# COMMISSION OF INQUIRY INTO FORENSIC DNA TESTING IN QUEENSLAND

Brisbane Magistrates Court Level 1/363 George Street, Brisbane

On Tuesday, 1 November 2022 at 9.30am

Before: The Hon Walter Sofronoff KC, Commissioner

Counsel Assisting: Mr Michael Hodge KC

Ms Laura Reece Mr Joshua Jones Ms Susan Hedge <CATHERINE JANET ALLEN, recalled, on former oath:</pre>

#### <EXAMINATION BY MR HUNTER:</pre>

THE COMMISSIONER: Mr Hunter.

MR HUNTER: Ms Allen, can you hear me and see me? A. Yes, I can.

Q. I'm going to return to the topic of the re-works on mixtures. Could we please have brought up on the screen, Mr Woolridge, [FSS.0001.0085.4217]. You were asked some questions about this email yesterday; correct?

A. Yes.

- Q. And this was an email drafted by you in response to the problem of mixtures being reported in one way initially only for the reported conclusion to be withdrawn and a different opinion substituted; correct?
- A. That's right. Yes. That had been a topic of discussion with QPS, yes.

Q. Sure. I accept that you say in that email that there's an ability for people to seek authorisation from you, but do you accept that the terms of that email are to effectively discourage scientists from re-working mixtures? A. I understand that that may be the way that it's read now. However, as Mr Howse and Ms Brisotto were aware of the lead-up to this particular email and the context around reviewing the spreadsheet, to provide confidence to the QPS that particular mixtures, you know, wouldn't change, they were aware of QPS's position on this, and so that was one of the things that I was saying to them, is that how can we - which we talked about, how can we ensure that the results that we're putting out to QPS won't necessarily change and how can we make that process better rather than thinking that the first option might be to re-work.

- Q. My question to you was do you accept that the terms of that email are to effectively discourage people from re-working mixtures?
- A. If you read that email on face value without the context, yes, that's how it appears.

Q. All right. So do I understand that the solution you devised was one that would avoid the issue of a result being withdrawn; correct?

A. No, because I couldn't guarantee the QPS that we wouldn't still amend results. So it was to try to ensure that we had better processes in the lead-up and then for me to be aware - an administrative process for me to be aware that results had been amended due to re-work so that I was across that and I was able to advise the QPS regarding why particular amendments had been made.

- Q. But you accept, don't you, that if a result is withdrawn then it's withdrawn because it's been detected by another scientist that an error has been made?
- A. Not necessarily. So there were there could be a human error within there. There was also other categories where additional reference samples had been submitted for the case after that particular mixture had been assessed, and also because there's a number of different staff members that may have reviewed or undertaken mixture interpretation within a particular case then there's also looking at the case as a whole and ensuring that all the mixtures were interpreted in a similar way, so from a case context perspective.

- Q. All right, let's forget about the word "error". If a result is withdrawn it's withdrawn because it's subsequently been found to be wrong?
- A. As I say, not necessarily --

Sorry, what was the --

Q. To be --

Α.

- Q. If a result is withdrawn it's withdrawn because
- someone subsequently looked at it and found that what had previously been reported was incorrect?
- A. As I say, not --

- Q. Surely you'd agree with that?
- A. Not necessarily, because of additional information that may have been provided.

- Q. Yes, but the additional information that's been provided has shown the earlier result to be incorrect; isn't it as simple as that?
- A. That's not necessarily the way that it's viewed. It's that additional information was provided which provides a different context to that particular mixture.

Q. Do you understand I'm not necessarily being critical

of the scientists who arrived at the first opinion?

 THE COMMISSIONER: Can we put it this way, Ms Allen: if a result is to be re-worked it's because, first, there's some doubt for some reason about the original result, and if the re-work results in a different result then that is because the first result is to be regarded as replaced by the second result; is that right?

A. Yes.

MR HUNTER: Thank you, Commissioner. Well, do you accept that, if your approach to this issue had the effect of discouraging scientists from undertaking re-works, you created a situation whereby incorrect results have gone

14 created a s 15 undetected? 16 A. No. be

A. No, because a reporting scientist would put forward the requests and the requests were always accepted, and they were able to undertake their re-work and provide information back to the QPS.

- Q. That assumes that the scientists ask you for permission to re-work in the first place. My question is premised upon the proposition that your attitude to re-works had the effect of discouraging scientists from doing them, and my question is: if it had that effect, do you accept that that created a situation whereby errors may have gone undetected?
- A. In a hypothetical situation as you've described, that could be true.

Q. All right. And you understood of course that the police would want to know if there were errors in the scientific opinions that had been provided to them?

A. Yes.

 Q. And in terms of a change in opinion having an adverse effect on a prosecution case do you accept that the police would want to know?

A. Yes.

Q. That the police would want to know if they had acted on a scientific opinion that might not have been correct? A. Yes. So that's why this process was put in place so that we could advise QPS at the earliest possible time regarding that.

Q. And isn't it the case that what the police were

- 1 seeking from you was that the initial advice perhaps be 2 couched in provisional terms, that is this is a provisional opinion? 3
  - I'm not sure I quite understand what you mean by that.

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- So when a mixture was first interpreted perhaps by one of the level 1 scientists that we know existed at the time isn't it the case that what the police asked is whether or not that report could be reported in a - sorry, that opinion could be reported in a cautionary way or a provisional way; that is, to effectively say to the police, "Here's a preliminary opinion, but if this is all you've got you shouldn't act on it"?
- That wasn't my understanding of the direction of QPS, that they were seeking assurances regarding the results wouldn't be amended so that they could act on those results.

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- Q. Did you see the EBN that went to the Police Commissioner?
- Α. No.

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Q. I showed you - you saw an extract from it yesterday? I think I only saw the extract. I don't remember seeing the EBN.

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See, it wasn't really possible for you to guarantee that results wouldn't change, was it? No, and that was part of the discussion that we had with the QPS, that they were wanting to tighten up that so

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Q. And so what the police wanted then, can I suggest, was that there be some moderation to the terms of the initial report so that they didn't go and lock someone up, to use the vernacular, on the basis of a DNA opinion that might subsequently change?

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Α. Yes, that's what they were seeking, yes.

that there were minimal amendments.

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- All right. Thanks, Mr Woolridge. I was asking you yesterday about the Options Paper, and I think we had got to the point where you agreed that the Options Paper might have been better if it had been worded differently, but nonetheless you are of the view that it placed before the police in a transparent way the pros and cons of what I'll
- call the DIFP process; that's your evidence? 46 47
  - That's what I remember. Yes, that's what I remember Α.

1 we got to yesterday. 2 Your evidence is that the Options Paper enabled the 3 QPS to make an informed decision about whether to adopt the 4 DIFP process? 5 Α. Yes. 6 7 8 What do you say to the proposition that the Options Paper was designed to induce the police to accept Option 2? 9 That's not the way that it was put forward to them. 10 It was put forward to them as, "We can continue to do the 11 process we're doing or there is another option should you 12 wish to undertake the other option." 13 14 15 But you were very keen, can I suggest, for the decision to be made by the police; correct? 16 Yes, because it had an impact on the QPS. 17 18 19 You wanted to make certain that if there were any adverse consequences from adopting the DIFP process they'd 20 21 be on the QPS, not you? No, that's not - that's not true. 22 Α. 23 Well, you knew when this inquiry commenced 24 All right. that the decision making around the Options Paper in DIFP 25 was going to be closely scrutinised; correct? 26 27 Α. Yes. 28 29 And your position, I take it, is today that the police, having been properly informed by the Options Paper, 30 made an informed decision; correct? 31 32 Α. Yes. 33 34 This was not a decision that was effectively forced on 35 them by the laboratory? Α. No. 36 37 38 Q. There had been a decision in May of 2013 with respect to volume crime, P3 crime, hadn't there? 39 40 Α. Yes. 41 42 Because initially volume crime was being processed using PP21? 43 44 Yes, that's right. Α. 45 46 And a decision was made to roll bulk crime back to Profiler Plus; correct? 47

1 Α. Yes, that's right. 2 And you in your position as managing scientist at the 3 laboratory were responsible for the collection and 4 collation of documents that were the subject of notices to 5 produce by the Commissioner? 6 For some of them, yes, and for ones that had been 7 8 given to me, yes. 9 One of the documents that was the subject of a notice 10 to produce was a document that related to that decision; 11 12 correct? 13 Α. Yes. 14 And it was a document that said this, amongst other 15 Q. 16 things: 17 In an effort to return to pre-Powerplex 18 results turnaround times, Forensic DNA 19 Analysis decided to return priority 3 20 21 volume samples to Profiler on 6 May 2013. 22 Correct? 23 Yes. 24 Α. 25 Can we perhaps put that document up on the screen, 26 [FSS.0001.0010.7078]. If you just scroll down a bit, please, Mr Woolridge, you can see in that second 27 28 29 paragraph - you can see that there, "In an effort to return to pre-Powerplex 21 results", in the second paragraph? 30 Yes. Yes. 31 Α. 32 That was a document that you found? 33 Q. 34 I believe it was a document that was collated and I was transferring it to MS Teams and checking that all of 35 the documents had transferred. 36 37 38 Q. And this was a historical document dating back to the time of this decision; correct? 39 I'm unsure. To be honest, I can't remember what the 40 date of the document was. 41 42 43 Well, it was created by someone other than you; 44 correct? 45 That's right. 46 47 THE COMMISSIONER: Mr Hunter, could you give me the number

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        of that document?
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        MR HUNTER:
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                      Sure. It's [FSS.0001.0010.7078].
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        THE COMMISSIONER:
                             Thank you.
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                      It was a document created by another staff
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        MR HUNTER:
        member, wasn't it? Didn't you look at the metadata?
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              Afterwards, yes.
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        Α.
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              Well, you understood, didn't you, that it was an
11
         historical document some years old that had been prepared
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         bv --
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        Α.
             Yes.
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              -- someone other than you; correct? Yes?
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        Q.
              Yes, that's right.
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        Α.
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        Q.
              And the version that you saw, was it in Word format?
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        Α.
              Yes.
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        Q.
              You changed it, didn't you?
              The original document was placed onto MS Teams --
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        Α.
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              Answer the question, please. You changed it, didn't
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         vou?
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              I --
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        Α.
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        THE COMMISSIONER:
                             You'd better be more specific what you
        mean by "change".
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        MR HUNTER:
                      You changed it by adding into that sentence
         that I've put to you - after the words "turn-around times,"
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         you added the words "the QPS requested forensic DNA
         analysis to return to priority 3 volume samples", didn't
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         you?
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              The following day I amended that and provided the
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         Α.
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         amended document.
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              In what universe did you think it was okay to amend a
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         document that was being asked for by this Commission?
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              My understanding was that putting forward an amended
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         document to Queensland Health legal, they would provide
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         legal advice around what next steps to take around that,
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        which is what I did.
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        Q.
              Why not just produce the document as you were required
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1 to do so?

A. I had produced the original document, and then I had amended the document and sent the amended document as well, so both documents were held by Queensland Health legal, and for me I was concerned that an error going forward in a document may be - I had felt I had an ethical dilemma around that because that wasn't correct.

- Q. Did you think it was ethical for you to amend a historical document?
- A. I amended the document and provided it to Queensland Health expecting legal advice regarding that.

Q. Did you think it was ethical for you to do that? A. Given that there was the original and an amended document, and I had put forward both.

- THE COMMISSIONER: What legal advice did you seek and who did you seek it from, and how did you seek it, orally or in writing?
- A. It was I believe that I had contacted Queensland Health legal contact the contact person, and then I had a phone call from Queensland Health legal team regarding the document and the amendment.

- Q. No, no, you said you asked for legal advice. What legal advice did you ask for let's stick with that for the moment or what advice did you need?
- A. I said that I had found an error in the document and that I needed to you know, it would be better if it was amended. They said, "Send the amendment." Then I was contacted by --

Q. Wait a minute. Wait a minute. What advice did you seek? You said you sought advice. What advice did you seek?

A. Around the amendment and the error in the document.

Q. But what did you ask for? What was the advice you asked for?

A. I was putting it forward that both of them were there, and my expectation was that legal advice would be of what I - what should go forward: the original document, the amended document, both documents with explanation.

Q. So you asked advice about whether both documents should go forward or only one of them; is that what you're

1 saying?

A. I said that I had found an error and that it should be amended and sent them that, and then that's when they contacted me regarding what the amendment was to discuss.

Q. Ms Allen, that's you telling them something. Your evidence a moment ago was that you asked for legal advice. I'm asking you what was the legal advice you asked for?

A. I guess that's where I'm saying that I put both of them forward expecting that they would provide me with that legal advice on what was the proper procedure.

- Q. So you didn't ask for legal advice; is that what you're now saying?
- A. Not specifically asking for it, but putting those two things to the legal team for their consideration.

Q. Mr Hunter.

 MR HUNTER: Thank you. Isn't what you did was to send a text message to someone who was then your - part of the Queensland Health legal team to say, "I've found an error in a document so I would like to fix the error and then replace the document"? Isn't that what you said?

A. I sent an MS Teams message to her, yes.

Q. Sorry, all right, but in that MS Teams message you said, "I've found an error in a document so I would like to fix the error and then replace the document"?

A. Yes.

- Q. Correct?
- A. That's right, yes.

Q. You didn't ask for advice, did you?

A. From my - from what I was trying to do was to amend that. The original document was already with the Queensland Health legal team, provide them with the amended document, and they would provide me with advice on what needed to - whether both documents went forward or just the original.

- Q. But why wouldn't you have just told the legal team,
  "Look, this is a document that we're required to disclose.
  Here it is, but it's wrong"? Why not just say that?
- A. And if I could go back and do it again that's exactly what would I do. But in the stress of the situation that's

1	obviously clouded my judgment because I was concerned about		
2	ensuring that we were providing accurate information to the		
3	Commission, and so I should have called Queensland Health		
4	legal to discuss with them before I did anything.		
5			
6	Q. But you weren't providing accurate information to the		
7	Commission by changing the document, were you? By handing		
8	over the amended document you would be representing that		
9	the records of the laboratory were different from the true		
10	position?		
11	A. No, my understanding is that the discussion with the		
12	QPS was around the turnaround times that were being		
13			
14	Q. No, no, I'm not asking you about that. I'm not asking		
15	you about that. What I'm saying is that you would, by		
16	changing this document and providing it to the Commission,		
17	be misleading the Commission because you would induce		
18	people at the Commission to think that that's how the		
19	document was when it was created, wouldn't you?		
20	A. That was not what I was attempting to do.		
21			
22	Q. Is this really emblematic of your sensitivity about		
23	decision making concerning DNA?		
24	A. No, it is not.		
25			
26	Q. You're so sensitive to the idea that your laboratory		
27	might be making decisions rather than the Queensland Police		
28	Service that you're prepared to amend a document, the		
29	production of which was compelled under notice?		
30	A. No, that's not true.		
31			
32	Q. Ms Allen, do you accept that it's imperative that the		
33	Queensland Police Service have faith in the laboratory that		
34	is testing its samples?		
35	A. Yes.		
36			
37	Q. You accept that the Police Service puts its trust in		
38	the laboratory to do its best work with the samples		
39	A. Yes.		
40			
41	Q that are provided to it?		
42	A. Yes.		
43			
44	Q. What do you say to the proposition that the Queensland		
45	Police Service cannot have any faith in the work of a		
46	laboratory with which you are in any way connected?		
47	A. I don't know what to say to that because the work that		

1	I've done with QPS has always been in good faith. I've			
2	worked with a number of different officers of different			
3	ranks regarding different forensic cases over a number of			
4	years. I've worked extensively with them to try to make an			
5	efficient process that allows them to obtain results. So			
6	from my perspective I have worked as best I can to ensure			
7	the QPS get the results that they need.			
8				
9	Q. And along the way you have lied or misled various			
10	members of the Queensland Police Service whenever it suited			
11	your purposes to do so?			
12	A. No, that's not true.			
13				
14	Q. Commissioner, I'll tender that document that's on the			
15	screen entitled "Volume crime processing in P+ instead of			
16	PP21".			
17				
18	THE COMMISSIONER: Exhibit 183			
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20	EXHIBIT #183 DOCUMENT ENTITLED "VOLUME CRIME PROCESSING IN			
21	P+ INSTEAD OF PP21"			
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23	MR HUNTER: I have no further questions. Thank you.			
24	Journal of the control of the			
25	THE COMMISSIONER: Yes. Mr Rice?			
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27	MR RICE: No, thank you.			
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29	THE COMMISSIONER: Ms McKenzie?			
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31	MS McKENZIE: No, thank you.			
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33	THE COMMISSIONER: Anybody else here?			
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35	MS FREEMAN: Yes, Commissioner.			
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37	THE COMMISSIONER: Yes.			
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39	MS FREEMAN: Thank you.			
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41	<examination by="" freeman:<="" ms="" td=""></examination>			
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43	MS FREEMAN: Ms Allen, can you see and hear me okay?			
44	A. Yes, I can.			
45				
46	Q. Great. My name is April Freeman. I act for			
47	Mr John Doherty. I just have a couple of questions to ask			

1 you in relation to exhibit 182. So if we could have that on the screen, please, Mr Operator. It's 2 [FSS.0001.0085.4217], just the email that Mr Hunter took 3 you to a short time ago. So it's an email that you sent to 4 Mr Howse on 25 January 2019; can you see that on your 5 6 screen? Yes, I can. 7 Α. 8 All right. And in that email you indicate there that 9 John Doherty had requested that you implement a process 10 where any reported sample is not re-worked without your 11 authorisation; can you see that there? 12 13 Α. Yes. 14 15 All right. Now, Mr Doherty started in the executive Q. director role in January 2019, didn't he? 16 17 Α. Yes. 18 19 Q. And that was executive director of the whole of FSS, 20 wasn't it? Yes, that's right. 21 Α. 22 And so that covers a broad range of areas, not just 23 24 the Forensic DNA Analysis Unit, doesn't it? 25 Yes. Α. 26 All right. And so you were reporting directly to 27 28 Mr Doherty at this particular time; is that right? 29 Α. Yes. 30 Along with a number of other people from other units 31 32 under the FSS banner; is that right? Yes, yes. 33 Α. 34 All right. And so in those first couple of weeks of 35 Q. Mr Doherty starting in this role as ED in January 2019 you 36 would have had some conversations with him about particular 37 38 issues within your unit; is that right? Α. Yes. 39 40 All right. And you had a conversation with him about 41 42 the re-working of reported samples in about his first week or two in that role, didn't you? 43 My recollection is that he attended a meeting with 44 myself and Superintendent McNab and Inspector David Neville 45 46 where this was discussed. 47

1	Q. All right. And it was your suggestion that you		
2	authorise any re-working of samples, wasn't it?		
3	A. My recollection is that Mr Doherty came to me - to my		
4	office and said that a process needed to be put in place		
5	regarding authorisations for re-works.		
6			
7	Q. All right. You'd been in the role of managing		
8	scientist for about 11 years up until this point, hadn't		
9	you?		
10	Ä. Yes.		
11			
12	Q. All right. I'm suggesting to you that it was your		
13	suggestion to Mr Doherty that you authorise any re-working		
14	of samples and he indicated he was happy for you to take		
15	that approach?		
16	A. My recollection is that Mr Doherty asked me to		
17	implement that process.		
18	rimpromorte that process.		
19	Q. All right. There was no formal direction issued to		
20	you about that, was there?		
21	A. No.		
22	A. NO.		
23	Q. I have nothing further. Thank you, Commissioner.		
23 24	Q. I have nothing further. Thank you, commissioner.		
	THE COMMISSIONED. Thank you Ma Engamon?		
25	THE COMMISSIONER: Thank you. Ms Freeman?		
26	MC FDFFMAN, Voc		
27	MS FREEMAN: Yes.		
28	THE COMMISSIONED. Con you namind me did Mn Debenty sive		
29	THE COMMISSIONER: Can you remind me did Mr Doherty give		
30	evidence about this point?		
31	MC EDEEMAN		
32	MS FREEMAN: He did, yes.		
33	THE COMMICCIONED AND A 15 LA		
34	THE COMMISSIONER: What did he say?		
35	NO EDEEMAN		
36	MS FREEMAN: His evidence was that it was a suggestion of		
37	Ms Allen's.		
38			
39	THE COMMISSIONER: Thank you very much. Yes. Yes,		
40	Mr Hickey?		
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42	MR HICKEY: Thank you, Commissioner.		
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44	<examination by="" hickey:<="" mr="" td=""></examination>		
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46	MR HICKEY: Ms Allen, I presume you can see and hear me		
47	all right?		

1 A. Yes, I can, thank you.

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- Q. Thank you. Could I ask you, first of all, why did you wish to work in this field in the first place?
- A. I liked the opportunity to be able to help the community without necessarily being close to, you know, offenders or victims of crime but being able to provide some type of service back to the community.

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Q. And you've been working in the lab for many, many years now. Presumably with your skills and qualifications and experience you could have sought employment elsewhere? A. Yes.

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- Q. Why have you stayed so long?
- A. Most people within forensics are very passionate about what they do and and I share that same passion with every single person within both of my teams, and I remain passionate about the service that we were delivering to the community, and I wanted to keep keep doing that role.

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- Q. Could I ask you, please, about the Options Paper that was delivered to QPS in 2018. From your perspective, what was the motivation for that process?
- QPS had always advised us that they didn't want us to do any unnecessary testing because they were aware that then that meant that turnaround times could be longer because we were spending time on items that weren't They provide us with information saying that items are no longer required for testing. So a review of a particular group of samples would show that 90 per cent of the time they weren't able to get any valuable information, 10 per cent of the time they were able to get some information, and then there was a 1.45 per cent regarding the NCIDD upload for that. The perspective that was put forward was, 'This is what we are currently doing and this is the outcome of that testing, or there can be a different option where testing is paused after quantitation, you can be advised of what that sample is, and then you can make a" - "QPS can make a decision regarding whether that sample's required, whether other samples will be submitted, or whether that sample can stay paused, and they may

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Q. Was that approach consistent or inconsistent with what you had understood to be the QPS preference for samples going back historically? Let me try it again --

revisit it later."

1 Α. Yes, my --2 Your evidence was that from time to time QPS would 3 tell you which samples they wanted to have tested? 4 Yes. 5 Α. 6 7 Q. And those which they no longer required to have 8 tested; is that right? 9 Α. Yes. 10 And so did you understand that what was being proposed 11 12 in the Options Paper was consistent with that approach? 13 Α. Yes. 14 15 Now, we've heard a lot of evidence about the way in Q. which the efficiency and efficacy of the lab is measured. 16 17 Is it the case that there are - that in your job you're 18 required to balance quality and quantity? Yes. 19 Α. 20 21 Q. Is it the case that scientists within the lab can 22 process samples at their leisure, or is there some 23 imperative to do it in an efficient way? We try to process - staff try to process samples in 24 25 the most efficient way possible to maximise the resources that we have. Reporting scientists can request for a 26 sample to be re-worked. If the final result hasn't gone to 27 28 QPS, they can undertake that at any time. 29 All right. Now, you mentioned maximising resources. 30 Can you tell us something about the nature of the resources 31 32 that were available to you from time to time? Additional funding was not forthcoming. 33 Requests that 34 I had put forward to the executive director - each 35 executive director regarding the budget I was managing and that it needed to be more than what I had, that wasn't 36 forthcoming. The Queensland Audit Office also undertook an 37 38 audit and showed within that document that over a period of five years \$1 million had been decreased from the Forensic 39 DNA Analysis budget for that. And then we went into a debt 40 and savings strategy from a whole of government. And in 41 this current financial year I've been asked to save 42 43 \$1.2 million from the Police Service's stream budget.

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All right. So am I right then in suggesting that the culture for you as a manager within which you were required to work was one of maximising savings rather than looking

1 for opportunity to spend more money? 2 Yes, that's right. Α. 3 And how long has that been the prevailing situation 4 within which you've worked? 5 For all the time that I've been the managing 6 scientist. 7 8 Now, can I ask you about the Options Paper itself. 9 Can you think of any personal benefit that flowed to you as 10 a consequence of QPS choosing one option over the other? 11 12 I gain no personal benefit whatsoever. 13 Was your personal remuneration linked to turnaround 14 Q. 15 time? No, it's not. I'm just a public servant. 16 Α. 17 Was your personal remuneration linked to the volume of 18 samples that the lab processed from time to time? 19 20 No, not at all. 21 22 Now, if I could ask you to turn your mind to the meeting that you had in February of 2018 with Mr Csoban, 23 Superintendent Frieberg and other members of the Queensland 24 Police; do you recall that meeting? 25 Α. Yes, I do. 26 27 28 To be clear, that's the meeting at which the 29 presentation was given about the Options Paper and its contents? 30 31 Α. Yes. 32 33 At the time of that meeting did you think the Q. 34 discussion was in any way controversial? 35 Α. No, I didn't. 36 37 Did you have any reason at that time to believe that it would become controversial in time? 38 No, I didn't. 39 Α. 40 Did you take notes of the kind that you would have 41 taken if you had expected the meeting would be 42 43 controversial? 44 Yes, I didn't take any notes because it was a free-flowing conversation; seemed positive. Yes, so I 45 46 didn't take any notes. 47

1	Q.	Now, chronologically that arose against the background			
2	of th	of the sperm microscopy controversy; you recall being asked			
3		some questions about that?			
4		Yes.			
5	,				
6	Q.	And in particular you were asked some extensive			
7	-	tions by my learned friend Mr Hodge about the			
	•	· · ·			
8		gement of ESR; do you recall that?			
9	Α.	Yes.			
10	•				
11	Q.	Now, that process included involvement by Clayton Utz,			
12	an ir	ndependent law firm?			
13	Α.	They were involved at the same time, yes.			
14					
15	Q.	And Crown Law?			
16	Α.	Yes.			
17					
18	Q.	And the HR department of Queensland Health?			
19	Α.	Yes.			
20					
21	Q.	Was that kind of body of external advice being			
22	-	ided something that happened regularly and routinely in			
23	-	experience within the lab?			
24	A.	No, it wasn't.			
	Λ.	NO, IC Wash C.			
25	0	In all the years that you've been the managing			
26	Q.	In all the years that you've been the managing			
27		ntist of the lab have you had regular contact with			
28	lawye				
29	Α.	There was contact in 2008 when the automated platforms			
30		an issue and we needed to engage with Crown Law			
31	•	ding advice around that, and then again in 2018			
32		n't recall off the top of my head if there was any			
33	other	time.			
34					
35	Q.	All right. Do you yourself have any legal training?			
36	Α.	No, I don't.			
37					
38	Q.	Do you regard the process of engaging with lawyers as			
39	being	g a natural one for you?			
40	Α.	No, not necessarily.			
41		<b>,</b>			
42	Q.	Is it a process that you found as a manager			
43		nidating?			
44	Α.	Yes.			
45					
46	Q.	And why is that?			
47		Scientists and lawyers speak different languages and			
<b>→</b> /	Α.	octentiata and lawyers speak utilierent languages and			

1	so I found it sometimes difficult to communicate the
2	scientific information to lawyers, and the legal process
3	and how that unfolds and the different parts of that
4	I definitely am not across. So I wouldn't necessarily know
5	what the next steps were for different parts of processes
6	because I'm not knowledgeable in that area.
7	O And in thinking about that CCD passes did you hald
8	Q. And in thinking about that ESR process did you hold
9	any concern about making inadvertent errors in your
10	engagement with the lawyers?
11 12	A. No. I provided as much information to them as possible.
13	possible.
14	Q. Was your concern to be responsive to their advices?
15	A. Yes.
16	Α. 163.
17	Q. Now, we've seen some evidence of your being involved
18	in the preparation of the terms of reference for ESR; do
19	you recall that?
20	A. Yes.
21	
22	Q. Have you ever before this occasion been called upon to
23	prepare a terms of reference of this kind?
24	A. Not to my recollection, no.
25	
26	Q. Were you given any assistance by people senior to you
27	within the department in the process of preparing the terms
28	of reference?
29	A. Not assistance in preparing, but Mr Csoban and
30	Mr Franklin reviewed the terms of reference that I put
31	together.
32	
33	Q. Do you recall Mr Csoban or Mr Franklin telling you,
34	whether orally or in writing, that any aspect of your
35	preparation of the terms of reference was deficient in any
36	way?
37	A. No, I don't have any recollection of that.
38	
39	Q. Do you recall there was a process by which Livingstons
40	were involved and engaged to provide some consultancy
41	services to the lab?
42	A. Yes.
43	
44	Q. Were you involved in preparing the terms of reference
45	for them?
46	A. No, I wasn't.

1 Now, you were asked some questions about what has been 2 described variously as the shredding party or bin gate; do you recall being asked about that? 3 4 Α. Yes. 5 The term "shredding party", was that something that 6 you came up with or was that a term described by somebody 7 8 else? That was the term that the staff member provided to 9 Α. 10 me. 11 12 All right. Did you have responsibilities around Q. document retention within the FSS lab? 13 14 Α. Yes. 15 What did you understand those to be? 16 Q. To ensure that the retention and destruction policy 17 was adhered to, particularly with documents that should be 18 retained either indefinitely or for specified periods of 19 20 time. 21 22 Now, that responsibility, was it something that you 23 only bore or did you have a responsibility to ensure others adhered to the document retention policy? 24 25 It's not just my responsibility. All staff members have that responsibility, and so I have oversight of that. 26 27 What did you think would happen if you turned a blind 28 29 eye to the concern that had been raised with you by that staff member about the shredding party? 30 That I could face some discipline action because 31 I hadn't acted and ensured that documents were either 32 retained or didn't find any. 33 34 And is that the reason you raised it with Mr Csoban 35 initially? 36 Yes. 37 Α. 38 And it came to be your evidence was that HR became 39 involved in the enquiries around that issue. Did anyone 40 from HR ever tell you that they weren't interested in the 41 42 issue? 43 No, they didn't. Α. 44 45 Did anyone from HR ever say to you, "Stop providing us assistance with this issue"? 46

I just didn't get any replies to my emails once

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Α.

1 2 3	I had undertaken the review of what was in the confidentia bin.
4 5 6 7	Q. Was that experience in things going cold at the HR end, was that a regular experience for you? A. Yes.
8 9 10	Q. Did anybody from HR at the time of these matters give you any reason at all to think that they regarded your conduct around the shredding party issue as being mad in any way?
12 13	A. No, they didn't provide any feedback on that, no.
14 15 16	Q. Did Mr Csoban give you any feedback to that effect? A. No, he didn't.
17 18 19 20 21	Q. Did you yourself have any reason at all at the time to think that your conduct was in any way aberrant in respect of that issue? A. No, I didn't.
22 23 24 25 26 27	Q. Now, could I ask you some questions, please, about the process of Workplace Edge having been engaged to provide some consultancy services around staff members in the lab; do you recall that? A. Yes.
-	Q. You attended a meeting at which their feedback was delivered to team members? A. Yes.
32 33 34	Q. Did you chair that meeting? A. No, I did not.
35 36 37 38	Q. Did you have input into the manner in which the feedback was delivered to the team members? A. No, I did not.
39 40 41 42 43 44 45 46	Q. Were you given any opportunity to modify the feedback that was to be given to team members before it was given? A. There was one opportunity where Mr Csoban asked me to amend a particular slide from the presentation. I don't remember what the modification was, but I remember supplying him with that, saying, "As discussed, amendment has been made," and then it was given to Mr Csoban and Mr Alan Holz from Workplace Edge.

- 1 Was your amendment one which Mr Csoban had directed or 2 was it one that you yourself composed?
  - No, it was one that Mr Csoban or Mr Alan Holz had asked to be made.

There's been some suggestion that you interfered with 6 the Workplace Edge consultancy processes. What do you say 8 to that?

- I did not interfere at all. I ensured that I remained out of those processes so that staff could independently talk with the Workplace Edge staff. Mr Alan Holz from Workplace Edge spoke to me one on one regarding, you know, issues that I may have had. So I spoke to him one on one. But I didn't interfere with any of the work that they were doing. I wanted them to be able to have free access to staff so that we could build a better culture moving forward.
- Was there any benefit to you in having a disenchanted staff within the lab?
- There was no benefit whatsoever to me.
- Were you aware that staff were feeling disenchanted within the lab?
- There had been discussions, and 2018 was a particularly difficult year. Some staff had felt that there was some low morale within their team, and the line managers were working with their teams to try to increase that.
- And that low morale, was that something that concerned you?
- Α. Yes.
- Q. What did you do about it?
- Within the Queensland Health each year there's a Working for Queensland survey. I encouraged staff to undertake the survey so that we could get results from that so that we could pinpoint particular areas where line managers could work with their teams on how we could improve particular areas that we may have low scores in to try to improve the workplace.
- 44 And did you discuss your concerns about the low morale with the line managers? 45
- 46 Yes. Α.

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- Q. In particular in first instance with Mr Howse and Ms Brisotto?
  - A. Yes.

- Q. Did you discuss with them strategies by which the morale might be improved?
- A. I don't have any independent recollection but that in my normal course of discussing things with them we would try to come up with strategies around how we could resolve situations. So, yes, I would have.

- Q. All right. Now, could I ask you to summarise, please, the breadth of your responsibilities as the managing scientist in the FSS lab?
- A. So from 2008 until 2013 the only team that I managed was Forensic DNA Analysis. Then from 2013 until now I now manage both Forensic DNA Analysis and also Forensic Chemistry. Forensic Chemistry is made up of three different groups. There's the illicit drug group, which test illicit drugs that are seized by QPS; there is the clandestine laboratory group, who test items that come from more of a manufacturing of the illicit drugs perspective; and there's the trace evidence group that does a diverse range of activities around chemical warfare testing, explosives testing, lubrication testing from sexual assault cases, those types of things. And then I still had the responsibilities within Forensic DNA Analysis. So overall I have responsibility for about 110 staff members within that.

- Q. All right. Now, Mr Howse has given some evidence that his experience in performing your role in an acting capacity was that your job is inherently stressful. Do you agree with his assessment?
- A. Yes. When I take leave and I ask each of my three direct reports, the chief chemist or either of the two team leaders from Forensic DNA Analysis, they're not overly keen to undertake higher duties in my role.

- Q. But, returning to you in particular, do you regard the role itself as being inherently stressful?
- A. Yes, it is.

- Q. And why is that?
- A. There's a number of different responsibilities across both of the different teams that require attention, and sometimes it's not just attention within one team, there's

attention that's required in both of the teams, and there's also responsibilities within the FSS leadership group as well about provision of information and meetings, meetings with the QPS, and those types of things. So there is a fair breadth of tasks to undertake every day.

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Q. Now, has that - has the pressure of the role - I'm sorry, and perhaps I should be clear about this. You do agree that it's an inherently stressful role? A. Yes, I do.

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- Q. Has the stress of the role become worse or better over the years you've been performing it?
- A. The stress has become worse.

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- Q. And why is that?
- The difficulty with trying to obtain resources for either of my two teams is extremely stressful. do more with less is really difficult. Trying to put forward, you know, additional - requests for additional funding is difficult. I've had a change of line manager over the past couple of years. From my experience within Pathology Queensland, they don't necessarily understand the forensic aspect of what we do, so that is difficult to communicate that. And over the last number of years there's been a change almost annually of the HR officer that I interact with, so that becomes difficult to then catch - you know, to explain the situation that may be going on, what advice had been previously given, and then the next lot of advice. And over the last couple of years we've also had a change in a finance officer, so then providing information to them around the types of consumables that we purchase, how expensive they are, that you can't necessarily try to reduce expenses in particular areas because it's inherent in the nature of the business.

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Q. Now, you've mentioned that you've had a number of line managers. I presume you mean --

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- THE COMMISSIONER: Sorry, Mr Hickey, just so that I don't forget. Ms Allen, you mentioned the finance officer. What's the split of responsibilities, if there is a split of responsibilities, between you as managing scientist and the finance officer? What is a finance officer's role in relation to you?
- A. The finance officer provides the budget to budget allocation, to police services stream. It's my

responsibility to manage that, and they will also review that and flag where expenses have gone over what was expected and help try to forecast if there are upcoming expenses, whether they're known expenses - like, if there was a particular police operation and there needed to be overtime, if we can forecast that we can put that in as a known expense. So liaising with them. They need to understand the types of consumables that we purchase, so why they are expensive, and, you know, the cost of those is usually international, so there's the international - the exchange rate, I should say, between the Australian dollar and the American dollar to take into consideration.

Q. Yes, thank you. Yes, Mr Hickey.

MR HICKEY: Thank you, Commissioner. Now, you mentioned a moment ago that you have responsibilities both for the DNA testing lab and also for Forensic Chemistry?

A. Yes.

Q. Forensic - the forensic chemistry side is also concerned with very serious crimes, isn't it?

A. Yes, that's right.

Q. Have you experienced anything like the kinds of problems which seem to have beset the DNA lab in that team? A. No.

Q. Now, you've given some evidence just a moment ago about funding and the difficulties in seeking additional funding. We've heard some evidence throughout the course of the Commission from Mr Drummond to the effect that if you had asked for additional funding it would have been given. Is that consistent with your experience as the long-time manager of the lab?

A. No.

Q. Because there's a process, isn't there, by which you could have made a formal request for further money?

A. Yes, that's right.

Q. You could propose a business case?A. Yes.

- Q. There doesn't seem to be any evidence of that having occurred; do you agree?
- 47 A. Yes, that's right.

- Q. And why is that?
- A. In consultation with the executive director about putting forward a business case, it would require different aspects around costings and those types of aspects. I've been through a costing exercise with the finance department over the years I think I've been through that on four different occasions but we've never finalised that, so I don't have the costing data to be able to provide within a business case for those types of things, and at the time the executive director didn't see that it would be worthwhile putting up a business case because he didn't see that it would be approved.

- Q. Now, when you say that costings weren't finalised, you didn't have those things finalised, was that something within your control or somebody else's?
- A. Within the finance department, it's within their control to finalise that.

Q. And how much of your time in your job do you suppose is --

THE COMMISSIONER: Mr Hickey, I just want to be clear so I understand this. What finance department? Can you just elucidate it?

MR HICKEY: Yes, of course. When you say that sat within the finance department, who precisely or what precisely do you refer to?

A. Within Queensland Health finance department --

Q. External to -A. -- or the - yes, external to FSS, or within the - what was previously called the health support Queensland finance group.

THE COMMISSIONER: So what stopped you asking for data that you would need to prepare a business case?

A. I had asked for that - for the data, but the actual costing around how much does it cost for a sample to go through evidence recovery, then process through - so a costing from end to end of a sample had never been completed, and that takes into account various factors, and that's what I had worked with the finance group on to actually get that so that we could see how much did it cost for a particular sample so we could base, you know, the

1 budget, et cetera moving forward on that.

- Q. So why didn't you do it? Why didn't you work up the costings?
- A. I don't have the expertise to do that, which is why I was working with the finance group to actually do that.

Q. Yes, Mr Hickey

 MR HICKEY: Thank you, Commissioner. How much of the time spent in your job is dealing with funding constraints?

A. It varies across the year. In the lead-up to the budget being set there is a fair amount of time put into reviewing what budget may be put forward, and then once the budget is provided it's reviewing what has been provided, and then on a monthly basis preparing a finance report for the FSS leadership team meeting.

Q. Would you describe it as a significant part of your role as managing scientist?

A. Yes.

- Q. Now, you said a few moments ago that you considered that the stress, the inherent stress, in your role had increased, not decreased, over time. Can I ask have you sought support from the department to scaffold you, given that situation?
- A. I've attempted to work with each of my line managers, the executive director, regarding that. I have at times sought HR support. But at different times I have gone outside that and sought the advice of my own two medical practitioners to assist me with that.

- Q. All right. To the extent you feel comfortable telling me about it, what's the nature of the assistance that you've sought from external medical practitioners?

  A So from my GP that I've been seeing for a long time
- A. So from my GP that I've been seeing for a long time I've sought advice and assistance from her both in 2018 and most recently, and in 2018 and most recently I've also been under the care of a psychologist.

Q. And to what extent do you regard the need for that care as being a consequence of your work environment?

A. One hundred per cent of that care is required due to work.

Q. All right. Now, you gave some evidence a few minutes

ago that you'd worked with a number of line managers over the years. I presume you were referring specifically to executive directors?

A. Yes, that's right.

Q. With how many executive directors have you worked?

A. I think it's six.

- Q. And has that number of executive directors with whom you've worked created a positive or a negative impact on your ability to do your job?
- A. Each of the people that I've worked with I've really enjoyed working with them, but it has made it difficult because some haven't come from a forensic background, so there is information needed to be provided around the nature of the work that's undertaken, some of the issues that we may face liaising with the QPS, how we operate in those two different teams. Within Forensic DNA Analysis there's a single point of contact with QPS, whereas in Forensic Chemistry there isn't a single point of contact, so there is different arrangements within those two teams and how they may and how the executive director may be involved in any of those meetings. So it's a lot of information transfer to bring them up to speed on where we're at at that particular time.

Q. I asked you a moment ago whether you had sought support from the department to scaffold you and your work, and you said that you had engaged with the EDs and with HR. To the extent that you did that, has the support that you've been provided been sufficient, in your mind?

A. No, that's why I've gone externally.

Q. And what could have been provided to you which hasn't been provided to you?

A. I think if we had a more stable HR person available at FSS rather than a number of changes, then you would be able to develop a relationship with them, plus also they would be across the issues and would also be working with you to resolve some of those issues. I understand that the executive director role, you know, can change, but that is also - can be also disruptive with change for that. So developing that relationship with them takes time for them to understand that. And from my perspective I have engaged with the employee assistance service previously, but I didn't find that that was of assistance, which is why I've gone outside and sought external help with that.

1 2 3 4	Q. All right. Now, we've heard quite a lot of evidence about the suggestion that there was a pervasive toxic culture within the lab; you're aware of that?			
5 6	A. Yes.			
7 8 9	Q. Was that your experience? A. In some teams it was, but in other teams, no, as in within Forensic DNA Analysis			
10 11 12 13 14	Q. In which A. Within Forensic DNA Analysis some of the teams had good culture and in other teams not as good culture.			
15 16 17 18	Q. Is there anything that you identify as being significant as to why some teams might have a good culture and others might have a bad culture?  A. From my perspective I think there are some staff			
19 20 21 22 23	members that have a strongly held belief regarding me, and any efforts that I have made over the years to change and grow as a person and a leader haven't necessarily been noted within that.			
24 25 26 27	Q. If you turn your mind to those particular people, do you regard yourself as having been supported by those staff? A. No, not at all.			
28 29 30 31 32	Q. Have you had any experience of being white-anted by those staff? A. Yes.			
33 34 35	Q. In what way? A. If there was an issue with something that I had done, a process that was attributed to me, they wouldn't			
36 37 38	necessarily come to talk to me about it. They would go to my line manager instead. So they wouldn't approach me so that we could discuss that, or even send me an email to			
39 40 41	discuss that, or ask a HR staff member to mediate a session with me to discuss that. They would go to my line manager, and in some instances my line manager hadn't come and			
42 43 44	talked to me about what those issues were, so I felt that I didn't get an opportunity to change or provide clarification around that because I was unaware of what the			
45 46	issue was.			

Q.

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And did you ask your line manager to give you clarity

around those complaints that had been received? Some of those managers, yes. Most recently I wasn't aware of the number of staff that had gone to my most recent line manager, Ms Keller.

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And what difference might it have made to you if you had been given detail and particulars about the nature of the concerns that had been expressed to your line manager? That I could have sought advice from my line manager regarding next steps on how to handle that in the best Possibly sought assistance from a coach or a possible wav. mentor. Work with another staff member within the wider group of Queensland Health on how to tackle particular issues. Work closer with the HR person. Perhaps also try to have a facilitated conversation with that staff member to try to reach a resolution.

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19 20 Q. Did knowing that complaints had been made to your line manager but not clearly articulated to you make you more or less trusting of the people that you worked with? Α. Less trusting.

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And what impact, if any, did that have upon your decision making and judgment as the manager of the FSS lab? It did have an impact, particularly around the Forensic DNA Analysis management team and sharing information with the management team because there wasn't necessarily a high level of trust within the team, and so I was uncertain about where some information may go. even if I said to them that it was confidential I didn't necessarily know whether it would remain confidential or not.

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And so did that cause you to change your practice in respect of the kind of information that you might or might not share from time to time?

Yes, I became more guarded and was more careful about sharing information and trying to share as much as I could but in the best possible way so that if it was shared with others my true meaning would be understood rather than, you know, a different view, that I was trying to ensure that they did understand what I was saying.

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- Now, were you aware that it had been suggested that you had described Ms Rika's team as "those fuckers over there"?
- Α. No, I was not.

1	0	the way you decembed them?
2		the way you described them? o recollection of that whatsoever.
4	A. I liave li	o recorrection of that whatsoever.
5	Q. And did	anybody ever raise that concern with you?
6	A. No, they	
7	,, e,	4.4
8	Q. And were	you made aware that Ms Rika and members of
9	her team desc	ribed themselves as "the FRIT fuckers"?
0	A. No, I wa	s not aware of that at all.
1		
2	Q. Now, you	were asked some questions about questions
3	•	d by email to your management team about staff
4	members who m	ight be contemplating becoming pregnant; do
5	you recall th	at yesterday?
6	A. Yes.	
7		
8	•	email said something about the consequences -
9		hrasing, but if people weren't identified
20	•	e dire consequences if they were to fall
21		they hadn't been identified; do you know what
22	I'm referring	
23	A. Yes, con	sequences on a budget, yes.
24	O Now sou	Id I ook you some things shout that . Was that
25 26		ld I ask you some things about that. Was that
27		that email was referring to about the funding of those who might become or were pregnant,
28		ding in respect of those who would backfill
29		ile they were taking leave?
30	•	n respect to funding for the maternity leave
31		the staff member may take, because the
32		heir recreation or long-service leave could be
33		thin the budget, but it was the maternity
34		that wouldn't be included in the budget, so
35	therefore the	position would have to remain vacant until
36	the maternity	leave period had finished.
37		
38	Q. I see.	Was there ever any intention on your part to
39	• •	members of your staff should not become
10	pregnant?	
11		at all. We have a high percentage of female
12		was just trying to ensure that we could get
13	•	1 that so that the position didn't remain
14		at period of time, which added stress to the
15	other team me	mbers.
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Did you ever suggest to anybody that if they should

Q.

1 become pregnant they wouldn't be entitled to parental 2 leave? Α. No, not at all. 3 4 5 Did you intend to suggest that? Q. Α. No, not at all. 6 7 8 All right. Now, you've mentioned that there's a high percentage of women who work within the laboratory. 9 you responsible for hiring scientists who come to work in 10 the reporting side of the lab? 11 No. I'm not usually on a recruitment panel for a HP4 12 13 reporting scientist, no. 14 15 All right. Who would be on a panel for those Q. positions? 16 17 It would be the senior scientist at a HP5 level or it could include a team leader at a HP6 level, and would 18 usually also include an external staff member, and 19 usually - given that it's a reporting staff member, we 20 21 would usually ask a QPS officer to be on the panel. 22 23 All right. Thank you. Now, could I ask you some Q. things about flexible work arrangements. Was it the case 24 that you had delegation to make decisions about flexible 25 work arrangements? 26 No, I don't. 27 Α. 28 29 Q. Who had that delegation? The delegation sits with the executive director. 30 Α. 31 32 Has that always been the case while you've been the manager of the lab? 33 34 Α. Yes. 35 36 Do you provide information to assist the executive director in making that decision? 37 38 Α. Yes. 39 Have you ever had any reason to consider that the 40 Q. executive directors did not make those decisions 41 42 independently of any views you might have expressed? 43 No, I don't. Α. 44 45 Did you ever bring pressure to bear on executive

ones?

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directors to make decisions that you thought were the right

1 Α. No. 2 Is it easy to fill roles of reporting scientists in 3 4 the lab generally? No, they're more difficult to fill a reporting 5 scientist role, and the training usually takes anywhere 6 between nine months and 12 months to complete before 7 8 they're able to become productive and issue statement of witness documents. 9 10 Now, if roles were left vacant for any 11 All right. period of time would that impact the ability of the lab to 12 13 do its job? Yes. 14 Α. 15 And ultimately is it your responsibility to ensure 16 17 that the lab's roles are filled? 18 In conjunction with the team leaders, yes. 19 20 Q. That is to say I'm not asking you about the process of 21 interviewing for them, but if staff can't be recruited to 22 those roles is that ultimately your problem? It is a problem that we will need to tackle, yes, 23 because if we don't have a staff member in that role then 24 there's more work for everybody else that's in the team. 25 26 And so, given that, is there any benefit to you that 27 you can identify in having dissatisfied staff in the lab? 28 29 There is no benefit to me whatsoever. 30 Now, you've been depicted, can I suggest to you, 31 32 through the questions you've been asked and the evidence of some of the others called in this Commission, as some kind 33 34 of Disney villain; is that your impression? Yes, that's how I feel, yes. 35 Α. 36 37 Q. What impact has that had on you? I find it quite - quite distressing. It upsets me, as 38 I'm just trying to do the best job I can 39 vou can see. because I care about the community and I want to try to 40 provide as many resources to the lab so that they can do 41 42 the best possible job that they can. I work really hard to do that. I'm a human being and I make mistakes, and it's 43 been really - these past 12 months has been really 44

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which I'm sure others can appreciate, because of

distressing, and it has had an enormous impact on my mental

and physical health. I am suffering mental health issues,

the stress. I've never been through a commission of inquiry before. I've been through the ministerial taskforce in 2005, and it was a very different experience than this one has been. I've found that this one has been quite personal to me, and I'm not quite sure how to take that because it has had an impact on me and has definitely affected my mental health.

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- Q. What impact has that had on your ability to give assistance to the Commission in answering the questions you've been asked over the last few days?
- you've been asked over the last few days? I've found it very challenging to provide information. I've been open and willing to provide information. I've provided as many documents as I possibly can. over different notices a number of times to ensure that I've provided everything. If I felt that there was something that was even remotely included, I've provided I've tried to do it in the fast possible manner that I've worked longer hours than required to try to I have. I've worked weekends as well to ensure that I can do that. provide everything possible. I don't want to hide anything from the Commission. I want to make sure that they have everything that they need from me. But I have found it

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Q. Now, you were suspended from your duties as the managing scientist some weeks ago now?

A. Yes, that's right.

challenging over the past couple of days.

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31 32 Q. At the time that occurred were you actively engaged in assisting the Commission of Inquiry with its investigations?

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Α.

A. Yes, I was.

No, I haven't been.

34 35

Q. Were you given any explanation for the reason for your suspension at the time?A. No, I wasn't.

36 37 38

Q. Have you been given any explanation since?

40 41 42

43

44

45 46

47

- Q. What effect did the suspension have upon you?
- A. I went into shock when I was told. I was given five minutes to leave the campus. I found it incredible that I would be given such distressing news and then allowed to drive a motor vehicle home. I was in shock for a good portion of the day I was told. A friend was concerned

about me, so she came to be with me because she was concerned what I might do. I lost two days from being able to help the Commission because when I was stood down they didn't advise me that I needed to continue working for the Commission, and then the following day was a public holiday, so they were the two days that I had lost to be able to help, and then it - only clarity was given on the Thursday night that I needed to continue to work on the Commission information from home.

- Q. Has the fact of your being suspended affected in any way your ability to give comprehensive evidence to this Commission?
- A. Yes.

- Q. How?
- A. It has been extremely stressful to be stood down not knowing why. To try to seek, you know, any information about that I've been excluded from the work place. So the normal support mechanism that I would have is no longer there. So I'm even more isolated than before. There's only limited people that I'm allowed to contact for support. So that has made it more difficult. And just the mechanism of working from home is difficult as well. So there's two different things on my mind. There's the Commission of Inquiry and the need to provide information, and then there's also the suspension on my mind as well.

 Q. Has anybody from the department reached out to provide you employee assistance since you were suspended?

A. Yes. Ms Lara Keller offered a session with the Queensland Ambulance Priority One service, and I've had one session with that staff member.

THE COMMISSIONER: Mr Hickey, would you prefer to continue or adjourn for morning tea?

MR HICKEY: No, let's adjourn, please, Commissioner.

THE COMMISSIONER: Yes. We'll adjourn for 20 minutes, Mr Hickey.

#### SHORT ADJOURNMENT

THE COMMISSIONER: Mr Hickey.

MR HICKEY: Thank you, Commissioner. Ms Allen, can you

1 still see and hear me? 2 Yes, I can. Α. 3 You're aware that we're now entering the sixth week of 4 hearings in the Commission and you're aware of all of the 5 people who have been called to give evidence so far, 6 I presume? 7 Yes. 8 Α. 9 Do you regard there as being people who ought to have 10 been called to give evidence to give the Commission the 11 12 complete picture who are employed by FSS but who have not been called? 13 Yes. 14 Α. 15 And who are they? 16 Q. 17 I'm aware that some of the management team members have supplied statements but haven't necessarily been 18 called, and some other staff members have reached out to me 19 to let me know that they would like to have been able to 20 21 put something forward but they're too scared because they 22 may be stood down or they may be treated a particular way 23 by the Commission, and so from that they won't come forward. 24 25 And in what way do you consider their evidence might 26 have assisted the Commissioner? 27 I think that it would have shown a balance, that 28 29 there's different issues within the different teams, and that there is cohesion across most of the teams and 30 provided some of the positives. 31 32 33 Now, my learned friend Mr Hunter asked you this 34 morning about the process of re-works, and I think in one of your answers you gave to him you suggested that you'd 35 never refused a request for re-work; is that so? 36 Yes, that's right. 37 Α. 38 Did anyone ever suggest to you that they or other 39 staff held the belief that they were not permitted to carry 40 out re-works? 41 42 Α. No. 43 44 Did anyone ever suggest to you that they or other staff felt as though re-works were discouraged? 45 46 Α. No.

1 I presume in the course of your work you spoke with Ms Brisotto and Mr Howse every day? 2 Yes. 3 4 Did either of them ever say anything to lead you to 5 Q. conclude that they thought re-works were not permitted? 6 7 Α. 8 Q. 9 Or that re-works were discouraged? Α. 10 11 12 Q. Now, in December 2021 the media began to take interest in the activities of the lab; do you recall that? 13 14 Α. Yes. 15 This wasn't the first time that you had experienced 16 the brutal glare of the media upon the lab's operations, 17 18 was it? That's right. 19 Α. 20 21 And so did the fact that the media were interested in 22 the lab's operations cause you any concern? 23 Yes. Α. 24 25 Q. Why was that? Because the lab operates in a fairly structured manner 26 with standard operating procedures that they adhere to. 27 All of the staff members do their very best every single 28 29 day to achieve good outcomes. So to have negative media regarding the lab is of a concern and stressful for all 30 staff members. 31 32 Did it have any effect on the stressfulness of your 33 Q. particular role? 34 35 Α. Yes, it increased the stress, yes. 36 And what effect, if any, did that increased stress 37 have upon your ability to make clear decisions? 38 It did have an impact. Now that I review that in 39 hindsight, the level of stress was much elevated because 40 the negative media continued cyclic, so it appeared 41 42 never-ending. 43 What do you say to the suggestion that you were like a 44 deer in the headlights at that point? 45 46 I don't think that that's an accurate description. The first day that I returned from leave and became more 47

fully aware of what had happened while I was on leave, briefings that had occurred, et cetera, I was quite shocked. But then the next day I turned my mind to, well, what are the types of things that we could actually do to help us in this situation to try to get onto the front foot.

- Q. Was the fact that you had a relatively new executive director to whom you answered a help or a hindrance, given the unfolding situation?
- A. Ms Keller was really trying to help, but I don't necessarily think that she understood the gravity of the situation because it related to forensics, and, having been through this before, my best action plan was, you know, to take some action to have the laboratory reviewed by another jurisdiction, if that could be possible, to have case files reviewed by other jurisdictions, that we try to work out how we can ensure that, you know, we are the best possible we are working in the best possible way to be able to keep continuing and to show staff that, you know, we have faith in them.

- Q. Was it business as usual from your perspective from December 2021 until the time you were suspended a month or so ago?
- A. Not necessarily business as usual. There was added pressure regarding the negative media. But we were trying to undertake business-as-usual activities. I'm not sure that those activities were able to be done at the same level once the Commission of Inquiry was announced, and staff required to produce statements or documents for notices, et cetera, reduced their available time for results or processing samples.

Q. Did you feel in June, when you were canvassed for information about how to undo the DIFP process, that you were obliged to make decisions quickly?

A. Yes, I was.

Q. And do you, with the benefit of hindsight, consider you ought to have had more time to make those decisions?

A. Yes, and, the benefit of hindsight, have a colleague from Forensic DNA Analysis review the options so that they

could spot my human error before it went to Ms Keller.

Q. Now, you were asked some questions yesterday about why Mr Howse's Update Paper did not become a project; do you

1	recall that?
2	A. Yes.
3	
	O To it the sees that all impostingtions understated in
4	Q. Is it the case that all investigations undertaken in
5	the lab proceed by way of project?
6	A. Not necessarily all, no.
7	
8	Q. And you said that you expected that the work that
	· · · · · · · · · · · · · · · · · · ·
9	might be undertaken through that Update Paper might also be
10	undertaken as part of an external review?
11	A. Yes.
12	
13	Q. Was it typical in the lab to duplicate work that was
14	proposed to be undertaken by an external consultant?
	·
15	A. No.
16	
17	Q. Now, you were asked some questions towards the end of
18	the day yesterday by the Commissioner about your
19	qualifications for the role you hold. Can I ask you some
20	questions about that. When you applied for the role was it
21	a process of competitive selection which led to your being
22	appointed?
23	A. Yes.
24	
25	Q. Do you recall there being other applicants for the
26	role?
27	A. Yes.
	Α. 103.
28	O D
29	Q. Do you recall who comprised the panel that interviewed
30	you?
31	A. Yes, I do. I remember two of the three people. One
32	was Greg Shaw, who was the executive director of FSS, one
33	of the other panel members was Inspector David Neville, and
34	I can't remember who the third panel member was.
	1 can t remember who the third paner member was.
35	O Did was marries and band area 5
36	Q. Did you receive any hand-over from the previous
37	managing scientist?
38	A. A small hand-over from her before she went on her
39	secondment, yes.
40	
41	Q. Do you consider that that was an adequate hand-over?
	·
42	A. Yes.
43	
44	Q. Has anybody ever suggested to you before yesterday
45	afternoon that you're not properly qualified for the role
46	you hold?
47	A. No, they have not.
• •	it its, they have her

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1
2
              Has anybody ever suggested to you that your
        performance as the managing scientist fell well short of
3
4
         expectations?
5
              No, they have not.
        Α.
6
              From time to time you've acted up as the executive
7
8
        director of the lab; is that so?
             Yes, I have.
9
        Α.
10
              Have you ever received any negative feedback about
11
        your performance in that acting position?
12
13
             No, I have not.
14
15
              Finally, Ms Allen, can I ask you this. We find
        ourselves here in a $6 million Commission of Inquiry,
16
        having sat here for six weeks exploring the entrails of the
17
         lab, and of course part of the Commissioner's role is to
18
        make findings about what has happened, but, importantly,
19
        part of his role is to make recommendations about what
20
21
         should happen in the future, and I want to ask you this:
22
         upon reflection, upon considering all of the things you now
         know, how did we end up here?
23
              I was given the team in 2013.
24
                                             That meant that I had
         extra responsibilities, regardless of my time and
25
         availability to do that. I think that change in 2008 when
26
        QPS changed the business model with us, that change staff
27
28
         didn't necessarily embrace.
                                      But from where I was standing
29
         that change was coming whether we embraced it or not.
        we went through the automated platform issues that we had,
30
        which, you know, created disharmony within the team.
31
32
        Moving forward, you know, trying to be innovative to assist
        QPS around the Options Paper, I see that staff didn't
33
34
         embrace that process and that more work was required within
         that process so that both the QPS and FSS were on the same
35
36
         page for that. To be really helpful, we tried to engage
         three different mechanisms to help with the culture,
37
38
         particularly within the management team, and that didn't
         necessarily work either. The first was with a psychologist
39
         that could help us, then with Workplace Edge, and then
40
         later with Tess Brook. But we didn't ever seem to have any
41
42
         success with maintaining the trust and the working
         relationship that we had with each other while we were
43
         going through that, and we'd gotten to a place where we
44
        weren't able to have robust scientific discussions because
45
46
         they would become - they turned into people taking it quite
         personally instead of trying to be able to discuss the
47
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science. And so as a group we are not cohesive and not moving in the same direction even though we may have the same passion for the work that we're doing.

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Q. And have you attempted to address those problems you've just described in good faith?

A. Yes, I have. Absolutely in good faith I have tried to do my best to work with everyone that I can to - because, for me, the ultimate goal is to help the community. So what can we do to ensure that we can provide good quality results to the QPS in a turnaround time that allows them to prevent and disrupt crime, that's the focus, and I understand that people view that I only look at the operational side of the work, but for me that's where I feel that I can be of most benefit, is on the operational side, to try to help the staff to actually achieve those

17 18 19 qoals.

Q. Thank you, Commissioner. Those are the questions.

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41 42 THE COMMISSIONER: Thank you. Ms Allen, I just wanted to ask you about the change that happened in 2013 that you just mentioned. I see from your curriculum vitae attached to your statement that you had been the managing scientist of the DNA Analysis Unit between 2008 and 2013, and then you were appointed managing scientist for the Police Services stream, and your responsibilities obviously increased substantially both in terms of the work that was being done by people under you and also a substantial increase in the number of people under your supervision. Can I ask you this: prior to your appointment to that position in 2013 who was running the other parts of the Police Services stream? What was the organisational structure before you were appointed in 2013? The Chemistry - so Forensic Chemistry had been under a chemical analysis group and had been headed up by a managing scientist for that chemical analysis group, and they had managed Forensic Chemistry and other chemistry That managing scientist took a areas within that. voluntary redundancy through the Newman era, and so all of the work groups within that chemical analysis group were then split up and given to different managing scientists, and I was given Forensic Chemistry.

43 44 45

Q. I understand. Yes, I understand. Thanks. Mr Hodge?

46 47

MR HODGE: Thank you. I did have some further questions.

1 2 THE COMMISSIONER: Yes. Go ahead. 3 <EXAMINATION BY MR HODGE:</pre> 4 5 Ms Allen, Mr Hickey asked you some questions 6 MR HODGE: about your involvement in the presentation of 7 8 the information that came back from Workplace Edge; do you recall that? 9 Yes. 10 Α. 11 And I just wanted to understand that and then show you 12 Q. some documents. What was your involvement in the 13 formulation of the presentation that was provided or 14 15 delivered to the staff members at that meeting in January of 2018? 16 17 I don't believe I was involved in the formulation, but I had been given a copy to review. 18 19 20 And tell me if you agree with this: you saw a couple 21 of versions of the presentation as it had been originally 22 done by Workplace Edge? Yes, I think I did see a couple of versions, yes. 23 24 25 Q. And then you edited versions of the presentation? I was asked to edit one version of the presentation, 26 and, as I say, I can't recall what that was, but then 27 forwarded that back to Mr Csoban and Mr Holz, and said to 28 29 them that, you know, I had edited it as they had requested. 30 31 And then when the presentation was delivered on the Q. 32 day you say it was just Mr Csoban who delivered the presentation? 33 Yes, that's my recollection. 34 Α. 35 And did you speak at all, as you remember? 36 Not as far as I remember, unless someone had asked me 37 38 a direct question, but I don't remember speaking, no. 39 40 I see. And before the PowerPoint presentation was created do you recall that Workplace Edge prepared a draft 41 42 report? Yes, they did a draft themed report. 43 44 45 And did you see a copy of that? Q. 46 Α. Yes. 47

1 Q. And did you have the opportunity to comment on that? 2 Not as far as I remember. The themes report then Α. became the presentation that was provided to the staff. 3

4 5

- You don't remember getting a copy of the report and responding with your comments by email?
- I may have, and so I may have forgotten that. I don't recall whether I did on the themes report.

8 9 10

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- And you don't remember tracking changes into the document prepared by Workplace Edge?
- I could have. As I say, I don't have a recollection around that.

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- Do you agree with this proposition: the Workplace Edge process wasn't something that happened independent of you but, rather, you were intimately involved in providing feedback on what came from Workplace Edge and with Mr Csoban formulating the way in which that would be presented to the staff?
- Α. I was asked to provide some feedback, but that was on the basis that they had written their report and I wasn't involved in the formulation of that report or the presentation.

24 25 26

27 28

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I'll show you some documents. So first can we bring up [FSS.0001.0083.4025]. So you see this is an email that somebody from Workplace Edge sent to Mr Lok and Mr Csoban on 10 January 2018 and then Mr Csoban forwards to you? Yes, the themes document, ves. Α.

30 31 32

33

34

35

- Yes. And was it the case that Workplace Edge weren't meeting with you but they were meeting with Mr Lok and Mr Csoban?
- That's probably right. I'm not sure. I don't I'm sorry, I don't recall.

36 37 38

39

And you see the version that Mr Csoban is sending to vou is a PDF?

Yes. 40 Α.

41 42

43

44

45

And he's sending that to you on 11 January. Nobody else is copied into that email; it's just Mr Csoban forwarding it to you. Was that common, that Mr Csoban would just send things to you and not copy anyone else in? It depended on what the topic was, whether it went to other people or myself.

_	
1 2	Q. And then if we - sorry, Commissioner, I'll tender that
3 4	email.
5 6	THE COMMISSIONER: The email of 11 January 2018 from Mr Csoban to Ms Allen is exhibit 184.
7 8	EXHIBIT #184 EMAIL OF 11 JANUARY 2018 FROM MR CSOBAN TO
9  0	MS ALLEN BARCODED [FSS.0001.0083.4025]
1   2	MR HODGE: Then if we could bring up [FSS.0001.0083.4017]. So this is the attachment to that email, the themes
3  4  5	document as you've referred to it? A. Yes.
6   7	Q. And I think you know you reviewed this at the time? A. Yes, I would likely have read it at the time, yes.
8  9 20	Q. I think you probably know you definitely read it at the time because you provided detailed comments on it, as
21 22	we'll come to in a moment.  A. Okay, yes.
23 24	Q. Do you remember that?
25 26	A. I don't have any independent recollection of it, no.
27 28 29	Q. Do you remember that one of the things that Workplace Edge recommended in their initial draft was a significant restructure of the way that the lab worked?
30 31	A. I don't necessarily remember that now, no.
32 33 34	Q. Do you recall that they described the model that was being used within the laboratory as the production line model?
35 36	A. Yes, it has been referred to as that, yes.
37 38 39	Q. And they - and I'll show you this. If we go to page 2 of the document, at the bottom of the page you see in $1(a)$ they say:
10 11 12 13	The production line model has not achieved the optimal delivery of services under the current structure with the existing systems
14	and processes, and resource allocation.
16	A. Yes.
39 40 41 42 43 44	The production line model has not achieved the optimal delivery of services under the current structure with the existing systems and processes, and resource allocation.

1 Q. And you see they also say: 2 The organisational structure does not fully 3 support the current operating model as 4 illustrated by comments provided by staff. 5 6 7 And then you see they go on to say: 8 The Team, as a whole is over-governed with 9 10 supervisors managing approximately 60 10 staff, giving a ratio of 1:6 actuals and 11 between 1:4 and 1:5 FTE. 12 13 Yes. 14 Α. 15 And then if we go over the page I just want to note 16 two more things. You see they raise an issue about 17 projects taking too long to establish and complete? 18 Yes. 19 Α. 20 21 And they raise an issue about the quality function; do you see that in the next dash? 22 Yes. 23 Α. 24 25 And then if you skip over the next one about pay, but Q. then you see they raise two further issues about the 26 production line concept? 27 28 Α. Yes. 29 And tell me do you remember them raising these 30 criticisms of the production line concept now? 31 32 Not specifically, no, but I see it in the document, Α. 33 yes. 34 35 If we then come to page 5, which is .4021, you see starting at the bottom of the page they have 36 recommendations? 37 38 Α. Yes. 39 And Option 1 is, "Process integrated team approach", 40 and they say this would involve a shift from the production 41 42 line model? Yes. 43 Α. 44 45 And then if we go over the page we can see - if we could keep scrolling, Mr Operator, thank you - we can see 46 they have a model for what they describe as the process 47

1 2 3	integrated team approach as a possibility? A. Yes.
5 5 6 7	Q. And then if we keep scrolling down the page we see they have what they describe as Option 2, which is an "Enhanced production line model"; and you see they say:
8 9 10	This option would involve structural and process changes to address many of the concerns expressed above.
11 12 13	A. Yes.
14 15 16 17	Q. And if we then go over the page you can see then they have this different form of structural chart that they propose? A. Yes.
19 20 21 22 23 24 25 26	Q. And then we can see if we go to the bottom of the page they have their recommendation 1, which is about considering options for operational model and structural change. Then if we go over the page we see recommendation 2 is, regardless of what operating model is chosen, that there needs to be strengthening of quality and projects? A. Yes.
27 28 29	Q. And they said the role of quality should report directly to the managing scientist? A. Yes.
30 31 32 33	Q. All right. Now, do you agree with me you rejected the possibility of structural change in the lab?  A. No, I don't recall rejecting change.
34 35 36 37	Q. Do you agree with me that you rejected changes to strengthen the quality function?  A. I don't recall rejecting that, no.
38 39 40 41 42 43 44 45 46 47	Q. I'll show you a document. Now, I'll just check if you've got this, Mr Operator. My version doesn't have a doc ID. Could we bring up, Operator, [FSS.1000.0079.8813]. That doesn't help. It may have just been emailed to you. Thank you. So if we could just scroll down first just so that Ms Allen can see. So you see you sent an email - if you can scroll up very slightly just so you can see the header. You sent an email on Thursday, 11 January, in the morning giving detailed feedback about that themes

1 document? 2 Α. Yes. 3 And if we scroll up to the top, and then a little 4 later - I'm not sure if you can see the header time, but 5 this was sent at 10.30 in the morning - you sent a further 6 email dealing with the recommendations and you see you say: 7 8 9 I support the spirit of recommendation 1, however I don't support either of the 10 proposed Org structures put forward. 11 12 13 Α. Yes. 14 Q. And then you see recommendation 2 you say: 15 16 By placing quality and projects under the 17 managing scientist, the expectation is that 18 the managing scientist is able to achieve 19 more things than others, which isn't a good 20 21 assumption to move forward with. I don't support this but I wouldn't oppose it if it 22 is framed very carefully with the staff. 23 24 25 Α. Yes. 26 And then you see at the bottom of your email there's a 27 Q. 28 line which is: 29 There hasn't been enough focus on fixing 30 the issue with reporting - as reporters 31 32 have advised Justin that they have laid everything on the line with Allan - and 33 34 this report won't satisfy them at all. 35 36 Α. Yes. 37 38 And tell me if you agree with this: the senior managers in reporting under Mr Howse were Ms Rika and 39 Ms Reeves at the time? 40 Α. Yes. 41 42 And your view was that they were the problem? 43 Q. No. I was trying to advise that the reporting staff 44 had put forward all of their ideas around this process and 45 46 that it hadn't necessarily addressed that. 47

1	Q. I see. I tender that email, Commissioner.
2	THE COMMISSIONER: What's the date of it, Mr Hodge?
4	MD HODGE: 11 January 2019
5 6	MR HODGE: 11 January 2018.
7 8	THE COMMISSIONER: Exhibit 185.
9	EXHIBIT #185 EMAIL SENT BY MS ALLEN DATED 11/01/2018
∣0 ∣1	MR HODGE: Now, then, then at some stage you were provided
2	with a Word document version of that themes report; do you
3	recall that?
4   5	A. I don't specifically recall it, but okay.
6	Q. And you went through and you tracked in comments into
7	the Word version of the report?
8	A. Okay.
19 20	Q. And if we bring up [WIT.0019.0016.1688], if you look
21	at the bottom of the page there's an email from Mr Csoban
22	to you where we can't see there's an attachment but
23	presumably there's an attachment and he says, "For review
24	please"?
25 26	A. Yes.
27	Q. And if we scroll up the page you see you email back a
28	version to Mr Csoban?
29	A. Yes.
30	
31	Q. And then if we go over the page, Mr Operator, do you
32 33	see the start of the document, so this is the themes document, and in blue we can see just the changes that
34	you're tracking to the document?
35	A. Yes.
36	
37	Q. And then if we go over the page. Now, this is a bit
38	confusing, but I want to suggest something to you. We see
39	both red and blue on this document. What I want to suggest
10	to you is both red and blue come from you.
11 12	A. Okay. Why would you say that?
13	Q. I'll show you why.
14	A. That both red and - okay.
15	· ·
16	Q. You see in the - sorry, can I just ask something.
17	This is a document that's actually attached to one of your

1 witness statements. Have you reviewed it any time in the 2 last couple of months? No, I haven't, I'm sorry; no. 3 4 5 Q. How would it get attached to one of your witness statements without you reviewing it? 6 I would have used that as some type of evidence for a 7 8 particular question. 9 Q. 10 I see. You see in the red it says: 11 12 It is not accurate to say that the production line model has not achieved the 13 optimal delivery of services. 14 15 Α. Yes. 16 17 And then, Mr Operator, could I get you to bring back 18 up that email that we were looking at, which was the last 19 20 exhibit. 21 22 OPERATOR: Side by side? 23 Yes, if you could do it side by side, that 24 MR HODGE: would be great. Thanks. And if you could scroll down to 25 the email that was sent at 9.20 in the morning, and just 26 stop there. It may be hard to zoom it in, but do you see, 27 Ms Allen, in that email on 11 January at 9.20 (a) says: 28 29 It is not accurate to say that the 30 31 production line model has not achieved the 32 optimal delivery of services. 33 34 Α. Yes. 35 And what it looks like has happened is that you have 36 copied your initial comments from 11 January over into the 37 document, and they are in red, and then you have in tracked 38 over them in blue and so the blue is also your 39 contribution; do you agree with that? 40 That's possible. I don't remember doing that, but 41 42 that's possible. 43 44 And it looks like - you tell me if you agree with this - you rejected the criticism of the production line 45 46 model?

Α.

47

I didn't necessarily think it was an accurate

1	port	rayal given that there had been an end to end process
2	that	had been conducted.
3		
4	Q.	And you see - I think we can take down the email now,
5	Mr O	perator, and that will allow us to blow up a bit better
6	what	's happening on the page. Could you just blow up the
7	last	section at the bottom of that page, that's right, the
8	-	highlighted in blue including the comment. So you see
9	what	Workplace Edge had said is:
10		
11		The Team as a whole is over-governed with
12		10 supervisors managing approximately 60
13		staff.
14	Α.	Yes.
15		
16	Q.	And then your comment is or the words you add is:
17		
18		Staff members highlighted that the
19		reporting teams were over-governed by
20		supervisors and proposed that the two
21		reporting teams could be merged into one
22		team with one supervisor.
23	_	
24	Α.	Yes.
25		
26	Q.	And then you see you have a comment to the side which
27	is:	
28		T
29		Its my understanding that the comments made
30		about over-governed were restricted to the
31		reporting teams, not the whole team.
32	۸	Voc
33	Α.	Yes.
34	0	And then you say
35	Q.	And then you say:
36 37		Its Workplace Edge's opinion that the whole
38		team is over-governed. This needs to be
39		made clearer.
40		made Crearer.
41	Α.	Yes.
42	Λ.	163.
43	Q.	And I want to just clarify some things about that.
44		supervisors in the reporting teams, that was Ms Rika
45		Ms Reeves?
46	A.	Yes.
47	, , ,	
• •		

- Q. And tell me if you agree with this: what you were trying to bring about was the abolition of one of their positions?
  - A. No. Sorry, I thought you were going to go on. No, not necessarily, because we would use that position somewhere else within the lab.

Q. Yes. But they each held a particular role as a supervisor of a reporting team, and you wanted to get rid of one of them as the supervisor of a reporting team?

A. If you merged the two teams then the other HP5 senior scientist position could be utilised somewhere else within Forensic DNA Analysis.

- Q. I understand. But I think if you just come back to my question. They were each the supervisor of a reporting team, and you wanted to get rid of one of those positions held by one of them?
- A. As I said, that was my understanding of what the team had put forward.

Q. And I just want to understand that. When you say "what the team had put forward" tell me if you agree with this: this was supposed to be anonymous feedback being reported to Workplace Edge as external consultants?

A. Yes.

Q. And you weren't sitting in on the interviews? A. No.

Q. That would have defeated the entire purpose of it? A. Yes.

Q. And so when you said staff wrote in the documents, "Staff members highlighted that the reporting teams were over-governed by supervisors and proposed that the two reporting teams could be merged into one team with one supervisor," that statement couldn't have come from you having actually participated in the interviews that were the basis of this document?

A. No.

Q. Do you honestly say that you thought it was legitimate for you to be seeking to manipulate the results of what was supposedly the anonymous feedback from a workplace survey?

A. I was not manipulating the results of the survey.

1 2 3 4 5	Q. Ms Allen, we can see it in writing what you did? A. That information had been - so what was the date of this email to me and the email regarding the presentation? Weren't they at about the same time?
6 7 8 9 10	Q. I don't understand. This email is 12 January 2018. The presentation I think from memory was 23 January 2018. A. That's when the presentation was given to the staff members, but I thought that presentation had been provided earlier.
12 13 14 15 16	Q. I see. If you're asking when was the first draft of the PowerPoint representation provided by Mr Csoban to you it was on 17 January, five days later.  A. Okay.
17 18 19 20 21	THE COMMISSIONER: Mr Hodge, is this document that we're looking at that was given to Ms Allen by Mr Csoban for her comments a document that was in due course supposed to become the report of Workplace Edge?
22 23	MR HODGE: Yes, as I understand it.
24 25	THE COMMISSIONER: Right.
26 27 28	MR HODGE: But, as it turned out, I think this is fair, the Workplace Edge engagement ended early.
29 30	THE COMMISSIONER: Yes. But at this stage
31 32	MR HODGE: So it was never finalised.
33 34 35	THE COMMISSIONER: At this stage it was a draft that was circulated for comment.
36 37	MR HODGE: Yes.
38 39 40 41	THE COMMISSIONER: And the alterations that were made to it were made to it with the aim in due course that that would be the form of the report that Workplace Edge would submit?
42 43 44	MR HODGE: Yes.
45 46 47	Q. So I'm sorry, Ms Allen, I understood you were explaining that you weren't trying to bring about a change in the opinion from Workplace Edge and you said - you asked

- the question about when the PowerPoint presentation was provided.
  - A. My recollection is that Workplace Edge had provided an overview to Mr Csoban and myself. I can't remember whether that I think that was a verbal overview which is where my comment had come from, because I wasn't involved in any of the interviews and this information came from Workplace Edge, which is what I was trying to say within my comment, that if this is what they had told us previously that this is what's in this document then it needs to be clearer around that.

- Q. So you're saying you thought Workplace Edge had told you at some earlier time that staff members had said that the reporting teams were over-governed and proposed that the two reporting teams could be merged into one team with one supervisor?
- A. That's my recollection, yes.

- Q. And when do you say that happened?
- A. At around this time. It was either late December or in the January around this time.

Q. I see. But you can't tell us when it was?

25 A. I'm sorry, I can't - I don't remember.

- Q. And was it a briefing, as you recall, just to you and Mr Csoban?
- A. And I think also Mr Peter Matthews may have also been there as well from Workplace Edge.

- Q. And this was something separate from the direct engagement that Workplace Edge was having with Mr Csoban and Mr Lok?
- A. Yes, that's my recollection; yes.

Q. And why - tell me if you can - was it necessary for Mr Csoban to be separately forwarding these documents on to you for your comment if you were involved in any way with the formal process by which Workplace Edge was engaging with him and Mr Lok from Queensland Health?

A. He asked for my opinion on the document. That's all I can say from that.

Q. And then I want to suggest to you - I don't want to go through them now, but I will if I need to - then from the 17th versions of PowerPoint presentations started being

1 prepared and you - I think you might have edited a couple of versions of the PowerPoint presentation for Mr Csoban? 2 Okay. I don't remember how many. 3 4 5 I don't want to suggest to you that you made any Q. significant changes other than in one respect. Do you 6 recall that there was an issue that Workplace Edge raised 7 8 about Project 181 taking too long? I don't specifically recall that, no. 9 10 Is it possible that you removed the reference to 11 12 Project 181 taking too long from the PowerPoint 13 presentation? I don't remember. So it's possible. I don't remember 14 15 what I did for those presentations. 16 17 THE COMMISSIONER: 181 was the spermatozoa? 18 If I could then just get the operator to 19 MR HODGE: Yes. bring back up that email chain once more that we were 20 21 looking at. Can we just scroll down again so we can see 22 Ms Allen's email. You see your opening line is: 23 24 My feedback is based on ensuring that adequate details have been supplied within 25 the document given that it is highly likely 26 to be requested through RTI? 27 28 29 Α. Yes. 30 And what was going on at the time or what had begun by 31 32 this time was that Ms Reeves was making RTI applications? I'm not sure whether it was at that time or whether 33 34 I was anticipating that there would be. I didn't think it was that early in the year, but I could be wrong because 35 I don't have the dates of the RTIs. 36 37 38 In any event when you refer to it being "highly likely to be requested through RTI" the person that you 39 anticipated requesting it through RTI was Ms Reeves? 40 She was most likely, but there were other staff that 41 42 could have also requested it. 43 44 But she was the one who was foremost in your mind, wasn't she, at the time? 45

Yes.

Α.

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- Q. And so what you were seeking to do I think you use the phrase "ensuring that adequate details have been supplied within the document", what you were seeking to do, though, was make it or get the document into a form where you would be content for it to be produced to Ms Reeves if she requested it by RTI?
  - A. No, that was not my intent. The intent was to ensure that there was adequate detail within the document so that it could, for want of a better word, stand alone because it had enough detail in it.

Q. But, Ms Allen, look at the words that you use:

My feedback is based on ensuring that adequate details have been supplied within the document given that it is highly likely to be requested through RTI.

THE COMMISSIONER: That's the point you're making, Mr Hodge?

MR HODGE: I'll put it more bluntly.

- Q. What you were seeking to do was to ensure that whatever opinions or statements were made by Workplace Edge were statements that you would be happy with if they were produced to Ms Reeves?
- A. No, that's not right.

- Q. You didn't want any statements that might be supportive of her criticisms of the operation of the laboratory?
- A. No, that's not right.

Q. And then I want to ask you then one last thing, which is about that meeting with Workplace Edge. Can we bring up another document which is [FSS.0001.0083.4210].

THE COMMISSIONER: At some point, Mr Hodge, can you make plain what point you're making about the foot of that amended report of Workplace Edge, the last two lines in blue relating to reporting scientists wanting to get rid of one of the positions?

MR HODGE: Yes, I thought I might have put it. But I can put it in a different way.

THE COMMISSIONER: I just need to understand the point and be sure that Ms Allen has had an opportunity to respond to it, whatever it is.

MR HODGE: I might just do it now.

Q. Ms Allen, what I'm putting to you is this: you wanted to get rid of Amanda Reeves; do you agree?

A. No.

- Q. And the problem with the feedback that Workplace Edge was saying they had received was that they were suggesting that as a whole the DNA lab had too many managers, and that wasn't feedback that you wanted?
- A. No, that's not true.

- Q. The feedback that you wanted was that one of the positions held by Ms Reeves or Ms Rika should be abolished?
- A. No, that's not true.

- Q. And so you set about adding as a comment the suggestion that that was in fact the feedback that had been received, even though that wasn't how Workplace Edge in writing had summarised the feedback?
- A. As I said, that was my understanding of what had occurred and so that they needed to clarify that so that there was understanding, because I was confused by it.

- Q. If the evidence you're giving now is true, tell me this: why would you not simply add a comment to the side saying, "Is this description of the feedback accurate, because I had thought that Workplace Edge had said that the feedback was something different"? Why would you take the trouble to edit the document to type in different words as the feedback?
- A. Because Mr Csoban had asked me to review that document and so my understanding was to add information to it. He didn't ask me just to make comments on it. He asked me to review it. So that's what I did.

- Q. I have to suggest this to you, Ms Allen. Once again this explanation that you're giving doesn't make sense and that's because it's not true.
- 45 A. I'm not lying.

Q. Now, you see this email that you sent to Mr Csoban and

1 Allan - you can't see who the other person is but it's Allan from Workplace Edge? 2 Α. Yes. 3 4 5 And you sent it on 25 January 2018, so this is two days after that all staff meeting? 6 7 Α. Yes. 8 9 And you see what you're setting out is some feedback that you'd received from Mr Howse about feedback that he in 10 turn had received from Ms Rika? 11 Α. Yes. 12 13 And you see it says the feedback was at least half the 14 Q. staff had said that the presentation didn't represent their 15 views or what they provided in the interviews? 16 17 Α. Yes. 18 Q. And then you see two dashes down it says: 19 20 21 It was put forward that Allan, Cathie and 22 Paul only put forward the things that they wanted to say and not the views of the 23 staff. 24 25 Α. Yes. 26 27 28 Now, I'm interested in that. How could it be that you were putting forward any views at this meeting if, as you 29 said in evidence a day ago, you were just sitting in the 30 audience? 31 32 That was their perception from the preparation, is my understanding of that dot point. The staff's perception 33 34 was that the views that had been put forward was mine and Paul's. 35 36 37 Q. And then you see the last paragraph says: 38 I'm concerned about this type of discussion 39 and the affect that it will have on the 40 staff member who have opened up. And 41 I would welcome any ideas on what we could 42 do to quell some of this nonsense. 43 44 45 Α. Yes. 46 47 Q. And by "quell some of this nonsense" you meant the

1	dissent that Ms Rika was expressing and saying that other
2	staff felt about the things that had been said at the
3	meeting?
4	A. No, it was about how all of that discussion would

A. No, it was about how all of that discussion would affect the team and we were trying to move forward with the team. So discussions like this needed to be quelled in some way or additional feedback provided from Workplace Edge on their specific points so that staff could understand that the comments had come from the staff and not come from others so that that could possibly help them move forward.

Q. You agree with me that what was presented on 23 January - we can leave aside for the moment who presented it - but what was presented was things that you and Mr Csoban were happy to have presented to staff?

A. What was presented to them was generated by Workplace Edge and that was the process that we undertook. It wasn't necessarily that I was happy about the content.

Q. I don't have any further questions. Sorry, I tender that email, Commissioner.

THE COMMISSIONER: Exhibit 186.

# EXHIBIT #186 EMAIL FROM MS ALLEN TO MR CSOBAN AND ALLAN DATED 25/01/2018

THE COMMISSIONER: Mr Hickey.

MR HICKEY: Yes, Commissioner.

THE COMMISSIONER: I think this line of questioning arose out of some of the evidence that you elicited about the efforts that were made to bring in external consultants and so on. But Mr Hodge's questions extended over a range. So I wondered whether you wanted to cover any of it.

MR HICKEY: No, thank you, Commissioner.

THE COMMISSIONER: All right. Thanks. That concludes your questioning, Ms Allen. You're free to cut the link now, if you wish.

A. Thank you.

<THE WITNESS WITHDREW

1 THE COMMISSIONER: Mr Hodge, what's next on the agenda? 2 MR HODGE: Commissioner, we now seamlessly transition to 3 module 5, which is the evidence of Ms Baker and Dr Kogios. 4 What I was going to do was give a short opening about that 5 evidence and then we were going to call them. 6 7 going to appear via video. 8 THE COMMISSIONER: 9 Yes. 10 I'm sorry, there's one other thing I have to do MR HODGE: 11 which is there's a large number of additional documents 12 which are to be tendered in line with what you indicated, 13 Commissioner, that the parties should have certainty about 14 15 what you're going to look at. 16 THE COMMISSIONER: 17 Yes. 18 19 MR HODGE: So can I hand up - these are broken down into for each different topic the additional documents that are 20 21 to be tendered. So I might just formally hand that up and 22 then in due course we'll do something where we allocate exhibit numbers. 23 24 Why don't you keep it until you're 25 THE COMMISSIONER: ready to make it part of the record in some form that can 26 be useful to anybody reading the transcript. 27 28 29 MR HODGE: I'll have a further discussion with my team over lunch about how that's to go up. 30 31 32 THE COMMISSIONER: All right. 33 So, Commissioner, I'll just say something, it 34 MR HODGE: 35 will probably take me about 10 minutes, I think, just to run through some things. 36 37 38 THE COMMISSIONER: And has everybody got the expert report that you're going to be speaking about? 39 MR HODGE: Commissioner, in the last five weeks of 41 Yes.

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hearing we've heard much about the past and present practices of the laboratory, and you have heard evidence from scientists, police and experts, and then over the last week from managers involved in the management of the In this final module, which will probably take laboratory. a day to a day and a half, you'll hear evidence from two

highly experienced forensic DNA scientists, Dr Rebecca Kogios and Ms Heidi Baker, who have conducted a review of the current operation of the lab. That review involved a thorough consideration of whether the lab's operations are consistent with international best practice, what needs to be done to rectify deficiencies that they identified, and where the lab should review or retest samples which have been dealt with in a way that has in the past been inconsistent with best practice.

I might bring up the expert report. Can we bring up [EXP.0007.0001.0001]. So this is the expert report that Ms Baker and Dr Kogios have prepared, and it's dated 28 October 2022. I might just tender it now, Commissioner, and it can go as an exhibit.

THE COMMISSIONER: Yes. The report is exhibit 187.

# EXHIBIT #187 EXPERT REPORT THAT MS BAKER AND DR KOGIOS BARCODED [EXP.0007.0001.0001]

MR HODGE: I might just tell you very briefly about the two experts, Commissioner, and then we won't need to traverse through that again when they give evidence.

THE COMMISSIONER: Yes.

MR HODGE: Ms Baker has a bachelor of science with honours in genetics. She's currently a forensic senior scientist at the highly regarded Institute of Environmental Science and Research, which is what is referred to as ESR, and you've heard references to ESR before, and ESR is in New Zealand. Ms Baker has over 20 years experience as a scientist and senior scientist in forensic DNA laboratories and is currently based in the forensic research and development team at ESR providing technical advice for genomics research and social systems projects.

 Dr Kogios has a PhD in molecular biology and bachelor degrees in science and law. She is the executive director of the forensic services department which is part of the Victorian Police and she has had a career of more than 20 years in forensic biology. She has responsibility for all forensic services provided by the Victorian Police, including forensic DNA.

Between them the two experts have worked at four

forensic service providers in three different countries and have a combined experience of more than 40 years in forensic DNA. They were provided with over 2,300 documents that had been compiled by the Commission to cover the current operations of the lab, and that included a full set of the current standard operating procedures, recent quality system documents and management team meetings, and they were also given the other expert reports that you obtained, Commissioner, for the purposes of this Commission.

THE COMMISSIONER: That is 2,300 document, not 2,300 pages.

MR HODGE: Correct.

THE COMMISSIONER: Each document maybe containing dozens or hundreds of pages.

MR HODGE: Sometimes many pages.

THE COMMISSIONER: Yes.

MR HODGE: Many pages. They visited the laboratory over four days between 21 and 27 September 2022, and that included tours and explanations of relevant parts of the laboratory and interviews with staff. They were involved in speaking to staff of the laboratory on 22 occasions from reporting, analytical and evidence recovery scientists to senior scientists, team leaders, the managing scientist, the quality manager of FSS and the executive director of FSS.

 They have in their report identified a number of features of the laboratory that fall below best practice. Some are high level and they affect the way that the lab as a whole functions and others are technical, and we will canvass a number of them in their oral evidence today and likely continuing into tomorrow.

Ms Baker and Dr Kogios also deal with a number of issues in relation to the governance and structure of the laboratory, and they identify some difficulties with the current structure for the management and governance of the lab. They have made 47 recommendations to bring the lab back into line with best practice.

Before we call them I wanted to just outline very briefly a few of the issues that they have discussed in their report. The first key issue is about workflow, something, Commissioner, that you've heard about a number of times over the course of the last five weeks. Ms Baker and Dr Kogios identify a number of features of the current workflow that create a potential issue, including that there is no dedicated case manager for cases other than those designated as P1. The lab tests everything it receives, and the lab uses a work list system so that reporting is done on a sample by sample basis without full case context.

No DNA and DIFP results when they were still reported were reported without a reporting scientist considering the results at all. A whole of case review would only occur when a statement was required for court, and they express the opinion that that for reasons they identify is problematic because the risks of this workflow are significant.

One risk is the missed opportunity to obtain all available forensic evidence. Another is the risk of not detecting contamination or unexpected results, and also the potential overservicing and damage to the relationships of the lab both internally in the sense of the trust of its scientists and externally in terms of the trust of the QPS.

Ms Baker and Dr Kogios consider that this workflow model, the production line model, presently in use falls below what is required to be best practice, and they consider it results in suboptimal triage and case review in some cases. Ultimately, as you'll hear from them, they recommend some changes that could be made or considered in terms of the operation of the lab.

In relation to the issue of thresholds, something that you've heard a lot of evidence about, Commissioner, Dr Kogios and Ms Baker consistently, with what other experts have said, have found that the lab's approach to thresholds falls below best practice. The no DNA threshold could be best practice if properly validated and properly explained to end users of the results, but neither is the case at present. The DIFP threshold is not supported by Ms Baker or Dr Kogios at all. Rectification of this issue will require a review of all samples reported as DIFP and immediate proper validation of the limited detection. In

terms of future thresholds Dr Kogios and Ms Baker suggest no threshold at all for serious or complex crimes or only a limited detection threshold which would be able to be overruled by a scientist at the scientist's discretion.

Ms Baker and Dr Kogios also found a number of other features of the lab's current operations to fall below best practice, including the lack of Y-STR testing, the reporting of incorrect results to police, the elution and auto-concentration practices, and the change in cleaning protocols for bones that was implemented in 2019. They also make a number of recommendations in areas where practice can be improved.

In relation to SAIKs they recommend the provision of feedback to collectors of SAIKs, accreditation of kits, and the formation of an interagency group to provide advice on best practice. They also recommend changes to the content of the SAIKs, including extra swabs, a swab to take a reference sample, materials to take fingernail clippings, and to make slides for microscopy.

 In quality management they are concerned by the current arrangements which involve the quality senior scientist, who is embedded in case work and has no power to overrule a manager, and the quality manager of FSS, who plays an advisory role only. They recommend a quality manager role which is dedicated solely to the forensic case work and a quality lead within all DNA analysis teams.

In the structure of the lab they recommend the division of managerial and technical roles, with the management role for sole responsibility for DNA analysis and a technical lead to serve as the custodian of scientific help. Ms Baker and Dr Kogios also recommend the development of research, development and innovation capabilities in the lab to ensure they keep apace of scientific developments in the field.

Ms Baker and Dr Kogios also identify significant problems in the culture of the laboratory, including strained relationships, inability to resolve scientific differences, disconnect from the other labs and opaque decision-making. However, as you will hear, Commissioner, I expect very early in their evidence they have a very high regard for the scientists who work in the laboratory and were very impressed by the quality and experience of those

1	scientists. They identify the features of a healthy
2	scientific workplace culture as being one in which you can
3	raise concerns without the fear of retribution, one that
4	encourages innovation, and one that is supported by a
5	values based culture.
6	
7	Dr Kogios and Ms Baker will give evidence
8	simultaneously and by video link; Dr Kogios in Victoria and
9	Ms Baker in New Zealand. Is it convenient, Commissioner,
10	if we call them now and commence their evidence?
11	The same time and commences their evidence.
12	THE COMMISSIONER: Of course. Yes, certainly.
13	THE COMMITCOTOMERY.
14	MR HODGE: I call Dr Kogios and Ms Baker. I'm told they
15	should be in the link. I can see Ms Baker. I should say,
16	Commissioner, just while we're waiting for Dr Kogios to
17	join, I'm going to address a few topics with the experts
18	and then I'll hand over to Ms Hedge and she'll address
19	other topics. Can we just adjourn for a moment and we'll
20	get the technology working?
21	THE COMMICCIONED. That Is all winds I had not it would not
22	THE COMMISSIONER: That's all right. Just get it working.
23	I'll wait.
24	MD HODGE O I I I I I I I I I I I
25	MR HODGE: Commissioner, I understand the operator has
26	asked us to adjourn so he can sort out the feedback.
27	THE COMMITTEE AND ADDRESS OF THE COMMITTEE AN
28	THE COMMISSIONER: All right.
29	WD 110005 0 16
30	MR HODGE: So if we can adjourn for five minutes.
31	
32	THE COMMISSIONER: Yes.
33	
34	LUNCHEON ADJOURNMENT
35	
36	THE COMMISSIONER: Ready to go, Mr Hodge?
37	
38	MR HODGE: I am. Just before we do that can I just hand
39	back up that list of documents to be tendered
40	
41	THE COMMISSIONER: Yes.
42	
43	MR HODGE: and I'll get you, Commissioner, to allocate
44	exhibit numbers.
45	
46	THE COMMISSIONER: Yes. And how do you want to describe
47	that bundle?

1	
2	MR HODGE: So each - if you have a look at the front page.
3	there's a series of topics that are identified and then
4	behind that there's a list for each topic, and the idea is
5	there will be a single exhibit number that you'll allocate
6	now to each topic.
7	
8	THE COMMISSIONER: I see what you've done. What you're
9	doing is you're tendering a list in relation to each matter
10	that we've been examining; is that it?
11	g,
12	MR HODGE: Yes. So if you allocate an exhibit number to
13	each one of them where there's the space on the right-hand
14	side. Then when they go up on the website and for the
15	purposes of the parties making submissions they will, for
16	example, be able to identify - I'm not sure what the next
17	exhibit number is.
18	
19	THE COMMISSIONER: The next number is 188.
20	
21	MR HODGE: So "General" will be exhibit 188, and then
22	within that list the very first item, which is the Forensia
23	DNA Analysis Unit team chart, that will be 188.1, and
24	that's how it will go up on the website and the parties
25	will be able to identify it.
26	
27	THE COMMISSIONER: All right. So what do you want me to
28	do with this?
29	WD 110005 - T 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -
30	MR HODGE: I just want you to - well, you could do it now
31	or we could just do it later, but No.1 will be 188 and it
32	will continue through. So if you just accept that as
33	formally tendered we'll allocate the numbers or
34	THE COMMISSIONED: Sorry I'm not with the schome yet
35 36	THE COMMISSIONER: Sorry, I'm not with the scheme yet.
37	MR HODGE: Sorry, if you look at the front page.
38	TIK HODGE. SOLLY, IT YOU TOOK At the ITOIT page.
39	THE COMMISSIONER: Yes.
40	THE COMMISCIONER. 100.
41	MR HODGE: You see where it says "Exhibit number"?
42	
43	THE COMMISSIONER: Yes.
14	
45	MR HODGE: So "General", if you write "188" there.
46	
<b>47</b>	THE COMMISSIONER: Yes.

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1
2
         MR HODGE:
                     Options Paper will be 189, et cetera.
 3
 4
         THE COMMISSIONER:
                              Yes.
 5
         MR HODGE:
                     And that will be it. That then will be the
6
7
         formal tendering of all of those documents.
8
         THE COMMISSIONER:
                              I see. All right. Well, that's what
9
         I'll do. Take that as a given, from 188, No.1, to --
10
11
         MR HODGE:
                      I'm sorry, just before you do that - yes, if
12
         you just do it as 188 and onwards, and then - so it will be
13
         188, 189, et cetera.
14
15
         THE COMMISSIONER:
16
                              Yes.
17
                     And then the list itself we'll make 188.0.
18
         MR HODGE:
19
         THE COMMISSIONER:
                              I don't understand any of that. We can
20
21
         mark it later and then read it into the record, Mr Hodge.
22
                     Thank you, Commissioner.
23
         MR HODGE:
24
25
         THE COMMISSIONER:
                            All right? So that I don't add to any
         confusion.
26
27
28
         MR HODGE:
                     I'll deal with that later.
29
         THE COMMISSIONER:
30
                              All right.
31
32
         MR HODGE:
                     Thank you.
33
34
         THE COMMISSIONER:
                              Are we ready to proceed?
35
36
         MR HODGE:
                     We are.
                               Thank you.
37
         <REBECCA JUSTINE KOGIOS, sworn:</pre>
38
39
40
         <HEIDI MIRANDA RUTH BAKER, affirmed:</pre>
41
42
         THE COMMISSIONER:
                              Thank you, both. Mr Hodge.
43
44
         MR HODGE:
                     Thank you.
45
         <EXAMINATION BY MR HODGE:</pre>
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1
                    I might just have each of you just identify
 2
         yourselves.
                      Dr Kogios, could you state your full name for
         the Commission?
 3
 4
 5
         DR KOGIOS:
                      Yes, my name is Rebecca Justine Kogios.
6
        MR HODGE:
7
                     And what is your occupation?
8
9
         DR KOGIOS:
                      I'm the executive director of the Victoria
         Police Forensic Services Department. I should say that for
10
         the purposes of this Commission the views that I express
11
         are personal views and should not be perceived as an
12
         official commitment or a view of Victoria Police or of the
13
        Victorian government.
14
15
         THE COMMISSIONER:
                             Yes, I'm acting upon the basis that
16
         each of you has been retained in your personal capacity to
17
         give your personal expert opinions and that nothing that
18
         you say is to be taken by anybody as reflecting an official
19
         view on the part of anyone. Is that what you're saying?
20
21
        Yes?
22
23
        DR KOGIOS:
                      Yes, thank you.
24
25
        MS BAKER:
                     Yes.
26
27
        THE COMMISSIONER:
                             Thank you, Mr Hodge.
28
29
        MR HODGE:
                     And, Ms Baker, could you state your full name
         for the Commission?
30
31
32
        MS BAKER:
                     Yes, it's Heidi Miranda Ruth Baker.
33
34
        MR HODGE:
                     Thank you. And what is your occupation?
35
        MS BAKER:
                     I'm a senior forensic scientist at ESR in New
36
        Zealand.
37
38
                                 Now, I'll just bring up, so that we
39
        MR HODGE:
                     Thank you.
40
         all at least have it on your screen, your expert report,
        which is [EXP.0007.0001.0114]. I'm hoping that both of you
41
         can see that as well?
42
43
44
        MS BAKER:
                     Yes.
45
46
        DR KOGIOS:
                      Yes.
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1 MR HODGE: Great. That's the report that the two of you 2 have prepared for the Commission? 3 4 MS BAKER: It is. 5 DR KOGIOS: 6 It is. 7 8 MR HODGE: And, insofar as it expresses opinions, those 9 opinions are ones that you genuinely hold? 10 MS BAKER: 11 Yes. 12 13 DR KOGIOS: Yes, that's correct. Now, we've already formally tendered it --14 15 THE COMMISSIONER: Just before you go on, because we're on 16 video with each of you and because you're not together 17 shall we just adopt the process that, Dr Kogios, if you 18 wouldn't mind speaking first in every case and, Ms Baker, 19 you speak second if the question is put to both of you, 20 21 subject to Mr Hodge asking or any of the counsel here 22 asking for some different sequence, and then you won't interrupt each other. 23 24 25 DR KOGIOS: Yes. 26 Yes, Mr Hodge. 27 THE COMMISSIONER: 28 29 MR HODGE: Thank you. What I want to start with is just inviting you to express your general observations about the 30 positive qualities of the lab based on your review, and 31 32 I wonder, Dr Kogios, if you might start? 33 More than happy to do that. 34 DR KOGIOS: Yes. It's fair 35 to say that Ms Baker and I did conduct quite a deep dive in the limited time that we had, both through our site visit 36 and through the extensive access to SOPs, to standard 37 38 operating procedures, that we were given, and on the whole overwhelmingly we found a highly functioning performing 39 laboratory with a very skilled cohort of staff across all 40 levels, staff who were very professional, very 41 knowledgeable, very experienced and incredibly committed to 42 their work and to providing an excellent service to the 43 State of Queensland. We found the facilities themselves to 44 be first rate, and on balance the methodology, the 45 protocols in use are very consistent with the methodology 46

and the protocols in use in the broader forensic sector in

this country and beyond. So we had many, many positive observations from our time with the staff of the Queensland Health Forensic and Scientific Services laboratory.

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MR HODGE: Thank you. Ms Baker, did you want to add anything to that?

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MS BAKER: I would, thank you. I certainly would describe the staff as incredibly passionate, very dedicated to the people of Queensland. Most of them have dedicated their whole careers to forensic science. I would like to express my thanks for how helpful the staff were when we did our visit, and I appreciate it was an incredibly challenging time for those staff, and their demeanour and the way they carried themselves during that visit was to be commended. So that would be a huge thankyou from me. I also noticed the really well-written extensive standard operating procedures, so those are like the manuals that people So there was certainly good evidence of a worked towards. high level of documentation there as well.

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MR HODGE: Now, is it fair to say, and I'll direct this first to Dr Kogios, what you also observed was within the work group fragmentation and issues of trust?

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Yes. Yes, that is fair to say, and that won't come as any surprise, given some of the evidence that has been before the Commission. I think it's probably an observation that I would make that as a general rule forensic work groups tend to be fairly fixed. It takes a long time to develop expertise in forensic science, particularly to the point that enables a person to give evidence in court. So what you end up often is a work group that has been there for a very long period of time, and this is a blessing because obviously you develop an enormous amount of skill and expertise in that work cohort. But it can also mean that there are difficulties that can arise potentially through longstanding issues in the workplace, and we certainly saw some evidence of that in our time in the laboratory.

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MR HODGE: And I wonder, Ms Baker, are you able to just describe for the Commissioner, if you're familiar with it, what might be thought of as a no-blame culture within a workplace?

MS BAKER: Yes, of course. So it's really important in forensics. You know, we're mostly human beings. We have some robotics as well, but we make mistakes, we're human, and it is really important to have a culture of no blame so that people feel able to raise any concerns that they may have. If you're in the middle of an item examination and you accidentally drop a tube, it is incredibly important that you feel able to own that and seek some support as to how best to rectify that situation.

We also are really sort of strongly advocating for that near-miss approach as well, so don't wait for something to go wrong. If you see a potential for something to go wrong in a process or the particular way that you're doing something, it's really important to flag that, and that's at all levels, and we really relish new staff coming into our laboratory who review our standard operating procedures as part of their initiation because we want to make sure that what we're saying we do we're actually doing, and we sort of really empower our new staff to sort of hold us to account, and if they hear us talking about something in a way that we're not describing it in our operating procedures we encourage them to speak out about that, and then that continues throughout people's forensic career.

I would just like to add in terms of the work culture the work we do is incredibly tough and challenging at times, mentally, emotionally as well, and it's incredibly important to have a support structure around you. A lot of our work is confidential. We can't talk about it with our friends or family. So to have work colleagues around you that you can trust and who understand the pressures that you're going through and some of the things that you find quite difficult to work with is really important, and I felt in some ways that was obviously lacking in certain areas within the laboratory, and that's a real shame for those staff.

 MR HODGE: And - I might direct this first to Dr Kogios - in terms of that idea of a no-blame culture, did you form a view about the extent to which that is reflected within the culture of the Queensland laboratory?

DR KOGIOS: Well, I mean, we did certainly see signs of a no-blame culture. A lot of people that we spoke to spoke about quality is everybody's business, everybody has a role

in calling out issues around quality. So, you know, there was evidence of a no-blame environment. We also heard from other staff members who had a very different experience. I think, you know, my personal view would be that the only way to really get the measure of a particular culture would be to be a part of that culture. You have to understand that Dr Baker and I, we had a view into the culture, but we've only seen certain perspectives. So whatever opinion we may offer would be limited, you know, when looking at it through that lens.

MR HODGE: I understand. What we might do is then move to the topic of workflow and start at a general level. One aspect of workflow I want to ask you about is in your experience are there ways in which different laboratories will go about trying to cap or limit the work that occurs at different stages of the forensic examination process?

DR KOGIOS: Yes, absolutely, and, look, I think it's important to understand that decisions are made right the way along the forensic continuum, if you like, from the start, from an event that takes place out at a crime scene, or what becomes a crime scene, right the way up to reporting the final results of a case. It's a series of decision making that takes place, and it's not just about efficiencies by any stretch. It is the case that not every scenario, not every case requires what we would consider to be a fully exhaustive approach to testing.

So, you know, to just give you some examples, at a crime scene it may well be that, you know, where do you draw the line? You could potentially collect every single item in the scene. It just wouldn't be necessary. So what you rely on is skilled experienced crime scene examiners to make good decisions based on what they observe and the information they have been given about the scene to zero in, if you like, on the exhibits that are going to be informative in that particular case. So there's an episode of sort of culling, if you like, from the start at the crime scene.

Then there's another round of decision making around which items will actually be examined. So you wouldn't necessarily in a major crime examine every item that was collected from a crime scene. Again, you're relying on experienced, skilled professionals to make judgment calls around setting an examination strategy that's going to give

you the most value in the case.

Then the next step down would be decision making around the samples that have been selected from those items that have been examined, which samples to process in which order, what level of re-work might be appropriate, and then at the end of that first initial round of testing a backward look, if you like, to say, "Well, what have we got so far? Is that sufficient in this case? Have we answered the key questions that we think forensic evidence are going to be able to provide answers to in this case?"

There may be new information as well that's come to the fore in the passage of time between the incident and, you know, this particular point in the sequence. So that might occasion a person to then go back and do some further item examination. Some of those exhibits or items that were collected are not examined in the first pass, you might want to go back and do further testing of those items. So it's a continual process of decision making to decide what gets carried forward at any particular point in time.

 MR HODGE: And there's two different aspects of that that I want to ask you some questions about. One is the way in which that ties into the workflow that's used, and the other is the way in which that ties into thresholds. So I'll start with the workflow. You make the observation in your report that the current process that is used within the Queensland laboratory is that cases are not routinely allocated to a case manager, except in the cases of the P1 cases. I wonder if you might just explain from your perspective how you would describe in its essence the type of workflow process that is used in the Queensland lab?

DR KOGIOS: Is that a question for me or for Ms Baker?

MR HODGE: I'll direct that to you, if that's all right, Dr Kogios

 DR KOGIOS: Sure. So I think we would describe it as a sort of end-to-end workflow that's split across two agencies. So, you know, you've got the Queensland Police, the QPS, responsible for the first part of that workflow. So that is not only the crime scene attendance but also the item - the setting of an examination strategy and the item examination. And then you've got a submission of samples

in an "in tube" model through to QHFSS - I'll use that acronym, I think, FSS, to describe the laboratory. So those samples are then submitted in tubes through to the laboratory for the subsequent analysis, the interpretation and the reporting stages of that workflow, with results then provided back to QPS. So I think we would think about this as being a split operating model between the two agencies with an "in tube" laboratory submission.

MR HODGE: And, Ms Baker, one of the expressions that sometimes's used is "robot ready". Could you just explain what that means?

MS BAKER: Yes. So to assist in processing large batches of samples and to I guess reduce the possibility of human error a lot of forensic laboratories do employ robots, so, for example, to do the extraction process, so actually getting the DNA out of the sample for quantification or measuring the amount of DNA in the sample, and also for amplification, so making lots of copies of them. And I guess being robot ready would mean that in this case the Queensland Police would submit a sample that's already in a - we call it a microcentrifuge tube, so a little tube that is ready to go straight onto the robot to be extracted.

 MR HODGE: And, again, I'll direct this to you, Ms Baker. The model that is used in Queensland, we've heard it referred to in different ways in the evidence, but sometimes there's something like a production line or something along those lines. Does that reflect the way, in your view, the process works within the Queensland laboratory, excluding P1 cases?

 MS BAKER: Yes. In some respects I guess the idea behind that is that samples are reported individually rather than as a case, and it's seen as a way of getting a high throughput and a relatively short turnaround time for a sample.

MR HODGE: Yes, and so I think - and I'll direct this to Dr Kogios - an advantage of that model is it is high throughput in terms of getting through samples?

DR KOGIOS: Yes, yes, absolutely, and I think one must - I guess I would caution against looking at high throughput as being necessarily a negative thing. In contemporary law

enforcement it's absolutely about intelligence-led policing. So police are really in some instances relying on a forensic link to give them a lead in a case that perhaps they wouldn't otherwise have, and having an early lead on a case can lead to significant positive outcomes in terms of the progression of that particular case. It might also be very useful in disrupting an active offender and therefore preventing further crimes from happening. So one of the benefits of this particular model is that it does have the potential to offer that high throughput, which then enables a rapid link and that information to be provided to police.

MR HODGE: Is it fair to say, though, in terms of where that kind of benefit accrues it's really for what's referred to as volume crime, or P3 cases in the Queensland parlance, rather than being - whilst it might occasionally be of significance, it's probably not normally of great significance in relation to murders and other kinds of serious offences?

DR KOGIOS: Broadly I would agree with you. I mean, as you say, there could be some instances of those serious crimes against the person where police just don't have a lead and they really are relying upon some forensic link to give them some sort of direction in a case. Forensic links are also vitally important in exculpating individuals from investigations as well, of course. So I think, yes, that I would agree that the high throughput and the ability to get a quick link is of particular benefit when you're dealing with high volume crime. It can be of benefit for some serious crimes against the person. But for those crimes it's often more the build of the full forensic picture that a court will subsequently rely on in that sort of evidential or that prosecutorial phase.

MR HODGE: The two of you identify a number of risks with this kind of process, so I just want to draw those out from you. One of those risks is the risk of over-servicing, and I wonder - and I'll direct this I think to you, Dr Kogios - could you just explain what that means in this context?

DR KOGIOS: Yes, Mr Hodge, if I can just take a moment just to explain. There is no such thing as a universally agreed international best practice for an operating model, and the operating model that is in place in Queensland is certainly within the range of accepted operating models in

Australia. So it is not that our evidence is that the operating model per se is the problem; rather, that under that particular operating model as opposed to other models, like a case manager led model, there is the potential for some risk through that lack of oversight, and that's where we say that, you know, safeguards are required to guard against some of those risks, and I think what you're asking me about now is one of those risks that we have identified.

MR HODGE: Yes. I think - let me say it back to you, and I'll direct it to both of you to make sure that I understand, and I was going to come to this point in due course, but your point is it's not that you don't want to be taken to be expressing the view that to have this kind of production line model is fundamentally wrong, and indeed it's a model that, as I understand it, at least aspects of it are reflected in New South Wales, but the point that you're making is it is a model like any model that comes with certain risks and if you're going to use that type of model then you need to make sure that you have risk mitigation strategies in place to address those risks?

THE COMMISSIONER: And that you don't use it in inappropriate circumstances.

DR KOGIOS: Yes. I'm terribly sorry, but the first part of that audio dropped out for me completely. So I caught the last part of what you said, and, yes, our evidence is that you need to have risk mitigation strategies in place really for any model, for any operating model. It's a question of identifying the weak points and putting some mitigation strategies in. For this particular model there are some specific risks that arise and we would say require a certain level of risk treatment.

MR HODGE: And we'll come to the detail of it, but is it fair to say that as an overall view your view is that the workflow model that's been used in Queensland doesn't have sufficient risk mitigation strategies around it to address the risks that come with that kind of production line model?

DR KOGIOS: It's fair to say we did see some evidence to support that statement through our case file review in particular.

MR HODGE: And, Ms Baker, what's your view about that?

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I would say I would agree with you and say the model itself isn't problematic and it does have advantages which we have talked about. What I would say is that for it to be successful it requires an incredibly strong collaborative relationship between those two providers; excellent communication, and that would be at all levels, between those two agencies; continual feedback as well is incredibly important to make sure there's no divergent evolution of practice; and I think, really importantly, any changes that either party make - and it could be as simple as changing which test you use to see whether something is blood or not, for example - they have to make the other party aware of that and consider the upstream and downstream impacts of any of those changes. just a number of really key areas where things could potentially fall through the cracks, and it's really important that this sort of working model has those ironed out and continually reviewed as well to make sure that there's not a slow creep of any issues.

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MR HODGE: And so then, Dr Kogios, to come back to some of the risks that you've identified with the current model, one of them is over-servicing. Could you just explain what you mean by that?

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41 42 DR KOGIOS: Yes. So through our case file review we observed a number of instances where the test - "we test everything we receive" approach potentially has led to a degree of unnecessary testing, and, look, I think it's fair to say here that really it's up to a court to decide what is important in any given case. So we are making some assumptions here, and the assumption that we're making is that if, for example, you had a sexual assault case and there were, you know, multiple swabs collected from the same part of the complainant's body and that all of those swabs were tested independently through the laboratory and all of them gave a sort of a very similar DNA result, if you like, let's say a two-person mixture with both the complainant and the accused not excluded from that two-person mixture, then we would say, you know, "Did you really need to test all nine of those swabs," for example, "You might have got that answer with just testing one of those swabs."

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Of course, it depends on the scenario, the particular scenario in question. Particularly if you had more than

one alleged offender, well, then of course you would need a more exhaustive testing regime. It would depend on what the accused person had said in terms of their version of what had taken place. But I think it's fair to say that we did see some evidence of what we felt to be multiple samples being put through the system at the same time, so a part of that first pass testing, all of which were giving very similar results, and that led us to believe that there was a degree of over-servicing in that process.

MR HODGE: And I'll come back in one moment to sexual assault investigation kits and ask Ms Baker a question about that. Could I just ask you to just address one more part of what you were talking about, Dr Kogios, which is outside the context of sexual assaults, and so looking at other kinds of offences of violence, does the same kind of issue arise; that is, does the same issue of potential over-servicing arise in relation to a murder or an assault?

 DR KOGIOS: Yes, potentially. So we didn't have the opportunity to take a deep dive into QPS protocols. was not part of our terms of reference. But what we do understand is that the submission for major crime like homicide, it's capped at 25 samples. So, you know, if you had a scenario where 25 samples were submitted - I'm not saying that in all cases 25 samples would be submitted, but the FSS, the scientists in the laboratory are really relying on the QPS to have made those decisions about which samples to submit. The sexual assault case work is slightly different in that the case - the scientists in the laboratory at FSS themselves are really the starter of that workflow process, whereas in homicide cases those decisions are likely being made by the QPS.

 But, you know, certainly from my experience and from Ms Baker's experience you wouldn't necessarily start off in a homicide case processing 25 samples in one go. You might start off, depending on the case scenario, processing a smaller number of samples and seeing what results you get back and then considering a second round and a third round, potentially even a fourth round of testing. I guess it's fair to say that in our experience we are used to a more sort of staged triage type approach.

MR HODGE: I understand, and just to explore that a little bit further, as you understand it within the way the Queensland laboratory works, if the QPS send over 25

samples, all of those samples will be tested by the laboratory regardless of whether, if after testing just two of those samples, they really have whatever the information is that could be useful to the case?

DR KOGIOS: That's our understanding of how the model has been established, yes.

 MR HODGE: And I'm going to come back to you in a moment, Dr Kogios, to get you to put that in the context of an examinations strategy. But can I just switch over to Ms Baker to ask you to talk about the context of testing all the samples in a SAIKs. Could you just explain are there risks involved with just automatically testing all of the samples within a SAIKs kit rather than looking at what results you get from some swabs, say, and then thinking about what other kinds of testing you might need to do?

So I would say that if you only have MS BAKER: Yes. standard DNA testing in your forensic toolbox then the risk is that you have used material from all of those samples, and we know that in sexual assaults we usually have an overwhelming amount of female DNA in those sort of intimate swabs, and if you've been unable to obtain a male DNA profile from that, so there might be cases where there's such a ratio that the female DNA swamps out any male DNA, you've actually already used quite a lot of material in that case, whereas potentially if at the outset you've identified there was no semen, for example, detected in that case, consideration could be given to submitting it for a more appropriate test, so perhaps a male-specific DNA test which has the ability to ignore all the female DNA that is in those samples and just target low levels of male DNA that you may not be able to sort of see or detect in a standard DNA test.

 MR HODGE: And is another issue with just taking the approach of testing every sample that is provided in a SAIKs kit without necessarily thinking in more detail about what exactly has occurred that you might have a different strategy for testing depending, for example, upon the nature of the allegation, whether, for example, it's digital penetration or penile penetration?

MS BAKER: Absolutely. So I would expect any sexual assault kits to have an examination strategy at the beginning to look at the information that's provided in the

case context and decide which samples to test and in what order. So not necessarily to test all the samples and, on top of that, not to necessarily test all the samples at once - all the ones you select for testing. Do a phased approach and see what information you get.

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MR HODGE: And what I might do is I want to ask each of you about examination strategy. I'll start with Dr Kogios. You refer in your report on a number of occasions to an examination strategy. Can you just start by explaining to the Commissioner what you mean by that?

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DR KOGIOS: So it's a term that I picked up when Sure. I first started working in the UK. It's certainly something that they refer to, an examination strategy. Other laboratories would call it triage or vetting. Ιt really is essentially looking at the case and deciding what is the best type of testing to perform on this particular So, to give you an example, it could be a homicide matter and it might be that there's, you know, 20 items that's probably unrealistic. It might be that there's, say, 50 items that have been collected from a crime scene. So you have the potential to examine 50 items. Your job in setting up the examination strategy is to say, "What's going to be most likely to give us the most valuable information in this case," and you need to have a bit of an understanding about the broader case context to be able to set the right strategy.

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So it might be that you decided, hypothetically, to say, "Right, well, we've got two items that we will start It's item 4. It's a knife (indistinct) and we need to look for trace DNA on the handle of the knife, and then examine the blade for blood. If blood is present, let's take a sample of that blood." So that would be an example of - an examination strategy that of all the clothing that has been submitted in this case - there's a particular allegation that seems to be under some sort of challenge. so it might be that, you know, of all of the pieces of clothing it's the T-shirt that's going to be most useful for us and we're interested in a particular type of blood pattern to see whether, you know, does the T-shirt show a particular type of blood staining that could be the vital piece of evidence in that case to help us refute/support the various hypotheses that have been put forward.

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So, instead of sort of a blanket testing regime, you

might end up with a very sort of tailored testing regime that says, "We're going to start with just two items, and we're going to treat those two items in this particular way and we're looking for specific things in those two items." That might be your first pass testing, and that might be all you need to do. So that would be an example of how you would go about setting an examination strategy for a complex case where you have potentially many, many exhibits.

MR HODGE: There's many elements of that that we'll deal with, but, just to focus on one part of it, what you are talking about is at the very beginning, before you undertake any DNA testing, first somebody thinking about what, given the samples we have, are the best things to start with testing, thinking about what is the most information that they will produce in the context of whatever the crime is that we're seeking to solve?

DR KOGIOS: That's right. That's right, and it's a very important step because it dictates what happens next. So if something is amiss in the examination strategy, well, you know, the horse has bolted, if you like, by the time it gets to a person to interpret those results.

The other benefit of a really well thought out examination strategy is at the time of reporting the case if the reporting scientist - they don't necessarily have to have been the same person who devised that strategy, but ideally they have access to that examination strategy, and then they have the opportunity to look at it and say, "Is anything amiss? Does anything look not right in the results that we've got," because that could be a sign of a contamination event, it could be the sign of something not working optimally in the process. So it gives them that opportunity. But it also gives them the opportunity to say - and here we are talking about those most complex crimes, those most serious crimes against a person where one would expect the community would be expecting, you know, law enforcement and forensic services to leave no stone unturned. So we're not talking about a theft of a bicycle here; we are talking about a serious crime against a person.

The reporting scientist having access to that examination strategy then gives them that ability to say, "Has everything been done that could be done in this case?

Have we left any stone unturned, and, if not, then we can go back and we can do some further testing." So that's another benefit of having a really well thought out examination strategy or a vetting strategy or a triage strategy.

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MR HODGE: Thank you, and I'll just turn to ask Ms Baker a question and then continue on in a different way, Dr Kogios. Ms Baker, are you able to, given some of the explanation you gave earlier in relation to SAIKs, just help us to think about an examination strategy for a SAIKs kit and what that would involve?

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So, for example, if there's been a sexual MS BAKER: Yes. assault and there's alleged to have been one male offender and there are sperm or semen detected on the microscope slides that are made as part of that process, I, for example, would not in that case feel the need to examine every swab that was submitted, and there's likely to be, you know, anywhere from three to perhaps nine, I think I've seen in particular case files at FSS. I would perhaps start with one of those samples because I would be confident in the process that I had that I should be able to obtain a male DNA profile from the semen that I could see on the microscope slide, so in that case you're looking at only processing one swab as opposed to multiple swabs, and perhaps if it turns out that there's only a small amount of semen or in fact it could be quite heavily degraded then you've got lots of material left to consider whether another type of DNA testing might be more appropriate for the next sample.

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39 40 MR HODGE: And so, just at a very basic level, if you had nine swabs from a SAIKs kit you might expect - and what you're dealing with was an allegation of penile penetration and you could identify semen on a slide in relation to a high vaginal swab, you would look to your high vaginal swab as the starting point because if you could identify semen that you could match to a reference sample from that swab it's not going to be necessary to test and use up all of the other swabs?

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MS BAKER: Exactly. Exactly. I think, you know, typically in that situation, to find semen that corresponds to an individual on someone's high vaginal swab would be seen as having a high probative value, and, again, it's a conversation that you would have with the police. They may

well be having conversations with the justice department to determine, "Is that sufficient in this case? Does this (indistinct) people's scenarios that they have suggested, and, if not, let's go back and test something else," or, "If so, that's great, let's move on to the next." Can I --

MR HODGE: And so --

MS BAKER: Sorry. One example of over-servicing, again, we talk about the swabs from a SAIK, but, for example, pieces of fabric that were submitted to the laboratory which were then sort of cut into, for example, two sections or four sections, and then all of those separate pieces were tested, and that to me is another example of over-servicing, where ideally you'd have a piece of fabric that's submitted that is small enough to be robot ready, as you've suggested, or an agreed procedure where you would just take one sample from that piece of fabric and run it through the system to see what you've got, as opposed to automatically processing, for one of the cases that I saw, all four subsamples, if you like, from a piece of fabric.

MR HODGE: I understand. So in both of those examples - that is, both the example of processing all nine swabs from a SAIKs kit and processing all four samples from a piece of cloth which had bloodstaining on it - it might well be that if you just processed one sample you would get sufficient information to not need to undertake testing of all of those other samples. So you've effectively - if you're talking about the case of nine swabs, you've done nine times the amount of work that was actually necessary in relation to that one case?

MS BAKER: Yes.

MR HODGE: And that is the issue of over-servicing that you're describing when you refer to one of the risks of the production line method that's used at the moment in Queensland, and, as I understand it, the point about having a good examination strategy is that's a way of trying to deal with that issue of over-servicing, amongst other things?

MS BAKER: Exactly.

MR HODGE: And then is having a good examination strategy also a way of dealing with another risk that you mention,

which is the missed opportunity to harvest all available forensic evidence?

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MS BAKER: It is, yes. I guess ideally and certainly from my own experience having that whole case overview means that you know which exhibits are available to you, you know what areas have been highlighted as potential for sampling or processing through DNA testing, and so you're able to make those sort of strategic decisions as to what is going to be in the first pass testing and what items or samples you have available should you need to go back and use in re-testing. Without having that sort of whole case overview it in my mind can be very difficult. You're expecting two different groups to have that covered, and they don't necessarily (indistinct) same amount of information from start to finish from that workflow.

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MR HODGE: And, Dr Kogios, does that issue or that issue of missing the opportunity to harvest all available forensic evidence, does that also tie into the issue of the strict application of thresholds?

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Yes, because if you're using a threshold to DR KOGIOS: make a decision not to proceed with testing then you're missing the opportunity to recover whatever DNA profile information might have been present in that sample. here I think again it's important to note that the use of thresholds is absolutely common practice in the forensic science sector. As part of our work in the Commission we had the opportunity to look at other Australasian forensic science providers and what they do with thresholds, and it's absolutely the case that the majority of jurisdictions in Australasia do apply a form of threshold. It's common practice. That threshold tends to be set at the limit of detection. So some jurisdictions will use a threshold at limit of detection only and they will only apply it to But I suppose what we're doing certain types of cases. here is pointing out that the use of thresholds per se is not unusual, and it is tied to the fact that we do know that you do get stochastic effect with low levels of DNA and in some case types and some instances it just may not be worth proceeding on with a particular type of testing. It's a decision, it's a policy decision, if you like, for each laboratory to set for themselves rather than it being a matter of science.

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MR HODGE: I understand, and I might just jump forward to

thresholds just to help everyone to understand your views about that. It's fair to say, isn't it, your view is that the way in which the Queensland lab is currently approaching thresholds is not best practice; you say that at paragraph 47 of your report?

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So 47 of my report, we're talking here about DR KOGIOS: their recent use of thresholds, and specifically we're talking about the so-called DIFP threshold. So we're not talking about, as we understand it, the current state, which is use of a limit of detection threshold, albeit that there are some questions over the current threshold and whether that has been appropriately obtained through appropriate validation. But, to make it really clear, what we're saying here is it would be considered broadly accepted practice, widespread accepted practice, to have a threshold beyond which you don't proceed set to the limit of detection. But the idea of having sort of that range, that DIFP range, so to speak, that's not something that we've seen in other jurisdictions as part of our review for this Commission.

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MR HODGE: I understand. Can I try to break that down into pieces so that we can all understand it.

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DR KOGIOS: Sure.

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33 34 MR HODGE: There's two thresholds that are apparent on the material to you that Queensland has been applying. One is a limit of detection, which they have set at  $0.001 ng/\mu L$ , and the other, until the changes made a few months ago, was what's referred to as the DIFP threshold, that if it's below  $0.0088 ng/\mu L$  they wouldn't as a matter of standard process continue to process the sample. So those are the two thresholds you were talking about?

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DR KOGIOS: That's correct.

(Day 23)

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MR HODGE: And I think your point is you and Ms Baker have the view that the use of the DIFP threshold is something that falls below best practice?

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DR KOGIOS: It's a very difficult question for a scientist to answer the binary is it best practice, is it not best practice. Ms Baker and I spent quite a bit of time thinking about this because we understand that that is exactly what the Commission is asking us to provide an

opinion on. So we came up with a bit of a framework that we have applied to our opinions around what is and what is not best practice, and our framework involved looking to see, you know, is there an ISO standard, let's say, that specifies that something must be done a certain way and is there some practice that we've seen that is inconsistent with that standard. If so, then we would say that that cannot be considered best practice.

There is no ISO standard that relates to the topic of threshold. So that is more of a policy decision for each laboratory to make. So what we did then was we looked at the responses from the other Australasian jurisdictions, and we looked for something to see whether it was, you know, significantly out of step with what the rest of the Australasian community was doing, and, if so, then we would say that falls below that definition of best practice. It's absolutely a subjective call that - you know, that we made with a degree of trepidation and with all of those caveats attached.

MR HODGE: I understand. If we just bring up paragraph 47 of your report. I know you've got it there, but I think it will just be helpful for everybody else. Sorry, it's not page 47. Paragraph 47. That's on page 23. You see in the first sentence where you say:

QHFSS recent approach to thresholds falls below best practice.

 What you're explaining to the Commissioner is what you mean by best practice and the point that you're making is one of the challenges for you in preparing your report was understanding or identifying what is the content of best practice given that we're lawyers and we try to do things in a very binary way and we've just said to you, "Is it best practice or not," and you had to then think about what does that even mean, what does it mean to say it is or isn't best practice, and your point is you started by asking is there a particular international standard that sets the practice in this particular way, and if it didn't accord with that standard then you would say, "Well, it falls below best practice."

But if there wasn't an international standard but you were able to, between the two of you, given your expertise, recognise a consistent practice around laboratories around

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Australia, and the practice within the Queensland lab fell below that consistent practice that you could identify, then that was the other circumstance when you would say, "Well, it falls below best practice"; is that a fair summary of your evidence?

DR KOGIOS: That is a fair summary of our evidence. We did find this particularly challenging. I think the other element to call out is forensic science is, like all science, constantly evolving, and there is a body of what we would call emergent best practice, and all forensic science providers need to keep an eye on that emerging body of knowledge, and so it's a process of continually reviewing and making amendments to practice to ensure that you maintain a level of contemporaneous practice.

What we have tried to do is draw a line between what we would consider to be emergent best practice and a journey that all forensic science providers are on, as opposed to what would be considered accepted best practice and has been the case for some period of time, and it's those instances where we felt that the Queensland lab was out of step that we called that below best practice.

You know, obviously you're showing on the screen one area of our report. We do make many references throughout our report to there being no such thing necessarily as, you know, a universally accepted best practice model. But with the right caveats around it we have made these observations on a number of instances - actually, really quite a small number of instances when you consider the vast amount of material that we've looked at from this laboratory and the deep dive that we've done. But we can't shy away from the fact that there were a number of instances where we did feel that we had to use that phrase "below best practice", albeit that they were not (indistinct) in total.

THE COMMISSIONER: I wonder if I could ask you to approach the issue in a slightly different way. I do understand that when I come to consider what to write in the report I'll have to take account of the scientific principles that have been explained to me by you and other experts, I'll have to take into account instances of what's called best practice, if there is such a thing in particular instances, and I'll have to take into account international or Australian standards where they exist.

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But there is another way of looking at it, which is that one has a look at the purpose of what a lab is doing and the ultimate purpose of what a lab is doing, and one can ask whether a particular process suits that end or not. So I understood that the purpose of the Options Paper that led to the DIFP category and the decision not to test DIFP automatically was justified by seeking to, as it was believed, improve the efficiency of the lab overall in terms of turnaround times and matters of that kind, and I understand your evidence to be that turnaround times are a factor - I'm correct, aren't I, that they're a factor?

DR KOGIOS: Yes.

THE COMMISSIONER: One scientific article that came to my attention as having been looked at at the time that these issues were being considered in the lab was from a number of Israeli scientists, and, Mr Woolridge, could you put that document up? Just for those in the hearing room, it's [FSS.0001.0011.2109], and if you go to the next page, please, it will be a scientific article, I think, that was attached to that document.

MR HODGE: I'll give you that doc ID.

THE COMMISSIONER: Yes.

MR HODGE:

THE COMMISSIONER: Thank you, Mr Hodge. Just while that's being obtained, this is an article that was explaining how scientists in Israel in a DNA lab were looking to make their lab more efficient by understanding where they can save time and effort, and the approach that they took was to apply a factor to every piece of work that was done and in that way to quantify the work that's done on each phase of the relevant process and to arrive at a result. But what I was interested in was that - yes, if you could go to the second page of that article, and go to the next page, yes, you'll see - can you see that on the screen now?

It's [FSS.0001.0011.2110\_0001].

DR KOGIOS: Yes. It's very small for me, but I can see it.

THE COMMISSIONER: We'll expand it a little. If you go to the preceding page, please, and go back to the next page, and would you go to the page after this, yes, you'll see

that in that first paragraph - if you could expand that first paragraph; thanks - they were looking at samples and what results they got, and they found that samples they regarded as insufficient DNA for amplification that had less than 250 picograms. So their whole study was to do with the amount of effort that was put in and the lack of results from low-quant samples.

But, Mr Woolridge, if you go back to the previous page, and if you highlight the top right-hand corner paragraph - that's it - you'll see that the authors there say that the work they're doing has got nothing to do with serious offences. What they're doing is they're analysing efficiency from those points of view to do with what we call in Queensland volume crime. But with the processing of homicide and sexual assaults cases they just test to the limits, subject to selection of samples and matters of that kind.

So it occurs to me that is this the perspective that you might advocate, that a threshold, whether it's the limit of detection or some other threshold, is a relevant factor to take into account as a guideline or a trigger for consideration as to what further steps, if any, should be taken and what those steps should be, but that there is, firstly, no place anywhere for an arbitrary guideline that would dictate in every case what is to be done, and, secondly, when dealing with the most serious offences one ought - that it is in general the practice of labs with which you are familiar to test samples that are to be tested within that particular case to the limit; what would you say to that? That is to say, thresholds should never be arbitrary, they should only be used to trigger a particular approach or to trigger considerations, and, secondly, samples in cases of serious crime should be tested to the hilt - I'm not saying every sample, but those samples that you choose to test along the lines that have been explained this afternoon should be tested to the hilt - and should not be ever excluded because of the lack of faith of getting a result; so what will you say to that as a proposition?

DR KOGIOS: Look, it's very difficult to be definitive because we can't - we don't know every laboratory, we don't - you know, there might be some laboratories that just aren't funded to do everything in every case, and so it makes perfect sense for them to prioritise the work

that's most likely to give them a result. My own personal view is that if funding is not an issue then having a threshold that can be overruled by a scientist on the basis of what they see in that particular sample is the way to go, that an empowered scientist furnished with the right level of information should be able to make a decision around the progression of a particular sample regardless of the crime type, and personally my view would be for serious crimes against the person that an approach where you leave no stone unturned would be preferable.

THE COMMISSIONER: Yes, thank you. Ms Baker, what would you say?

MS BAKER: Well, I think I would agree. It is really important that the scientist is empowered to make those decisions. They'll know what type of sample it is, what the substrate is, the sort of ease or not of recovering DNA from that particular type of sample. What I would say is it also comes down to what forensic tools you have available to you, and again we talked about if you only had standard DNA testing you may well consider that very low levels of DNA or a very degraded DNA sample may not benefit from that standard DNA testing, and consider outsourcing to another provider for more specific testing.

I guess in terms of thresholds, if you're looking at a sample that you want to be able to load up to your database and you know that you have a minimum number of pieces of information of DNA that you need to obtain in order to load to that database, that could be a consideration. But for a lot of cases they're not necessarily going to a database. You may well have reference DNA samples from individuals in that case for comparison. And I think it's something that shows again to have that really collaborative relationship with the police in understanding what stage their investigation is at, what particular areas they're looking to forensics to provide evidence or information to support their investigation, and really having those conversations throughout an investigative process is crucial for understanding what type of testing you do when.

THE COMMISSIONER: So what I've drawn from both of you then is this: that it's no part of the scientific approach of scientists in your field to adopt an arbitrary threshold to determine the course of sampling. That is a policy issue, not a scientific issue, and it will driven by

1 resource questions. That's the first thing, and the second thing is that the question - subject to that policy issue, 2 the question how samples ought to be treated ought be at 3 the discretion of the scientist. That is to say you ought 4 to trust your scientists to make the right decisions; you 5 can't do better than that. 6 7 8 DR KOGIOS: Yes. 9 THE COMMISSIONER: 10 You can't do worse than that, but you can't do better than that 11 12 13 That's right, and equip that scientist with DR KOGIOS: the right information that they need to be able to know, 14 15 you know, to make the best decision in the individual case. 16 THE COMMISSIONER: 17 And it seems to me then that a production line system of the kind that we've encountered 18 here for what were believed to be reasons of efficiency and 19 productivity and so on is prone to deny that level of 20 21 discretion to scientists? 22 23 DR KOGIOS: I think that it doesn't - the two are not mutually exclusive, but with the right level of 24 communication and engagement and collaboration between the 25 two agencies it could happen. I guess from our experience, 26 Ms Baker and I, our lived experience in the agencies that 27 28 we've worked in, this work is done inhouse, not necessarily 29 by the same person, but within the same agency. So, you know, these groups of people are sharing a tearoom, they're 30 31 sharing a car park, they're known to each other. They're 32 communicating frequently throughout the day, so it's very easy to share information in that kind of environment. 33 34 Splitting across a workflow across two agencies doesn't preclude that. It just - as Ms Baker has said a number of 35 times now, it just really does rely on that really 36 connected collaborative engagement between the 37 38 practitioners at the different - or stages of the work. THE COMMISSIONER: Yes, that's very helpful. 40 Did you want to add anything, Ms Baker? 41

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No, I think between the two of you you have it MS BAKER: covered. Thank you, sir.

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THE COMMISSIONER: Thank you. Mr Hodge?

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MR HODGE: Thank you. Ms Baker, could I just clarify one thing with you, and I think this is my fault. But when we've been referring to the production line system that's used in Queensland --

THE COMMISSIONER: For what it's worth, I'll mark that article exhibit 188.

## EXHIBIT #188 SCIENTIFIC ARTICLE BARCODED [FSS.0001.0011.2110\_0001]

THE COMMISSIONER: Sorry, Mr Hodge.

MR HODGE: Thank you. When we've been referring to the production line system in Queensland is it fair to say that there are two significant elements to it? One is what I think what most of - what we've been talking about seems to be about, which is that the collection of samples and the decision about what samples would be submitted for analysis is made by the QPS separate from decisions that are made in the DNA laboratory, but the second element of it is that within the laboratory the system that they use other than in relation to P1 samples is the work list system?

Yes, that's correct. So the police will decide MS BAKER: which - they'll do the evidence recovery, I guess, so looking, for example, for a piece of clothing and deciding which samples to submit. It will then come into the lab and go through the analytical process, and only when it comes out of the other side of that DNA processing will it be seen by a reporting scientist, who will then interpret and then have that result verified on a sample by sample basis, and each time one of those samples is verified, as I understand it, the result is made available in the forensic-register to the police. My understanding is only perhaps in about 10 per cent of cases are statements requested, and at that point a case manager will be assigned who will have oversight of all the samples in a particular case and draw those results together and put them into a statement format report.

 MR HODGE: Thank you. I'll come back to that aspect in a moment. Mr Operator, could you just bring back up the report and again go to page 23. I just want to cover off on a couple of other aspects of this. I'll direct this question first to you, Ms Baker. It's paragraph 49, if we

could just blow that up.

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This reflects the recommendations that you and Dr Kogios come to in relation to thresholds, and I just want to clarify that there's - I think this reflects what you've said to the Commissioner. Although it looks like it's (a), (b) and (c), there's actually two possibilities here that you're identifying. (a) is that there be no quantification threshold for serious or complex crimes, so that would mean not even a limit of detection threshold; whereas the alternative is (b) and (c) together, that is applying a lower limit threshold in the form of a limit of detection that's been validated, but then enabling scientists to overrule that limit; is that right?

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Yes, that's correct. So the scientists are MS BAKER: highly trained, they're very experienced and they will know what type of samples they have and also what extra testing is available to them. In terms of the FSS, unfortunately they're limited by only having that standard DNA testing. But they have certain clean-up or concentration steps that are available to them. So I would expect a scientist to look at a result and decide what their expectations are about getting a probative result from that if they do some more work or some different types of testing.

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33 34 MR HODGE: Thank you. Then to come back again to the question I was asking Dr Kogios earlier about two thresholds, which can get a bit confusing, I think, for all of us, just focusing on the higher threshold, the DIFP threshold that had been in use, is it right, Ms Baker, that your view and the view of you and Dr Kogios is that threshold, the DIFP threshold, is out of step with the practice of other laboratories in Australia and New Zealand?

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I think it is certainly with respect to those MS BAKER: serious or complex crimes. I can see a situation where for a volume crime case, sort of a property related offence, that that might be the only evidence in the case. the police's perspective they're interested in obtaining a profile that can be loaded up to the database. If you have sufficient validation to show the types of DNA results you get and based on their quantification or when you measure the amount of DNA in them, I can see a scenario where that would be considered okay. But it would worry me to be applying that for all case types.

MR HODGE: Thank you. Then if we blow up paragraph 48 of your report, in 48 you and Dr Kogios recommend that the Queensland laboratory should cease application of it says "the current threshold" - so, just to be clear about what that means, that mean this current limit of detection threshold of .001ng/ $\mu$ L?

MS BAKER: Yes.

MR HODGE: And the recommendation from the two of you is that they should cease applying that threshold until they have properly validated what the actual limit of detection is of the kit within the laboratory; is that right?

MS BAKER: Yes, exactly. If you're making that sort of binary decision of yes to processing or no to processing based on a limit of detection that hasn't been properly validated, that is inappropriate and we would recommend that that ceases.

DR KOGIOS: I was just going to add that here we are drawing upon the report of Commission expert Dr Duncan Taylor and his findings specifically around that particular validation, that piece of validation work. So we effectively have incorporated his finding into our report through this particular aspect.

MR HODGE: Yes, he's identified the point that the Queensland laboratory hasn't validated that limit of detection of .001.

DR KOGIOS: That is right. And it's fair to say that we don't expect there to be a significant difference between their current lower limit threshold and an appropriately determined limit of detection threshold because their threshold looks about right. I haven't got in front of me the thresholds that are in place and used in other Australasian jurisdictions, but it's certainly in the ballpark, and Dr Taylor himself has given that evidence as I understand it as well. So this is more a sort of a cease using the current threshold as a decision-making factor until such time as that work has been done, and once that work has been done then compare the two and essentially check that that is indeed the case, that the two are very similar.

MR HODGE: Thank you. Then I think just to tie off on one last thing, could we just blow back up paragraph 47 again. I think there might have been a slight confusion in some of the evidence earlier, but am I right in thinking the approach to both thresholds that Queensland has adopted recently is problematic but the limit of detection threshold is not problematic in a permanent way; it's just problematic in the sense that you shouldn't be applying a limit of detection without having validated what the limit of detection is; is that fair, Dr Kogios?

DR KOGIOS: Yes.

MR HODGE: Thank you. To then return back to where we were, which is in relation to the issues that arise where you're using the production line method, I think where we had started was I was referring to paragraph 37, subparagraph (b), subparagraph (iii), which is on page 18 of the report, and that's where you're identifying one of the risks of the production line method is this strict application of process and thresholds rather than the exercise of discretion of the kind that you've described in relation to thresholds. I'm just wondering given what we've already talked about is there anything you wanted to add about what that risk is, Dr Kogios?

DR KOGIOS: No. I mean, I think the application of thresholds, it's not in and of itself an essential part of that so-called production line model. It was more that in this section of our report we were detailing our observations around potential missed opportunity. And so that's why we've referenced the strict application of thresholds in this section of our report. You could certainly have that current operating model and not apply thresholds to that operating model. So the two things are related but they're not one and the same.

MR HODGE: Thank you. And then another risk that the two of you have identified of this kind or the current kind of model is the missed opportunity to detect contamination or other unexpected results, and I might direct this one to Ms Baker. Could you just explain what that risk is?

MS BAKER: I suppose if you're not proceeding with a result - I mean, there's two aspects of that. One is to do with processing the negative control, and I'm not sure if that's what you are referring to here, but the other one is

actually progressing some of those samples through and just checking whether or not you do actually have low levels of DNA detected and whether those are popping up in your controls.

MR HODGE: I'll bring it up just so the two of you can see it. If we bring up page 19 and subparagraph (c) of paragraph 37. I had understood this as referring to the issue of where a result arises which is strange or doesn't quite make sense in the context of the case and the missed opportunity to be able to identify that. So I don't know whether maybe you want to expand upon that, Ms Baker, and then I might just ask you there's a reference to the Jama case. I don't know, Dr Kogios, whether you're able to speak about that or whether I should ask Ms Baker to speak about that.

DR KOGIOS: I'm happy to speak about that, Ms Baker, if that's --

MS BAKER: Yes.

So, look, we included a reference to this DR KOGIOS: particular case - it is a Victorian case - really as a bit of a cautionary tale. So, without going into detail, it certainly is the case that that particular case has ultimately been found to have had a miscarriage of justice It was an allegation of sexual assault where a contamination had happened upstream of the laboratory but was not detected at the time. So really we highlight that case as being, you know, a case to provide support for the notion that the reporting biologist, the more equipped with information that they can be about the circumstances of the case and that ability to look more broadly at the result rather than just, you know, a DNA result on a page, but to understand the context of the case, it might be helpful then in, I suppose, raising an alarm that something doesn't look quite right.

Here we draw on the UK's Forensic Science Regulator, and there's a reference that we have provided to one of the Forensic Science Regulator's report from the UK that really encourages reporting officers to take that time to consider anything in terms of a result that doesn't fit with the case circumstances. You know, we've said this a couple of times in our evidence. When you put humans into systems mistakes will happen and we can't shy away from that.

Contamination happens in forensic laboratories in every forensic laboratory. The best defence that we in the business of forensic science provision have to that is, you know, accepting it and being on the hunt for it, and then having appropriate risk mitigation strategies.

This is a vital risk mitigation strategy, to have a reporting scientist who can look across that case holistically and see whether there is anything that just doesn't look right on the circumstances. It could be what stands between, you know, a result going out the door or that result being caught, and the opportunity for a deeper dive to look into what might have happened to explain this result. This could have happened as a result of a contamination event.

MR HODGE: Does the Queensland system at present outside of P1 cases have that kind of overview?

DR KOGIOS: Well, I think it's fair to say that the scientists at FSS are very limited in their (indistinct) ability to perform this kind of check (indistinct) because they just don't have (indistinct) perhaps QPS have that ability to (indistinct) in any great detail, but I guess the point --

THE COMMISSIONER: Dr Kogios, we lost the last 30 seconds.

DR KOGIOS: My apologies.

THE COMMISSIONER: I'm sorry, could you start again.

DR KOGIOS: Yes. So the point that I'm making here is that the scientists at QHFSS are limited in their ability to perform this type of check because they don't have that whole of case visibility. It may well be that the staff, the scientists at QPS are performing this whole of case check. We didn't do a deep dive into QPS's function really in any great sense. That was outside of our terms of reference.

But the point we make is where you have a workflow that straddles the two agencies and you have people in both of those agencies whose got some sort of knowledge you really do need to have those safeguard mechanisms in place so that you can join up the various pieces of the puzzle so that things don't fall through the cracks.

MR HODGE: Thank you. And then, Ms Baker, I might just then ask you to talk about one last risk that the two of you raise under the current model, and that is the loss of trust or the relationship, and I was wondering if you could in particular just explain the view that the two of you have informed about the use of incorrects and the term "unintended human error".

 MS BAKER: Yes, I'm happy to. Just to add to that last point as well it's really important that if there are any events such as that that Dr Kogios has just described that they need to be identified and remedied as early on in the process as possible. I think that's what we're advocating for by the case managers can have more overall case visibility. But, yes, more than happy to talk to this.

So we noted that because the results are reported sample by sample through the work list, so they're made available to QPS, and then, for example, say there's 10 samples in a case, they get reported perhaps by different individuals, they're verified on the way out, but they go sample by sample. At some point the case manager is asked So they bring together all of to produce the statements. those results. It's a completely normal scientific forensic phenomenon that we don't always agree with the interpretation of DNA results. That's not to say that one person is right and another person is wrong. It can be down to different experiences, different perspectives. Sometimes two people just need to sit down in the same room together and nut it out and listen to each other's points of view, and that can be sufficient.

But what we noticed at the FSS is that the (indistinct) looking at the statement disagreed with the original interpretation, they were forced to sort of by way of retraction issue an incorrect notice and to say that the original result was as a result of unintended human error. I think from my perspective that gives the opinion that somebody's done something wrong; whereas what we're really talking about is a perfectly normal difference of opinion between forensic scientists that can either be resolved through conversation or perhaps through additional testing as examples. So to label them as an incorrect and to say it's an unintended human error, I personally would feel quite uncomfortable if that was a label that was applied to some work that I had done, and I feel it goes against that

no blame culture that we discussed earlier and that people should be able to have those differences of opinion and have those robust discussions and do a bit of extra testing to gain some more confidence in the result. But the person delivering that statement and going to court and providing evidence has to have the belief and the confidence in the result that they're reporting.

MR HODGE: Thank you. Is it fair to say - and I direct this to Dr Kogios - one of the things you'd say that's problematic about describing that type of situation where the reporting side - where the scientist ultimately doing a review at the end comes to a different view is that to describe it as error is incorrect or maybe incorrect in the sense that it may not involve any error at all; it's just a matter of differing professional judgment.

 DR KOGIOS: Yes, that's right. That is right. sort of put ourselves in the shoes of the QPS members receiving, you know, notification of the amendment to the result and we sort of thought about from their perspective that must be - you know, it might perhaps cause them to lose faith a little bit in the laboratory. "What's going on in that laboratory? They keep changing their mind." As Ms Baker said, these are not errors per se. It's just the natural consequence of the complexity of DNA evidence and the fact that in some instance, not all instances but in some instances, two highly trained individuals will come to a difference of opinion.

In a case manager allocation type model those differences of opinions tend to get nutted out before a result goes out the door. So that might be one of the things that's playing out with the Queensland model, that police are being brought into the process at an earlier stage perhaps before that difference of opinion has had a chance to be nutted out. So we were concerned about what that might mean for QPS in terms of their ability to rely on the results that are coming from FSS and for the general faith that they have in the people who are providing, you know, forensic DNA services to them. It's very important of course that the parties involved have that level of trust and can rely on the results.

MR HODGE: Thank you. I think that's a really important point. Can I give an example of something that might occur, which is that it might be that one scientist looks

at the profile or looks at the result that they get and thinks, "That is something where I can see two distinct profiles," but another scientist might look at it and think, "Actually, it's more than two contributors," and that changes their interpretation of the result that they can draw from the sample; that's the kind of situation you're talking about?

DR KOGIOS: Yes, that's exactly right. The number of contributors can be one of the most vexed issues. So, you know, what would you do in that circumstance? Well, ideally the two individuals would sit down, nut it out. Potentially they might reach agreement at that point. If not, then perhaps a third person can be brought in to adjudicate. We did see evidence of that process at QHFSS. So that certainly is the practice.

 I guess, the difference is because of the operating model numerous scientists are coming into contact with the samples in the case. Let's say it's a large case with lots of samples in it. You've got many, many hands touching that case. So you've got that potential for divergence of opinion in a much greater scale than you would have if you had a whole of case, dedicated case manager reporting that particular case. So that was why we thought this might be happening more often in Queensland; that coupled with the fact that, you know, the results are being made available to police earlier on in the piece.

Again, how might you put in a safeguard as sort of a checker balance operating model? You could consider perhaps use of some sort of flag so that if you're reporting to police on a sample by sample basis rather than having done whole of case review perhaps some way of communicating to police that, you know, "This is a preliminary result, perhaps even an interim result, and it may be subject to change. So if you are intending to rely on this result perhaps to arrest an individual or to, you know, do something in your investigation, let us know and we might pull this one out and look at it from a holistic point of view to make sure that you are ultimately going to be able to rely on that result in court." That's an example of a safety net, if you like, that could be applied to this type of operating model.

MR HODGE: I understand that as an idea for a safeguard. Can I just ask about that. Would that be, in your view, a

good practice to adopt for a major crime?

DR KOGIOS: I think that's a good practice to adopt if that's your operating model. If your operating model is you have this split between two agencies then, yes, I would see that as a vital safeguard.

MR HODGE: I suppose, sorry, what I meant was presumably even within the model of having the split between two agencies you could still have some kind of case management so that you weren't delivering results on a sample by sample basis and you were waiting until there had been a holistic review at the end before the results were delivered back from the DNA laboratory back to the police?

DR KOGIOS: Yes, that's true. You could invoke a case manager allocation model within that split workflow; yes, that's absolutely correct.

MR HODGE: And I was just interested in whether your view is in terms of good practice whether for major crime it would be good practice to be continuing to provide those kinds of provisional sample by sample results to police or whether you think it would be better for major crime to have case management so that you're not delivering interim results; you're only delivering final results after you've had a holistic review. It may be there's no right answer, but I'm just interested in your views about it

DR KOGIOS: Yes, look, I think Ms Baker and I, our lived experience has absolutely been in the case manager allocation model. We mike that model. It's served us well over the years. Neither of us has ever worked in a model like the model in Queensland, and I think it's important that we flag that. There are labs and people who work within that model who sing the praises of that model. So I don't think it's for us to say the model itself is wrong. All we can do is call out some of the perhaps gaps that we've seen around the way that model is currently operating.

MR HODGE: And then just to tie off on the significance of this description "unintended human error" am I right in thinking your points are in relation to scientists within the lab the problematic aspect of it is it conveys the idea that if one person, the final person, has a different view from you that that must mean that you are in error or you

are wrong in the original view that you took, which is not something that the two of you would agree with as an approach to scientific reasoning?

DR KOGIOS: Yes, that's correct. So the reality is you can never really know - unless, you know, a construct, an artificial construct that you've set up and you have that ground truth knowledge, you know exactly how many people have contributed to that mixture because you've constructed that mixture; absence of that, which of course is not the reality in a case work scenario, you can never know. All you can do is come to an opinion on the basis of the evidence. So you can never say that one person was wrong and the other person was right.

MR HODGE: And then the second aspect of it is, which I'll direct to Ms Baker, but the second aspect of it is it gives police the misleading impression that if somebody else comes to a different - another scientist comes to a different view from the first scientist, then that means that the first scientist was wrong as opposed to this is just a matter of professional judgment about which judgments will differ.

MS BAKER: Yes, I'd agree with that. I think it's an incredibly poor choice of words. I can also see a situation where it could act as a deterrent from a scientist wanting to come to a different conclusion about a particular DNA result. We're all different. lot of people who work in forensics who perhaps wouldn't appreciate the sort of confrontation that would need to be involved in having those discussions or wouldn't want to (indistinct) an incorrect or an unintended human error on one of their colleagues. I think that as a barrier is concerning and certainly not ideal. Just the negative impact on people's own morale. Like I said, possibly just a very poor choice of words, where what we're actually talking about is a difference in scientific opinion.

MR HODGE: Thank you. And then I want to then move to talk about the possible changes that might be made. So can we bring up paragraph 39 on page 20 first. This is where you talk about the in tube model that's presently used by the QPS and the FSS, and you explain the advantages of that which is high throughput and fast turnaround times. You make the point there that there are risks, and you've talked already about the risks and we've talked about them,

because you don't have case management oversight.

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 Then I want to just move to paragraph 40, which is over the page, where you explain the kinds of safeguards that are required for sexual assaults and other complex cases where you're using that kind of in tube model. This comes back to the point I was asking you about earlier, which is that there's really two different aspects of the Queensland model. One is the separation between QPS and FSS, but the other of it is the work list system. Perhaps at a general level am I right in thinking the safeguards that you identify are really ones that would mean not applying the work list system to these kinds of complex crimes even if you are using the in tube model? I might start with - I can see Ms Baker nodding, but I might start with Dr Kogios

DR KOGIOS: Yes, I think that's right. I think that's right. I mean, ultimately it's a matter for each jurisdiction to determine their own operating model. So I guess we're working from the sort of assumption that the operating model exists. As we've said, we didn't look at QPS. So we've got, you know, limited line of sight into how the model works from their perspective. If the model persists these are the types of safeguards that we thought could be put in place at FSS because, remember, that was of course the focus of the work that we were there to do, to look at FSS per se. So these aspects would be, I suppose, modifications in the work list model.

MR HODGE: Well, I mean, in saying that, is it fair to say it's much more like using the model that's presently used for P1 samples but applying it to all complex cases? So that is treating it as a whole case, somebody having an overview of it, somebody interacting directly with QPS, somebody watching the progress of it through?

DR KOGIOS: Yes, I think that's fair to say. Certainly (a) there, case manager allocation at the point of entry, yes, that's as we understand it what happens in the P1 category of cases. So applying that to P2, yes. (b) as well, whole of case review prior to reporting. We didn't see whole of case review prior to reporting for P1s. So I don't think it would be as simple as picking up the P1 model and broadening that out to P2s. That would work in terms of 40(a), case manager allocation at point of entry. But, 40(b), it would be new for the P1 category of cases as

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MR HODGE: Yes. Am I right in thinking - I might just direct this one to Ms Baker - your proposition is not that any one of these safeguards would be sufficient by itself; it's that you need all of these safeguards to deal with the kind of risks that arise from using that separation production line model of in tube samples coming from QPS?

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MS BAKER: So in this case we sort of talk about the Yes. Swiss cheese effects where all the holes have to line up for something not great to happen at the end, and it's a way of how do you plug all of those gaps so that you're avoiding or minimising any risk of that end result I would like to note as well those P1 cases, happening. I did review some of those as part of our work and I noted that there was really good collaboration and communication between the FSS and the QPS for those cases. That gave me great faith that this particular sort of case manager model that we're suggesting is something that the scientists are more than capable of and that they have already demonstrated the ability to work well with that kind of collaboration and communication throughout the lifecycle of those cases.

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31 32 MR HODGE: And, Dr Kogios, I suppose then one of the things that it will be important for the Commissioner to grapple with is you and Ms Baker obviously have a lot of experience with the case management model. Is it the case that you can use the case management model whilst maintaining the separation between of function lying with QPS and some function lying with FSS or does it really require a reconsideration of where you draw the line?

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DR KOGIOS: I think you can apply the principles that underpin the case manager-led model to a split workflow. It just might mean that in some instances you get your QPS member and your FSS member together to conduct a whole of case review, for example, because the knowledge might be split between those two individuals. So from a principle based approach, yes, it can be applied to the current operating model.

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MR HODGE: Just to unpack that a little bit, in terms of the kind of model that you and Ms Baker are familiar with as a case management model, in that model would the examination strategy be devised by the same person or preferably be devised by the same person who also then has a holistic overview of the entire case?

DR KOGIOS: Look, it's certainly not the case that it must be the same person. There are instances where, you know, you might have a person who's on maternity leave or who has left the organisation who started the case and then you've got someone else coming in to finish off the case and, you know, there would be some crimes against the person where there's no problem at all with it being two separate people.

For me I think the key is that the person at the back end, the person who is reporting, has access, full access, to that examination strategy so that they can see and understand the basis for the examination strategy having been set up the way that it was. So, you know, coupling those two people together or providing that level of information, sharing that level of information, you know, you could achieve the right outcome albeit with those two people not being the same - there's a level of extra difficulty if those two people don't belong to the same agency, but these are not insurmountable problems.

 MR HODGE: I understand. They're surmountable if you have adequate levels of communication between at one end the person who is devising the examination strategy and at the other end the person who is undertaking the holistic review of the DNA testing?

DR KOGIOS: Yes, that's right. As Ms Baker said, we did see evidence of that in P1 cases from both QPS side and from FSS side.

MR HODGE: And, Ms Baker, in your experience of this kind of case management model could you just maybe give the Commissioner some sense of how common would it be for the person who is undertaking the devising of the examination strategy to be the same person who ultimately is going to have the holistic responsibility for the case?

MS BAKER: I would say that would be the ideal and that would certainly be sort of normal practice from my perspective from the laboratories where I've worked. There are occasions, and I can think of one being where there's actually quite a large backlog at the sort of reporting end and that you don't want your labs (indistinct) twiddling

their thumbs while your reporting staff are heavily trying to get through a backlog of work for any particular reason. So in that situation you may allocate the person to do the triaging or the examination strategy for cases coming in to make sure that those cases are progressing through the laboratory in a timely way, and perhaps even if there is a backlog at the reporting end you are having those conversations with the police, with the Crown prosecutor or, sorry, the OPDD in Queensland and you can prioritise work based on that. But at least the work has been done. So that's the situation. But, ideally, you would have a case start to finish. It's a much smoother process because it takes time for another scientist to go back and review from the start to the point where they entered that case.

MR HODGE: At the moment under the Queensland model whatever examination strategy there is is one that's devised within the QPS?

MS BAKER: Yes, with the exception of the sexual assault kit.

MR HODGE: And you haven't undertaken a review of the QPS procedures, but one of the things I gather you haven't seen is what SOP the QPS have for devising an examination strategy?

DR KOGIOS: That's right.

 MR HODGE: And if we go to page 24 of your report you deal there with bias, and I just wanted to get you to explain that a little bit to the Commissioner because it seems to be one of the points that is inherent in your opinion from the stages of the workflow you set out is that under your case management model you're not assuming that the same scientist will devise the examination strategy, conduct the collection analysis plate reading, and effectively undertake all stages of the analysis; that's right, isn't it?

DR KOGIOS: Yes, that's correct. That's correct.

MR HODGE: And in fact you think it would be - in fact you're not even suggesting that would be a good model. To you a good model would be one where you do have different people at different stages because you want to blind some of them to certain information so they're not biased in

relation to what's going on?

DR KOGIOS: Yes, that's right. That's right. So we've included this because, you know, we understand that there may be some concerns or some challenge offered to the so-called case manager allocation model, which is the model that Ms Baker and I are used to, in that, well, then how do you manage bias. Well, this section of our report speaks to what we consider to be, you know, international emergent best practice around how to handle the topic of bias in DNA case work in particular. But in actual fact it does have broader application right across forensics.

 So if we think about bias really as being, you know, the shortcuts taken by the human brain unconsciously, and there are two types of bias that can arise in the forensic context. One is around context bias. So that might be where information around, you know, case information not relevant to the task at hand might be biasing the analyst in some of the decisions that they're making. Then the second form of bias is confirmation bias. So that might be where you have a view about something because of something that you've already seen and so you're unconsciously focused on that information that then would confirm what you expect to see.

 So the international recommended gold standard approach to this we looked to a paper by Krane in the Journal of Forensic Sciences that speaks to a concept called sequential unmasking. Essentially what it does is and you can see it up there on the screen - it sets out the various stages of the workflow, five in total; the first one being really, you know, assessing the case and setting your examination strategy, and you've heard us use that language today; the second one being the actual sort of analysis phase; then through into interpretation; through to reporting; and through to final review.

What this best practice calls out would be, you know, if you think about it as a bookend approach at the start and at the end of the process having that full picture so that you can set the right examination strategy, you can make sure all the right testing has been done and nothing looks odd in the case, but in between the people who are involved in the workflow don't have all of that additional information. So they just know, "I'm to pick up item 4 and examine the handle of the knife for trace and look for

1 blood on the knife blade." They don't need to know anything about the broader context of the case. That would 2 be what we would consider to be that emergent best practice 3 4 approach. 5 You know, can we say we think that the substructure of 6 the FSS laboratory in that they have got groupings aligned 7 8 to tasks, they have that evidence recovery team, they have their analysis team, and they have their reporting team, it 9 actually lends itself really well to this style of 10 approach, provided you've got the ability to sort of mask 11 12 from individuals at certain stages in the process, you know, key pieces of information masked at certain stages, 13 and that's where we make a recommendation around changes to 14 the forensic-register to support that level of sequential. 15 16 17 SHORT ADJOURNMENT 18 Mr Hodge. 19 THE COMMISSIONER: 20 21 MR HODGE: Thank you. Dr Kogios, can you see and hear me? 22 DR KOGIOS: 23 I can, yes. 24 25 MR HODGE: Thank you. And, Ms Baker, can you see and hear me? 26 27 MS BAKER: Yes, I can, Mr Hodge. 28 29 Great. Thank you. Now, could we just scroll MR HODGE: 30 down a little bit on that report that we've got up there. 31 32 I might start with you, Ms Baker. I wanted to ask you about recommendation 2. So you see in recommendation 2 the 33 34 recommendation you and Dr Kogios make is that: 35 QPS/FSS retrospectively review all sexual 36 assault and complex cases falling outside 37 the "hot jobs" and "major incident" 38 categories. 39 40 I want to just start with understanding some terms there. 41 42 When you say complex cases what do you mean by that? 43 44 So I think as Dr Kogios gave us an example MS BAKER: earlier, not all homicides could be considered a complex 45

handle for trace DNA and the blade for blood, then we

If your submission is a knife, that you look at the

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wouldn't necessarily call that a complex case.

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 Most volume crime cases or sort of property crime cases might be considered simple. But you may well have a situation where a gang of individuals go from one end of the street to another smashing windows in cars, being cut in the process and depositing blood. Whilst it's a property crime, that may well be considered complex because you have multiple scenes in effect in terms of each vehicle; potential multiple offenders as well. So I think we're just highlighting that the complex cases aren't necessarily defined by the type of crime that has been committed but more so the number of samples that are required and the number of crime scenes, the number of individual victims or complainants in a case, or the number of offenders as examples.

MR HODGE: And then where you refer to hot jobs and major incident categories can you just explain what you mean by that?

MS BAKER: Honestly I might struggle. Those are terms used by the QPS that I'm aware of. We refer to the report of Anna Davey when she was describing those types of cases, though I wouldn't give my opinion as to what they mean when they're a QPS term as far as I'm aware.

 MR HODGE: Maybe I should approach this in a slightly different way. You're recommending a retrospective review of all of the sexual assault and complex cases other than those that were classified as hot jobs and major incidents. So maybe you could just explain to us why not review the hot jobs and major incident ones?

MS BAKER: Yes, of course. So my understanding from Anna Davey's report is that that sort of holistic case overview has been carried out for those hot jobs or major instances - sorry, major categories. My understanding was that Ms Davey didn't find evidence to suggest that that holistic case overview to make sure there's no potential other avenues for evidence or processing or different types of testing, and that she didn't see evidence that that's been carried out for other types of cases. We felt that given that there is the potential for some of that missed opportunity to fall through the cracks, if that holistic overview isn't carried out, that that would be a really good idea just to make sure that there was nothing missing

for individual cases which require further testing or fall into that missed opportunity category.

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MR HODGE: So is the reason for or is the reasoning behind recommendation 2 then that sexual assault and complex cases are the ones most likely to have been detrimentally affected, not in terms of the ultimate conclusion necessarily but in terms of the process of gathering and analysing DNA evidence by virtue of the system that was in place in Queensland without the kinds of safeguards that you and Dr Kogios have recommended?

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MS BAKER: Yes, I think that would be fair. We were also, I guess, taking a risk based approach of the impact of not - or missing evidence in those types of cases, or missing the potential for more testing. That's not to say it will necessarily bring a different outcome, but just the missed opportunity for potential work.

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MR HODGE: And are you able to - I'll perhaps direct this to Dr Kogios first. Are you able to help us or help - yes, help us with understanding what kind of criteria might be applied to determine whether a case is a complex case, or is it a matter of judgment depending upon each case?

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DR KOGIOS: Well, I think it is a matter of judgment. I think as a starting point - and Ms Baker is right; in this recommendation we are highly reliant on the findings of Commission expert Anna Davey and we do quote from her report on page 18 of paragraph 37(b)(ii). She made a finding as a result of having a good look at the QPS process that this process of holistic case review was missing in certain categories of cases. So I think a starting point would be to be guided by that and look at those categories of cases where that holistic review hadn't happened. I think the criteria would be and the reason why we've sort of listed this as a sort of sequential QPS first and then FSS would be that the police themselves would be best placed to understand, yes, you know, perhaps there were some cases there that hadn't been holistically reviewed, but those cases may not be proceeding for reasons unrelated to the forensics, for example. So there perhaps is no point to go back and look at those particular cases if there's no realistic chance of a prosecution for other reasons, reasons not related to the forensics. than a blanket go back and test all of those samples and also, to be frank, FSS wouldn't necessarily know, one would assume, which cases fell outside of the ones that had been holistically reviewed.

So I think as a starting point it would be Anna Davey's report to look at the category of cases that fell outside of those that had been holistically reviewed, and then a systemic review of each case by QPS to ascertain, you know, whether there was more work to be done given the circumstances of the case, and then at that point engaging the lab to facilitate the progression of further testing for those cases that had been pulled out and identified as being in scope.

MR HODGE: I see. And just, though, in terms of maybe then for the QPS and thinking about what constitutes a complex case, how would you suggest they go about thinking about that?

 DR KOGIOS: Well, we really have defined "complex case" as really being a case involving multiple items, multiple persons of interest, multiple complainants, for example, multiple accused. So I really think there's no simple answer to this. Each case turns on its individual merits. But, because we were guided by Ms Davey's report, her report would be a starting point for QPS - I mean, assuming that this is accurate and is in fact the case, for QPS to start with those jobs that hadn't been holistically reviewed and to systematically work their way through those cases.

MR HODGE: And presumably in the absence of the kind of case management framework that you and Ms Baker have described where you've got a DNA scientist who's involved in the examination strategy at the beginning, the person who is or the group that is going to need to make a judgment about whether a case is complex or not will have to be the QPS?

DR KOGIOS: Yes, yes. And our understanding is QPS are outside of SAIKs, QPS are setting examination strategies. So there's nothing in the materials for us to suggest that they're not entirely capable of and, you know, well placed to perform that work.

MR HODGE: Thank you. And then if we go over the page to recommendation 3, so this is where you make a recommendation - I'll direct this to you, Dr Kogios - for a

1 change to the process that is used in Queensland at the moment to establish what you describe as fit for purpose 2 work streams for the different type of case work received; 3 4 that's right? 5 Yes, that's right, and again prefaced with 6 DR KOGIOS: there is no one universal accepted best practice model. 7 Ιf 8 this operating model is retained, this recommendation speaks to some safeguards and some checks and balances that 9 we think could be helpful in terms of those complex cases. 10 11 12 MR HODGE: I understand. I think is it fair to say your 13 point is the recommendation as to what ought to be done is that there needs to be the introduction of fit for purpose 14 work streams within QHFSS; it may be that there are 15 different models that might be used that are fit for 16 purpose depending upon whether or not, for example, the QPS 17 continues to be the controller and decider of examination 18 strategies? 19 20 21 DR KOGIOS: Yes, that's right. 22 MR HODGE: And, if the QPS continues to be the controller 23 and decider of examination strategies, then that will 24 require particular safeguards to be introduced to deal with 25 the risks that arise from that kind of situation? 26 27 28 DR KOGIOS: Yes. From the situation that involves people 29 in the workflow split between two different agencies. 30 MR HODGE: And in general, regardless of whether QPS or 31 FSS - that is regardless of whether it is somebody within 32 the police or somebody within the DNA lab who decides what 33 34 the examination strategy is going to be - your 35

recommendation is that for sexual assaults and complex cases that there be a case management approach to those kinds of cases?

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I mean, there's many different ways to DR KOGIOS: Yes. define a case manager approach, but this recommendation details those specific elements. They're sort of principles that underpin that case manager model that we think would be helpful.

MR HODGE: Thank you. Now, I was then going to move to talk about or to ask Ms Baker about the toolkit that's available to the FSS. I just wondered whether either of

you wanted to say anything finally in relation to the workflow model before we move to that other topic.

MS BAKER: No, nothing from me.

MR HODGE: I'll take silence as no. Thank you. So then can we move to the toolkit that is available to FSS at the moment, and if we go to paragraph 74 on page 36. The point that you identify at the start of paragraph 74 - I'm directing this to Ms Baker - is that FSS offers standard DNA testing only; it doesn't have Y-STR testing?

MS BAKER: That's correct, yes.

MR HODGE: And there's a number of other kinds of technology or methodologies that you then list in the following sentence which are also things that are not presently available to FSS?

MS BAKER: Yes.

MR HODGE: And I think the two of you also observe that there seems to be very limited subcontracting of samples, that is the sending out of samples, by FSS to other laboratories around Australia?

MS BAKER: Yes. My understanding is it's the QPS that will decide what samples to outsource in that respect, and the FSS provide them the samples to send out.

MR HODGE: And if we then go over the page to page 37 in paragraph 80 the two of you make the point that the lack of Y-STR capability places FSS outside of best practice in terms of the provision of service. I was interested in understanding, or maybe for the Commissioner to understand, what are the benefits of Y-STR testing for sexual assault investigations?

MS BAKER: I'm happy to speak to that. So I will say Y-STR capability is revolutionary when it comes to sexual assault investigations. So we know that in many cases sexual assault may involve touching or, for example, penile or digital penetration. There's not always semen deposited as a result of those assaults. So, as I spoke to earlier, having the capability to detect really low levels of male DNA against an overwhelming background of female DNA, so, for example, in those intimate body swabs, it is

revolutionary now that we have that male specific testing. It's also really helpful in determining (indistinct) providing more confidence when you're determining the number of contributors of DNA. So you use Y-STR for that to say, "Well, actually, there was a low level of male DNA. It could have been one person. It could have potentially been two." We've run that sample through this Y-STR testing and we've detected, for example, two paternally unrelated males in that sample. MR HODGE: And is it right to say from your report most other Australian labs have had Y-STR for the last five years? Yes, my understanding is that the final MS BAKER: forensic service provider implemented Y-STR approximately five years ago. MR HODGE: And that's why, if we go over the page to page 38, in recommendation 15 you and Dr Kogios recommend that the: QPS/FSS retrospectively review all sexual assault casework to identify cases with samples suitable for Y-STR testing.

MS BAKER: Yes. I think we both feel this is incredibly important for those people who are experiencing sexual assault in that just because no male DNA was detected using standard DNA testing that's not to say in fact that incident didn't occur and there could have in fact been low levels of male DNA that just weren't able to be detected using that standard test.

That's a matter of great concern to me THE COMMISSIONER: because anybody who practises criminal law in Queensland knows that sexual offences are a large proportion of the offences that are tried in court. Were you able to get any sense of why it's taken FSS so long to undertake the validation process? Were there some troubles they were facing that need to be addressed?

I might speak to that one, if you can hear me DR KOGIOS: over the rain that's just unfolding above my office. I would say it's very hard to quickly operationalise new capabilities when you don't have a dedicated research,

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development and innovation group or when you don't have staff members who are dedicated to research and development. So the lab has rightly been trying to implement and validate and implement Y-STR for many years. I would say that they're hamstrung by the pressures that you know, the constant grind of getting cases out the door and not having that dedicated research capability to support them in rapidly turning on a new capability. I think that that is a significant part of what has contributed to it taking them quite so long.

THE COMMISSIONER: Ms Baker?

MS BAKER: Yes, so I share your concern as well, Commissioner, in that I understand that sexual assault cases are highly underreported and, sort of, of those that are reported very few make it through to a court process.

THE COMMISSIONER: Yes

MS BAKER: So it really does have a significant impact. I think the statistics of the number of people, not just women but men and children as well, who experience sexual assault, this capability as I said is revolutionary for those types of cases.

 THE COMMISSIONER: Yes. So the trouble that they were facing in the lab is that it seems that scientists have to do this sort of work in effect part-time, whether it's a project or whether it's Y-STR validation. Nobody's allocated to it as a matter of sole interest and priority. They do it when they can find time in amongst all their other work; is that an exaggeration, what I've put?

No, I think that would be fair. I think you've also got the backdrop of a focus on high throughputs, rapid turnaround as well is also going to be a factor in that. I'm quite sure the lab very much wanted to have implemented I will say it is a more complex DNA test. Y-STR by now. The interpretation and the way in which it's reported and the statistical assessment of Y-STR is different to So it's not just a straightforward standard DNA testing. validate a particular kit. It has knock-on effects as well in terms of the reporting of those results. So for that reason it would take longer than, for example, just introducing a new standard DNA kit.

 THE COMMISSIONER: Yes. Thank you.

DR KOGIOS: I might just add it's also certainly the case that not every forensic science provider can have every tool, every technique, particularly some of the smaller ones. It's just not possible. The volume of work that they get wouldn't necessarily justify everything. So outsourcing is entirely appropriate. We have seen instances of outsourcing of this particular type of case work. As we understand it, that's not a decision that's made by FSS as to whether a particular case gets an outsource or not. So I just think it's important to offer that as well in the interests of, you know, a full balanced view.

THE COMMISSIONER: Thank you for mentioning that because I saw in your report - and it's obvious when you think about it - you may not be able to do everything in a particular lab, in Australia anyway

DR KOGIOS: Yes.

THE COMMISSIONER: Mr Hodge, you carry on, but just keep an eye on the time, which is almost there.

 MR HODGE: I will. Thank you, Commissioner. Could I just tie off on three things from what both of you have just said. The first is the significance of Y-STR is the reason why - and I'll direct this to you, Ms Baker. If we go to the bottom of page 37, recommendation 13, you recommend as a priority the validation and implementation of Y-STR profiling, and I think that reflects the things that you've already said to the Commissioner about its significance.

MS BAKER: Yes, and I would like to say that this isn't something that the laboratory should have to do in isolation. I would hope that the Australasian forensic community will support the laboratory in doing that because certainly many of the other laboratories have significant experience in Y-STR testing.

MR HODGE: And then the second thing is you were talking a little earlier about recommendation 15, if we could just scroll down again. One thing I just wanted to clarify about that is if it's been the practice of FSS to test all swabs in a SAIKs kit will that pose any difficulty for going back to perform Y-STR testing?

MS BAKER: There may well be some loss of material if samples, for example, have gone through that microcon concentration to full, which I know the Commission has spoken about. What I will say is there is all the potential to re-extract the swabs, and we do have successful Y-STR from re-extraction of those swabs or the sample remains, if there's differential lysis being carried out, it's not ideal but we could always go back to that (indistinct) as well. So there are sort of several bites at that cherry, I guess is what I'm trying to say. So just because all the samples were tested doesn't mean that there is nothing left to look at for Y-STR. It is a more sensitive DNA test than standard DNA testing, which is helpful.

MR HODGE: And so, given your answer, if samples haven't been microconned to full and depending on whether there's been re-works of samples or not, there's likely to be sufficient material still available from original extracts to be able to perform Y-STR testing?

MS BAKER: Yes, there should be. Like I said, the options to go back to sample remains can also be available. Sometimes we find that the original extraction has taken out a lot of the female DNA and in fact when the sample is re-extracted we actually get almost get a better ratio of male DNA from those.

MR HODGE: And the other part of that is, as I think you know, Queensland has ceased the DIFP process and will have to go back and retrospectively review cases that were designated as DIFP over the course of the last more than four years. Is it fair to say you would expect that as part of that review good practice would be to consider whether in reviewing those samples Y-STR should be used for them?

MS BAKER: Yes, absolutely. I think it has to be a case of best science approach and what is most likely to give a probative result given the amount of extract that's available. I wouldn't necessarily limit the Y-STR retrospectively to just four years.

MR HODGE: No, I understand. You wouldn't just stop four years ago. You would go back further than that. But you would agree, I think, that given that you are going to be

reviewing DIFP cases it's particularly important to take this into account

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MS BAKER: Absolutely, yes.

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MR HODGE: And then, Dr Kogios, you in answering one of the Commissioner's questions, you were referring to the decision to outsource to other laboratories being made by the QPS rather than FSS, and I was interested in whether you have a view about whether it will be better for FSS to be independently considering whether or not to outsource to other laboratories?

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DR KOGIOS: Not necessarily. I mean, I think that ultimately, you know, it is the remit of QPS to decide how to build their case, and they are perhaps better placed to have the fulsome knowledge of the case context to know whether a particular sample would benefit from that approach. So QPS taking carriage of that decision doesn't trouble me.

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MR HODGE: I understand. And then in making that decision what are the kinds of factors that you would expect to be taken into account, or does it depend on the case?

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DR KOGIOS: Well, the case circumstances, absolutely. So it does depend on the case. The case scenario, the availability - I mean, are we talking specifically about I guess any kind of outsourcing depends on the availability to procure those particular services. Sometimes it might be a question of having conversations with the scientist because the scientists are best placed to know about emerging techniques. The forensic science community is watching with great interest; there are developments in relation to the forensic investigative genetic genealogy as another examine of new capability that we've not had traditionally. So I think, broadly speaking, the DNA scientists would be best placed to provide advice to QPS on the potential for further testing more broadly than Y-STR, for example. But ultimately I would see that that would be a decision for QPS.

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MR HODGE: Is it fair to say that some of the factors that you expect would need to be taken into account would include particular information about the scientific qualities of the material, for example, the quant value, level of degradation.

45 46 47 DR KOGIOS: Yes, I think that would be fair enough. So again it does call for that conversation between the QPS and FSS.

 MR HODGE: Thank you. And then I just want to ask you about one last thing before we finish. Could we just go to page 13 of your report, and here you have recommendation No.1, which is:

Consideration be given to the establishment of a Forensic Science Advisory Board to assist with the coordination and accountability for managing forensic services across agencies.

 I think a little earlier in your report you say if the provision of forensic services is to remain within the Department of Health then you would recommend the formation of this kind of board. But I wonder if you might just step back - and I'll ask you to do it first, Dr Kogios - and explain to us what kind of board you envisage and what the purpose of that kind of board would be?

DR KOGIOS: Really, this recommendation was grounded in the fact that QHFSS is funded by QPS and Queensland Health, but its output has, you know, much broader implications for police, for the criminal justice system, the broader criminal justice system, and for the Queensland community. So really it's about having that broader frame of reference that brings in to play all the different stakeholders across that broader criminal justice system.

You know, ultimately the operating model, it's a matter for each jurisdiction, as we have said. But particularly where you've got forensic science provision happening outside of the traditional criminal justice system and, I guess, in that sense we're saying Health perhaps sits to the side of that system, we felt that it could be of benefit to have some sort of overarching board that could help connect the different players in this space, provide a place of guidance perhaps to sense check things like a DIFP policy so that it's not just sort of left to the lab to make a decision in isolation or just by checking in with QPS.

I mean, ideally you take into account a broader

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         spectrum of considerations when you're setting those sorts
         of policies. But in practical reality how do you do that?
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         It's not always easy to do that. It's not always easy to
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         know where to go and in what format to go.
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                                                      So this was
        where we felt that some sort of overarching forensic
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         science advisory board to connect the agencies together to
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         help with that sort of development of a whole of system
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         sort of lens, perhaps, sort of understanding of what risk
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         appetite might look like across that broader criminal
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         justice system could be really helpful for a lab like FSS
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         sitting as they currently do within Health.
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        MR HODGE:
                     Thank you.
                                 Commissioner, that was all I wanted
         to ask this afternoon, and then tomorrow when we resume
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        Ms Hedge will be continuing the examination.
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                             Yes.
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        THE COMMISSIONER:
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        MR HODGE: So is that a convenient time to adjourn?
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        THE COMMISSIONER:
                             It is.
                                     Ms Hedge, how long do you think
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        you'll be?
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        MS HEDGE:
                     Maybe half a day.
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        THE COMMISSIONER:
                             All right. You might not have
26
         formulated your views yet, the rest of you, but, Mr Hunter,
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         do you have a great deal for the two experts?
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        MR HUNTER:
                      Very little.
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        THE COMMISSIONER:
                             Mr Rice?
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        MR RICE:
                    Not a great deal, Commissioner.
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        THE COMMISSIONER:
                             Ms McKenzie?
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        MS McKENZIE:
                        No.
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                             Mr Hickey?
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        THE COMMISSIONER:
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        MR HICKEY:
                      No, Commissioner.
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        THE COMMISSIONER:
                             All right.
                                         So it looks like we'll
         finish with this part of it tomorrow at some point; is that
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         right?
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        MR HODGE:
                    Yes.
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        THE COMMISSIONER:
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                             That's what you think?
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        MR HODGE:
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                     That's what I expect.
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        THE COMMISSIONER:
                             That's good. Thank you very much for
         this afternoon and for your help. New Zealand is ahead of
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         us, isn't it?
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        MR HODGE:
                     Three hours ahead in New Zealand and one hour
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         ahead in Victoria.
                             So if we resume at 10 then it will be
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         1 pm for Ms Baker and 11 am for --
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        THE COMMISSIONER:
                             So we could resume at nine?
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                     I think they have been told 10 am Brisbane
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        MR HODGE:
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         time.
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        THE COMMISSIONER:
                             What suits you two?
                                                   Ms Baker, what
21
         time suits you tomorrow?
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                     Well, I'm probably - I'm happy starting earlier
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         just because it's probably not got the same impact on me.
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         But I'll leave it up to Dr Kogios.
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26
        THE COMMISSIONER:
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                             Thank you.
                                         Dr Kogios?
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         DR KOGIOS:
                      Look, I'm fine as well. I'm just mindful that
         for Ms Baker the finish time in Brisbane is later on in the
30
         day for her. So that opportunity to start earlier that
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        might be appreciated. But from my perspective I can fall
         in with whatever works with you, Commissioner.
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        THE COMMISSIONER:
                             All right.
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                     I think just from feedback at the Bar table we
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         couldn't start before 9.30.
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        THE COMMISSIONER:
                             All right. We'll make it 9.30 Brisbane
         time.
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                     Thank you, Commissioner.
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        MR HODGE:
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        THE COMMISSIONER:
                             See you then. Thank you.
46
        <THE WITNESSES WITHDREW
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         AT 4.38PM THE COMMISSION WAS ADJOURNED TO WEDNESDAY,
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         2 NOVEMBER 2022 AT 9.30AM
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