

## FOR THE RECORD

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# Population Data of ABO Gene Applicable to Human Identification in Koreans\*

**POPULATION:** A total of 100 unrelated healthy individuals living in Seoul, Korea

**KEYWORDS:** forensic science, ABO gene, human identification, Koreans

Genomic DNA was isolated from buccal swabs by the QIAamp DNA Mini Kit (QIAGEN). The primers used for amplification and sequencing of exon 6 and 7 of ABO gene were suggested by Lee and Chang (1), and Ogasawara et al. (2), respectively. About 1–2 ng of genomic DNA was used for PCR in 25  $\mu$ L reaction volume. PCR mixture contained 10 mM Tris-HCl pH 8.5, 50 mM KCl, 1.5 mM MgCl<sub>2</sub>, 200  $\mu$ M of each dNTP, 0.25  $\mu$ M of each primer, and 1 U of AmpliTaq Gold DNA Polymerase (Applied Biosystems). Thermal cycling was performed initially at 95°C for 10 min, then 35 cycles consisting of 95°C for 1 min, 60°C for 1 min for exon 6 and 63°C for 1 min for exon 7, 72°C for 1 min, followed by 10 min extension at 72°C in a GeneAmp PCR System 9600 (Perkin Elmer).

PCR products were purified by QIAquick PCR Purification kit (QIAGEN). The BigDye Terminator Cycle Sequencing kit (Applied Biosystems) and an ABI PRISM 310 Genetic Analyzer (Applied Biosystems) were used for direct DNA sequencing. Sequences were analyzed with Sequence Navigator software (Applied Biosys-

tems). ABO genotype was determined by comparing those with the reported nucleotide sequences of ABO alleles (3,4).

Statistical analysis was performed using the Genetic Data Analysis software, (<http://lewis.eeb.uconn.edu/lewishome/software.html>) and the PowerStat (<http://www.promega.com/geneticid-tools/powerstats/>) program of Promega Corporation. In case of mean exclusion chance (MEC), it was calculated according to Krüger et al. (5). We did not find significant deviation from Hardy-Weinberg equilibrium at ABO gene.

The complete data are available to any interested researcher upon request from the corresponding author.

## References

1. Lee JC, Chang JG. ABO genotyping by polymerase chain reaction. *J Forensic Sci* 1992;37:1269–75.
2. Ogasawara K, Bannai M, Saitou N, Yabe R, Nakata K, Takenaka M et al. Extensive polymorphism of ABO blood group gene: three major lineages of the alleles for the common ABO phenotypes. *Hum Genet* 1996; 97:777–83.
3. Yamamoto F. Molecular genetics of ABO. *Vox Sang* 2000;78:91–103.
4. Yip SP. Sequence variation at the human ABO locus. *Ann Hum Genet* 2002;66:1–27.
5. Krüger J, Fuhrmann W, Lichte KH, Steffens C. Zur Verwendung der sauren Erythrocytenphosphatase bei der Vaterschaftsbegutachtung. *Dtsch Z Gerichtl Med* 1968;64:127–46.

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TABLE 1—*Allele frequencies and forensic efficiency of ABO gene in Koreans.*

Allele	A101	A102	B101	B102	O01	O02	O04	Number	Frequency
A101	3	1	1	...	2	2	...	12	0.060
A102		7	10	...	10	9	...	44	0.220
B101			8	1	11	5	...	44	0.220
B102				...	...	...	...	1	0.005
O01					9	13	1	55	0.275
O02						6	1	42	0.210
O04							...	2	0.010
Observed heterozygosity			0.670						
Expected heterozygosity			0.784						
Polymorphism information content (PIC)			0.744						
Power of discrimination (PD)			0.915						
Mean exclusion chance (MEC)			0.564						