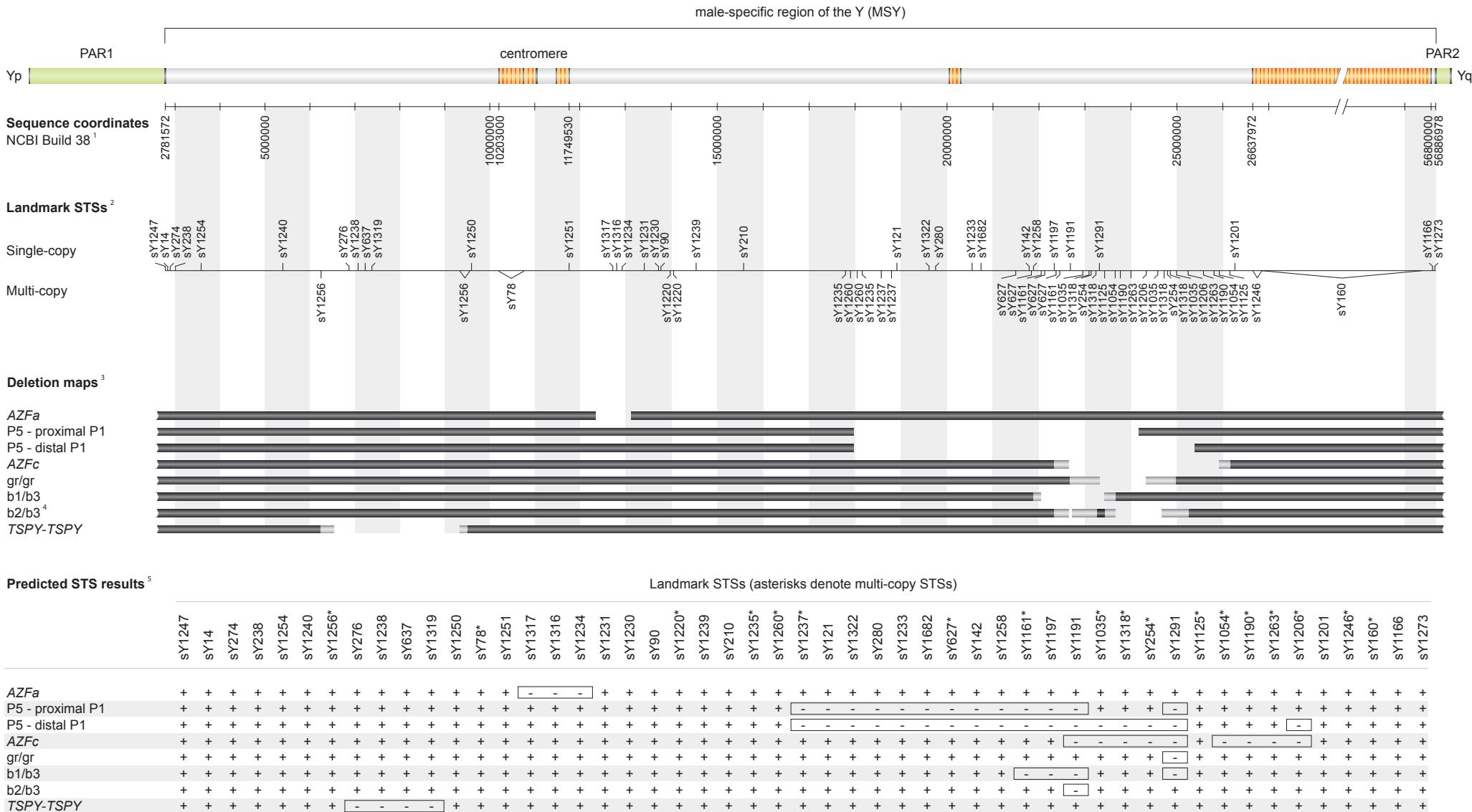


STS maps of common interstitial deletions

Reported in Sun et al., 2000; Kamp et al., 2000; Blanco et al., 2000 (AZFa), Repping et al., 2002 (P5 - proximal P1 and P5 - distal P1), Kuroda-Kawaguchi et al., 2001 (AZFc), Repping et al., 2003 (gr/gr and b1/b3), Fernandes et al., 2004; Repping et al., 2004 (b2/b3), Jobling et al., 2007 (TSPY-TSPY)

MSY Breakpoint Mapper

<http://breakpointmapper.wi.mit.edu>



Notes

¹ NCBI Build 38: GRCh38/hg38 (December 2013)

²Landmark STSs consist of sY1247 and sY1273, located at the boundaries between MSY and pseudoautosomal (PAR) sequences on Yp and Yq, respectively, and a previously published panel of 49 STS (see Supplementary Methods Table SM-2 in Repping et al., 2006).

³ For each deletion, solid black bars encompass STSs expected to be present, gray bars indicate breakpoint intervals that cannot be further narrowed due to cross-amplification at other loci.

⁴The b2/b3 deletion is contiguous but appears in the deletion map as shown because the deletion arises in Y chromosomes that carry an inversion in this region.

⁵Landmark STSs are shown in order of first appearance from left to right on the Y chromosome. Although each deletion is contiguous, multi-copy STSs can appear to be present within a deletion due to amplification from sites located outside that deletion.