

研究成果の刊行に関する一覧表レイアウト

書籍

著者氏名	論文タイトル名	書籍全体の編集者名	書籍名	出版社名	出版地	出版年	ページ
なし							

雑誌

発表者氏名	論文タイトル名	発表誌名	巻号	ページ	出版年
Ieda D, Negishi Y, Miyamoto T, Johmura Y, Kumamoto N, Kato K, Miyoshi I, Nakanishi M, Ugawa S, Oishi H, <u>Saitoh S</u>	Two mouse models carrying truncating mutations in Magel2 show distinct phenotypes.	PLoS One	15	e0237814	2020
Hara-Isono K, Matsubara K, Fukube T, Yamazawa K, Satou K, Murakami N, <u>Saitoh S</u> , Nakabayashi K, Hata K, Ogata T, Fukami M, Kagami M.	Genome-wide methylation analysis in Silveira-Russell syndrome, Temple syndrome, and Prader-Willi syndrome.	Clin Epigenetics	12	159	2020
Isobe K, Ieda D, Miya F, Miyachi R, Otsuji S, Asai M, Tsunoda T, Kosaki K, Hattori A, <u>Saitoh S</u> , Mizuno M.	Hemorrhagic shock and encephalopathy syndrome in a patient with a de novo heterozygous variant in KIF1A.	Brain Dev	44	249-253	2022
Hayakawa K, Kawase K, Fujimoto M, Nakamura Y, <u>Saitoh S</u> .	Utility of breakpoint-specific nested polymerase chain reaction for the diagnosis of Emanuel syndrome.	Pediatr Int	63	1534-1536	2021
Negishi Y, Aoki Y, Itomi K, Yasuda K, Taniguchi H, Ishida A, Arakawa T, Miyamoto S, Nakashima M, Saitsu H, <u>Saitoh S</u> .	SCN8A-related developmental and epileptic encephalopathy with ictal asystole requiring cardiac pacemaker implantation.	Brain Dev	43	804-808	2021

Ohashi K, Fukuhara S, Miyachi T, Asai T, Imaeda M, Goto M, Kurokawa Y, Anzai T, Tsurusaki Y, Miyake N, Matsumoto N, Yamagata T, <u>Saitoh S.</u>	Comprehensive Genetic Analysis of Non-syndromic Autism Spectrum Disorder in Clinical Settings.	J Autism Dev Disord	51	4655-4662	2021
Hori I, Ieda D, Ito S, Ebe S, Nakamura Y, Ohashi K, Aoyama K, Hattori A, Kokubo M, <u>Saitoh S.</u>	Peripheral nerves are involved in hypomyelinating leukodystrophy-3 caused by a homozygous AIMP1 variant.	Brain Dev	43	590-595	2021