Supplemental Information

Identification and characterization of a novel *DGAT1* missense mutation associated with congenital diarrhea

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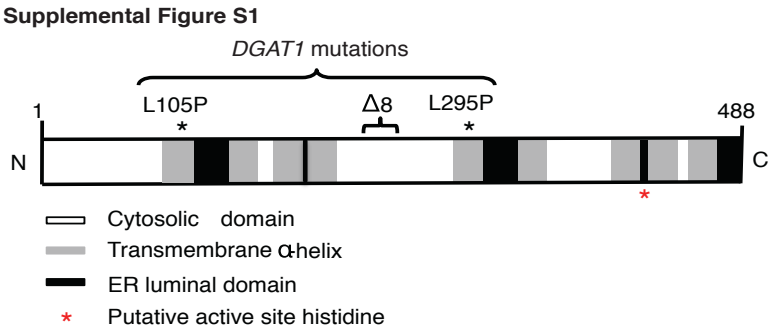

Supplemental Fig. S1. Schematic model of the DGAT1 protein showing nine predicted transmembrane domains, a histidine required for activity, and the location of amino acid changes due to the three human *DGAT1* mutations. Adapted from adapted from TMHMM Server v. 2.0 and McFie et al 2010. Δ8: chromosome 8 145541756 A->G.