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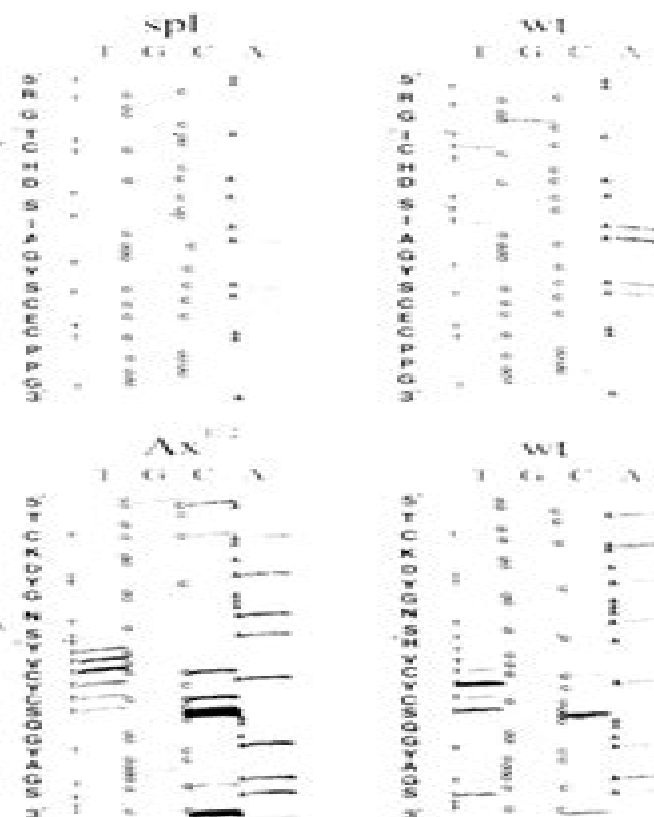


Fig. 6. (A) A diagrammatic representation of the 2703 amino acid putative Notch protein. The position of the *spl* (*spl*) mutation, a thymine to cytosine transition which causes the isoleucine at amino acid 578 in the 14th EGF-like repeat to change to a threonine, and the position of the *Abn* (*Abn*) mutation, a cytosine to thymine transition causing a histidine (1167 in the 29th EGF-like repeat) to change to a tyrosine are indicated. SP, signal peptide; TMD, transmembrane domain; P, A, nucleotide phosphate binding sequence homology; opa, opa repeat; EGF-like repeats, the epidermal growth factor-like repeats; Notch Repeats, cysteine-rich repeat present in Notch and lin-12. (B) Autoradiographs of sequencing gels showing part of the *spl* chromosome sequence, the *Abn* chromosome sequence and the corresponding wild-type (*wt*) chromosome sequence. (*) indicates the nucleotides or amino acids which differ between the mutant and wild-type genes. The autoradiograph of the *spl* sequence shows the nucleotide change responsible for the single amino acid alteration we detect and in addition a silent T-C transition. The single nucleotide change between 6 kb of wild-type and *Abn* sequence is shown in the autoradiograph to the right. The differences between the *spl* and wild-type sequence we have determined (Wharton *et al.*, 1985b) are as follows: 2474 (T-C), 2487 (T-C), 2889 (C-G), 2901 (T-C), 3447 (C-G), 4932 (C-T), 5307 (C-T), 5322 (C-T), 5454 (C-T), 5493 (A-G), 5712 (G-A), 5890 (C-A), 6000 (C-A), 6045 (A-G), 6096 (G-A), 6132 (C-T), 6453 (T-G), 6468 (T-A), 6489 (G-A). As a result of re-examining the autoradiographs of wild-type sequence corresponding to this region, we have discovered a number of errors in our published sequence (Wharton *et al.*, 1985b). They are at nucleotides 2811, 3360, 3616, 6654 and 6655. They change respectively an A to C, A to C, A to G, C to G and C to G. The latter four change the published amino acid sequence at 873 (Arg to Ser), 959 (Arg to Gly), 1971 (Phe to Leu) and 1972 (Ileu to Asp).

previously determined wild-type sequence (Wharton *et al.*, 1985b). Eighteen of these changes do not alter the deduced amino acid sequence (one results in the *Sa*I polymorphism we had

previously discovered). The 19th changes an isoleucine to a threonine at amino acid position 578, within the estimated limits of the *spl* mutation in the coding region. Figure 6 shows the

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