Legends to Supplemental Figures

Figure S1: *SRp40 increases the PLP/DM20 ratio.* Representative RT-PCR analysis of PLP and DM20 products derived from PLP-neo and PLP-neo-ISEMT2 amplified from Oli-neu cell transfected with 0.5 µg DNA of either pSRp40 or pcDNA3. The PLP/DM20 ratio derived from PLPneo and PLP-neo-ISEMT2 was ~4-fold higher in SRp40 transfected Oli-neu cells. Western blot of cell lysates of Oli-neu cells co-transfected with plasmid expressing SRp40 and probed with anti-SRp40 antibody shows increased expression of SRp40 vs. control cells.

Figure S2: *hnRNPH and F regulate the PLP/DM20 ratio in non-glial cells*. Representative RT-PCR analysis of PLP-neo derived PLP and DM20 products amplified from RNA isolated from N2A (**A**) and L cells (**B**) treated with siH3, siF3, siH3+siF3 and siF/H (30 PCR cycles) (n=2). Mock are cells treated with scrambled siRNA.

Figure S3: *hnRNPF restores the* X_L/X_S *ratio derived from Bcl-x minigene after knock down of hnRNPH/F.* **A**. Schematic drawing of the Bcl-x minigene construct, showing the X_L and X_S 5' splice sites. The arrowheads indicate the position of the primers used for PCR amplification. **B**. Representative RT-PCR analysis of plasmid derived X_L and X_S products amplified from RNA isolated from untreated Oli-neu cells (lane 1), treated with siF/H (lane 2), treated with siF/H and transfected with pFlag-hnRNPF DNA (1 µg, lane 3) (PCR cycles= 30). The data are expressed as the X_L/X_S ratio.

Figure S4: *Expression of MS2hnRNPH and F fusion proteins*. Western blot analysis of cell lysates of Oli-neu cells co-transfected with plasmids expressing the fusion

proteins and probed with myc-tag antibody. The hnRNPH and F fusion proteins are expressed in similar amount.







В.

Α.

Bcl-x	+	+	+	-
siF/H	-	+	+	-
Flag-hnRNPF	-	-	+	-



pcDNA-MS2-H pcDNA-MS2-F

