

An AKT3-FOXP1-reelin network underlies defective migration and cortical malformations of cortical development

Nature Medicine

21, 1445-1454

DOI: [10.1038/nm.3982](https://doi.org/10.1038/nm.3982)

Citation Report

#	ARTICLE	IF	CITATIONS
1	Sending Mixed Signals: The Expanding Role of Molecular Cascade Mutations in Malformations of Cortical Development and Epilepsy. <i>Epilepsy Currents</i> , 2016, 16, 158-163.	0.4	5
2	Canonical and Non-canonical Reelin Signaling. <i>Frontiers in Cellular Neuroscience</i> , 2016, 10, 166.	1.8	89
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6	Identification of Gene Loci That Overlap Between Schizophrenia and Educational Attainment. <i>Schizophrenia Bulletin</i> , 2017, 43, sbw085.	2.3	56
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8	Focal Cortical Dysplasia: Gene Mutations, Cell Signaling, and Therapeutic Implications. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2017, 12, 547-571.	9.6	98
9	AKT2 Regulates Pulmonary Inflammation and Fibrosis via Modulating Macrophage Activation. <i>Journal of Immunology</i> , 2017, 198, 4470-4480.	0.4	62
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18	Evolutionary conservation and conversion of Foxg1 function in brain development. <i>Development Growth and Differentiation</i> , 2017, 59, 258-269.	0.6	77

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39	Malformations of Cerebral Cortex Development: Molecules and Mechanisms. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2019, 14, 293-318.	9.6	71
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