Inherited disorders of gamma-aminobutyric acid metabolism and advances in ALDH5A1 mutation identificatio

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#	Paper	IF	Citations
31	mTOR, Autophagy, Aminoacidopathies, and Human Genetic Disorders. **2016**, 143-166		0
30	Succinic semialdehyde dehydrogenase deficiency (SSADHD): Pathophysiological complexity and multifactorial trait associations in a rare monogenic disorder of GABA metabolism. *Neurochemistry International*, **2016**, 99, 72-84	4.4	41
29	mTOR inhibitors rescue premature lethality and attenuate dysregulation of GABAergic/glutamatergic transcription in murine succinate semialdehyde dehydrogenase deficiency (SSADHD), a disorder of GABA metabolism. *Journal of Inherited Metabolic Disease*, **2016**, 39, 877-886	5.4	19
28	Acute Infantile Encephalopathy as Presentation of Succinic Semialdehyde Dehydrogenase Deficiency. *Pediatric Neurology*, **2016**, 58, 113-5	2.9	5
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2	Gamma-Aminobutyric Acid Transaminase (GABA-T) Deficiency in a Consanguineous Saudi Family: A Case Report and Literature Review.		0
1	The presence and severity of epilepsy coincide with reduced aminobutyrate and cortical excitatory markers in succinic semialdehyde dehydrogenase deficiency.		0