RFLP for a DNA clone which maps to 19q13.2 - 19qter (D19S63)

J.D.Brook, H.G.Harley, S.A.Rundle, K.V.Walsh and D.J.Shaw

Institute of Medical Genetics, University of Wales College of Medicine, Heath Park, Cardiff CF4 4XN, Wales, UK

Source/Description: pD10 is a 0.85kb fragment with BamHI ends cloned in pSP64. It was sub-cloned from $\lambda D10$ which was isolated from a genomic library constructed in \(\lambda EMBL3 \) from a rodent human somatic cell hybrid (Stallings et al., 1988).

Frequency: Studied in 80 unrelated individuals.

7.1kb = 0.58

6.8kb = 0.27

6.5kb = 0.15

Not Polymorphic For: BanI, BglII, DraI, HinfI, TaqI, XbaI.

Chromosomal Localisation: Probe localises to 19q13.2 - 19qter on a panel of rodent human somatic cell hybrids.

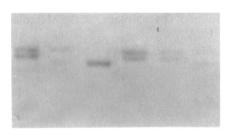
Mendelian Inheritance: Co-dominant segregation shown in 32 families, 280 individuals.

Probe Availability: Available for collaborative work on Myotonic Dystrophy. Freely available for other studies.

Other Comments: Wash to 0.1×SSC at 65°C for 30 mins. The probe also detects a 2 allele polymorphism with HincII.

References: Stallings et al. (1988) Am. J. Hum. Genet. 43, 144-151. Brook et al. (1987) Genomics 1, 320-328.

Acknowledgements: This work was supported by the Muscular Dystrophy Group of Great Britain and the Muscular Dystrophy Association, USA.



A HindIII polymorphism identified by a DNA clone which maps to chromosome 17 (D17S245)

J.D.Brook, M.Upadhyaya, W.Broadhead, S.A.Rundle, K.V.Walsh, H.G.Harlev and D.J.Shaw

Institute of Medical Genetics, University of Wales College of Medicine, Heath Park, Cardiff CF4 4XN. Wales, UK

Source/Description: $\lambda D13$ is an 18kb fragment with EcoRI ends cloned in \(\lambda EMBL3 \) from rodent human somatic cell hybrid DNA (Stallings et al. 1988).

Polymorphism: A polymorphism with two alleles at 4.2kb and 4.4kb is detected with HindIII.

Frequency: Studied in 28 unrelated individuals.

4.2kb = 0.5

4.4kb = 0.5

No other enzymes tested.

Chromosomal Localisation: Probe localises to chromosome 17 using a panel of rodent human somatic cell hybrids (Worwood et al. 1985).

Mendelian Inheritance: Co-dominant segregation shown in 15 families, 90 individuals.

Probe Availability: Freely available.

Other Comments: A 1.7kb fragment was used as probe following digestion of $\lambda D13$ with BamHI and HindIII. Filters were washed to 0.1×SSC at 65°C for 30 minutes.

References: Stallings et al. (1988) Am. J. Hum. Genet. 43, 141-151. Worwood et al. (1985) Hum. Genet. 69, 371-374.

