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The nob2 mouse, a null mutation in Cacna1f: anatomical and functional abnormalities in the outer retina and their consequences on ganglion cell visual respons

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#	Paper	IF	Citations
190	Ribbon synapses of the retina. 2006 , 326, 339-46		108
189	Reduced synaptic vesicle density and aberrant synaptic localization caused by a splice site mutation in the Rs1h gene. <i>Visual Neuroscience</i> , 2006 , 23, 887-98	1.7	19
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187	Molecular genetics and protein function involved in nocturnal vision. 2007 , 2, 467-485		33
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