

Judith H Ford

List of Publications by Year in descending order

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35
papers

822
citations

471509

17
h-index

501196

28
g-index

35
all docs

35
docs citations

35
times ranked

541
citing authors

#	ARTICLE	IF	CITATIONS
1	Simultaneous detection of X- and Y-bearing human sperm by double fluorescence in situ hybridization. <i>Molecular Reproduction and Development</i> , 1993, 34, 308-313.	2.0	76
2	Saturated fatty acid metabolism is key link between cell division, cancer, and senescence in cellular and whole organism aging. <i>Age</i> , 2010, 32, 231-237.	3.0	71
3	Detection of chromosome 17 and X-bearing human spermatozoa using fluorescence in situ hybridization. <i>Molecular Reproduction and Development</i> , 1992, 33, 189-194.	2.0	59
4	Detection of X- and Y-bearing human spermatozoa after motile sperm isolation by swim-up. <i>Fertility and Sterility</i> , 1993, 60, 1046-1051.	1.0	50
5	The impact of maternal serum screening on the birth prevalence of Down's syndrome and the use of amniocentesis and chorionic villus sampling in South Australia. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 2000, 107, 1453-1459.	2.3	44
6	Chromosome abnormalities in chronic lymphocytic leukemia revealed by TPA as a mitogen. <i>Cancer Genetics and Cytogenetics</i> , 1983, 10, 87-93.	1.0	43
7	PALS (pregnancy and lifestyle study): association between occupational and environmental exposure to chemicals and reproductive outcome. <i>Mutation Research - Environmental Mutagenesis and Related Subjects Including Methodology</i> , 1994, 313, 153-164.	0.4	43
8	VACTERL with hydrocephalus: Family with X-linked VACTERL-H. <i>American Journal of Medical Genetics Part A</i> , 1998, 76, 74-78.	2.4	42
9	A fluorescent <i>in situ</i> hybridization analysis of the chromosome constitution of ejaculated sperm in a 47, XYY male. <i>Clinical Genetics</i> , 1994, 45, 67-70.	2.0	40
10	Chromosome errors at mitotic anaphase. <i>Genome</i> , 1992, 35, 702-705.	2.0	35
11	Spindle microtubular dysfunction in mothers of Down syndrome children. <i>Human Genetics</i> , 1984, 68, 295-298.	3.8	33
12	Reduced quality and accelerated follicle loss with female reproductive aging - does decline in theca dehydroepiandrosterone (DHEA) underlie the problem?. <i>Journal of Biomedical Science</i> , 2013, 20, 93.	7.0	33
13	A13-Year Cytogenetic Study of Spontaneous Abortion: Clinical Applications of Testing. <i>Australian and New Zealand Journal of Obstetrics and Gynaecology</i> , 1996, 36, 314-318.	1.0	30
14	Cytogenetic Basis of Acute Myeloid Leukemia. <i>Journal of the National Cancer Institute</i> , 1975, 55, 761-765.	6.3	24
15	Diploid complete hydatidiform mole, mosaic for normally fertilized cells and androgenetic homozygous cells. Case report. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 1986, 93, 1181-1186.	2.3	21
16	ULTRASTRUCTURAL AND CHEMICAL STUDIES OF POLLEN WALL DEVELOPMENT IN THE EPACRIDACEAE. , 1971, , 130-173.		21
17	Hesx1, a Homeobox Gene Expressed by Murine Embryonic Stem Cells, Maps to Mouse Chromosome 14, Bands A3-B. <i>Genomics</i> , 1993, 18, 464-466.	2.9	20
18	A complex translocation in acute promyelocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 1985, 16, 45-48.	1.0	19

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19	Assignment of the interleukin-2 receptor $\hat{\alpha}$ 2 chain gene (IL-2rb) to band E on mouse chromosome 15. Genomics, 1992, 12, 179-180.	2.9	16
20	Translocations of chromosome 12. Human Genetics, 1981, 58, 279-281.	3.8	12
21	Translocations involving chromosome 12. Human Genetics, 1981, 58, 144-148.	3.8	11
22	Novel translocations in acute nonlymphocytic leukemia. Cancer Genetics and Cytogenetics, 1990, 44, 99-105.	1.0	11
23	Preconception risk factors and SGA babies: Papilloma virus, omega 3 and fat soluble vitamin deficiencies. Early Human Development, 2011, 87, 785-789.	1.8	11
24	Overlap between mutagens and teratogens. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 1997, 396, 1-8.	1.0	10
25	Interactions between C-bands of chromosomes 1 and 9 in recurrent reproductive loss. Human Genetics, 1983, 63, 58-62.	3.8	9
26	Does aneuploidy per se cause developmental abnormalities?. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 1997, 396, 195-203.	1.0	8
27	Analysis of fatty acids in early mid-life in fertile women: Implications for reproductive decline and other chronic health problems. American Journal of Human Biology, 2010, 22, 134-136.	1.6	7
28	Contribution of reciprocal translocations to an understanding of chromosome displacement: Inferences for studies of spatial order at metaphase. Human Genetics, 1984, 66, 302-305.	3.8	6
29	Protraction of anaphase B in lymphocyte mitosis with ageing: possible contribution to age-related cancer risk. Mutagenesis, 2013, 28, 307-314.	2.6	5
30	A Model for the Mechanism of Aneuploidy Involving Chromosome Displacement. , 1985, 36, 291-295.		4
31	Lack of correlation between conceptual karyotype and maternal response to placentation. Reproduction, Fertility and Development, 1997, 9, 271.	0.4	4
32	Unusual segregation in a family with a 11/21 translocation. Clinical Genetics, 1988, 33, 449-453.	2.0	2
33	Aneuploidy in Humans. , 1990, , 97-114.		2
34	Segregation of Univalents on Mini Spindles. Nature, 1971, 229, 570-571.	27.8	0
35	Aneuploidy Studies in Sperm: Post-Meiotic Selection against Aneuploid Sperm. , 1994, , 305-310.		0