

Marjorie Whitfield

List of Publications by Year in descending order

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Version: 2024-02-01

14
papers

582
citations

933447

10
h-index

996975

15
g-index

15
all docs

15
docs citations

15
times ranked

649
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Sperm Ion Transporters and Channels in Human Asthenozoospermia: Genetic Etiology, Lessons from Animal Models, and Clinical Perspectives. <i>International Journal of Molecular Sciences</i> , 2022, 23, 3926. | 4.1 | 11 |
| 2 | The sodium/proton exchanger <scp>SLC9C1</scp> (<scp>sNHE</scp>) is essential for human sperm motility and fertility. <i>Clinical Genetics</i> , 2021, 99, 684-693. | 2.0 | 26 |
| 3 | Tubulin glycylation controls axonemal dynein activity, flagellar beat, and male fertility. <i>Science</i> , 2021, 371, . | 12.6 | 84 |
| 4 | Identification and Characterization of the Most Common Genetic Variant Responsible for Acephalic Spermatozoa Syndrome in Men Originating from North Africa. <i>International Journal of Molecular Sciences</i> , 2021, 22, 2187. | 4.1 | 5 |
| 5 | A missense mutation in IFT74, encoding for an essential component for intraflagellar transport of Tubulin, causes asthenozoospermia and male infertility without clinical signs of Bardetâ€Biedl syndrome. <i>Human Genetics</i> , 2021, 140, 1031-1043. | 3.8 | 20 |
| 6 | Bi-allelic truncating variants in CFAP206 cause male infertility in human and mouse. <i>Human Genetics</i> , 2021, 140, 1367-1377. | 3.8 | 23 |
| 7 | Bi-allelic DNAH8 Variants Lead to Multiple Morphological Abnormalities of the Sperm Flagella and Primary Male Infertility. <i>American Journal of Human Genetics</i> , 2020, 107, 330-341. | 6.2 | 111 |
| 8 | TTC12 Loss-of-Function Mutations Cause Primary Ciliary Dyskinesia and Unveil Distinct Dynein Assembly Mechanisms in Motile Cilia Versus Flagella. <i>American Journal of Human Genetics</i> , 2020, 106, 153-169. | 6.2 | 46 |
| 9 | Mutations in TTC29, Encoding an Evolutionarily Conserved Axonemal Protein, Result in Asthenozoospermia and Male Infertility. <i>American Journal of Human Genetics</i> , 2019, 105, 1148-1167. | 6.2 | 44 |
| 10 | Mutations in DNAH17, Encoding a Sperm-Specific Axonemal Outer Dynein Arm Heavy Chain, Cause Isolated Male Infertility Due to Asthenozoospermia. <i>American Journal of Human Genetics</i> , 2019, 105, 198-212. | 6.2 | 116 |
| 11 | Comprehensive overview of murine epididymal mononuclear phagocytes and lymphocytes: Unexpected populations arise. <i>Journal of Reproductive Immunology</i> , 2018, 126, 11-17. | 1.9 | 39 |
| 12 | Slc26a3 deficiency is associated with epididymis dysplasia and impaired sperm fertilization potential in the mouse. <i>Molecular Reproduction and Development</i> , 2018, 85, 682-695. | 2.0 | 21 |
| 13 | Liver X Receptors (LXRs) Alpha and Beta Play Distinct Roles in the Mouse Epididymis1. <i>Biology of Reproduction</i> , 2016, 94, 55. | 2.7 | 7 |
| 14 | Posttesticular sperm maturation, infertility, and hypercholesterolemia. <i>Asian Journal of Andrology</i> , 2015, 17, 742. | 1.6 | 28 |