Creation of clinical developmental labeling documents

Extensive proof reading and labeling specialist review reduced the error rates to less than 1%

Scope

- Creation of labeling documents for US, EU, ASEAN, LATAM, ROW Market (USPI, SmPC etc.)
- QC of labeling documents (involves review of editorial changes, compliance per guidance documents and formatting checks)
- **Supportive documents:** The labeling content development using Investigator's Brochure (IB), Core Safety Information (CSI), Target Product Profile (TPP), Development Product Label (DPP) provided by client etc.