

Supplementary information for
DDX39B promotes translation through regulation of
pre-ribosomal RNA levels

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Table S1. Primers used in qRT-PCR for various investigations

Gene Name	Sequence (5' to 3')	Investigation
DDX39B-FP	GAGCAAAGAGATCCGTCCAG	DDX39B mRNA level
DDX39B-RP	TTCCGGTTCTTCTCGTTGTC	DDX39B mRNA level
DDX49-FP	CTTCTTCTGGGAAGCACAGG	DDX49 mRNA level
DDX49-RP	TTCATCATGGAGTGCAGAGC	DDX49 mRNA level
GAPDH-FP	TCACCAGGGCTGCTTTTAAC	qPCR normalization
GAPDH-RP	TGACGGTGCCATGGAATTTG	qPCR normalization
47S-FP	CTGTCCTCTGGCGACCTG	47S rRNA level
47S-RP	GAGAGAACAGCAGGCCCG	47S rRNA level
H42-FP	GCACCGTTTGTGTGGGGTTGG	Chromatin Immunoprecipitation (ChIP)
H42-RP	CGAGACAGATCCGGCTGGCAG	
H0-FP	GGAGGTATATCTTTCGCTCCGAG	
H0-RP	GACGACAGGTCGCCAGAGGA	
H13-FP	ACCTGGCGCTAAACCATTTCGT	
H13-RP	GGACAAACCCTTGTGTCGAGG	
H18-FP	GTTGACGTACAGGGTGGACTG	
H18-RP	GGAAGTTGTCTTCACGCCTGA	
YY2-FP	TATAGCGGCTGCGAAAAGAT	
YY2-RP	CTTTGCCACATTCTGCACAT	
CALML3-FP	GGCCTTCTCCCTGTTTGAC	
CALML3-RP	TCCGTGTCCTTCATCTTCCT	

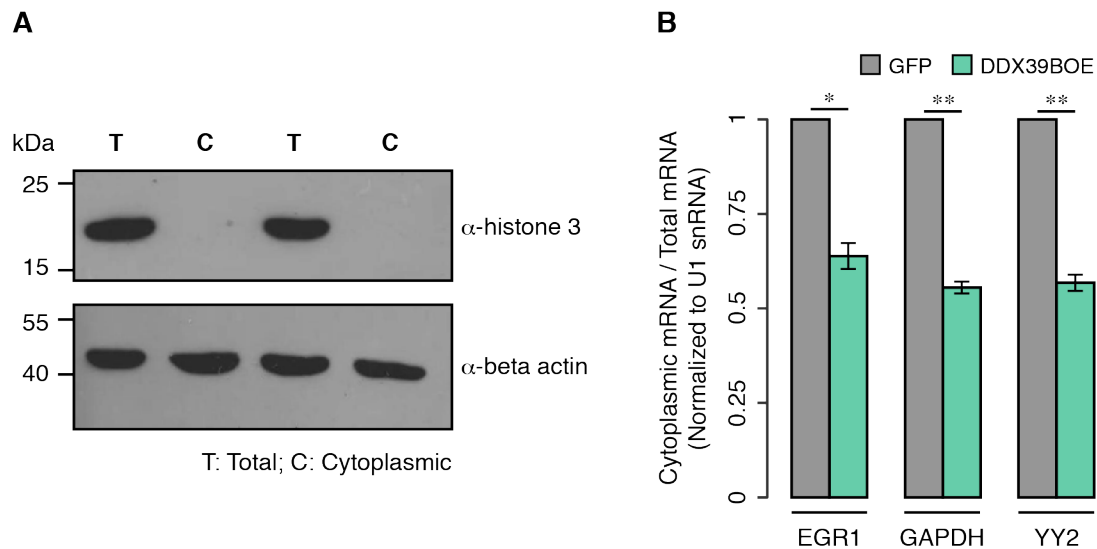


Figure S1: Overexpression of DDX39B inhibits mRNA export. HEK293 cells were transfected with pEGFP-C1 vector or pEGFP-DDX39B and cytoplasmic RNA and total RNA were isolated after 60 hours of transfection. **(A)** Western blot analysis of total (T) and cytoplasmic (C) extracts, used for RNA isolation. The immunoblotting was performed with histone 3 (nuclear protein) or beta actin antibody. The absence of histone 3 signal in cytoplasmic extracts confirms that there is no nuclear leakage in the cytoplasmic fractions. **(B)** Subcellular distribution of mRNAs of the indicated genes was quantified using qRT-PCR. The ratios of cytoplasmic to total mRNA were normalized to U1 small nuclear RNA (snRNA) levels and are presented relative to the control sample. Data are represented as mean of three independent experiments, with error bars representing standard deviations. The statistical significance was assessed by two tailed t-Test: paired two samples for means. * represents P-value < 0.05 and ** P-value < 0.01.

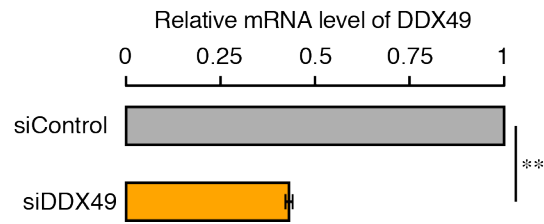


Figure S2: Knockdown efficiency of DDX49 siRNA. HEK293 cells were transfected with control siRNA or DDX49 and transcript levels of DDX49 were quantified using qRT-PCR. DDX49 transcripts levels were normalized to GAPDH expression and are presented relative to the control sample. Data are represented as mean of three independent experiments, with error bars representing standard deviation. Statistical significance was assessed by two tailed t-Test: paired two samples for means. ** represents P-value < 0.01.