FOR THE RECORD

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Population Data on Eight Short Tandem Repeat Loci in the Barbadian Population

POPULATION: Barbadian (*n* = 186).

KEYWORDS: forensic science, DNA typing, short tandem repeats, polymerase chain reaction, population genetics, Barbados, D16S539, D7S820, D13S317, D5S818, CSF1PO, TPOX, TH01, vWA

Whole blood drawn in EDTA vacutainer tubes from 200 unrelated individuals was kindly donated from the Queen Elizabeth Hospital blood bank and stored on Ultrastain Cards (Whatman[®] Inc., Newton Center, MA). DNA extraction was performed using the QIAamp DNA mini kit (Qiagen, Valencia, CA) dried blood spot protocol, according to the manufacturer's instructions. The Quantiblot[®] Human DNA Quantitation kit (Applied Biosys-

tems, Foster City, CA) was used to determine the quantity of extracted DNA, as per manufacturer's instructions. Amplification was performed using the reagents contained in the PowerPlex[®] 1.1 System (Promega, Madison WI), according to the manufacturer's instructions. Postelectrophoresis amplified products were detected using the Hitachi FMBIO[®] II Fluorescent Scanner (Hitachi Software Engineering Ltd., San Francisco, CA). Results were

TABLE 1—Allele frequency data on eight STR loci in a sample population from Barbados.

Allele	D16S539 $(n = 167)$	D7S820 ($n = 182$)	D13S317 ($n = 184$)	D5S818 $(n = 183)$	CSF1PO (<i>n</i> = 173)	$\begin{array}{c} \text{TPOX} \\ (n = 183) \end{array}$	TH01 $(n = 181)$	vWA $(n = 184)$
5	× ,	. ,	. ,	. ,	. ,	· /	0.000	. ,
5	_		_	_	0.003	0.00	0.009	_
0	_	0 000		0.006	0.005	0.09	0.109	
8	0.045	0.009	0 019	0.000	0.075	0.358	0.378	
9	0.186	0.110	0.017	0.000	0.075	0.330	0.202	
03	0.100	0.110	0.014	0.022	0.027	0.107	0.088	
10	0 177	0 363	0 035	0.087	0 251	0 082	0.000	
10	0.200	0.303	0.033	0.243	0.257	0.002	0.017	0 009
12	0.277	0.082	0.205	0.245	0.257	0.036		0.007
12	0.217	0.002	0.174	0.224	0.035	0.050		0 022
14	0.024	0.0157	0.073	0.014	0.009			0.022
15	0.024	_	0.073	0.014	0.007	_	_	0.171
16			0.003					0.285
17			0.005					0.193
18								0.153
19								0.065
20								0.030
20								0.009
MP	0.069	0.094	0.124	0.096	0.087	0.084	0.100	0.058
PD	0.931	0.906	0.876	0.904	0.913	0.916	0.900	0.942
PIC	0.77	0.72	0.68	0.72	0.75	0.74	0.72	0.79
PE	0.608	0.542	0.491	0.535	0.575	0.571	0.538	0.638
HWE	01000	01012	01191	0.000	01070	01071	0.000	01020
Exact test	0.050	0.735	0.134	0.790	0.342	0.275	0.370	0.629
γ^2	0.644	0.998	0.522	0.999	0.995	0.934	0.984	0.206
Homozygosity test	0.001	0.546	0.371	0.933	0.225	0.033	0.690	0.412

MP, match probability; PD, power of discrimination; PIC, polymorphism information content; PE, power of exclusion.

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analyzed using the STaR CallTM Allele Calling Software. The genotypes obtained from the STaR CallTM Software were imported into the Power Stats V12 program for each locus.

Possible divergence from Hardy–Weinberg expectations (HWE) was tested by calculating the unbiased estimate of the expected homozygote/heterozygote frequencies (1) and the exact test (2), based on 2,000 shuffling experiments. An interclass correlation criterion (3) for two-locus associations was used for detecting disequilibrium between the STR loci.

The program to perform these tests (DNA type) was developed and kindly provided by R. Chakraborty (University of Cincinnati). Table 1 displays the results of the study. Based on the exact test there were no detectable departures from HWE.

The complete data are available to any interested researcher upon request.

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