Else Marie Vestergaard

List of Publications by Year
in descending order

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	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
2	Clinical aspects of altered glycosylation of glycoproteins in cancer. Electrophoresis, 1999, 20, 362-371.	2.4	170
3	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
4	Influence of Lewis Î±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
5	Trefoil factors in inflammatory bowel disease. World Journal of Gastroenterology, 2014, 20, 3223.	3.3	85
6	Plasma Levels of Trefoil Factors are Increased in Patients with Advanced Prostate Cancer. Clinical Cancer Research, 2006, 12, 807-812.	7.0	70
7	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
8	A gradient of TFF3 (trefoil factor familyï¿½3) peptide synthesis within the normal human gastric mucosa. Cell and Tissue Research, 2004, 316, 155-165.	2.9	56
9	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
10	Development and Evaluation of an ELISA for Human Trefoil Factor 3. Clinical Chemistry, 2002, 48, 1689-1695.	3.2	49
11	Chromosomal microarray in fetuses with increased nuchal translucency. Ultrasound in Obstetrics and Gynecology, 2015, 45, 95-100.	1.7	49
12	Promoter hypomethylation and upregulation of trefoil factors in prostate cancer. International Journal of Cancer, 2010, 127, 1857-1865.	5.1	48
13	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
14	Chromosomal microarray as primary diagnostic genomic tool for pregnancies at increased risk within a populationâ€based combined firstâ€trimester screening program. Ultrasound in Obstetrics and Gynecology, 2018, 51, 480-486.	1.7	46
15	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
16	Serum Trefoil Factors in Patients with Inflammatory Bowel Disease. Digestion, 2006, 74, 33-39.	2.3	42
17	Trefoil factors in human milk. Early Human Development, 2008, 84, 631-635.	1.8	33
18	Pharmacokinetics of trefoil peptides and their stability in gastrointestinal contents. Peptides, 2007, 28, 1197-1206.	2.4	30

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19	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
20	Prenatal diagnosis: array comparative genomic hybridization in fetuses with abnormal sonographic findings. Acta Obstetricia Et Gynecologica Scandinavica, 2013, 92, 762-768.	2.8	29
21	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
22	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
23	Development and evaluation of an ELISA for human trefoil factor 3. Clinical Chemistry, 2002, 48, 1689-95.	3.2	21
24	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
25	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
26	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
27	Codon Y791F Mutations in a Large Kindred: Is Prophylactic Thyroidectomy Always Indicated?. World Journal of Surgery, 2007, 31, 996-1001.	1.6	16
28	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
29	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
30	Fetal Heart Defects and Measures of Cerebral Size. Journal of Pediatrics, 2019, 210, 146-153.	1.8	15
31	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.	4.5	11
32	National data on the early clinical use of nonâ€invasive prenatal testing in public and private healthcare in Denmark 2013â€“2017. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 100, 884-892.	2.8	11
33	LEWIS ANTIGEN MEDIATED ADHESION OF FRESHLY REMOVED HUMAN BLADDER TUMORS TO E-SELECTIN. Journal of Urology, 1999, 161, 1316-1323.	0.4	9
34	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
35	First reported case of Simpsonâ€“Golabiâ€“Behmel syndrome in a female fetus diagnosed prenatally with chromosomal microarray. Clinical Case Reports (discontinued), 2017, 5, 608-612.	0.5	8
36	Placental mosaicism in the era of chromosomal microarrays. European Journal of Medical Genetics, 2020, 63, 103778.	1.3	7

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37	Ocular albinism with infertility and lateâ€onset sensorineural hearing loss. American Journal of Medical Genetics, Part A, 2018, 176, 1587-1593.	1.2	6
38	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
39	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
40	Transposition of the great arteries - a phenotype associated with 16p11.2 duplications?. World Journal of Cardiology, 2017, 9, 848-852.	1.5	5