Daisuke Sato

List of Publications by Year
in descending order

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27 papers	4,203 citations	567281 15 h-index	526287 27 g-index
28 all docs	28 docs citations	28 times ranked	7677 citing authors

#	Article	IF	Citations
1	Metastatic pulmonary nodule after a seventeenâ€year diseaseâ€free interval resected through thoracoscopic subsegmentectomy: A case report. Thoracic Cancer, 2022, , .	1.9	1
2	Mediastinal mature teratoma with chest pain onset and subsequent perforation: A case report. International Journal of Surgery Case Reports, 2021, 81, 105807.	0.6	4
3	Optimal timing of echocardiography for heart failure inpatients in Japanese institutions: OPTIMAL Study. ESC Heart Failure, 2020, 7, 4213-4221.	3.1	4
4	Evaluation of echocardiographic parameters in Japanese patients agedâ€‰overâ€‰90Âyears at a single institution. Journal of Medical Ultrasonics (2001), 2020, 47, 555-563.	1.3	0
5	Effectiveness and safety of oral direct factor Xa inhibitors for the treatment of venous thromboembolism in patients with cancer and/or older age. Heart and Vessels, 2019, 34, 678-687.	1.2	3
6	Comparison of the effects of edoxaban, an oral direct factor Xa inhibitor, on venous thromboembolism between patients with and without cancer. Journal of Cardiology, 2018, 72, 120-127.	1.9	7
7	Synthesis, structure-activity relationships, and mechanistic studies of 5-arylazo-tropolone derivatives as novel xanthine oxidase (XO) inhibitors. Bioorganic and Medicinal Chemistry, 2018, 26, 536-542.	3.0	13
8	A cross-sectional survey of methods for controling hand-foot syndrome in patients receiving capecitabine treatment. Molecular and Clinical Oncology, 2018, 9, 443-448.	1.0	4
9	Dual emissive bispyrene peptide probes for highly sensitive measurements of trypsin activity and evaluation of trypsin inhibitors. Bioorganic and Medicinal Chemistry, 2018, 26, 3468-3473.	3.0	10
10	SMCHD1 mutations associated with a rare muscular dystrophy can also cause isolated arhinia and Bosma arhinia microphthalmia syndrome. Nature Genetics, 2017, 49, 238-248.	21.4	131
11	Efficient near infrared fluorescence detection of elastase enzyme using peptide-bound unsymmetrical squaraine dye. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 4024-4029.	2.2	10
12	Structureâ€“activity relations of rosmarinic acid derivatives for the amyloid Î² aggregation inhibition and antioxidant properties. European Journal of Medicinal Chemistry, 2017, 138, 1066-1075.	5.5	51
13	Radical pulmonary resection for metastatic lung tumor from hepatocellular carcinoma. The Journal of the Japanese Association for Chest Surgery, 2017, 31, 439-445.	0.0	1
14	Novel fluorescent substrates for detection of trypsin activity and inhibitor screening by self-quenching. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 5736-5740.	2.2	16
15	Meta-analysis of SHANK Mutations in Autism Spectrum Disorders: A Gradient of Severity in Cognitive Impairments. PLoS Genetics, 2014, 10, e1004580.	3.5	501
16	A genome-wide copy number association study of osteoporotic fractures points to the 6p25.1 locus. Journal of Medical Genetics, 2014, 51, 122-131.	3.2	36
17	Rare exonic deletions implicate the synaptic organizer Gephyrin (GPHN) in risk for autism, schizophrenia and seizures. Human Molecular Genetics, 2013, 22, 2055-2066.	2.9	139
18	A Discovery Resource of Rare Copy Number Variations in Individuals with Autism Spectrum Disorder. G3: Genes, Genomes, Genetics, 2012, 2, 1665-1685.	1.8	175

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19	Rare Deletions at the Neurexin 3 Locus in Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 133-141.	6.2	182
20	SHANK1 Deletions in Males with Autism Spectrum Disorder. American Journal of Human Genetics, 2012, 90, 879-887.	6.2	292
21	Culprit segments identified by optical coherence tomography in patients with acute myocardial infarction: two case reports. Cardiovascular Intervention and Therapeutics, 2012, 27, 47-51.	2.3	1
22	Alleviation of cisplatin-induced acute kidney injury using phytochemical polyphenols is accompanied by reduced accumulation of indoxyl sulfate in rats. Clinical and Experimental Nephrology, 2011, 15, 820-830.	1.6	27
23	Functional impact of global rare copy number variation in autism spectrum disorders. Nature, 2010, 466, 368-372.	27.8	1,803
24	A genome-wide scan for common alleles affecting risk for autism. Human Molecular Genetics, 2010, 19, 4072-4082.	2.9	538
25	Disruption at the PTCHD1 Locus on Xp22.11 in Autism Spectrum Disorder and Intellectual Disability. Science Translational Medicine, 2010, 2, 49ra68.	12.4	178
26	A ZRS duplication causes syndactyly type IV with tibial hypoplasia. American Journal of Medical Genetics, Part A, 2009, 149A, 816-818.	1.2	35
27	A syndactyly type IV locus maps to 7q36. Journal of Human Genetics, 2007, 52, 561-564.	2.3	35