Isolation and mapping of a polymorphic DNA sequence (pCMM65) on chromosome 16 [D16S84]

Y.Nakamura, C.Martin, K.Krapcho, P.O'Connell, M.Leppert, G.M.Lathrop, J.-M.Lalouel and R.White

The Howard Hughes Medical Institute, University of Utah Medical School, Salt Lake City, UT 84132, USA

SOURCE/DESCRIPTION: A 3.2 kb RsaI fragment from cosmid CMM65 isolated by myoglobin-2 oligonucleotide (1) was subcloned into the HincII site of pUCl8.

<u>POLYMORPHISM</u>: MspI identifies a 2 allele VNTR polymorphism with a band at 1.5 kb(M1) or 1.3 kb(M2) without a constant band. TaqI, RsaI, BglII, PstI, EcoRI, BamHI HindIII and PvuII also identify the same polymorphism.

FREQUECY: Estimated from 111 unrelated Caucasians.

M1: 0.37 M2: 0.63

NOT POLYMORPHIC FOR: Not known

CHROMOSOMAL LOCALIZATION: pcMM65 has been assigned to chromosome 16 by linkage analysis(2) with a locus (HBZP1) known to span this region.

MENDELIAN INHERITANCE: Co-dominant segregation of the MspI RFLP was observed in 56 three generation families.

PROBE AVAILABILITY: Contact Y.N.

OTHER COMMENTS: RFLPs were observed under normal hybridization and wash stringencies.

REFERENCES:

- 1. Y. Nakamura et al., Science 235: 1616-1622 (1987)
- 2. G.M. Lathrop et al., Am. J. Hum. Genet. 37:482-498 (1985)