

Supplementary Table 1: Overview of DVL knock-out mouse phenotypes

Genotype \ Phenotype	<i>DVL1</i> ^{-/-}	<i>DVL2</i> ^{-/-}	<i>DVL3</i> ^{-/-}	<i>DVL1</i> ^{-/-} ; <i>DVL2</i> ^{-/-}	<i>DVL2</i> ^{+/-} ; <i>DVL3</i> ^{-/-}	<i>DVL2</i> ^{-/-} ; <i>DVL3</i> ^{+/-}	<i>DVL2</i> ^{-/-} ; <i>DVL3</i> ^{-/-}	<i>DVL1</i> ^{-/-} ; <i>DVL3</i> ^{-/-}	<i>DVL1</i> ^{-/-} ; <i>DVL2</i> ^{+/-} ; <i>DVL3</i> ^{-/-}	<i>DVL1</i> ^{-/-} ; <i>DVL2</i> ^{-/-} ; <i>DVL3</i> ^{+/-}	<i>DVL1</i> ^{-/-} ; <i>DVL2</i> ^{-/-} ; <i>DVL3</i> ^{-/-}
abnormal social behavior	+										
sensorimotor gating defects	+										
neuronal differentiation and function	+										
congenital heart defects		+	+	+	+	+					
skeletal defects posterior					+	+					
skeletal defects vertebrae and ribs		+		+							
neural tube defects		+ (2-3%)		+ (>90%)	+	+					
PCP defect in organ of Corti			+	+	+						
Nodal cilia defects									+	+	
cilia defects in ependymal cells (hydrocephalus)										+	(cond.)
gastrulation defects											+
anterior-posterior shortening				+					+		
embryonic lethal						+	(E9.5)	+	(E8.5)	+	(E13.5-15.5)

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