List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Large Intra-Age Group Variation in Chromosome Abnormalities in Human Blastocysts. Dna, 2021, 1, 91-104.	0.4	1
2	Clinical outcomes after the transfer of blastocysts characterized as mosaic by high resolution Next Generation Sequencing- further insights. European Journal of Medical Genetics, 2020, 63, 103741.	0.7	82
3	First PGT-A using human in vivo blastocysts recovered by uterine lavage: comparison with matched IVF embryo controlsâ€. Human Reproduction, 2020, 35, 70-80.	0.4	38
4	Clinical error rates of next generation sequencing and array comparative genomic hybridization with single thawed euploid embryo transfer. European Journal of Medical Genetics, 2020, 63, 103852.	0.7	23
5	Current and Novel Methods for Chromosome Testing. , 2019, , 603-612.		0
6	Controlled ovarian hyperstimulation (COH) parameters associated with euploidy rates in donor oocytes. European Journal of Medical Genetics, 2019, 62, 103707.	0.7	14
7	25 historic papers: an ASRM 75th birthday gift from Fertility and Sterility. Fertility and Sterility, 2019, 112, e2-e27.	0.5	5
8	Preimplantation genetic testing for aneuploidy versus morphology as selection criteria for single frozen-thawed embryo transfer in good-prognosis patients: a multicenter randomized clinical trial. Fertility and Sterility, 2019, 112, 1071-1079.e7.	0.5	379
9	The cytogenetic constitution of human blastocysts: insights from comprehensive chromosome screening strategies. Human Reproduction Update, 2019, 25, 15-33.	5.2	87
10	Next generation sequencing for preimplantation genetic screening improves pregnancy outcomes compared with array comparative genomic hybridization in single thawed euploid embryo transfer cycles. Fertility and Sterility, 2018, 109, 627-632.	0.5	99
11	Genetic Selection of the Human Embryos: From FISH to NGS, Past and Future. , 2018, , 227-242.		4
12	Forty years of IVF. Fertility and Sterility, 2018, 110, 185-324.e5.	0.5	211
13	Status of preimplantation genetic testing and embryo selection. Reproductive BioMedicine Online, 2018, 37, 393-396.	1.1	50
14	Response: how PGS/PGT-a laboratories succeeded in losing all credibility. Reproductive BioMedicine Online, 2018, 37, 247-249.	1.1	4
15	Optimal euploid embryo transfer strategy, fresh versus frozen, after preimplantation genetic screening with next generation sequencing: a randomized controlled trial. Fertility and Sterility, 2017, 107, 723-730.e3.	0.5	120
16	Detection of mosaicism at blastocyst stage with the use of high-resolution next-generation sequencing. Fertility and Sterility, 2017, 107, 1085-1091.	0.5	164
17	Advanced maternal age patients benefit from preimplantation genetic diagnosis of aneuploidy. Fertility and Sterility, 2017, 107, 1145-1146.	0.5	19
18	Comment on: Gleicher N et al., 2016. Reprod biol endocrinol Sep 5;14(1). Reproductive Biology and Endocrinology, 2017, 15, 24.	1.4	1

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19	Detailed investigation into the cytogenetic constitution and pregnancy outcome of replacing mosaic blastocysts detected with the use of high-resolution next-generation sequencing. Fertility and Sterility, 2017, 108, 62-71.e8.	0.5	219
20	Differences in pregnancy outcomes in donor egg frozen embryo transfer (FET) cycles following preimplantation genetic screening (PGS): a single center retrospective study. Journal of Assisted Reproduction and Genetics, 2017, 34, 71-78.	1.2	26
21	Discrepant diagnosis rate of array comparative genomic hybridization in thawed euploid blastocysts. Journal of Assisted Reproduction and Genetics, 2016, 33, 893-897.	1.2	31
22	Why do euploid embryos miscarry? A case-control study comparing the rate of aneuploidy within presumed euploid embryos that resulted in miscarriage or live birth using next-generation sequencing. Fertility and Sterility, 2016, 106, 1414-1419.e5.	0.5	154
23	Causes and estimated incidences of sex-chromosome misdiagnosis in preimplantation genetic diagnosis of aneuploidy. Reproductive BioMedicine Online, 2016, 33, 550-559.	1.1	5
24	The why, the how and the when of PGS 2.0: current practices and expert opinions of fertility specialists, molecular biologists, and embryologists. Molecular Human Reproduction, 2016, 22, 845-857.	1.3	116
25	Mosaicism: "survival of the fittest―versus "no embryo left behind― Fertility and Sterility, 2016, 105, 1146-1149.	0.5	102
26	Altered Levels of Mitochondrial DNA Are Associated with Female Age, Aneuploidy, and Provide an Independent Measure of Embryonic Implantation Potential. PLoS Genetics, 2015, 11, e1005241.	1.5	253
27	Detection and phasing of single base de novo mutations in biopsies from human in vitro fertilized embryos by advanced whole-genome sequencing. Genome Research, 2015, 25, 426-434.	2.4	49
28	Deliveries from trophectoderm biopsied, fresh and vitrified blastocysts derived from polar body biopsied, vitrified oocytes. Reproductive BioMedicine Online, 2015, 31, 210-216.	1.1	4
29	The human sex ratio from conception to birth. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E2102-11.	3.3	206
30	Validation of next-generation sequencing for comprehensive chromosome screening of embryos. Reproductive BioMedicine Online, 2015, 31, 760-769.	1.1	107
31	Live births following Karyomapping of human blastocysts: experience from clinical application of the method. Reproductive BioMedicine Online, 2015, 31, 394-403.	1.1	61
32	Biomarkers for infertility and recurrent pregnancy loss. Reproductive BioMedicine Online, 2014, 29, 1-2.	1.1	2
33	A greater number of euploid blastocysts in a given cohort predicts excellent outcomes in single embryo transfer cycles. Journal of Assisted Reproduction and Genetics, 2014, 31, 667-673.	1.2	11
34	Genome-wide karyomapping accurately identifies the inheritance of single-gene defects in human preimplantation embryos in vitro. Genetics in Medicine, 2014, 16, 838-845.	1.1	126
35	Clinical utilisation of a rapid low-pass whole genome sequencing technique for the diagnosis of aneuploidy in human embryos prior to implantation. Journal of Medical Genetics, 2014, 51, 553-562.	1.5	200
36	Diminished effect of maternal age onÂimplantation after preimplantation genetic diagnosis with array comparative genomic hybridization. Fertility and Sterility, 2013, 100, 1695-1703.	0.5	284

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37	Live Birth from Previously Vitrified Oocytes, after Trophectoderm Biopsy, Revitrification, and Transfer of a Euploid Blastocyst. Clinical Medicine Insights Reproductive Health, 2013, 7, CMRH.S11919.	3.9	2
38	Microarrays and CGH for PGD of Chromosome Abnormalities and Gene Defects. , 2013, , 425-438.		0
39	Microarrays and CGH for PGD of Chromosome Abnormalities and Gene Defects. , 2013, , 303-316.		0
40	Preimplantation Genetic Diagnosis for Aneuploidy and Translocations Using Array Comparative Genomic Hybridization. Current Genomics, 2012, 13, 463-470.	0.7	73
41	ldiopathic recurrent miscarriage is caused mostly by aneuploid embryos. Fertility and Sterility, 2012, 98, 675-680.	0.5	160
42	Validation of array comparative genome hybridization for diagnosis of translocations in preimplantation human embryos. Reproductive BioMedicine Online, 2012, 24, 621-629.	1.1	73
43	Array CGH analysis shows that aneuploidy is not related to the number of embryos generated. Reproductive BioMedicine Online, 2012, 24, 614-620.	1.1	191
44	Intra-age, intercenter, and intercycle differences in chromosome abnormalities in oocytes. Fertility and Sterility, 2012, 97, 935-942.	0.5	19
45	The effect of timing of embryonicÂprogression on chromosomal abnormality. Fertility and Sterility, 2012, 98, 876-880.	0.5	61
46	Dynamic blastomere behaviour reflects human embryo ploidy by the four-cell stage. Nature Communications, 2012, 3, 1251.	5.8	260
47	Microarrays and CGH for PGD of Chromosome Abnormalities and Gene Defects. , 2012, , 483-490.		1
48	Validation of microarray comparative genomic hybridization for comprehensive chromosome analysis of embryos. Fertility and Sterility, 2011, 95, 953-958.	0.5	272
49	Culture-induced chromosome abnormalities: the canary in the mine. Reproductive BioMedicine Online, 2011, 22, 506-508.	1.1	11
50	Preimplantation genetic diagnosis (PGD) improves pregnancy outcome for translocation carriers with a history of recurrent losses. Fertility and Sterility, 2010, 94, 283-289.	0.5	133
51	Technology requirements for preimplantation genetic diagnosis to improve assisted reproduction outcomes. Fertility and Sterility, 2010, 94, 408-430.	0.5	86
52	Clinical application of comprehensive chromosomal screening at the blastocyst stage. Fertility and Sterility, 2010, 94, 1700-1706.	0.5	293
53	Comprehensive chromosome screening of polar bodies and blastocysts from couples experiencing repeated implantation failure. Fertility and Sterility, 2010, 94, 875-887.	0.5	147
54	Overview of preimplantation genetic diagnosis. Expert Review of Obstetrics and Gynecology, 2010, 5, 403-408.	0.4	1

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55	Rapid mapping of chromosomal breakpoints: from blood to BAC in 20 days Folia Histochemica Et Cytobiologica, 2010, 47, 367-75.	0.6	11
56	Chromosomal Rainbows detect Oncogenic Rearrangements of Signaling Molecules in Thyroid Tumors. , 2010, 2, 13-22.		7
57	Pregnancy Complicated by Triploidy: A Comparison of the Three Karyotypes. American Journal of Perinatology, 2009, 26, 641-645.	0.6	22
58	DNA Probe Pooling for Rapid Delineation of Chromosomal Breakpoints. Journal of Histochemistry and Cytochemistry, 2009, 57, 587-597.	1.3	24
59	Selection of embryos by morphology is less effective than by a combination of aneuploidy testing and morphology observations. Fertility and Sterility, 2009, 91, 943-945.	0.5	26
60	Preimplantation aneuploidy testing for infertile patients of advanced maternal age: a randomized prospective trial. Fertility and Sterility, 2009, 92, 157-162.	0.5	136
61	Effect of infertility, maternal age, and number of previous miscarriages on the outcome of preimplantation genetic diagnosis for idiopathic recurrent pregnancy loss. Fertility and Sterility, 2009, 92, 288-295.	0.5	62
62	The impact of LH-containing gonadotropin stimulation on euploidy rates in preimplantation embryos: antagonist cycles. Fertility and Sterility, 2009, 92, 937-942.	0.5	26
63	Preimplantation genetic diagnosis of single-gene disorders: experience with more than 200 cycles conducted by a reference laboratory in the United States. Fertility and Sterility, 2009, 92, 1544-1556.	0.5	66
64	The role of preimplantation genetic diagnosis in diagnosing embryo aneuploidy. Current Opinion in Obstetrics and Gynecology, 2009, 21, 442-449.	0.9	27
65	The impact of LH-containing gonadotropins on diploidy rates in preimplantation embryos: long protocol stimulation. Human Reproduction, 2008, 23, 499-503.	0.4	48
66	Improving pregnancy outcome for IVF patients with preimplantation genetic screening. Expert Review of Obstetrics and Gynecology, 2008, 3, 635-646.	0.4	3
67	Preimplantation genetic diagnosis for infertility. , 2008, , 381-401.		0
68	In Vitro Fertilization with Preimplantation Genetic Screening. New England Journal of Medicine, 2007, 357, 1769-1771.	13.9	49
69	Removal of 2 cells from cleavage stage embryos is likely to reduce the efficacy of chromosomal tests that are used to enhance implantation rates. Fertility and Sterility, 2007, 87, 496-503.	0.5	206
70	Increased efficiency of preimplantation genetic diagnosis for infertility using "no result rescue― Fertility and Sterility, 2007, 88, 53-61.	0.5	124
71	Lack of association between polycystic ovary syndrome and embryonic aneuploidy. Fertility and Sterility, 2007, 88, 900-905.	0.5	71
72	Substandard application of preimplantation genetic screening may interfere with its clinical success. Fertility and Sterility, 2007, 88, 781-784.	0.5	104

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73	Maternal age-related differential global expression profiles observed in human oocytes. Reproductive BioMedicine Online, 2007, 14, 700-708.	1.1	181
74	Derivation of Human Embryonic Stem Cells in Xeno-Free Conditions. Methods in Molecular Biology, 2007, 407, 1-10.	0.4	18
75	Chromosomal status of human embryos. Reproductive Medicine and Assisted Reproductive Techniques Series, 2007, , 209-234.	0.1	3
76	Cryopreservation of unfertilized human oocytes. Reproductive BioMedicine Online, 2006, 13, 222-227.	1.1	43
77	Chromosome abnormalities and their relationship to morphology and development of human embryos. Reproductive BioMedicine Online, 2006, 12, 234-253.	1.1	217
78	Preimplantation genetic diagnosis significantly improves the pregnancy outcome of translocation carriers with a history of recurrent miscarriage and unsuccessful pregnancies. Reproductive BioMedicine Online, 2006, 13, 869-874.	1.1	123
79	Preimplantation genetic diagnosis significantly reduces pregnancy loss in infertile couples: a multicenter study. Fertility and Sterility, 2006, 85, 326-332.	0.5	138
80	Reply: PGD—a model to evaluate efficacy?. Fertility and Sterility, 2006, 85, 535-536.	0.5	2
81	Recurrent abortion and live birth rate per patient. Fertility and Sterility, 2006, 85, 1071.	0.5	1
82	Preimplantation genetic diagnosis for translocations. Human Reproduction, 2006, 21, 839-840.	0.4	13
83	Nonviable Human Pre-Implantation Embryos as a Source of Stem Cells for Research and Potential Therapy. Stem Cell Reviews and Reports, 2005, 1, 337-344.	5.6	14
84	Preimplantation genetic diagnosis for chromosome abnormalities. , 2005, , .		0
85	Fluorescence In Situ Hybridization and Spectral Imaging Analysis of Human Oocytes and First Polar Bodies. Journal of Histochemistry and Cytochemistry, 2005, 53, 269-272.	1.3	12
86	Patterns of ovarian response to gonadotropin stimulation in female carriers of balanced translocation. Fertility and Sterility, 2005, 83, 1504-1509.	0.5	29
87	Preimplantation genetic diagnosis reduces pregnancy loss in women aged 35 years and older with a history of recurrent miscarriages. Fertility and Sterility, 2005, 84, 331-335.	0.5	229
88	Genetic testing of embryos: a critical need for data. Reproductive BioMedicine Online, 2005, 11, 667-670.	1.1	26
89	Cryopreservation of biopsied cleavage stage human embryos. Reproductive BioMedicine Online, 2005, 11, 711-715.	1.1	27
90	Negligible interchromosomal effect in embryos of Robertsonian translocation carriers. Reproductive BioMedicine Online, 2005, 10, 363-369.	1.1	30

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91	Self-correction of chromosomally abnormal embryos in culture and implications for stem cell production. Fertility and Sterility, 2005, 84, 1328-1334.	0.5	146
92	Reliability of comparative genomic hybridization to detect chromosome abnormalities in first polar bodies and metaphase II oocytes. Human Reproduction, 2004, 19, 2118-2125.	0.4	74
93	PGD analysis for aneuploidy in a patient heterozygous for a polymorphism of chromosome 16(16qhâ^'). Prenatal Diagnosis, 2004, 24, 741-744.	1.1	24
94	Increased rate of aneuploid embryos in young women with previous aneuploid conceptions. Prenatal Diagnosis, 2004, 24, 638-643.	1.1	101
95	Differences in chromosome susceptibility to aneuploidy and survival to first trimester. Reproductive BioMedicine Online, 2004, 8, 81-90.	1.1	128
96	The status of preimplantation genetic diagnosis in Japan: a criticism. Reproductive BioMedicine Online, 2004, 9, 258-259.	1.1	8
97	Over a decade of experience with preimplantation genetic diagnosis: A multicenter report. Fertility and Sterility, 2004, 82, 292-294.	0.5	204
98	Chromosomal abnormalities in embryos derived from testicular sperm extraction. Fertility and Sterility, 2003, 79, 30-38.	0.5	162
99	Predictive value of sperm fluorescence in situ hybridization analysis on the outcome of preimplantation genetic diagnosis for translocations. Fertility and Sterility, 2003, 79, 1528-1534.	0.5	102
100	Preimplantation genetic diagnosis as both a therapeutic and diagnostic tool in assisted reproductive technology. Fertility and Sterility, 2003, 80, 467-468.	0.5	93
101	Questions concerning the suitability of comparative genomic hybridization for preimplantation genetic diagnosis. Fertility and Sterility, 2003, 80, 871-872.	0.5	25
102	Preimplantation genetic diagnosis of numerical abnormalities for 13 chromosomes. Reproductive BioMedicine Online, 2003, 6, 226-231.	1.1	41
103	Improved implantation after preimplantation genetic diagnosis of aneuploidy. Reproductive BioMedicine Online, 2003, 7, 91-97.	1.1	306
104	Spectral karyotyping of fresh, non-inseminated oocytes. Molecular Human Reproduction, 2002, 8, 580-585.	1.3	114
105	Preimplantation genetic diagnosis. Current Opinion in Obstetrics and Gynecology, 2002, 14, 239-244.	0.9	27
106	Preimplantation genetic diagnosis for advanced maternal age and other indications. Fertility and Sterility, 2002, 78, 234-236.	0.5	80
107	First clinical application of comparative genomic hybridization and polar body testing for preimplantation genetic diagnosis of aneuploidy. Fertility and Sterility, 2002, 78, 543-549.	0.5	201
108	Blastomere fixation techniques and risk of misdiagnosis for preimplantation genetic diagnosis of aneuploidy. Reproductive BioMedicine Online, 2002, 4, 210-217.	1.1	104

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109	Chromosome mosaicism in cleavage-stage human embryos: evidence of a maternal age effect. Reproductive BioMedicine Online, 2002, 4, 223-232.	1.1	186
110	Preimplantation genetic diagnosis of numerical and structural chromosome abnormalities. Reproductive BioMedicine Online, 2002, 4, 183-196.	1.1	169
111	Pregnancies from single normal embryo transfer in women older than 40 years. Reproductive BioMedicine Online, 2001, 2, 98-101.	1.1	26
112	Preimplantation genetic diagnosis of structural abnormalities. Molecular and Cellular Endocrinology, 2001, 183, S55-S58.	1.6	28
113	Detection of chromosome translocation products in single interphase cell nuclei. Methods in Cell Biology, 2001, 64, 97-114.	0.5	13
114	Preimplantation genetic diagnosis of pericentric inversions. Prenatal Diagnosis, 2001, 21, 760-766.	1.1	29
115	Preimplantation genetic diagnosis (PGD), a collaborative activity of clinical genetic departments and IVF centres. Prenatal Diagnosis, 2001, 21, 1086-1092.	1.1	31
116	Female gamete segregation in two carriers of translocations involving 2q and 14q. , 2000, 20, 235-237.		16
117	Selection of the most common chromosome abnormalities in oocytes prior to ICSI. Prenatal Diagnosis, 2000, 20, 582-586.	1.1	29
118	Analysis of chromosome abnormalities in sperm and embryos from two 45,XY,t(13;14)(q10;q10) carriers. Prenatal Diagnosis, 2000, 20, 599-602.	1.1	15
119	ls there an interchromosomal effect in reciprocal translocation carriers? Sperm FISH studies. Human Genetics, 2000, 106, 517-524.	1.8	64
120	Chromosome abnormalities in 1255 cleavage-stage human embryos. Reproductive BioMedicine Online, 2000, 1, 17-26.	1.1	188
121	Outcome of preimplantation genetic diagnosis of translocations. Fertility and Sterility, 2000, 73, 1209-1218.	0.5	278
122	Pregnancy after polar body biopsy and freezing and thawing of human embryos. Fertility and Sterility, 2000, 73, 645-647.	0.5	25
123	Analysis of chromosome abnormalities in sperm and embryos from two 45,XY,t(13;14)(q10;q10) carriers. Prenatal Diagnosis, 2000, 20, 599-602.	1.1	78
124	Preimplantation genetic diagnosis of aneuploidy: were we looking at the wrong chromosomes?. Journal of Assisted Reproduction and Genetics, 1999, 16, 176-181.	1.2	71
125	Advantages of day 4 embryo transfer in patients undergoing preimplantation genetic diagnosis of aneuploidy. Journal of Assisted Reproduction and Genetics, 1999, 16, 170-175.	1.2	45
126	Patient-specific probes for preimplantation genetic diagnosis of structural and numerical aberrations in interphase cells. Journal of Assisted Reproduction and Genetics, 1999, 16, 182-191.	1.2	35

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127	Multicolor fluorescence in situ hybridization analysis of the spermatozoa of a male heterozygous for a reciprocal translocation t(11;22)(q23;q11). Human Genetics, 1999, 104, 412-417.	1.8	34
128	Preimplantation diagnosis for aneuploidies in patients undergoing in vitro fertilization with a poor prognosis: identification of the categories for which it should be proposed. Fertility and Sterility, 1999, 72, 837-844.	0.5	439
129	Sperm quality may adversely affect the chromosome constitution of embryos that result from intracytoplasmic sperm injection. Fertility and Sterility, 1999, 72, 1113-1115.	0.5	38
130	Molecular cloning of translocation breakpoints in a case of constitutional translocation t(11;22)(q23;q11) and preparation of probes for preimplantation genetic diagnosis. Reproduction, Fertility and Development, 1999, 11, 17.	0.1	12
131	Spontaneous abortions are reduced after preconception diagnosis of translocations. Journal of Assisted Reproduction and Genetics, 1998, 15, 290-296.	1.2	121
132	Case report: chromatid exchange and predivision of chromatids as other sources of abnormal occytes detected by preimplantation genetic diagnosis of translocations. , 1998, 18, 1450-1458.		27
133	Preimplantation diagnosis of the aneuploidies most commonly found in spontaneous abortions and live births: XY, 13, 14, 15, 16, 18, 21, 22. Prenatal Diagnosis, 1998, 18, 1459-1466.	1.1	212
134	Impaired development of zygotes with uneven pronuclear size. Zygote, 1998, 6, 137-141.	0.5	85
135	Case report: chromatid exchange and predivision of chromatids as other sources of abnormal oocytes detected by preimplantation genetic diagnosis of translocations. , 1998, 18, 1450.		2
136	Preimplantation genetic diagnosis increases the implantation rate in human in vitro fertilization by avoiding the transfer of chromosomally abnormal embryos. Fertility and Sterility, 1997, 68, 1128-1131.	0.5	175
137	Aneuploidy 16 in human embryos increases significantly with maternal age. Fertility and Sterility, 1996, 66, 248-255.	0.5	98
138	Reduction in signal overlap results in increased FISH efficiency: Implications for preimplantation genetic diagnosis. Journal of Assisted Reproduction and Genetics, 1996, 13, 149-156.	1.2	70
139	Embryo morphology, developmental rates, and maternal age are correlated with chromosome abnormalities. Fertility and Sterility, 1995, 64, 382-391.	0.5	729
140	Male and Female Genomes Associated in a Single Pronucleus in Human Zygotes. Biology of Reproduction, 1995, 52, 653-657.	1.2	63
141	Formation of Male Pronuclei in Partitioned Human Oocytes. Biology of Reproduction, 1995, 53, 209-213.	1.2	12
142	Assessment of numeric abnormalities of X, Y, 18, and 16 chromosomes in preimplantation human embryos before transfer. American Journal of Obstetrics and Gynecology, 1995, 172, 1191-1201.	0.7	139
143	Micromanipulation in Clinical Management of Fertility Disorders. Seminars in Reproductive Medicine, 1994, 12, 151-168.	0.5	17
144	Monospermic polyploidy and atypical embryo morphology. Human Reproduction, 1994, 9, 506-510.	0.4	61

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145	Healthy deliveries from biopsied human embryos. Human Reproduction, 1994, 9, 912-916.	0.4	72
146	The human zygote inherits its mitotic potential from the male gamete. Human Reproduction, 1994, 9, 1220-1225.	0.4	225
147	Sex determination of human embryos using the polymerase chain reaction and confirmation by fluorescence in situ hybridization. Fertility and Sterility, 1994, 61, 111-117.	0.5	45
148	A fast and efficient method for simultaneous X and Y in situ hybridization of human blastomeres. Journal of Assisted Reproduction and Genetics, 1993, 10, 82-90.	1.2	113
149	Origin of single pronucleated human zygotes. Journal of Assisted Reproduction and Genetics, 1993, 10, 276-279.	1.2	35
150	Sex distribution in arrested precompacted human embryos. Zygote, 1993, 1, 155-162.	0.5	11
151	Microinjection of FITC-dextran into mouse blastomeres to assess topical effects of zona photoablation. Zygote, 1993, 1, 43-48.	0.5	18
152	Fertilization and early embryology: Diagnosis of major chromosome aneuploidies in human preimplantation embryos. Human Reproduction, 1993, 8, 2185-2191.	0.4	517
153	Unsuitability of multinucleated human blastomeres for preimplantation genetic diagnosis. Human Reproduction, 1993, 8, 1120-1125.	0.4	99
154	PGD for Chromosomal Anomalies. , 0, , 643-656.		0