

Table S3: Human orthologs and biomedical relevance of characterized *Drosophila* genes.

Genes	Biological process	Ct and/or Kn	Human ortholog; Biomedical relevance	References
<i>RpL7</i> <i>RpL36A</i>	ribosomal regulatory function	Ct/Kn	<i>RpL7</i> <i>RpL36A</i> Autism; ribosomopathies	Klauck et al. 2006; Narla & Ebert, 2010; Armistead & Triggs-Raine, 2014; Zhou et al. 2015
<i>dmn</i>	neurogenesis, microtubule-based transport	Ct	<i>DCTN2</i> ; exacerbate tauopathies (e.g. Alzheimer disease)	Butzlaff et al. 2015
<i>T-cpl</i> <i>CCT2</i>	chaperonin activity	Kn	<i>TCPI</i> <i>CCT2</i> Huntington disease; Gaucher disease; hypoparathyroidism-retardation-Dysmorphism (HRD) syndrome; Kenny-affey syndromes	Tam et al 2006; Parvari et al. 2002
<i>wdb</i>	cytoskeletal regulation, autophagy	Ct/Kn	<i>PPP2R5D</i> ; autosomal dominant mental retardation	Deciphering Developmental Disorders Study, 2015; Shulman & Feany, 2003; Hannan et al. 2016
<i>Ank2</i>	cytoskeletal regulation	Kn	<i>Ank3</i> ; autosomal recessive mental retardation	Iqbal et al 2013; Bi et al. 2012

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