Saturnino LujÃ;n

List of Publications by Year in descending order

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1306789 1199166 14 149 7 12 citations g-index h-index papers 16 16 16 213 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Sperm DNA Methylation Epimutation Biomarkers for Male Infertility and FSH Therapeutic Responsiveness. Scientific Reports, 2019, 9, 16786.	1.6	53
2	Evaluation of Adult Acute Scrotum in the Emergency Room: Clinical Characteristics, Diagnosis, Management, and Costs. Urology, 2016, 94, 36-41.	0.5	20
3	Effect and in silico characterization of genetic variants associated with severe spermatogenic disorders in a large Iberian cohort. Andrology, 2021, 9, 1151-1165.	1.9	12
4	Are Urology Residents Ready to Treat Premature Ejaculation After Their Training?. Journal of Sexual Medicine, 2012, 9, 404-410.	0.3	10
5	Local Treatment of Penile Prosthesis Infection as Alternative to Immediate Salvage Surgery. Sexual Medicine, 2016, 4, e255-e258.	0.9	10
6	Evaluation of Male Fertility-Associated Loci in a European Population of Patients with Severe Spermatogenic Impairment. Journal of Personalized Medicine, 2021, 11, 22.	1,1	10
7	Intronic variation of the SOHLH2 gene confers risk to male reproductive impairment. Fertility and Sterility, 2020, 114, 398-406.	0.5	9
8	Polyorchidism: the case in a young male and review of the literature. Asian Journal of Andrology, 2015, 17, 511.	0.8	8
9	Common genetic variation in <i>KATNAL1</i> nonâ€coding regions is involved in the susceptibility to severe phenotypes of male infertility. Andrology, 2022, 10, 1339-1350.	1.9	5
10	Protocol for developing a core outcome set for male infertility research: an international consensus development study. Human Reproduction Open, 2022, 2022, hoac014.	2.3	4
11	Long-standing hemospermia in a patient with megacava associated to a circumaortic renal vein. Urology Annals, 2015, 7, 405-7.	0.3	2
12	Paternal Age and Fertility Concerns. , 2018, , 301-306.		1
13	Ejaculatory Dysfunction: Evaluation and Pathophysiology. , 2018, , 177-183.		0
14	Common Variation in the PIN1 Locus Increases the Genetic Risk to Suffer from Sertoli Cell-Only Syndrome. Journal of Personalized Medicine, 2022, 12, 932.	1.1	0