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Manufacturer's Information Test No. 22-5803: DNA-Blood

Each sample pack consisted of two known bloodstains on FTA cards (Items 1 & 2), and two questioned stains on clean, colored fabric (Item 3 & 4). Participants were requested to analyze these items using their existing protocols.

SAMPLE PREPARATION: The stains in Items 1, 2, 3, and 4 were prepared from human whole blood which was either drawn into citric acid preservative bags or EDTA tubes. Items 1 (75 μ I) and 3 (50 μ I) were created using blood from the same male donor. Item 2 (75 μ I) was created using blood collected from a male donor, but different than that of Items 1 and 3. Item 4 (50 μ I) was created using blood collected from a male donor, but different than that of Items 1, 2, and 3. Stains from different sources were prepared at separate times and were packaged once they were thoroughly dried. Completed sample sets were stored at -20°C until shipment on April 19, 2022 following completion of verification stage.

SAMPLE SET ASSEMBLY: For each sample set, all four Items (1-4) were packed into separate envelopes and then placed together in a pre-labeled sample pack envelope and sealed. The sealed sample pack envelopes were then packaged in pre-labeled heat seal envelopes and sealed. This process was repeated until all of the sample sets were prepared.

VERIFICATION: Laboratories that conducted predistribution analysis of the samples reported consistent allelic results and confirmed the manufacturer's expected associations.

The information presented here is that received from the sample manufacturer. It presents details of the design specification for the test samples and/or details of how they were prepared. This information does not necessarily represent the answers that should or could be obtained from an examination of the sample. Final interpretation of the results should be deferred until the summary report is available.

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Manufacturer's Information, continued Test No. 22-5803: DNA-Blood

Amelogenin and STR Results													
	Results compiled from predistribution laboratories and a consensus of at least 10 participants.												
ltem	D1S1656	D2S1338	D2S441	D3S1358	D5S818	D6S1043							
	D7S820	D8S1179	D10S1248	D12S391	D13S317	D16S539							
_	D18S51	D19S433	D21S11	D22S1045	Amelogenin	CSF1PO							
	FGA	Penta D	Penta E	SE33	TH01	TPOX							
	vWA	DYS391	DYS570	DYS576	Y Indel								
1	12,17	17,20	10,14	16,17	11,11	*							
	8,11	10,15	15,15	15,22	8,12	10,11							
	11,13	13,14	31,32.2	11,15	X,Y	10,10							
	24,24	11,11	7,14	12,22.2	6,9	8,8							
	16,18	11	20	18	2								
2	12,17	17,24	10,11	16,16	9,13	*							
	11,11	10,14	13,16	20,20	12,12	10,12							
	14,14	13.2,14	30,32	15,15	X,Y	10,12							
	24,26	9,13	10,12	16,16.2	6,9	8,11							
	16,18	11	18	18	2								
3	 12,17	17,20	10,14	16,17	 11,11	*							
	8,11	10,15	15,15	15,22	8,12	10,11							
	11,13	13,14	31,32.2	11,15	X,Y	10,10							
	24,24	11,11	7,14	12,22.2	6,9	8,8							
	16,18	11	20	18	2								
4	16,17.3	20,21	14,14	14,14	12,12	*							
	9,10	11,15	13,17	18,21	8,12	10,12							
	15,19	14,15	28,32.2	15,15	X,Y	10,13							
	20,23	11,11	12,20	18,28.2	9,9	8,11							
	15,18	10	17	19	2								

YSTR Results Results compiled from predistribution laboratories and a consensus of at least 10 participants.												
	DYS437 DYS518	DYS438 DYS533	DYS439 DYS549	DYS448 DYS570	DYS449 DYS576	DYS456 DYS627	DYS458 DYS635	DYS460 DYS643	DYS481 YGATAH4			
		D13333					D13033	D13043	IGAIAN4			
1	38,38	15	11,15	13	29	24	11	11	13			
	14	11	10	19	33	16	15	11	23			
	42	12	*	20	18	16	24	*	13			
2	36,37	14	11,14	13	29	24	11	13	13			
	15	12	12	19	32	15	15	11	23			
L	37	12	*	18	18	19	23	*	12			
3	38,38	15	11,15	13	29	24	11	11	13			
	14	11	10	19	33	16	15	11	23			
	42	12	*	20	18	16	24	*	13			
4	40,40	15	14,17	13	29	23	10	11	12			
	15	9	12	21	32	16	16	10	22			
	37	11	*	17	19	21	21	*	11			

 $^{^{}st}$ Results were not received by a minimum of 10 participants for the loci indicated.

The information presented here is that received from the sample manufacturer. It presents details of the design specification for the test samples and/or details of how they were prepared. This information does not necessarily represent the answers that should or could be obtained from an examination of the sample. Final interpretation of the results should be deferred until the summary report is available.

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