# Yoram Cohen

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1072946/publications.pdf

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| 42 | 2,541 | 471371 | 302012 |
|---|---|---|---|
| papers | citations | 17 h-index | 39 g-index |

| 46 | 46 | 46 | 3509 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Associations between breast implants and postpartum lactational mastitis in breastfeeding women: retrospective study. BJOG: an International Journal of Obstetrics and Gynaecology, 2022, 129, 267-272. | 1.1 | 3 |
| 2 | Female fragile X premutation carriers are at increased risk for metabolic syndrome from early adulthood. Nutrition, Metabolism and Cardiovascular Diseases, 2022, 32, 1010-1018. | 1.1 | 1 |
| 3 | P-124â€ƒThe impact of Fragile x premutation carrier state on embryo morphokinetic development, A comparison between genetically normal and abnormal embryos. Human Reproduction, 2022, 37, . | 0.4 | 0 |
| 4 | Dysregulation of anti-Mullerian hormone expression levels in mural granulosa cells of FMR1 premutation carriers. Scientific Reports, 2021, 11, 14139. | 1.6 | 1 |
| 5 | Does the presence of AGG interruptions within the CGG repeat tract have a protective effect on the fertility phenotype of female FMR1 premutation carriers?. Journal of Assisted Reproduction and Genetics, 2020, 37, 849-854. | 1.2 | 6 |
| 6 | FMRpolyG accumulates in FMR1 premutation granulosa cells. Journal of Ovarian Research, 2020, 13, 22. | 1.3 | 16 |
| 7 | Single-Nucleotide Polymorphisms in IL23R-IL12RB2 (rs1495965) Are Highly Prevalent in Patients with Behcetâ€™s Uveitis and Vary Between Populations. Ocular Immunology and Inflammation, 2019, 27, 766-773. | 1.0 | 6 |
| 8 | Absence of AGG Interruptions Is a Risk Factor for Full Mutation Expansion Among Israeli FMR1 Premutation Carriers. Frontiers in Genetics, 2018, 9, 606. | 1.1 | 14 |
| 9 | Fragile X Associated Primary Ovarian Insufficiency (FXPOI): Case Report and Literature Review. Frontiers in Genetics, 2018, 9, 529. | 1.1 | 26 |
| 10 | Preimplantation genetic diagnosis versus prenatal diagnosisâ€”decision-making among pregnant FMR1 premutation carriers. Journal of Assisted Reproduction and Genetics, 2018, 35, 2071-2075. | 1.2 | 9 |
| 11 | Choice of anaesthesia for categoryâ€1 caesarean section in women with anticipated difficult tracheal intubation: the use of decision analysis. Anaesthesia, 2017, 72, 156-171. | 1.8 | 37 |
| 12 | In Vitro Exposure of Human Luteinized Mural Granulosa Cells to Dibutyl Phthalate Affects Global Gene Expression. Toxicological Sciences, 2017, 160, 180-188. | 1.4 | 10 |
| 13 | Category-1 caesarean section, airways and Julius Caesar. A reply. Anaesthesia, 2017, 72, 1154-1155. | 1.8 | 0 |
| 14 | Fragile X Premutation Carrier Epidemiology and Symptomatology in Israelâ€”Results from a Tertiary Child Developmental Center. Cerebellum, 2016, 15, 595-598. | 1.4 | 8 |
| 15 | An antibody to amphiregulin, an abundant growth factor in patientsâ€™ fluids, inhibits ovarian tumors. Oncogene, 2016, 35, 438-447. | 2.6 | 33 |
| 16 | Mutational analysis of PI3K/AKT and RAS/RAF pathway activation in malignant salivary gland tumours with a new mutation of PIK3CA. International Journal of Oral and Maxillofacial Surgery, 2016, 45, 721-725. | 0.7 | 6 |
| 17 | FMR6 may play a role in the pathogenesis of fragile X-associated premature ovarian insufficiency. Gynecological Endocrinology, 2016, 32, 334-337. | 0.7 | 19 |
| 18 | Interleukinâ€2 Production by Cultured Human Granulosa Cells. American Journal of Reproductive Immunology, 2015, 74, 392-397. | 1.2 | 6 |

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|---|---|---|---|
| 19 | Elevated Levels of FMR1 mRNA in Granulosa Cells Are Associated with Low Ovarian Reserve in FMR1 Premutation Carriers. PLoS ONE, 2014, 9, e105121. | 1.1 | 57 |
| 20 | Interleukin-2 and SOCS-1 proteins involvement in the pathophysiology of severe ovarian hyperstimulation syndrome-a preliminary proof of concept. Journal of Ovarian Research, 2014, 7, 106. | 1.3 | 17 |
| 21 | Genetic Mutation Screen in Early Nonâ€“Small-Cell Lung Cancer (NSCLC) Specimens. Clinical Lung Cancer, 2014, 15, 159-165. | 1.1 | 17 |
| 22 | Evaluation of EGFR, KRAS, and TP53 mutations as predictive of disease recurrence in resected early non-small cell lung carcinomas (NSCLCs). Memo - Magazine of European Medical Oncology, 2014, 7, 10-15. | 0.3 | 0 |
| 23 | Exposure of fallopian tube epithelium to follicular fluid mimics carcinogenic changes in precursor lesions of serous papillary carcinoma. Gynecologic Oncology, 2014, 132, 322-327. | 0.6 | 87 |
| 24 | BRCA1/2 mutations and FMR1 alleles are randomly distributed: a case control study. European Journal of Human Genetics, 2014, 22, 277-279. | 1.4 | 9 |
| 25 | Tamoxifen co-administration during controlled ovarian hyperstimulation for inÂvitro fertilization in breast cancer patients increases the safety of fertility-preservation treatment strategies. Fertility and Sterility, 2014, 102, 488-495.e3. | 0.5 | 95 |
| 26 | Surgically treated ovarian endometriosis association with BRCA1 and BRCA2 mutations. Pathology Research and Practice, 2014, 210, 250-255. | 1.0 | 6 |
| 27 | Association between a common CYP17A1 haplotype and anxiety in female anorexia nervosa. Archives of Women's Mental Health, 2013, 16, 423-428. | 1.2 | 2 |
| 28 | JAK2V617F allele burden is associated with transformation to myelofibrosis. Leukemia and Lymphoma, 2012, 53, 2210-2213. | 0.6 | 22 |
| 29 | Niche-modulated and niche-modulating genes in bone marrow cells. Blood Cancer Journal, 2012, 2, e97-e97. | 2.8 | 6 |
| 30 | BRAF and GNAQ mutations in melanocytic tumors of the oral cavity. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2012, 114, 778-784. | 0.2 | 14 |
| 31 | Mutational analysis of PTEN/PIK3CA/AKT pathway in oral squamous cell carcinoma. Oral Oncology, 2011, 47, 946-950. | 0.8 | 54 |
| 32 | High-Throughput Mutation Profiling in Intraductal Papillary Mucinous Neoplasm (IPMN). Journal of Gastrointestinal Surgery, 2011, 15, 503-511. | 0.9 | 31 |
| 33 | AKT1 pleckstrin homology domain E17K activating mutation in endometrial carcinoma. Gynecologic Oncology, 2010, 116, 88-91. | 0.6 | 48 |
| 34 | PI3K/Akt Pathway Mutations in Retinoblastoma. , 2009, 50, 5054. | | 16 |
| 35 | AKT1 E17 K pleckstrin homology domain mutation in urothelial carcinoma. Cancer Genetics and Cytogenetics, 2009, 191, 34-37. | 1.0 | 30 |
| 36 | A new MALDI-TOF-based assay for monitoring JAK2 V617F mutation level in patients undergoing allogeneic stem cell transplantation (allo SCT) for classic myeloproliferative disorders (MPD). Leukemia Research, 2008, 32, 421-427. | 0.4 | 30 |

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|---|---|---|---|
| 37 | Hypermethylation of CpG island loci of multiple tumor suppressor genes in retinoblastoma. Experimental Eye Research, 2008, 86, 201-206. | 1.2 | 41 |
| 38 | High-throughput, sensitive and quantitative assay for the detection of BCR-ABL kinase domain mutations. Leukemia, 2007, 21, 1318-1321. | 3.3 | 39 |
| 39 | Spindle imaging: a new marker for optimal timing of ICSI?. Human Reproduction, 2004, 19, 649-654. | 0.4 | 103 |
| 40 | BRAF Mutation in Papillary Thyroid Carcinoma. Journal of the National Cancer Institute, 2003, 95, 625-627. | 3.0 | 849 |
| 41 | Lack of BRAF Mutation in Primary Uveal Melanoma. , 2003, 44, 2876. | | 117 |
| 42 | Comparison of Factors Associated With 30-Day Mortality After Coronary Artery Bypass Grafting in Patients With Versus Without Diabetes Mellitus. American Journal of Cardiology, 1998, 81, 7-11. | 0.7 | 633 |