## Else Marie Vestergaard

List of Publications by Year in descending order

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40 papers

1,732 citations

331670 21 h-index 276875 41 g-index

42 all docs 42 docs citations

42 times ranked 2806 citing authors

#	Article	IF	CITATIONS
1	National data on the early clinical use of nonâ€invasive prenatal testing in public and private healthcare in Denmark 2013–2017. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 100, 884-892.	2.8	11
2	Mosaicism for copy number variations in the placenta is even more difficult to interpret than mosaicism for whole chromosome aneuploidy. Prenatal Diagnosis, 2021, 41, 668-680.	2.3	6
3	Placental mosaicism in the era of chromosomal microarrays. European Journal of Medical Genetics, 2020, 63, 103778.	1.3	7
4	Prevalence of mosaicism in uncultured chorionic villus samples after chromosomal microarray and clinical outcome in pregnancies affected by confined placental mosaicism. Prenatal Diagnosis, 2020, 40, 244-259.	2.3	23
5	Fetal Heart Defects and Measures of Cerebral Size. Journal of Pediatrics, 2019, 210, 146-153.	1.8	15
6	Case of successful IVF treatment of an oligospermic male with 46,XX/46,XY chimerism. Journal of Assisted Reproduction and Genetics, 2018, 35, 1325-1328.	2.5	15
7	Chromosomal microarray as primary diagnostic genomic tool for pregnancies at increased risk within a populationâ€based combined firstâ€ŧrimester screening program. Ultrasound in Obstetrics and Gynecology, 2018, 51, 480-486.	1.7	46
8	Ocular albinism with infertility and lateâ€onset sensorineural hearing loss. American Journal of Medical Genetics, Part A, 2018, 176, 1587-1593.	1.2	6
9	Isochromosome 21q is overrepresented among false-negative cell-free DNA prenatal screening results involving Down syndrome. European Journal of Human Genetics, 2018, 26, 1490-1496.	2.8	16
10	On the road to replacing invasive testing with cellâ€based NIPT: Five clinical cases with aneuploidies, microduplication, unbalanced structural rearrangement, or mosaicism. Prenatal Diagnosis, 2017, 37, 1120-1124.	2.3	47
11	First reported case of Simpson–Golabi–Behmel syndrome in a female fetus diagnosed prenatally with chromosomal microarray. Clinical Case Reports (discontinued), 2017, 5, 608-612.	0.5	8
12	Comprehensive Evaluation of TFF3 Promoter Hypomethylation and Molecular Biomarker Potential for Prostate Cancer Diagnosis and Prognosis. International Journal of Molecular Sciences, 2017, 18, 2017.	4.1	20
13	Transposition of the great arteries - a phenotype associated with 16p11.2 duplications?. World Journal of Cardiology, 2017, 9, 848-852.	1.5	5
14	17q12 deletion and duplication syndrome in Denmarkâ€"A clinical cohort of 38 patients and review of the literature. American Journal of Medical Genetics, Part A, 2016, 170, 2934-2942.	1.2	53
15	Genome-wide copy number analysis on DNA from fetal cells isolated from the blood of pregnant women. Prenatal Diagnosis, 2016, 36, 1127-1134.	2.3	68
16	Non-invasive prenatal testing offered as part of a combined first-trimester screening program identifies tetrasomy 18p in a high-risk pregnancy. Prenatal Diagnosis, 2016, 36, 1112-1114.	2.3	5
17	Prenatal diagnosis of Nager syndrome in a 12-week-old fetus with a whole gene deletion of SF3B4 by chromosomal microarray. European Journal of Medical Genetics, 2016, 59, 48-51.	1.3	19
18	Aggressive Medullary Thyroid Carcinoma in a Ten-Year-Old Patient with Multiple Endocrine Neoplasia 2B Due to the A883F Mutation. Thyroid, 2015, 25, 139-140.	<b>4.</b> 5	11

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19	Chromosomal microarray in fetuses with increased nuchal translucency. Ultrasound in Obstetrics and Gynecology, 2015, 45, 95-100.	1.7	49
20	Trefoil factors in inflammatory bowel disease. World Journal of Gastroenterology, 2014, 20, 3223.	3.3	85
21	Prenatal diagnosis: array comparative genomic hybridization in fetuses with abnormal sonographic findings. Acta Obstetricia Et Gynecologica Scandinavica, 2013, 92, 762-768.	2.8	29
22	DNA Methylation Signatures for Prediction of Biochemical Recurrence After Radical Prostatectomy of Clinically Localized Prostate Cancer. Journal of Clinical Oncology, 2013, 31, 3250-3258.	1.6	117
23	Absolute immature platelet count may predict imminent platelet recovery in thrombocytopenic children following chemotherapy. Pediatric Blood and Cancer, 2013, 60, 1198-1203.	1.5	30
24	Trefoil factor family peptides in human saliva and cyclical cervical mucus. Method evaluation and results on healthy individuals. Clinical Chemistry and Laboratory Medicine, 2011, 49, 861-868.	2.3	28
25	Promoter hypomethylation and upregulation of trefoil factors in prostate cancer. International Journal of Cancer, 2010, 127, 1857-1865.	5.1	48
26	High-resolution melting analysis using unlabeled probe and amplicon scanning simultaneously detects several lactase persistence variants. Scandinavian Journal of Clinical and Laboratory Investigation, 2010, 70, 535-540.	1.2	8
27	Trefoil factors in human milk. Early Human Development, 2008, 84, 631-635.	1.8	33
28	Independent Introduction of Two Lactase-Persistence Alleles into Human Populations Reflects Different History of Adaptation to Milk Culture. American Journal of Human Genetics, 2008, 82, 57-72.	6.2	301
29	Pharmacokinetics of trefoil peptides and their stability in gastrointestinal contents. Peptides, 2007, 28, 1197-1206.	2.4	30
30	Codon Y791F Mutations in a Large Kindred: Is Prophylactic Thyroidectomy Always Indicated?. World Journal of Surgery, 2007, 31, 996-1001.	1.6	16
31	Plasma Levels of Trefoil Factors are Increased in Patients with Advanced Prostate Cancer. Clinical Cancer Research, 2006, 12, 807-812.	7.0	70
32	Serum Trefoil Factors in Patients with Inflammatory Bowel Disease. Digestion, 2006, 74, 33-39.	2.3	42
33	A gradient of TFF3 (trefoil factor family�3) peptide synthesis within the normal human gastric mucosa. Cell and Tissue Research, 2004, 316, 155-165.	2.9	56
34	Immunoassays of human trefoil factors 1 and 2: measured on serum from patients with inflammatory bowel disease. Scandinavian Journal of Clinical and Laboratory Investigation, 2004, 64, 146-156.	1.2	45
35	Development and Evaluation of an ELISA for Human Trefoil Factor 3. Clinical Chemistry, 2002, 48, 1689-1695.	3.2	49
36	Development and evaluation of an ELISA for human trefoil factor 3. Clinical Chemistry, 2002, 48, 1689-95.	3.2	21

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37	Clinical aspects of altered glycosylation of glycoproteins in cancer. Electrophoresis, 1999, 20, 362-371.	2.4	170
38	LEWIS ANTIGEN MEDIATED ADHESION OF FRESHLY REMOVED HUMAN BLADDER TUMORS TO E-SELECTIN. Journal of Urology, 1999, 161, 1316-1323.	0.4	9
39	Increased concentrations of genotype-interpreted Ca 19-9 in urine of bladder cancer patients mark diffuse atypia of the urothelium. Clinical Chemistry, 1998, 44, 197-204.	3.2	20
40	Influence of Lewis $\hat{l}\pm 1$ -3/4-L-Fucosyltransferase (FUT3) Gene Mutations on Enzyme Activity, Erythrocyte Phenotyping, and Circulating Tumor Marker Sialyl-Lewis a Levels. Journal of Biological Chemistry, 1996, 271, 32260-32268.	3.4	94