Isolation and mapping of a polymorphic DNA sequence pMLJ14 on chromosome 14 [D14S13]

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SOURCE/DESCRIPTION: This clone is a cosmid pJB-8 derivative with a 40kb insert in the EcoRI site identified with the myoglobin oligonucleotide (1).

<u>POLYMORPHISM</u>: <u>Rsa</u>I optimally resolves a >20 allele VNTR polymorphism with bands between 4.0 - 15.0 kb. With <u>Rsa</u>I, numerous constant bands are observed below the size range of <u>Msp</u>I, <u>Tag</u>I, <u>Pst</u>I, <u>Bgl</u>II, <u>BamH</u>I, <u>Pvu</u>II, the polymorphism. HindIII, and EcoRI also detect the same polymorphism.

HETEROZYGOSITY: With RsaI, 95% heterozygosity was observed in 108 unrelated Caucasians.

pMLJ14 has been assigned to distal CHROMOSOMAL LOCALIZATION: chromosome 14q by multipoint linkage analysis(2) with loci (D14S1, PI, GM) known to span this region (3,4).

MENDELIAN INHERITIANCE: Co-dominant segregation of the VNTR polymorphism has been observed in 54 three generation families.

PROBE AVAILABILITY: Contact Y.N.

OTHER COMMENTS: RFLPs observed under normal stringeny. The whole cosmid can be used as a probe without pre-association with excess human DNA.

REFERENCES:

- Y. Nakamura et al., Science <u>235</u>: 1616-1622 (1987)
 G.M. Lathrop et al., Am. J. Hum. Genet. <u>37</u>:482-498 (1985) 3. D.R. Cox and T. Gedde-Dahl, Cytogenet. Cell Genet. 40:206-
- 242 (1985) 4. Y. Nakamura et al., abstract submitted to Human Gene Mapping Workshop 9. Cytogenet Cell Genet., in press