An anonymous DNA segment (Π 227) maps to the long arm of human chromosome 5 and identifies a BstXI polymorphism (D5S26)

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SOURCE/DESCRIPTION: Recovered from the library of Lawn et al. cloned into TTAN7. 900 bp HindIII/EcoR1 insert. POLYMORPHISM: BstXI allele 1:2.7kb; allele 2:2.3kb. Others for PstI (4.3 and 3.0); MboI (550bp and 450bp) and BcII (3.0 and 1.8, 1.2) are in disequilibrium with BstXI polymorphism. There are no constant bands. FREQUENCY: BstXI polymorphism: Allele 1, 0.3; Allele 2, 0.7 based on 34 chromosomes. NOT POLYMORPHIC FOR: ApaI, BamHI, BanII, BgII, BgIII, EcoRI, EcoRV, HaeII, HincII, HinfI, HindIII, KpnI, MspI, NciI, PvuII, RsaI, Sau96A, StyI, TaqI, XbaI, XmnI CHROMOSOMAL LOCALISATION: 5q by somatic cell hybrids, provisionally 5q14-5q32. MENDELIAN INHERITANCE: Yes, based on 3 single child families. PROBE AVAILABILITY: Freely available. OTHER COMMENTS: Gel conditions: BstXI and BclI -90V in 0.8% agarose -50V in 1.2% agarose MboI All 16 hours in TBE -70V in 0.8% agarose PstI **REFERENCE:** Lawn RM, Fritsch EH, Parker RC, Blake G, and Maniatis T. Cell 15:1157-1174 (1978). ACKNOWLEDGEMENTS: NIH HD20118 and Howard Hughes Medical Institute

