Tristan Hardy

List of Publications by Year
in descending order

13	169	1478505 6	1281871 11
papers	citations	h-index	g-index

14	14	14	300
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Gene selection for the Australian Reproductive Genetic Carrier Screening Project (â€œMackenzieâ€™s) Tj ETQq1 1 0.784314 rgBT /Ove	2.8	60
2	Isolation of Circulating Fetal Trophoblasts Using Inertial Microfluidics for Noninvasive Prenatal Testing. Advanced Materials Technologies, 2018, 3, 1800066.	5.8	32
3	A Reappraisal of Circulating Fetal Cell Noninvasive Prenatal Testing. Trends in Biotechnology, 2019, 37, 632-644.	9.3	21
4	The role of prenatal diagnosis following preimplantation genetic testing for singleâ€gene conditions: A historical overview of evolving technologies and clinical practice. Prenatal Diagnosis, 2020, 40, 647-651.	2.3	18
5	Paternal mosaicism for a novel <scp>PBX1</scp> mutation associated with recurrent perinatal death: Phenotypic expansion of the <scp>PBX1</scp>â€related syndrome. American Journal of Medical Genetics, Part A, 2020, 182, 1273-1277.	1.2	12
6	Aberrant Splicing of SDHC in Families With Unexplained Succinate Dehydrogenase-Deficient Paragangliomas. Journal of the Endocrine Society, 2020, 4, bvaa071.	0.2	9
7	Genetic Testing in Endocrinology. Clinical Biochemist Reviews, 2018, 39, 17-28.	3.3	7
8	Preimplantation Genetic Testing for Monogenic Conditions: Is Cell-Free DNA Testing the Next Step?. Molecular Diagnosis and Therapy, 2021, 25, 683-690.	3.8	4
9	Concurrent maternal malignancy and fetal trisomy detected using genomeâ€wide noninvasive prenatal screening. Prenatal Diagnosis, 2021, 41, 1273-1276.	2.3	2
10	Higher Doses of FSH Used for Superovulation Do Not Adversely Affect Embryonic Ploidy: A Randomized Controlled Trial (STimulation Resulting in Embryonic Aneuploidy using Menopur) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 377	0.2	2
11	Recurrent pneumothorax in a case of t enascinâ€X deficient Ehlersâ€“Danlos syndrome: Broadening the phenotypic spectrum. American Journal of Medical Genetics, Part A, 2022, , .	1.2	2
12	The next generation of noninvasive fetal genotyping and the return to single gene conditions. BJOG: an International Journal of Obstetrics and Gynaecology, 2019, 126, 1475-1475.	2.3	0
13	OR34-6 A Novel Mechanism of SDH-Deficient Tumorigenesis and Implications for Genetic Testing in Patients with Pheochromocytoma-Paraganglioma. Journal of the Endocrine Society, 2019, 3, .	0.2	0