

# Hongyu Sun

## List of Publications by Year in descending order

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50  
papers

2,378  
citations

430874

18  
h-index

206112

48  
g-index

53  
all docs

53  
docs citations

53  
times ranked

2421  
citing authors

#	ARTICLE	IF	CITATIONS
1	Maternal Plasma DNA Sequencing Reveals the Genome-Wide Genetic and Mutational Profile of the Fetus. <i>Science Translational Medicine</i> , 2010, 2, 61ra91.	12.4	878
2	Non-invasive prenatal assessment of trisomy 21 by multiplexed maternal plasma DNA sequencing: large scale validity study. <i>BMJ: British Medical Journal</i> , 2011, 342, c7401-c7401.	2.3	641
3	Genetic polymorphisms and mutation rates of 27 Y-chromosomal STRs in a Han population from Guangdong Province, Southern China. <i>Forensic Science International: Genetics</i> , 2016, 21, 5-9.	3.1	76
4	Future directions of forensic DNA databases. <i>Croatian Medical Journal</i> , 2014, 55, 163-166.	0.7	47
5	Improved pairwise kinship analysis using massively parallel sequencing. <i>Forensic Science International: Genetics</i> , 2019, 38, 77-85.	3.1	46
6	Polymorphism analysis and evaluation of 19 STR loci in the Han population of Southern China. <i>Annals of Human Biology</i> , 2013, 40, 191-196.	1.0	42
7	Predicting Human Age with Bloodstains by sjTREC Quantification. <i>PLoS ONE</i> , 2012, 7, e42412.	2.5	42
8	Haplotype analysis of the polymorphic 40 Y-STR markers in Chinese populations. <i>Forensic Science International: Genetics</i> , 2015, 19, 255-262.	3.1	34
9	Null alleles of the X and Y chromosomal amelogenin gene in a Chinese population. <i>International Journal of Legal Medicine</i> , 2012, 126, 513-518.	2.2	33
10	Detection and quantification of the age-related sjTREC decline in human peripheral blood. <i>International Journal of Legal Medicine</i> , 2011, 125, 603-608.	2.2	29
11	Increased Expression and Altered Methylation of HERVWE1 in the Human Placentas of Smaller Fetuses from Monozygotic, Dichorionic, Discordant Twins. <i>PLoS ONE</i> , 2012, 7, e33503.	2.5	28
12	Genetic polymorphisms of twelve X-chromosomal STR loci in Chinese Han population from Guangdong Province. <i>Forensic Science International: Genetics</i> , 2011, 5, e114-e116.	3.1	26
13	Comparison of southern Chinese Han and Brazilian Caucasian mutation rates at autosomal short tandem repeat loci used in human forensic genetics. <i>International Journal of Legal Medicine</i> , 2014, 128, 1-9.	2.2	25
14	Epigenome-wide DNA methylation assay reveals placental epigenetic markers for noninvasive fetal single nucleotide polymorphism genotyping in maternal plasma. <i>Transfusion</i> , 2014, 54, 2523-2533.	1.6	23
15	SNP typing using the HID-Ion AmpliSeq <sup>®</sup> Identity Panel in a southern Chinese population. <i>International Journal of Legal Medicine</i> , 2018, 132, 997-1006.	2.2	22
16	Characterizing stutter variants in forensic STRs with massively parallel sequencing. <i>Forensic Science International: Genetics</i> , 2020, 45, 102225.	3.1	21
17	Identification of sequence polymorphisms at 58 STRs and 94 iSNPs in a Tibetan population using massively parallel sequencing. <i>Scientific Reports</i> , 2020, 10, 12225.	3.3	20
18	Developmental validation of the MGIEasy Signature Identification Library Prep Kit, an all-in-one multiplex system for forensic applications. <i>International Journal of Legal Medicine</i> , 2021, 135, 739-753.	2.2	20

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19	Detection of the deletion on Yp11.2 in a Chinese population. <i>Forensic Science International: Genetics</i> , 2014, 8, 73-79.	3.1	19
20	Complete paternal uniparental isodisomy for Chromosome 2 revealed in a parentage testing case. <i>Transfusion</i> , 2013, 53, 1266-1269.	1.6	18
21	Development of a five ChX STRs loci typing system. <i>International Journal of Legal Medicine</i> , 2008, 122, 261-265.	2.2	17
22	Noninvasive fetal genotyping of paternally inherited alleles using targeted massively parallel sequencing in parentage testing cases. <i>Transfusion</i> , 2017, 57, 1505-1514.	1.6	16
23	Identification and sequencing of 59 highly polymorphic microhaplotypes for analysis of DNA mixtures. <i>International Journal of Legal Medicine</i> , 2021, 135, 1137-1149.	2.2	15
24	Investigation of the Application of miR10b and miR135b in the Identification of Semen Stains. <i>PLoS ONE</i> , 2015, 10, e0137067.	2.5	12
25	Pairwise kinship testing with microhaplotypes: Can advancements be made in kinship inference with these markers?. <i>Forensic Science International</i> , 2021, 325, 110875.	2.2	12
26	Genetic polymorphism and population structure of Torghut Mongols and comparison with a Mongolian population 3000 kilometers away. <i>Forensic Science International: Genetics</i> , 2019, 42, 235-243.	3.1	11
27	An SNP panel for the analysis of paternally inherited alleles in maternal plasma using ion Torrent PGM. <i>International Journal of Legal Medicine</i> , 2018, 132, 343-352.	2.2	10
28	A paternity case with mutations at three CODIS core STR loci. <i>Forensic Science International: Genetics</i> , 2012, 6, e61-e62.	3.1	9
29	Genome-wide screen for universal individual identification SNPs based on the HapMap and 1000 Genomes databases. <i>Scientific Reports</i> , 2018, 8, 5553.	3.3	9
30	Analysis of 17 Y-STR loci haplotype and Y-chromosome haplogroup distribution in five Chinese ethnic groups. <i>Electrophoresis</i> , 2015, 36, 2546-2552.	2.4	8
31	Detecting hypermethylated fetal RASSF1A sequences in maternal plasma: implications for noninvasive paternity testing in pregnancy. <i>Transfusion</i> , 2013, 53, 1856-1858.	1.6	7
32	Analysis of 24 Y-STR haplotype data in a Chinese Han population from Guangdong Province. <i>International Journal of Legal Medicine</i> , 2016, 130, 689-691.	2.2	7
33	The novel HLA-DQB1*03:01:01:12 allele, identified by next-generation sequencing in a Chinese family. <i>Hla</i> , 2018, 91, 221-222.	0.6	7
34	Revisiting the potential power of human leukocyte antigen (HLA) genes on relationship testing by massively parallel sequencing-based HLA typing in an extended family. <i>Journal of Human Genetics</i> , 2019, 64, 29-38.	2.3	7
35	Noninvasive Prenatal Paternity Testing with a Combination of Well-Established SNP and STR Markers Using Massively Parallel Sequencing. <i>Genes</i> , 2021, 12, 454.	2.4	7
36	Polymorphism analysis of 15 STR loci in a large sample of the Han population in southern China. <i>Forensic Science International: Genetics</i> , 2009, 4, e27-e29.	3.1	6

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37	Identification of the sequence variations of 15 autosomal STR loci in a Chinese population. <i>Annals of Human Biology</i> , 2014, 41, 524-530.	1.0	6
38	Next-generation sequencing-based typing of a new allele <i>HLA-DQB1*03:01:01:20</i> in a Chinese family. <i>Hla</i> , 2018, 91, 223-224.	0.6	6
39	Integrated massively parallel sequencing of 15 autosomal STRs and Amelogenin using a simplified library preparation approach. <i>Electrophoresis</i> , 2018, 39, 1466-1473.	2.4	5
40	<i>HLA-A*02:01:01:28</i> , a novel HLA allele identified by next-generation sequencing in a Chinese family. <i>Hla</i> , 2018, 91, 195-196.	0.6	5
41	Genetic analysis of 39 Y-STR loci in a Han population from Henan province, central China. <i>International Journal of Legal Medicine</i> , 2019, 133, 95-97.	2.2	4
42	Genetic diversity of 17 autosomal STR loci in Meizhou Hakka population. <i>International Journal of Legal Medicine</i> , 2021, 135, 443-444.	2.2	4
43	Identification of the new <i>HLA-DRB5*01:01:01:02</i> allele in a Chinese individual. <i>Hla</i> , 2019, 93, 56-57.	0.6	3
44	Characterization of genetic polymorphisms in Nigerians residing in Guangzhou using massively parallel sequencing. <i>Forensic Science International: Genetics</i> , 2020, 48, 102323.	3.1	3
45	Development and validation of a new 18 STR typing assay for forensic applications. <i>Electrophoresis</i> , 2021, 42, 766-773.	2.4	3
46	Development of New mRNA Markers for the Identification of Menstrual Blood. <i>Annals of Clinical and Laboratory Science</i> , 2018, 48, 55-62.	0.2	3
47	The new allele <i>HLA-DRB1*14:54:01:04</i> in a Chinese family identified using next-generation sequencing. <i>Hla</i> , 2018, 92, 320-321.	0.6	2
48	Comparative evaluation of autosomal STRs and X-chromosome STRs as a complement of autosomal STRs in kinship testing in Southern Han Chinese. <i>Annals of Human Biology</i> , 2021, 48, 66-69.	1.0	2
49	Linkage and linkage disequilibrium among the markers in the forensic MPS panels. <i>Journal of Forensic Sciences</i> , 2021, 66, 1637-1646.	1.6	2
50	Identification of a rare off-ladder allele of the D13S325 locus during paternity testing. <i>Legal Medicine</i> , 2014, 16, 48-51.	1.3	0