Satellite II DNA of human lymphocytes: tandem repeats of a simple sequence element

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Submitted November 10, 1987

Accession no. X06199

A 250bp fragment of satellite II DNA was cloned from sonicated and renatured (Cot=\(^1\)) human lymphocyte DNA. This clone (AB6) was sequenced using standard methods (1). The sequence contains multiple \(\frac{\text{Hinf1}}{\text{and}}\) and \(\frac{\text{Taq1}}{\text{restriction}}\) restriction endonuclease sites and corresponds to the simple sequence component (satellite 2) of classical satellite II DNA (2).

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CqTTt
GATTCCĂTTtGATGtT 21
GATTCCATTC
GAGTCCATTCGATGAT 47
a A T T C C A T T C
GATTCttTgCGAT---
GATTCCATŤC
ctTTCCATTtGA-GAT 95
GATTCCATTC
GAGACCATTCGAT---
GAŤTgCATTC
a A T T Č - A T T C G A T G A c
GATTCqATTC
a A T T C C g T T C a A T - - -
GATTCCĂTTC
GATTCCA a T t GATGAT ^{192}
GATTCCATTC
GATTCCATT t GATGAT ^{218}
GATTCCATgC
GATTCCATTCGATGAT
GACTCC
       250
```

tcqa

Taq1

The sequence presented here consists entirely of tandem repeats of a 26bp element which has been seen in other satellite II sequences such as the 48bp and 59bp genomic sequences of Prosser et al (2) and the pPD17 clone of Deininger et al (3). The 26bp repeating unit is formed from alternating 10bp and 16bp units which have common core consensus of (GATTCCATTC). This core is related to the 5bp repeat unit (ATTCC) of satellite 3 DNA and may be the original segment which was amplified to form the satellite II DNA.

The sequence shown is continuous and is aligned to show the repeating units. Highly conserved bases are indicated by capital letters and deleted bases are shown as hyphens. The <u>Hinfl</u> and <u>Taql</u> sites are shown below the sequence and the numbering refers only to bases sequenced.

REFERENCES

g a(n)t c

Hinf1

- 1) Hindley, J. (1982) <u>DNA Sequencing</u>, Laboratory Techniques in Biochemistry and Molecular Biology 10.
- Prosser, J., Frommer, M., Paul, C. and Vincent, P.C. (1986) <u>J. Mol. Biol.</u> 187, 145-155.
- 3) Deininger, P.L., Jolly, D.J., Rubin, C.M., Friedmann, T. and Schmid, C.W. (1981) J. Mol. Biol. 151, 17-33.