

**Cathie Allen**

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**From:** Cathie Allen  
**Sent:** Tuesday, 11 February 2020 11:55 AM  
**To:** Krosch.MattN[OSC]  
**Cc:** Keatinge.DavidJ[OSC]; John Doherty; Allison Lloyd  
**Subject:** RE: DNA success rates manuscript

Hi Matt

Thanks for your time on Friday to discuss the manuscript.

I've discussed with the Team Leaders from Forensic DNA Analysis regarding an appropriate FSS staff member, and Allison Lloyd is very happy to assist with this. Allison is currently acting in the role of Senior Scientist for the Intelligence team, so is suitably placed to assist with DNA success rates, given NCIDD is within her portfolio. I've included Allison on this email, but will email her the manuscript on a separate email.

We look forward to working with you on this and other projects in the future.

Cheers  
 Cathie

**Cathie Allen**

Managing Scientist

**Police Services Stream, Forensic & Scientific Services**  
 Health Support Queensland, Queensland Health

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**From:** Krosch.MattN[OSC] [REDACTED]@police.qld.gov.au>  
**Sent:** Wednesday, 5 February 2020 3:13 PM  
**To:** Cathie Allen [REDACTED]@health.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED]@police.qld.gov.au>; John Doherty [REDACTED]@health.qld.gov.au>  
**Subject:** Re: DNA success rates manuscript

Hi Cathie,  
 I should be at my desk all tomorrow and Friday, when will be a good time to speak with you about this paper?

Matt

**From:** Krosch.MattN[OSC]  
**Sent:** Friday, 31 January 2020 3:05:36 PM  
**To:** Cathie Allen [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Cc:** Keatinge.DavidJ[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>; John Doherty [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Subject:** RE: DNA success rates manuscript

Hi Cathie,

Sorry, I must not have been in reception in the depths of the building. My apologies. Wednesday it is. Enjoy your weekend.

Matt



**Dr. Matt Krosch**  
Research Officer  
Quality Management Section, Forensic Services Group  
Queensland Police Service  
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**From:** Cathie Allen [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Sent:** Friday, 31 January 2020 14:59  
**To:** Krosch.MattN[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>  
**Cc:** Keatinge.DavidJ[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>; John Doherty [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Subject:** RE: DNA success rates manuscript

Hi Matt

I tried your mobile, as suggested, but it went straight to message bank.

I'll give you a call on Wednesday sometime to discuss the manuscript.

Cheers  
Cathie

**Cathie Allen**  
Managing Scientist  
**Police Services Stream, Forensic & Scientific Services**  
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**From:** Krosch.MattN[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>  
**Sent:** Friday, 31 January 2020 2:42 PM  
**To:** Cathie Allen [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Cc:** Keatinge.DavidJ[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>; John Doherty [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Subject:** RE: DNA success rates manuscript

Hi Cathie,

Sorry I missed your call earlier, I had to return to the lab to finish off the morning's experiments. I'm about to head off for the day, but back at the desk on Wednesday. Let's try to arrange a time to speak then.

Matt

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**From:** Cathie Allen [REDACTED] <[\[REDACTED\]@health.qld.gov.au](mailto:[REDACTED]@health.qld.gov.au)>  
**Sent:** Thursday, 30 January 2020 14:37  
**To:** Krosch.MattN[OSC] [REDACTED] <[\[REDACTED\]@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED] <[\[REDACTED\]@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>; John Doherty [REDACTED] <[\[REDACTED\]@health.qld.gov.au](mailto:[REDACTED]@health.qld.gov.au)>  
**Subject:** RE: DNA success rates manuscript

Hi Matt

Sorry for not getting back to you, I've had a few other priorities and some [REDACTED] leave.

I'll give you tomorrow at some stage, if that's ok?

Cheers  
 Cathie

**Cathie Allen**

Managing Scientist

**Police Services Stream, Forensic & Scientific Services**

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**From:** Krosch.MattN[OSC] [REDACTED] <[\[REDACTED\]@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>  
**Sent:** Wednesday, 29 January 2020 9:36 AM  
**To:** Cathie Allen [REDACTED] <[\[REDACTED\]@health.qld.gov.au](mailto:[REDACTED]@health.qld.gov.au)>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED] <[\[REDACTED\]@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>; John Doherty [REDACTED] <[\[REDACTED\]@health.qld.gov.au](mailto:[REDACTED]@health.qld.gov.au)>  
**Subject:** RE: DNA success rates manuscript

Hi Cathie,

We are keen to progress with submission of this manuscript. Can you please let me know if you would still like to meet to discuss the paper or chat over the phone. I'm available all this week and Wednesday-Friday next week.

Regards,

Matt

**Dr. Matt Krosch**



Research Officer  
 Quality Management Section, Forensic Services Group  
 Queensland Police Service  
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**From:** Krosch.MattN[OSC]  
**Sent:** Monday, 13 January 2020 09:59  
**To:** Cathie Allen [REDACTED]@health.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED]@police.qld.gov.au>; John Doherty [REDACTED]@health.qld.gov.au>  
**Subject:** RE: DNA success rates manuscript

Hi Cathie,

Certainly happy to meet with you to discuss the paper. I'm free all week, but Insp Keatinge has limited time this week so if he was to join us for a face-to-face then meeting here would be preferable. Anytime this week works for me at this stage.

Alternatively, I'm happy to discuss over the phone if that helps to save travel time?

Cheers  
 Matt



**Dr. Matt Krosch**  
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 Quality Management Section, Forensic Services Group  
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**From:** Cathie Allen [REDACTED]@health.qld.gov.au>  
**Sent:** Monday, 13 January 2020 09:16  
**To:** Krosch.MattN[OSC] [REDACTED]@police.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED]@police.qld.gov.au>; John Doherty [REDACTED]@health.qld.gov.au>  
**Subject:** RE: DNA success rates manuscript

Hi Matt

Thanks for the email and the opportunity to review the manuscript.

It would be great if we could meet to discuss the paper and the data used within it. I'm happy to host you at FSS or alternatively, I'm happy to meet with you at QPS HQ. Please let me know your preference and availability.

Cheers  
 Cathie

**Cathie Allen**

Managing Scientist

**Police Services Stream, Forensic & Scientific Services**

Health Support Queensland, Queensland Health

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e [Cathie.Allen@health.qld.gov.au](mailto:Cathie.Allen@health.qld.gov.au) w [www.health.qld.gov.au/healthsupport](http://www.health.qld.gov.au/healthsupport)*Queensland Health acknowledges the Traditional Owners of the land, and pays respect to Elders past, present and future.***From:** Krosch.MattN[OSC] [REDACTED] [@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>**Sent:** Tuesday, 7 January 2020 1:02 PM**To:** Cathie Allen [REDACTED] [@health.qld.gov.au](mailto:[REDACTED]@health.qld.gov.au)>**Cc:** Keatinge.DavidJ[OSC] [REDACTED] [@police.qld.gov.au](mailto:[REDACTED]@police.qld.gov.au)>**Subject:** DNA success rates manuscript

Dear Cathie,

Over the latter months of last year I spent some time summarising FR data for DNA results with a view to establish percentage successes for common items/substrates and collection methods. This was essentially a self-driven project that grew out of conversations with SOCOs and OICs and so the focus was on our side of the process to ensure we're making the best decisions on sampling to maximise success in the lab. In a nutshell it involved pulling information on the DNA results for every exhibit that was submitted over a set time period and searching the item description/location fields for keywords that allowed extraction of specific items/substrate results. The aim was to develop an evidence base on the success rates of sampling certain items to inform procedures and make recommendations to our officers on which collection methods were most effective for specific items based on recent data from actual casework.

I've now completed the analysis and have written the results up as a short paper that I hope to submit to AJFS as I believe this information is important to communicate to the forensic community. However, because the paper necessarily contains information about DNA profiling in Queensland we wish to offer you the opportunity to review the draft manuscript before submission to ensure that you and QHFSS are happy for the contents to be published. Please find attached the draft manuscript as a word document and the tables both at the end of the manuscript and as a separate excel file on individual sheets.

If you would like any further explanation on the methods or outcomes, please don't hesitate to get in touch.

Kind regards,

Matt

**Dr. Matt Krosch**

Research Officer

Quality Management Section, Forensic Services Group

Queensland Police Service

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RTI Release

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**Variation in forensic DNA profiling success rate among sampled items and collection methods: a Queensland perspective.**

Matt N. Krosch<sup>a\*</sup>

*<sup>a</sup>Quality Management Section, Forensic Services Group, Queensland Police Service, 200 Roma Street, Brisbane, QLD 4000*

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DOH DISCLOSURE LOG

## **Variation in forensic DNA profiling success rate among sampled items and collection methods: a Queensland perspective.**

Understanding the relative success rates of recovering DNA profiles from different touched evidentiary items/substrates and between different methods of collection is critical for optimal targeting of forensic sample collection and triaging for analysis. Further, reporting of such success rates allows comparison between jurisdictions that can drive improvements and prompt discussion between stakeholders. This study analysed success rates of DNA sampling from major and volume crimes attended by the Queensland Police Service, Australia, from January 2017 to September 2019. In total, 61 344 total records were analysed, representing the most comprehensive analysis of its kind to date. Success rates were determined for various sample types and items, including those that are commonly encountered or have high probative value. Results suggested that, overall, around 10% of trace DNA samples returned full profiles, but with some disparity between swabs (13.45%) and tapelifts (7.01%). Despite this, tapelifts provided nearly 25% of total suspect identifications compared with 17% for trace swabs. Substantial variation in profiling success among items/substrates was observed, as there was between swabs and tapelifts taken from the same item. These data contribute significantly to our understanding of DNA prevalence and recovery and provide a critical evidence base to inform changes to operational procedures.

**Keywords:** swabs, tapelifts, full profile, mixed profile, suspect identification

DOH DISCLOSURE LOG

## Introduction

DNA sampling, particularly of touched objects and surfaces, has become an increasing focus for forensic analysts globally<sup>1,2</sup>. Resolution of DNA profiles from such items can be highly probative and thus understanding the relative success rates of recovering profiles from items is important for targeting sample collection and triaging for analysis. Such success rates should be considered in the context of the specific collection and analysis methods used by a given jurisdiction. Comparing data generated from different extraction and profiling methods may not necessarily represent a like-for-like comparison and must be considered with some caution. Nevertheless, there can be great value in comparing between jurisdictions to determine whether substantial differences are apparent and where improvements could be made. Moreover, sampling of putatively touched items can be a point of friction between investigators and forensic scientists who may have contrasting anecdotal experience concerning a questioned item. Finally, where jurisdictions use multiple collection methods for similar items (because of officer preference or simply what consumables are available at the time), it is important to assess whether one method outperforms another to ensure operational procedures follow best practice. Therefore, there is a need for additional data to inform decision-making and assist forensic scientists in optimally targeting sampling effort.

There have been sporadic attempts over the last twelve years to address this issue in a range of national and state jurisdictions from New Zealand<sup>3</sup>, Switzerland<sup>4</sup>, Canada<sup>5</sup>, Netherlands<sup>6</sup>, Singapore<sup>7</sup>, and Australia<sup>8</sup>, including a comparative analysis of experimental and casework samples from Western Switzerland<sup>9</sup>. These studies analysed success rates for various types of casework samples; either those most commonly collected, restricted to volume crime cases, or other items of interest. Generally speaking, these studies were consistent in suggesting that, as expected, biological fluid traces (blood, saliva, semen) provided the greatest proportions of full profiles (up to 87.5%<sup>9</sup>), whereas touch samples were far less successful overall (<30%). Worn or touched items that often returned above average proportions of full profiles include hats/caps, gloves, adhesive tape, clothing, door handles and steering wheels<sup>3-9</sup>, though in some cases these may represent victim profiles.

This study aimed to analyse success rates of DNA sampling from major and volume crime for the Queensland Police Service, Queensland, Australia over a period of roughly 20 months. Success rates were determined for sample types over the entire period, as well as broken down to selected items of interest, including those that are commonly encountered or have high probative value. Queensland data are then discussed in the context of previous literature.

## Methods

Samples included in this analysis were collected from exhibits related to both major and volume crime between the 1<sup>st</sup> January 2017 and 11<sup>th</sup> September 2019. Methods of collection included swabbing with a rayon swab (Medical Wire, UK) pre-moistened with 70% ethanol, tapelifting with a custom 3M adhesive tape kit (Lovell Surgical Supplies, Australia), excision (e.g., fabric, cigarette butts), and scraping. All samples were processed at Queensland Health Forensic Scientific Services (QHFSS) following standard procedures: DNA extraction conducted using the DNA IQ™ Casework Pro Kit for Maxwell®16 (Promega Corp., Melbourne, Australia) on a Maxwell® 16 MDx (Promega Corp.); quantification using Quantifiler® Trio (ThermoFisher Scientific, Melbourne, Australia) on the 7500 Real Time PCR System (Applied Biosystems™, ThermoFisher Scientific), and STR amplification using PowerPlex® 21 (Promega Corp.). DNA quantification results determined progression to profiling, according to QHFSS standard procedures: samples of concentration <0.0088ng/μL were considered to have insufficient DNA and were thus categorised as 'no DNA'. Samples that yielded sufficient DNA (>0.0088ng/μL) proceeded to STR profiling.

Data was extracted from the in-house laboratory information management system (LIMS) for all DNA samples sent for processing between the 1<sup>st</sup> January 2017 and 11<sup>th</sup> September 2019. The LIMS was queried in such a way to return sample type (e.g., swab/tapelift) and exhibit description information, as well as STR profiling results categorised as 'full' (all 42 alleles present), 'partial/mixed' (less than 42 alleles, or more than one contributor), or 'no DNA' (DNA quantification insufficient for profiling). In some cases, profiling results could include multiple categories; for example, full+partial/mixed profile results may indicate full profiles deconvoluted from mixtures, or no DNA+full or

partial/mixed where sub-threshold information (<150rfu) was present, or where the original quantification was insufficient, but the sample was profiled following investigator request. Profiles were also recorded for whether they matched a suspect/offender reference sample. This master spreadsheet was queried using Windows Powershell to extract lines in which the exhibit description matched specific text strings. All resulting sub-sheets were manually reviewed to ensure only relevant data was included. Despite this, inconsistencies in spelling and terminology in the exhibit description limited the completeness of the analysis; however, this is unlikely to impact dramatically on the interpretation of DNA success rates. Percentages of each profile result category were calculated for the total dataset, each collection method across all items, and then broken down for collection method from each selected item. Success rates were also assessed for porous versus non-porous substrate surfaces. Sample metadata allowed separation of swabs from biological fluid stains (blood, saliva, semen) to be separated from those taken from putative touched areas or handled objects.

## Results

In total, 61 344 total records (representing 60 332 unique exhibits) were analysed, the majority of which were swabs or tapelifts (Table 1). Swabs collected from biological fluids represented a much smaller proportion than those from touched areas/objects. Overall, 25.85% of samples returned full profiles: the greatest proportion of full profiles was obtained from samples of obvious stains of biological fluids, with the most successful being swabs of bloodstains (73.96%, Table 2). Partial/mixed profiles were rarely obtained from non-sexual assault kit semen swabs (1.96%), but otherwise ranged up to 28.04% of DNA results from other sample types. Percentages of suspect identifications ranged from 13.49% (hair) to 41.55% (blood swabs). Both swabs and tapelifts of touched objects/surfaces returned suspect identifications from ~15% of samples, but there was a significant disparity between full profile results (swabs = 13.45%; tapelifts = 7.01%). Despite this, tapelifts provided nearly 25% of total suspect identifications compared with 17% for trace swabs (Table 1), suggesting that the success of tapelifting is often reliant on partial profiles or deconvolution of mixtures.

Individual items/surfaces showed great variation in their percentage success. The greatest success for exhibits where no visible stain was observed was for swabs and excised sections from drinking straws, which produced full profiles in ~47% of samples taken, whereas tapelifts from straws were slightly less successful at 33.3%. Bedding (swab), waistbands of lower garments (swab), discharged cartridge cases (tapelift), underwear (both), zip/cable ties (both), and drinking vessels (both) all produced full profiles in >20% of samples. The least successful items (no full profiles recorded) included: swabs of cigarette packets, rocks, helmets, firearm barrels, shirt collars, power cords, rubber key handles, and several tools; tapelifts of external car door handles, sweat smears on cars, and glovemarks; and both swabs and tapelifts of public phones and fingermarks. Despite this, several of these items did return suspect identifications based on partial profiles; including, external car door handles, shirt collars, and rubber key handles. Among sexual assault-related samples, breast swabs identified the greatest percentage of suspects after penis swabs (suspect reference samples), no suspect identifications were recorded from perineum samples. The highest percentage of full profiles were reported from oral swabs (most likely complainant profiles, though 8.41% were identified a suspect), whereas the lowest proportion of full profiles were from breast swabs.

Some distinct differences in the recovery of full profiles from swabs and tapelifts of trace samples were observed for specific items. Swabs were at least twice as successful as tapelifts for car doors, car door handles, seatbelt straps & buckles, adhesive tapes, drinking vessels, firearm handles, sweat smears on cars, waistbands of lower garments, sledgehammers, mattock/pickaxes, torches, and bedding. In contrast, tapelifts were more successful for discharged car airbags, gearsticks, motorcycles (including handlebars), cigarette packets, power cords, flyscreen, rubber and metal keys, cartridge cases (both discharged and live), firearm barrels, mobile phones, shirt collars, helmets, hats, rocks, and several tools. In contrast to conventional wisdom, tapelifts of non-porous surfaces recovered slightly more full profiles than swabs, whereas swabs were better for porous surfaces (Table 3). Furthermore, porous surfaces returned a greater percentage of full profiles and suspect identifications than non-porous surfaces.

### ***Data caveats***

A small number of samples were recorded as returning results in more than one category: 256 records were categorised as both partial/mixed and full (likely representing full profiles deconvoluted from mixtures), representing 2% of partial/mixed records and 1.6% of full profile results; 614 samples were categorised as both partial/mixed and no DNA, representing 1.7% of no DNA results and 4.8% of partial/mixed results; 3001 samples were categorised as both no DNA and full, representing 8.2% of no DNA results and 19% of full profile results; and 92 samples were categorised across all three categories. The vast bulk of such multiple categorisations are due to sub-threshold information present in otherwise full, partial or mixed profiles, or samples that fell below the internal quantification threshold for profiling but were processed following investigator request. In the context of the total dataset these multiple categorisations are not considered to substantially impact on the interpretation of profiling success rates. Manually reviewing every record was outside the scope of this project.

### **Discussion**

The analysis presented here of over 18 months of DNA sampling data, representing more than 60 000 individual exhibits, from the Queensland Police Service has revealed some interesting patterns that can inform operational procedures. Averaged over all items/surfaces, trace swabs recovered more full profiles than tapelifts; however, there was substantial variation noted among exhibit types, including many for which tapelifts were the more successful method of collection. Increasing the granularity of the analysis therefore provided a deeper insight into DNA profiling success rates among items and methods of collection. Interestingly, percentage profiling successes for swabs and tapelifts from porous and non-porous surfaces were opposite to conventional wisdom.

It is difficult to compare the data presented here with previous studies from other jurisdictions. The specifics of collection technique, consumables, DNA extraction and STR profiling procedures and kits between organisations and over time are likely to have significant influence on profiling success. In addition, there has been variation across studies

in the exhibit categorisation strategy used and hence granularity of data analysed. For example, some studies lump all clothing samples together<sup>4,7,9</sup>, whereas others separate them into subcategories for specific clothing types<sup>3,5,6</sup>. Further, some studies were deliberately restricted to samples taken from volume crime scenes<sup>8,9</sup>, whereas others either were from all crime scenes or did not specify<sup>3-7</sup>. This limits the ability to make truly like-for-like comparisons between studies. Nevertheless, some general trends deserve discussion.

Overall, trace DNA success was similar for Queensland as for most jurisdictions compared here (Table 4). Interestingly, profiling success for many items included in the comparison was poorer than that reported from other jurisdictions, despite the current use in Queensland of a more sensitive DNA profiling kit than that used in many of these previous studies. This suggests that there were many other more successful items sampled by Queensland that made up the shortfall (possibly including SAIK swabs, for example). Alternatively, it could be because of different collection, storage, submission and triage procedures in other regions, or a factor of analysing total sample data rather than smaller, selected subsets. Trace DNA profile success was also relatively high for items from cars (airbags, seatbelts), drinking straws, chewing gum, cartridge cases, underwear and waistbands, and bedding. The majority of comparisons with previous literature related to swabbed items (Table 4); however, tapelift sampling of many of these items in fact returned more full profiles than swabs (11 out of 19 items). Perhaps the most striking discrepancies were for swabs from hats/caps, inside of gloves, and collars compared with the results of Mapes et al<sup>6</sup>. Within the Queensland data, clear differences in profiling success were observed between collection methods which will contribute toward updated operational procedures.

These data provide valuable insight into DNA profiling success of one of Australia's largest police jurisdictions. Additional research is required to determine whether differences between Queensland and other published data stem from consumables used, collection technique, environmental effects (e.g., increased degradation), or some other factor. Some recent work has suggested that rayon swabs are not ideal for recovering maximum DNA from collected samples<sup>10</sup>, although this appears to contradict other research that supports rayon as



among the most effective swab materials<sup>11,12</sup>. Additional research is still required here to inform better consumables choice for forensic practitioners. Pleasingly, there is good support in the data presented here for the efficacy of forensic tapelifts, particularly in preference to swabs for many non-porous items. This accords with existing literature that supports tapelifting as a highly effective collection method<sup>13,14</sup>, including for the specific tape product used by QPS forensic officers<sup>15</sup>. Future research and reporting by other agencies into their success rates would benefit from a consistent approach to item and profile success categorisation, to maximise comparability between studies. This study demonstrates that increasing the granularity of data captured can reveal important trends that can inform best practice at the crime scene and laboratory.

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## DOH DISCLOSURE LOG

## Tables

**Table 1.** Number of records included for analysis separated into major sample types (minor sample types or those not subsequently analysed are not shown). Percentages of total records, suspect identifications, full or partial/mixed profiles, and no DNA records provided for each sample type.

Sample type	Number of exhibit records	Percentage of total records	Percentage of total suspect identifications (N=14267)	Percentage of total full profiles (N=15855)	Percentage of total partial/mixed profiles (N=12784)	Percentage of total no DNA (N=36484)
Cigarette butts	2633	4.29	7.46	9.16	6.31	1.75
Fabric	1865	3.04	4.56	5.00	3.83	2.50
Hair	289	0.47	0.27	0.52	0.21	0.53
Scraping	922	1.50	2.28	2.34	0.82	1.53
Swab (blood)	7248	11.82	21.10	33.81	9.05	4.00
Swab (saliva)	4769	7.77	12.93	12.17	10.46	4.97
Swab (semen)	51	0.08	0.10	0.09	0.01	0.11
Swab (trace)	16518	26.93	17.18	14.01	20.24	34.14
Tapelift	22576	36.76	24.45	9.97	38.40	45.74
All trace	39067	63.69	41.63	23.99	58.64	79.88

# DOH DISCLOSURE LOG

**Table 2.** DNA profiling results for samples collected by QPS forensic officers between 1 January 2017 and 11 September 2019.

Item	Collection method	Total results	Percentage suspect identification	Percentage full profile	Percentage partial/mixed profile	Percentage no DNA	
All	All	61344	23.26	25.85	20.84	59.47	
	Fabric	1865	34.91	42.52	26.27	48.90	
	Hair	289	13.49	28.72	9.34	67.47	
	Scrapings	922	35.25	40.24	11.39	60.74	
	Swab (blood)	7247	41.55	73.96	15.97	20.15	
	Swab (saliva)	4769	38.69	40.45	28.04	38.04	
	Swab (semen)	51	27.45	29.41	1.96	76.47	
	All trace	39066	15.20	9.73	19.19	74.60	
	Swab	16518	14.84	13.45	15.66	75.40	
	Tapelift	22548	15.47	7.01	21.77	74.01	
Steering wheel	Swab (blood)	40	67.50	62.50	25.00	27.50	
	All trace	3676	16.29	6.41	22.52	73.07	
	Swab	696	12.36	4.17	17.96	79.60	
	Tapelift	2980	17.21	6.95	23.59	71.54	
Cars	Airbags	Swab (blood)	53	69.81	84.91	13.21	15.09
		Excised	14	57.14	78.57	14.29	28.57
		All trace	236	31.78	18.64	27.12	61.44
		Swab	12	25.00	8.33	16.67	83.33
		Tapelift	224	32.14	19.20	27.68	60.27
Gear stick	Swab (blood)	9	55.56	55.56	44.44	11.11	
	All trace	761	10.91	5.65	15.24	82.00	
	Swab	241	6.64	2.90	9.54	88.38	
	Tapelift	520	11.73	5.96	16.73	78.85	

	Swab (blood)	110	58.18	79.09	11.82	19.09
All doors	All trace	164	12.80	6.71	14.02	80.49
	Swab	94	10.64	10.64	8.51	82.98
	Tapelift	70	15.71	1.43	21.43	77.14
	Swab (blood)	50	62.00	74.00	14.00	28.00
Internal door handle	All trace	104	14.42	7.69	15.38	78.85
	Swab	55	14.55	12.73	10.91	80.00
	Tapelift	49	14.29	2.04	20.41	77.55
	Swab (blood)	32	59.38	87.50	12.50	9.38
External door handle	All trace	39	7.69	5.13	12.82	82.05
	Swab	25	0.00	8.00	4.00	88.00
	Tapelift	14	21.43	0.00	28.57	71.43
	Swab (blood)	2	0.00	100.00	0.00	100.00
Seatbelt strap	Fabric	1	0.00	0.00	0.00	100.00
	All trace	154	6.49	3.25	10.39	87.66
	Swab	7	28.57	14.29	28.57	71.43
	Tapelift	147	5.44	2.72	9.52	88.44
	All trace	96	8.33	5.21	11.46	88.54
Seatbelt buckle	Swab	32	6.25	9.38	6.25	90.63
	Tapelift	64	9.38	3.13	14.06	87.50
	Swab (blood)	14	57.14	100.00	0.00	7.14
Motorcycles	All trace	83	8.43	3.61	12.05	86.75
	Swab	26	0.00	0.00	3.85	96.15
	Tapelift	57	12.28	5.26	15.79	82.46
	Swab (blood)	2	50.00	100.00	0.00	0.00
	All trace	73	8.22	4.11	12.33	86.30
	Swab	22	0.00	0.00	4.55	95.45
	Tapelift	51	11.76	5.88	15.69	82.35
	Cigarette butt	Excised (majority)	2633	40.41	55.15	30.65

Cigarette packet	Swab (blood)	5	40.00	100.00	0.00	0.00	
	All trace	12	8.33	8.33	33.33	58.33	
	Swab	4	0.00	0.00	25.00	75.00	
	Tapelift	8	12.50	12.50	37.50	50.00	
Cigarette lighter	All trace	185	7.57	4.32	11.89	84.32	
	Swab	141	8.51	4.26	11.35	84.40	
	Tapelift	44	4.55	4.55	13.64	84.09	
Bindings	All	421	9.50	10.93	14.73	77.91	
	Rope	Tapelift (majority)	87	4.60	13.79	18.39	72.41
	Zip/cable ties	All trace	70	22.86	21.43	14.29	68.57
		Swab	45	17.78	22.22	8.89	71.11
		Tapelift	25	32.00	20.00	24.00	64.00
	Power cords	Swab (blood)	7	42.86	42.86	28.57	57.14
		All trace	183	4.92	3.83	10.38	87.43
		Swab	89	1.12	0.00	6.74	93.26
		Tapelift	94	8.51	7.45	13.83	81.91
	Tapes	All trace	150	10.00	8.00	13.33	82.67
		Swab	87	9.20	11.49	13.79	80.46
		Tapelift	63	11.11	3.17	12.70	85.71
Deceased scenes	Tapelift (majority)	37	2.70	32.43	35.14	45.95	
Door handles (premises)	Swab (blood)	66	51.52	66.67	25.76	22.73	
	All trace	519	3.47	2.12	10.21	88.44	
	Swab	278	2.88	1.44	8.99	90.29	
	Tapelift	241	4.15	2.90	11.62	86.31	
Window frames/sills	Swab (blood)	174	51.72	78.74	11.49	16.09	
	All trace	126	8.73	7.14	6.35	88.89	
	Swab	73	8.22	8.22	6.85	87.67	
	Tapelift	53	9.43	5.66	5.66	90.57	
Flyscreen mesh	Swab (blood)	37	59.46	81.08	8.11	13.51	

	Excised	7	28.57	14.29	14.29	71.43	
	All trace	1117	5.01	4.57	10.92	85.50	
	Swab	159	2.52	1.89	6.29	92.45	
	Tapelift	958	5.43	5.01	11.69	84.34	
Mouth/rim of drinking vessel	All trace	4578	35.23	37.70	27.09	41.50	
	Swab	4423	36.08	38.68	27.36	40.29	
	Tapelift	155	10.97	9.68	19.35	76.13	
Drinking straw	Excised	68	55.88	47.06	33.82	32.35	
	All trace	506	50.20	46.44	28.66	33.79	
	Swab	494	49.80	46.76	28.34	34.01	
	Tapelift	12	66.67	33.33	41.67	25.00	
Drug pipe/bong	Swab (majority)	215	26.98	11.16	30.23	61.40	
Chewing gum	Whole item (majority)	47	14.89	63.83	12.77	31.91	
	All trace	425	5.88	2.35	11.29	87.29	
	Swab	238	4.20	1.68	6.30	92.86	
	Tapelift	187	8.02	3.21	17.65	80.21	
Keys	Rubber	All trace	12	8.33	8.33	16.67	83.33
		Swab	4	25.00	0.00	25.00	75.00
		Tapelift	8	0.00	12.50	12.50	87.50
	Metal	All trace	166	5.42	1.81	8.43	90.36
		Swab	106	2.83	0.94	4.72	94.34
		Tapelift	60	5.00	3.33	15.00	83.33
Plastic	All trace	161	6.21	3.73	11.80	85.09	
	Swab	70	4.29	2.86	4.29	92.86	
	Tapelift	91	7.69	4.40	17.58	79.12	
Cartridge cases	All trace	212	8.96	9.91	3.77	89.62	
	Swab	127	6.30	5.51	2.36	92.91	
	Tapelift	85	12.94	16.47	5.88	82.35	



Discharged	All trace	70	5.71	11.43	2.86	88.57	
	Swab	41	4.88	4.88	0.00	95.12	
	Tapelift	29	6.90	20.69	6.90	79.31	
Live	All trace	130	10.77	9.23	3.85	89.23	
	Swab	80	7.50	6.25	3.75	91.25	
	Tapelift	50	16.00	14.00	4.00	86.00	
Firearm	Swab (blood)	18	44.44	83.33	11.11	22.22	
	All trace	831	9.15	2.65	10.83	87.48	
	Swab	444	7.66	2.03	9.68	89.86	
	Tapelift	387	10.85	3.36	12.14	84.75	
	Handle	All trace	232	8.62	2.16	10.78	88.36
		Swab	92	7.61	4.35	11.96	86.96
		Tapelift	140	9.29	0.71	10.00	89.29
	Barrel	All trace	31	6.45	3.23	12.90	87.10
		Swab	19	5.26	0.00	10.53	94.74
		Tapelift	12	8.33	8.33	16.67	75.00
	Trigger	All trace	273	8.79	2.56	10.99	87.55
		Swab	174	8.62	2.87	10.34	87.93
		Tapelift	99	9.09	2.02	12.12	86.87
	Knife	Swab (blood)	363	34.71	50.69	34.16	26.45
		All trace	1329	15.65	7.22	18.96	77.20
Swab		792	14.52	7.20	17.55	78.28	
Tapelift		537	17.32	7.26	21.04	75.61	
Handle		All trace	986	16.33	4.97	20.08	77.79
		Swab	523	14.72	3.63	17.97	80.50
		Tapelift	463	18.14	6.48	22.46	74.73
Blade		All trace	236	13.56	14.83	17.37	72.03
		Swab	219	13.70	14.61	17.35	72.15
		Tapelift	17	11.76	17.65	17.65	70.59

Gloves		Swab (blood)	14	57.14	50.00	21.43	35.71	
		Excised	12	50.00	8.33	41.67	50.00	
		All trace	1686	20.23	6.47	24.67	70.82	
		Swab	384	13.02	5.99	16.67	79.69	
		Tapelift	1302	22.35	6.61	27.04	68.20	
		All trace	1076	20.72	7.53	26.02	68.59	
		Inside surfaces	Swab	223	15.25	8.07	18.39	75.34
		Tapelift	853	22.27	7.39	28.02	66.71	
Fingermarks		Swab (blood)	10	20.00	40.00	20.00	40.00	
		All trace	102	2.94	0.00	5.88	94.12	
		Swab	85	3.53	0.00	7.06	92.94	
		Tapelift	17	0.00	0.00	0.00	100.00	
Glovemarks		All trace	140	2.14	0.71	2.86	97.14	
		Swab	121	0.83	0.83	0.83	98.35	
		Tapelift	19	10.53	0.00	15.79	89.47	
Sweat smears	Premises	All trace	181	3.87	4.42	2.76	94.48	
		Swab	157	3.82	4.46	3.18	94.27	
		Tapelift	24	4.17	4.17	0.00	95.83	
	Cars	All trace	40	0.00	5.00	2.50	95.00	
		Swab	37	0.00	5.41	2.70	94.59	
		Tapelift	3	0.00	0.00	0.00	100.00	
Phones	Mobile phone	Swab (blood)	32	43.75	65.63	34.38	18.75	
		All trace	174	13.79	4.02	23.56	74.14	
		Swab	119	11.76	1.68	21.85	77.31	
		Tapelift	55	18.18	9.09	27.27	67.27	
	Public phone	Swab (blood)	2	100.00	100.00	0.00	100.00	
		All trace	10	0.00	0.00	0.00	100.00	
		Swab	6	0.00	0.00	0.00	100.00	
		Tape	4	0.00	0.00	0.00	100.00	

Keypad (eg., safe/alarm)	Swab (majority)	26	3.85	7.69	7.69	88.46	
Computer keyboard	Swab (blood/trace)	5	20.00	60.00	0.00	40.00	
Fingernails	Scrapings	549	56.83	39.89	47.91	30.42	
	Clippings	71	25.35	67.61	26.76	22.54	
Condom	Swab (majority)	253	50.59	23.72	45.45	46.25	
Sexual assault-related	All	4586	22.50	48.95	22.55	41.95	
	High vaginal	629	25.60	54.05	30.84	30.21	
	Low vaginal	615	20.81	53.33	25.20	33.33	
	Hymen	11	9.09	63.64	9.09	36.36	
	Vaginal other	65	26.15	64.62	20.00	18.46	
	Vulval	980	16.73	54.39	18.88	37.55	
	Labial	202	13.86	63.37	17.33	31.19	
	Perineum	28	0.00	50.00	0.00	50.00	
	Perianal	442	14.03	35.75	17.19	56.79	
	Anal	147	10.88	42.18	9.52	59.18	
	Rectal	216	10.65	40.28	12.50	56.94	
	Breast	46	39.13	6.52	41.30	67.39	
	Oral	309	8.41	72.17	5.18	32.04	
	Penis	450	55.56	26.44	36.67	49.78	
	Clothing	Swab (blood/saliva)	5	60.00	40.00	40.00	40.00
Collar		Fabric	18	38.89	33.33	38.89	33.33
		All trace	409	27.14	7.33	34.23	61.86
Tapelift		Swab	11	27.27	0.00	36.36	63.64
		Tapelift	398	27.14	7.54	34.17	61.81
Beanie		Tapelift (majority)	89	34.83	6.74	38.20	57.30
Balaclava		Tapelift (majority)	90	31.11	18.89	21.11	66.67
Helmet		Swab (blood)	12	41.67	91.67	8.33	16.67
	All trace	148	29.05	8.11	31.76	62.84	

	Swab	12	0.00	0.00	0.00	100.00
	Tapelift	136	31.62	8.82	34.56	59.56
Hat/cap	Swab (blood)	37	48.65	48.65	35.14	27.03
	All trace	888	28.83	10.47	33.45	60.02
	Swab	42	14.29	2.38	19.05	78.57
	Tapelift	846	29.55	10.87	34.16	59.10
	Excised/scraped	189	44.44	39.68	40.74	83.07
Underwear	All trace	324	40.43	25.62	66.36	68.52
	Swab	13	53.85	38.46	61.54	46.15
	Tapelift	311	39.87	25.08	66.56	69.45
	Excised/scraped	29	20.69	41.38	17.24	72.41
Waistband shorts/pants	All trace	196	20.41	5.61	35.71	62.76
	Swab	5	60.00	20.00	60.00	40.00
	Tapelift	191	19.37	5.24	35.08	63.35
	All trace	939	11.40	4.37	15.65	81.36
Screwdriver	Swab	469	9.81	4.05	12.15	84.86
	Tapelift	470	12.98	4.47	19.15	77.87
	Swab (blood)	4	0.00	75.00	0.00	50.00
Sledge hammer	All trace	75	9.33	2.67	12.00	85.33
	Swab	22	4.55	4.55	4.55	90.91
	Tapelift	53	11.32	1.89	15.09	83.02
	Swab (blood)	22	27.27	63.64	13.64	59.09
Hammer	All trace	356	10.39	3.65	13.48	83.71
	Swab	116	9.48	3.45	11.21	85.34
	Tapelift	240	10.83	3.75	14.58	82.92
	Swab (blood)	5	20.00	100.00	0.00	0.00
Spanner	All trace	104	8.65	2.88	8.65	89.42
	Swab	55	7.27	3.64	5.45	92.73
	Tapelift	49	10.20	2.04	12.24	85.71

Chisel	All trace	66	16.67	3.03	16.67	81.82
	Swab	25	0.00	0.00	0.00	100.00
	Tapelift	41	26.83	4.88	26.83	70.73
Shovel	Swab (blood)	2	0.00	100.00	0.00	100.00
	All trace	66	10.47	4.65	8.14	87.21
	Swab	25	7.14	0.00	7.14	92.86
	Tapelift	41	12.07	6.90	8.62	84.48
Crow bar	All trace	268	5.97	2.99	7.09	91.79
	Swab	108	3.70	1.85	5.56	94.44
	Tapelift	160	7.50	3.75	8.13	90.00
Axe	Swab (blood)	3	33.33	66.67	33.33	33.33
	All trace	114	12.28	3.51	13.16	84.21
	Swab	24	4.17	0.00	8.33	91.67
	Tapelift	90	14.44	4.44	14.44	82.22
Mattock/Pickaxe	All trace	41	4.88	2.44	9.76	87.80
	Swab	7	0.00	14.29	14.29	71.43
	Tapelift	34	5.88	0.00	8.82	91.18
Torch	All trace	376	19.95	10.11	19.68	72.87
	Swab	163	14.11	13.50	12.88	78.53
	Tapelift	213	24.41	7.51	24.88	68.54
Brick/rock	All	527	8.73	10.82	7.40	89.18
	Swab (blood)	14	14.29	64.29	7.14	28.57
	All trace	287	3.83	3.48	5.92	91.29
	Swab	21	0.00	0.00	4.76	95.24
	Tapelift	266	4.14	3.76	6.02	90.98
	Swab (blood)	29	41.38	79.31	3.45	20.69
	All trace	227	9.25	6.61	8.81	87.22
Brick/paver	Swab	18	0.00	5.56	0.00	100.00
	Tapelift	209	10.05	6.70	9.57	86.12

Clip-seal plastic bag	All trace	267	14.98	8.24	13.48	81.27	
	Swab	213	15.02	7.51	13.15	81.69	
	Tapelift	54	14.81	11.11	14.81	79.63	
Bedding	All	1440	25.76	28.47	23.19	60.97	
	Excised	491	28.11	38.29	24.85	57.64	
	Scraping	348	25.00	8.91	28.74	49.43	
	Other	278	28.42	41.37	12.95	83.45	
	Swab (blood)	96	31.25	56.25	27.08	27.08	
	All trace	226	16.37	9.73	22.12	73.01	
	Swab	5	0.00	40.00	20.00	60.00	
	Tapelift	221	16.74	9.05	22.17	73.30	
	Mattress	All	158	11.39	31.01	12.66	71.52
	Mattress protector	All	63	52.38	19.05	39.68	63.49
Sheets	All	679	28.57	27.54	24.30	58.62	
Blanket	All	403	21.09	31.27	20.35	62.03	
Pillow	All	179	23.46	24.02	25.70	60.89	

**Table 3.** Comparison of percentage success in DNA sampling between porous and non-porous items/surfaces from Table 2.

Surface	Collection method	Total results	Percentage suspect identification	Percentage full profile	Percentage partial/mixed profile	Percentage no DNA
Non-porous	All trace	23234	12.35	7.61	14.30	80.38
	Swab	11836	9.87	6.88	11.11	83.99
	Tapelift	11398	13.97	7.33	17.08	77.95
Porous	All trace	3125	20.82	11.67	26.52	71.74
	Swab	134	20.44	13.41	25.59	72.78
	Tapelift	2991	21.20	9.93	27.46	70.71

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**Table 4.** Comparison of Queensland DNA profiling success data for specific items against equivalent data from the literature.

Exhibit category	Profile Collection	This study	Netherlands <sup>6</sup>	Singapore <sup>7</sup>	Switzerland <sup>4</sup>	Switzerland <sup>9</sup>	New Zealand <sup>3</sup>	New South Wales <sup>8</sup>
		Full	Single	Single	Full/partial>5 loci	Single	Full	Full/partial>12 loci
		Cigarette butt	Excised	55	84	81		70.6
Hat/cap	Swab	2	42					
	Tapelift	11					25	
Collar	Swab	0*	34					
Glove (inside)	Swab	8	25a	11		18.8b		
	Tapelift	7					25	
Torch	Swab	14	27					
Drinking vessels	Swab	39	57	34		55.6	21c	
Knife handle	Swab	4*	19					
Lighter	Swab	4*	17					
Firearm grip	Swab	4	6					
Firearms (other) Handle	Swab	2*						15
motorcycle	Swab	0*	9					
Cartridge cases	Swab	6*	6					
Tape	Swab	11	9	16				
Keys	Swab	2*	12					
Hair	Excised	29		21.1				
Drug apparatus	Swab	11		15			21c	
Thrown stones	Swab	0*			7	7.5		
Cables/power cords	Swab	0*			29	12.2		
Tools	Swab	4*d	5e	10	22			15



Clothing	Swab	29f		5	18.8b		
	Tapelift	12g				15h	
	Excised	38i					
Blood	Swab	74	68		87.5		
Dataset average	All trace	15j	25k	12	12k	16	14

\*greater percentage full profiles from tapelifts where relevant

a combined here from latex & fabric glove results

b combined category clothing/gloves

c combined category drinking vessels/drug pipes

d averaged over all tools analysed in Table 2

e combined here from screwdriver/crowbar/hand-tools (other)

f averaged over underwear and waistband shorts/pants in Table 2

g averaged over collar/beanie/balaclava/helmet/hat/cap/underwear/waistband shorts/pants in Table 2

h combined here from underwear/socks/upper garments results

i averaged over collar/underwear/waistband shorts/pants in Table 2

j average profiling success for trace samples only (i.e., excludes biological fluids, hair, cigarette butts)

k included bloodstain profiling results

s.47(3)(b)

---

**From:** Neville.DavidH[OSC] [REDACTED]@police.qld.gov.au>

**Sent:** Friday, 14 February 2020 2:42 PM

**To:** Cathie Allen [REDACTED]@health.qld.gov.au>

**Cc:** Krosch.MattN[OSC] [REDACTED]@police.qld.gov.au>; McNab.BruceJ[OSC]

[REDACTED]@police.qld.gov.au>; Keatinge.DavidJ[OSC] <[REDACTED]@police.qld.gov.au>

**Subject:** FW: DNA success rates manuscript

Hi Cathie

Matt has forwarded me the below email and we have had a discussion in relation to this. Thanks for taking the time to review his work. This paper is aimed at crime scene examiners to help them better focus their sampling methodology. It is not aimed at the laboratory and the introduction of additional lab factors might unnecessarily complicate the matter. It is important that the possible the impact of micron be covered in the discussion, however I don't think it is necessary for us to rerun the data. In this instance we were looking to provide QHFSS an acknowledgement in the paper, however it was not anticipated that the article would be become lab focused. As a result, a general review is probably all that is needed, if possible please.

Regards

David Neville

---

**From:** Allison Lloyd [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Sent:** Friday, 14 February 2020 10:31  
**To:** Krosch.MattN[OSC] [redacted] <[\[redacted\]@police.qld.gov.au](mailto:[redacted]@police.qld.gov.au)>  
**Subject:** FW: DNA success rates manuscript

Hi Matt,

I've been asked to go through your manuscript. I've given it a good read and have a few questions/comments... I'm more than happy to meet up or talk on the phone, whatever suits you better.

My number is [redacted] or [redacted].

Looking forward to working with you on this.

Kind regards,



**Allison Lloyd**  
A/Senior Scientist - Intelligence Team

**Forensic DNA Analysis, Police Services Stream**  
Forensic & Scientific Services, Health Support Queensland, Queensland Health

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e [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)> w [www.health.qld.gov.au/healthsupport/businesses/forensic-and-scientific-services](http://www.health.qld.gov.au/healthsupport/businesses/forensic-and-scientific-services)



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**From:** Cathie Allen [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>  
**Sent:** Tuesday, 11 February 2020 12:04 PM  
**To:** Allison Lloyd [redacted] <[\[redacted\]@health.qld.gov.au](mailto:[redacted]@health.qld.gov.au)>

Cc: Justin Howes [redacted]@health.qld.gov.au>; Paula Brisotto [redacted]@health.qld.gov.au>  
Subject: FW: DNA success rates manuscript

Hi Allison

Thanks so much for agreeing to be the FSS collaborator on this paper – I really appreciate it, given your busy role.

Attached is the manuscript and also the raw data.

I've discussed with Matt that the Government would be expecting a collaboration on this, given the significant investment they have made in the Forensic DNA Analysis lab to undertake DNA testing solely for the purpose of the QPS. I appreciate that Matt has driven this work himself and has focussed on sampling, however my perspective is that the lab has tailored it's processes to ensure success for a sample that's submitted, so it's a collaboration and Matt readily agreed. Matt has done all of the evaluation of the data to date, so I suggested that perhaps the FSS rep (as we spoke on Friday, prior to offering you the opportunity so wasn't able to name you) would be able to review some data, as I believe he hasn't taken into account any microcons that we've done to achieve the profiles. So they may need to run the report in the FR again, to capture the post extraction techniques so that we can review them to see if they have affected the outcome. If the report needs to be re-run, Matt will be able to achieve that, given he's within the QPS.

Please let me know if you have any questions. I'm excited that we're able to collaborate with the QPS on this and am excited for you to be given this opportunity, given your vast experience with profiles and NCIDD.

Cheers  
Cathie

**Cathie Allen**  
Managing Scientist

**Police Services Stream, Forensic & Scientific Services**  
Health Support Queensland, Queensland Health

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**From:** Krosch.MattN[OSC] [redacted]@police.qld.gov.au>  
**Sent:** Tuesday, 7 January 2020 1:02 PM  
**To:** Cathie Allen [redacted]@health.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [redacted]@police.qld.gov.au>  
**Subject:** DNA success rates manuscript

Dear Cathie,

Over the latter months of last year I spent some time summarising FR data for DNA results with a view to establish percentage successes for common items/substrates and collection methods. This was essentially a self-driven project that grew out of conversations with SOCOs and OICs and so the focus was on our side of the process to

ensure we're making the best decisions on sampling to maximise success in the lab. In a nutshell it involved pulling information on the DNA results for every exhibit that was submitted over a set time period and searching the item description/location fields for keywords that allowed extraction of specific items/substrate results. The aim was to develop an evidence base on the success rates of sampling certain items to inform procedures and make recommendations to our officers on which collection methods were most effective for specific items based on recent data from actual casework.

I've now completed the analysis and have written the results up as a short paper that I hope to submit to AJFS as I believe this information is important to communicate to the forensic community. However, because the paper necessarily contains information about DNA profiling in Queensland we wish to offer you the opportunity to review the draft manuscript before submission to ensure that you and QHFSS are happy for the contents to be published. Please find attached the draft manuscript as a word document and the tables both at the end of the manuscript and as a separate excel file on individual sheets.

If you would like any further explanation on the methods or outcomes, please don't hesitate to get in touch.

Kind regards,

Matt



**Dr. Matt Krosch**

Research Officer

Quality Management Section, Forensic Services Group

Queensland Police Service

Ph: (07) [redacted] | M: [redacted] | Email: [redacted]@[police.qld.gov.au](mailto:police.qld.gov.au)

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# DOH DISCLOSURE LOG

**Allison Lloyd**

---

**From:** Allison Lloyd  
**Sent:** Friday, 14 February 2020 1:04 PM  
**To:** Krosch.MattN[OSC]  
**Subject:** RE: DNA success rates manuscript

Hi Matt,

Monday is better for me due to on and off meetings all day. Have a good weekend.

Thanks,

Allison

---

**From:** Krosch.MattN[OSC] [REDACTED]@police.qld.gov.au>  
**Sent:** Friday, 14 February 2020 10:59 AM  
**To:** Allison Lloyd [REDACTED]@health.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED]@police.qld.gov.au>  
**Subject:** RE: DNA success rates manuscript

Hi Allison,

I'm out of the office today, but on email. Otherwise I'll be back at the desk on Monday. Happy to hear your thoughts.

Cheers  
 Matt

**Dr. Matt Krosch**

Research Officer

Quality Management Section, Forensic Services Group

Queensland Police Service

Ph: (07) [REDACTED] | M: [REDACTED] | Email: [Krosch.MatthewN@police.qld.gov.au](mailto:Krosch.MatthewN@police.qld.gov.au)

---

**From:** Allison Lloyd [REDACTED]@health.qld.gov.au>  
**Sent:** Friday, 14 February 2020 10:31  
**To:** Krosch.MattN[OSC] [REDACTED]@police.qld.gov.au>  
**Subject:** FW: DNA success rates manuscript

Hi Matt,

I've been asked to go through your manuscript. I've given it a good read and have a few questions/comments... I'm more than happy to meet up or talk on the phone, whatever suits you better.

My number is [REDACTED] or [REDACTED].

Looking forward to working with you on this.

Kind regards,



## Allison Lloyd

A/Senior Scientist - Intelligence Team

### Forensic DNA Analysis, Police Services Stream

Forensic & Scientific Services, Health Support Queensland, Queensland Health

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**From:** Cathie Allen [REDACTED]@health.qld.gov.au>

**Sent:** Tuesday, 11 February 2020 12:04 PM

**To:** Allison Lloyd [REDACTED]@health.qld.gov.au>

**Cc:** Justin Howes [REDACTED]@health.qld.gov.au>; Paula Brisotto [REDACTED]@health.qld.gov.au>

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Cheers  
Cathie



**Cathie Allen**

Managing Scientist

**Police Services Stream, Forensic & Scientific Services**  
Health Support Queensland, Queensland Health

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**To:** Cathie Allen [REDACTED]@health.qld.gov.au>  
**Cc:** Keatinge.DavidJ[OSC] [REDACTED]J@police.qld.gov.au>  
**Subject:** DNA success rates manuscript

Dear Cathie,

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If you would like any further explanation on the methods or outcomes, please don't hesitate to get in touch.

Kind regards,

Matt

# DOH DISCLOSURE LOG

**Dr. Matt Krosch**  
Research Officer  
Quality Management Section, Forensic Services Group



Queensland Police Service

Ph: (07) [redacted] | M: [redacted] | Email: [redacted]@police.qld.gov.au

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**Allison Lloyd**

---

**From:** Allison Lloyd  
**Sent:** Monday, 17 February 2020 11:57 AM  
**To:** Krosch.MattN[OSC]  
**Subject:** Thresholds

Hi Matt,

One thing I forgot to mention... Sub threshold peaks for PowerPlex 21 are under 40 RFU but for Profiler Plus ( all of 2017 for volume crime) were less than 50 RFU.

Kind regards



**Allison Lloyd**

A/Senior Scientist - Intelligence Team

**Forensic DNA Analysis, Police Services Stream**

Forensic & Scientific Services, Health Support Queensland, Queensland Health

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**Allison Lloyd**

---

**From:** Allison Lloyd  
**Sent:** Tuesday, 18 February 2020 1:34 PM  
**To:** Cathie Allen  
**Cc:** Justin Howes  
**Subject:** QPS Manuscript Feedback given via phone call 17/02/2020

Hi Cathie,

Here is a breakdown of the feedback I gave to Dr Matt Krosch yesterday morning via phone call regarding the 'Variation in Forensic DNA profiling...' manuscript.

Points discussed:

1. Page 4/Methods – We were still using Profiler Plus for volume crime samples for the most part of 2017 which was not mentioned. This also had implications for results obtained in the second paragraph of Methods as we were using a binary method of interpretation which would affect the profiles that could be counted as a 'successful' profile.
2. Page 4/Methods – QIA symphony was not mentioned
3. Page 5/Methods, line 1: Sub-threshold information (<150rfu) was incorrect and different for the different kits used.
4. Page 4/Methods, bottom line: (no DNA+full) what does this mean? This is when a 'NO DNA' result line was released and most likely the quant was not right, the batch requanted and corrected results released. I said these results should be considered as being the 'updated/corrected' results.
5. General discussion on what was considered a 'full' profile/mixed/partial. These results were taken directly from result lines in the FR. I offered to go through them in more detail. In my opinion, Dr Krosch did not have a particularly strong understanding of the results or what they meant.
6. General discussion on what was considered a 'successful' profile. I said that in my opinion, obtaining a profile regardless of whether it was able to be interpreted would be considered successful. It is my understanding that the QPS version of a successful result was obtaining a suspect identification/LR favouring contribution for a suspect (Page 6, 1<sup>st</sup> paragraph). I suggested that some definitions around 'success' and even the types of results such as 'full/mixed/partial etc' were put in the manuscript to avoid ambiguity.
7. General discussion that the processes/reworking strategies that DNA Analysis used were not vital to the manuscript as this was generally looking at different sampling methods and the different types of results obtained from those sampling methods and substrates and the point of the paper was for SOCOs to have some printed advice to take to Investigators for discussions as to why certain samples might not be as worthwhile as others (as per the anecdotal experience mentioned on page 3/Introduction). I expressed enthusiasm for this as I could see that might be less complex or uninterpretable profiles and our analysts could focus more time on potentially meaningful samples which would benefit us all. The impression I got from this was that we were both on the same wavelength.
8. General discussion on the success of the tape lifts (page 7/Discussion). I gave anecdotal stories of where I had seen unexpected profiles obtained on objects such as tapelifted rocks/bricks and that the success of the tapelifts was pleasantly surprising.

Thanks,

# DOH DISCLOSURE LOG



**Allison Lloyd**

A/Senior Scientist - Intelligence Team

**Forensic DNA Analysis, Police Services Stream**

Forensic & Scientific Services, Health Support Queensland, Queensland Health

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S.47(3)(b)

DOH RTI 2960

## DOH DISCLOSURE LOG

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