Supplementary materials.

I. Primers used in the study

All are listed 5' to 3':

TAP2-1-5' GAGAATCACTTGAACCTGGGAG TAP2-1-3' AAGGAAACCCCAGGAAGAAATA TAP2-2-5' TTGTCCACAGTGTACCACATGA TAP2-2-3' ATCTTGCAATTCAGCATGACAC TAP2-3-5' TATTTCTTCCTGGGGTTTCCTT TAP2-3-3' ACCACCTCCCAAACTAAAAACA TAP2-4-5' CATGATGTGTCATGCTGAATTG TAP2-4-3' CGGGCTCTCCTAAATAGAAGGT TAP2-5-5' ATAGAACAAGAACCAAAGCCCA TAP2-5-3' GGCGGTTACTCTGGGATATAAA TAP2-6-5' GGACAACAGATAAAGTTGCCCT TAP2-6-3' CTGTCAACGGATACGAGATGAG TAP2-7-5' AAGTGACTACCCACTCCACGCT TAP2-7-3' AGTAAAGCCGCGTCCACCAGC TAP2-8-5' CATCTCGTATCCGTTGACAGAG TAP2-8-3' AGGGCCTAGAAATGGAGTTAGG TAP2-9-5' CCTCACTATTCTGGTCGTGTGA TAP2-9-3' TGAAATGGTTTTCGAGGGTACT TAP2-10-5' GGTGAATGTAGTCACCATTCTG TAP2-10-3' ATGACTGCACCTAACACTGTGG TAP2-11-5' GATGGTGGACCCACCAGATTAT TAP2-12-5' GGCAGGGAATTTATTATTGCAG TAP2-12-3' CCTCTGAACATCCTCCTTCACT TAP2-13-5' CATCTCCCTCCCCTCTTATTCT TAP2-13-3' TATACCTTCTCCCCTAACGGCT TAP2-14-5' CGGATACCACCCTGATGAGTAA TAP2-14-3' GGTTGGGGGTAGTAGGCAGTTA TAP2-15-5' ATAGGAAGGGGAGATGATAGGC TAP2-15-3' TCTTTTGTAAATTGCCCGATCT TAP2-16-5' CTCCGTTGTACCTTCTGTCTCC TAP2-16-3' CAGAAAGTGGTGTTGATGAGGA TAP2-17-5' AGATCGGGCAATTTACAAAAGA TAP2-17-3' CTTGGATATAACACCAAACGCA TAP2-18-5' TCTTTGCCCATTTTCTGATTTT TAP2-18-3' TGTTCAAGGGGCTCTTTATAGC TAP2-19-5' CCTTCCTTCCTGATTCAGACAC TAP2-19-3' GAAGGCCACAAAGAAAAAGAGA TAP2-20-5' GCTATAAAGAGGCCCTTGAACA TAP2-20-3' CTCATACACATACACTACATGG

II. Polymorphism data used for analyses.

Labels refer to the western chimpanzees listed in Methods. The genotypes are given as pseudohaplotypes, *i.e.* the phase is unknown. Positions are given with respect to the sequence from Jeffreys *et al.* (2000), available at http://www.le.ac.uk/ge/ajj/tap2/SEQdata.html/.

Positions:

536 552 556 557 568 703 1114 1211 1613 1808 2090 2176 2417 2669 3271 3319 3491 3600 3804 3841 4006 4062 4075 4196 4274 4415 4451 4452 4466 4572 4613 4887 4927 5197 5503 5569 5775 5789 5854 5868 6342 6388 6796 7429 7978 8089 8113 8377 8621 8824 8974 9141 9159 9501 9649 9680 9759

> 311

ATACCCTGGAGGCAAGAATCTTCCGATAGACGCCAGTCCCTAGTTGTCCCGACCTC

A T A C C C T G G A G G C A A A A A T C T T C T G A T A G A C G C C A A T C C T A G T T G T C C T C G A C A T C

> 312

G C A C C C T G G A G G C A A G A G T C T T C C G A T A G A C G C C A G T C C C T A G T T G C C C C G A C C T C

ATGTCTTGAAAATAAAGATCTTTCGATAGACCTCAGTCCCTAGTTGTACCCGATATC

> 313

G C A C C C T G G A G G C A A G A G T C T T C C G A T A G A C G C C A G T C C C C A G T C C C C G G C C T C

ATGTCTTGAAAAAAAATCTTCCGATAGACCCCAGTCC

> 314

G C A C C C T G G A G G C A A G A G T C T T C C G A T A G A C C C C A G T C C C C A G T C

ATGTCTTGAAAAAAAATCTTCCGATAGACCCCAGTCC

> 317

> 320

A T G T T T T G A A A A T G A G A A T C T T C C A A T A T A T C C T A G T G T T C A A T T C T C C C A A C C A T

> 321

ATACCCTGGAGGCAAGAATCTTCCGATAGACGCCGGTCCCTAGTTGCCCCCGACCTC

A T A C C C T A G A G G C A A A A A T C T T C C G A T A G A C G C C A G T C C C T A G C T C C C C G A C A T C

> 322

G C A C C C T G G A G G C A A G G G T C T T C C G A T A G A C G C C G G T C C C T A G T T G C C C C G A C C T C

ATGTCTTGAAAATAAAATCTTCCGATAGACCCCAGTCC CTAGCTGTCCCCGACATC

> 325

ATACCCTGGAGGCAAGGATCTTCCGATAGACGCCAGTCCCTAGTTGTCCCGGCCTC

ATACCCTAGAGGCAAAAATCTTCTGATAGACGCCAGTC CCTAGTTGTCCTCAACAAT

> 326

G C A C C C T G G G G G C A A G A G T C T T C C G A T A G A C G C C A G T C C C C A G T C C C C G G C C T C

ATGTCTTAAAAATAAAGATCTTCCAATAGACCCCAGTCC CTAATTGTACCCAACAAT

> 328

ATACCCTGGAGGCGAGGACCCCCGGGGAGGCGCCAGCCCCCGGGTTGTCCCCGACCTC

> 329

ATACTCCGGAGGCAAGGACCCCCGGGAAGGCCAGCCCCCGGGAAGGCCAGCC

> Annaclara

G C A C C C T G G A G G C A A G A G T C T T C C G A T A G A C G C C A G T C C C C A G T C C C C G G C C T C

ATGTCTTAAAAAAAATCTTCCAATAGACCCCAGTCC CTAATTGTCCCCAACAAT

> Frits

ATACCCTGGAGGCAAAATCTTCCGATCGACGCCAGTC

A T A C C C T A G A G G C A A A A A T C T T C T G A T A G A C G C C A G T C C C T A G T C C T C G A C A T C

> Hilko

G C G T C T T G A A A A C A G G A G T C T T C C G A T A G A C C C C A G T C C C C A G T C

G C G T C T T G A A A A C A A G A G T C T T C C A A T A G A C C C C A G T C C C T A A T T G T C C C C A A T A A T

> Liesbetha

ATACCCTGGAGGCAAAATCTTCTGATAGACGCCAGTC CCTAGTTGTCCTCGACATC

ATACCCTGGAGGCAAAATCTTCTGATAGACGCCAGTC

> Louise

A T A C C C T G G A G G C A A A A A T C T T C C G A T A G A C G C C G G T C C C T A G T C C C C G A C A T C

A T A C C C T A G A G G C A A A A A T C T T C C G A T A G A C G C C A G T C C C T A G C T C C T C G A C A T C

> Marco

A T A C C C T G G A G G C A A A A A T C T T C C G A T C G A C G C C A G T C C C T A G T T G T C C C C G A C C T C

A T A C C C T G G A G G C A A A A A T C T T T T G A T A G A C G T C A G T C C C T A G T T G T A C T C G A C A T C

> Oscar

ATACCCTGGAGGCAAGGACCCCCGGGAAGGCCAGCCCCCGGGTCGTCCCCAGCC

A T A C T C T G G A G A T G A A A A T C T T C T A A T A T A T C C T A G T G T T T A G T T C T C T C G A C A A C

> Regina

ATACCCTGGAGGCGAGGACCCCCGGGGAGGCGCCAGCCCCCGGGTCGTCCCCGACCTC

A T A C T C T A G A G A T A A G A A T C T T C C A A T A T A T C C T A G T G T T T A G T T C T C C G A C A A C

> Socrates

G C A C C C T G G A G G C A A G A G T C T T C C G A T A G A C G C C A G T C C C T A G T T G T C C C C G A C A T C

ATGTCTTAAAAATAAAATCTTCCGATAGACCCCAGTCC

> Sonia

ATACCCTGGAGGCAAAATCTTCCGATAGACGCCGGTCCCTAGTTGTCCCGACCTC

A T A C C C T A G A G G C A A A A A T C T T C C G A T A G A C G C C A G T C C C T A G C T C C C C G A C A T C

> Yoran

GTACCCTGGAGGCAAGAGTCTTCCGATAGACGCCGGTC CCCAGTTGTCCCCGGCCTC

A C G T C T T G A A A A T A A A A A T C T T C C A A T A G A C C C C A G T C C C T A A C T G T C C C C A A C C A T

> Yvonne

A T G T T T T G A A A A T G A G A A T C T T C C A A T A T G T C C T A G T G T T C A A T T C T C T C C A A C C A T

III. Search for sequence motifs previously associated with recombination hotspots.

Unless indicated otherwise, all searches used the interface available from http://bioweb.pasteur.fr/seqanal/interfaces/scan_for_matches.html and were run with the following tolerance levels: 0 for a mismatch, 0 for an indel.

The locations of the motifs are indicated by []. Positions are given with respect to the sequence from Jeffreys *et al.* (2000), available at http://www.le.ac.uk/ge/ajj/tap2/SEQdata.html/.

In the consensus sequences, the following equivalencies are used:

```
R = (A,G)
```

Y = (T,C)

W = (A,T)

S = (G,C)

M = (A,C)

K = (G,T)

H = (A,T,C)

B = (G,C,T)

V = (G,A,C)

D = (G, A, T)

N = (A,G,C,T)

1) Search for motifs listed in Smith et al. (1998).

Possible location where Pur element promotes duplex opening: ATATATTTT

>humantap2:[5330,5338]

>chimptap2:[5330,5338]

 $(ATTTT)_n$

none

WAWTTDDWWWDHWGWHMAWTT

none

2) Search for additional motifs listed in Badge et al. (2000).

CHI: GCTGGTGG

>humantap2:[3527,3534]

>humantap2:[3794,3801]

>chimptap2:[3527,3534]

>chimptap2:[3794,3801]

ade6-M26 heptamer: ATGACGT

none

LTR-IS: TGGAAATCCCC

none

retroransposon LTR: TCATACACCACGCAGGGGTAGAGGACT none

XY32 homopurine-pyrimidine H-palindrome: AAGGGAGAARGGGTATAGGGRAAGAGGGAA none

human minisatellite core sequence: GGGCAGGARG GGGCAGGAAG:

>humantap2:[5547,5556] >chimptap2:[5547,5556]

human hypervariable minisatellite sequences: GGAGGTGGGCAGGARG,AGAGGTGGGCAGGTGG none

pur: GGNNGAGGGAGARRRR >humantap2:[8784,8799] GGAAGAGGGAGAGAAA

translin: GCNC[A/T][G/C][G/C][A/T] N(0-2) GCCC[A/T][G/C][G/C][A/T] -> GCNCWSSW N(0-2) GCCCWSSW GCNCWSSWGCCCWSSW GCNCWSSWNGCCCWSSW GCNCWSSWNNGCCCWSSW none

human replication origin consensus: WAWTTDDWWWDHWGWHMAWTT none

S.cerevisiae ARS: WTTTATRTTTW

none

S.pombe ARS: WRTTTATTTAW

none

scaffold attachment regions: AATAAAYAAA: none

TWWTDTTWWW or TTWTWTTWTT: TTTTTTTTT

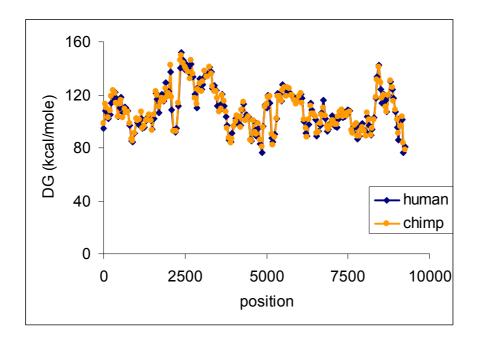
>humantap2:[2301,2310] >humantap2:[3201,3246] >humantap2:[3247,3256]

```
>chimptap2:[3201,3246]
>chimptap2:[8105,8114]
>chimptap2:[8278,8287]
WADAWAYAWW:
AAGAAACATT
>humantap2:[2749,2758]
>chimptap2:[2749,2758]
TAGAAACATA
>humantap2:[5323,5332]
>chimptap2:[5323,5332]
AAAATACAAA
>humantap2:[9736,9745]
>chimptap2:[9736,9745]
topoisomerase II binding site: GTNWAYATTNATNNR
none
3) Search for additional motifs listed in Petes (2001).
β hotspot:
(CCGNN)<sub>12</sub>:
none
(CGGATCCG)<sub>4</sub>:
none
GC content: human 46.8%, chimp 47.0%
(GT)_n:
none (no more than 3 repeats)
CoHR: -AG-T-GAACAA-ATAATC-AAAAAAAAAAAAAACTGT--C----C-A-----
Cf. Blumental-Perry et al. (2000)
none
4) Search for additional motifs listed in Wall et al. (2003).
(RY)_n:
```

WEB-THERMODYN: http://wings.buffalo.edu/gsa/dna/dk/ no difference

none (no more than 6 repeats)

Helical stability of the DNA sequence.



References:

Badge RM, Yardley J, Jeffreys AJ, Armour JA (2000) Crossover breakpoint mapping identifies a subtelomeric hotspot for male meiotic recombination. Hum Mol Genet 9:1239-1244

Blumental-Perry A, Zenvirth D, Klein S, Onn I, Simchen G (2000) DNA motif associated with meiotic double-strand break regions in *Saccharomyces cerevisiae*. EMBO Rep 1:232-238

Jeffreys AJ, Ritchie A, Neumann R (2000) High resolution analysis of haplotype diversity and meiotic crossover in the human TAP2 recombination hotspot. Hum Mol Genet 9:725-733

- Petes TD (2001) Meiotic recombination hot spots and cold spots. Nat Rev Genet 2:360-369
- Smith RA, Ho PJ, Clegg JB, Kidd JR, Thein SL (1998) Recombination breakpoints in the human beta-globin gene cluster. Blood 92:4415-4421
- Wall JD, Frisse LA, Hudson RR, Rienzo AD (2003) Comparative linkage disequilibrium analysis of the β-globin hotspot in primates. Am J Hum Genet 73:1330-1340.