

Human apolipoprotein CI (ApoC1) gene locus: BglII dimorphic siteP.M.Frossard, D.W.Lim, R.T.Coleman, H.Funke¹, G.Assmann¹, M.J.Malloy² and J.P.Kane²

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SOURCE/DESCRIPTION 0.3kb human apolipoprotein CI cDNA fragment isolated from a human liver cDNA library and subcloned into the EcoRI site of pBR329 (1).

POLYMORPHISM BglII (GCCNNNNNGGC) (International Biotechnologies, Inc.) detects a single two-allele polymorphism with bands at either 6.2kb or 8.7kb.

FREQUENCY Studied in 61 unrelated Germans and 79 unrelated Northern Americans.

	Northern Americans	Germans
8.7kb allele	0.025±0.012	0.016±0.011
6.2kb allele	0.975±0.012	0.984±0.011

CHROMOSOMAL LOCALISATION Human apolipoprotein CI gene located on chromosome 19 by Southern blot hybridization of DNA from human-rodent somatic cell hybrids (2,3).

MENDELIAN INHERITANCE Co-dominant segregation observed in 3 families (19 individuals).

PROBE AVAILABILITY Requests for probe to P.F. at the above address.

OTHER COMMENTS Regular background under usual stringency conditions.

REFERENCES 1- Appleby,V.A. and Levy-Wilson,B. (unpublished)
 2- Knott,T.J. *et al.* (1984) *B.B.R.C.* 125, 299-306
 3- Tata,F. *et al.* (1985) *Hum. Genet.* 69, 345-349

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