Isolation and mapping of a polymorphic DNA sequence (pEFZ31) on chromosome 22 [D22S32]

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

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

SOURCE/DESCRIPTION: A 3.1 kb MspI fragment from the cosmid EFZ31 was subcloned into the AccI site of pUC18.

POLYMORPHISM: MspI identifies a 2 allele polymorphism ( M1 : 3.5 kb, M2 : 2.6 kb).

FREQUENCY: Estimated from 48 unrelated Caucasians.

M1 : 0.60 MspI M2: 0.40

NOT POLYMORPHIC FOR: RsaI, BglII, PstI, PvuII, BamHI, HindIII and EcoRI

CHROMOSOMAL LOCALIZATION: This probe is assigned to chromosome 22 by linkage analysis (1) with a locus (D22S1) known to be on this chromosome (2).

MENDELIAN INHERITANCE: Co-dominant segregation has observed in 24 three generation families.

PROBE AVAILABILITY: Contact Y.N..

OTHER COMMENTS: RFLPs were observed under normal conditions of hybridization and washing.

## REFERENCES:

- 1. G.M. Lathrop et al., Am. J. Hum. Genet. 37:482-498 (1985)
  2. P. Tippett and J.-C. Kaplan, Cytognet. Cell Genet. 40:268-295(1985)