# 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 | 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 |
| 2 | Gene selection for the Australian Reproductive Genetic Carrier Screening Project (â€œMackenzieâ€™s) Tj ETQq0 0 0 rgBT /Overlock 10 | 2.8 | 60 |
| 3 | Concurrent maternal malignancy and fetal trisomy detected using genomeâ€wide noninvasive prenatal screening. Prenatal Diagnosis, 2021, 41, 1273-1276. | 2.3 | 2 |
| 4 | 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 |
| 5 | 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 ETQq1 1 0.784314 rgBT /Overlock 10 T | | 2 |
| 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 | 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 |
| 8 | 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 |
| 9 | 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 |
| 10 | A Reappraisal of Circulating Fetal Cell Noninvasive Prenatal Testing. Trends in Biotechnology, 2019, 37, 632-644. | 9.3 | 21 |
| 11 | 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 |
| 12 | Isolation of Circulating Fetal Trophoblasts Using Inertial Microfluidics for Noninvasive Prenatal Testing. Advanced Materials Technologies, 2018, 3, 1800066. | 5.8 | 32 |
| 13 | Genetic Testing in Endocrinology. Clinical Biochemist Reviews, 2018, 39, 17-28. | 3.3 | 7 |