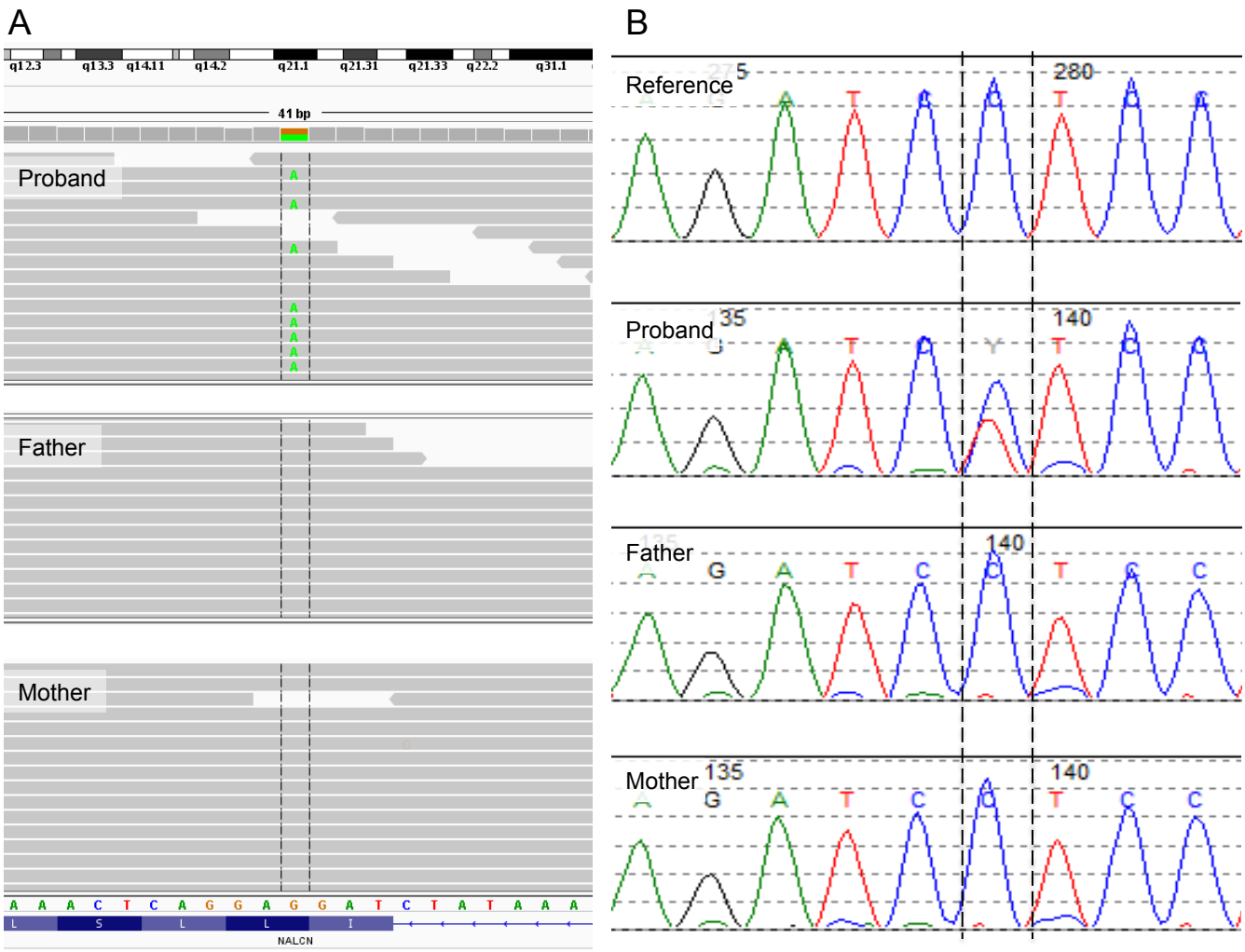

Figure e-1. Identification and confirmation of the *NALCN* variant. (A) Integrative Genomics Viewer (IGV) showing the chromosome region and NALCN exome sequencing results in the proband and the two unaffected parents. (B) Sanger sequencing shows the confirmation of the de novo change in *NALCN* (c.1768C>T; p.Leu590Phe).