



Fig. S3. Nucleotide and amino acid variant analysis of LAIR1-containing transcripts. (A) Outline of data processing. Paired-end reads mapping to *LAIR1* locus were extracted from samples and merged. Resulting reads were aligned to the consensus contig created by the main pipeline. Somatic hypermutations were assigned if a mismatch was present in more than 5% of reads. (B) Variants of *LAIR1* transcripts profiled for two donors of which one displayed LAIR1-containing antibodies in the serum, and another donor identified by suppression PCR (215). Two monoclonal cell lines (MMJ5, MGO3) expressing LAIR1-containing antibodies were used as controls. (C) Schematic representation of transcripts derived from monoclonal cell lines and donor 215. Black arrows indicate positions of nucleotide exchanges. The arrow in MMJ5 indicates a splice variant.