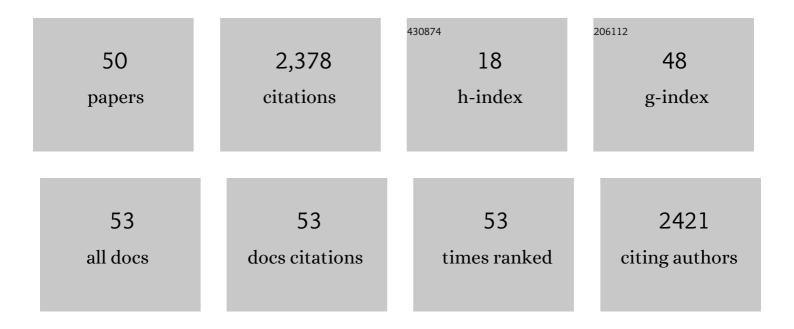
## Hongyu Sun

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Genetic diversity of 17 autosomal STR loci in Meizhou Hakka population. International Journal of Legal Medicine, 2021, 135, 443-444.	2.2	4
2	Comparative evaluation of autosomal STRs and X-chromosome STRs as a complement of autosomal STRs in kinship testing in Southern Han Chinese. Annals of Human Biology, 2021, 48, 66-69.	1.0	2
3	Developmental validation of the MGIEasy Signature Identification Library Prep Kit, an all-in-one multiplex system for forensic applications. International Journal of Legal Medicine, 2021, 135, 739-753.	2.2	20
4	Noninvasive Prenatal Paternity Testing with a Combination of Well-Established SNP and STR Markers Using Massively Parallel Sequencing. Genes, 2021, 12, 454.	2.4	7
5	Linkage and linkage disequilibrium among the markers in the forensic MPS panels. Journal of Forensic Sciences, 2021, 66, 1637-1646.	1.6	2
6	Pairwise kinship testing with microhaplotypes: Can advancements be made in kinship inference with these markers?. Forensic Science International, 2021, 325, 110875.	2.2	12
7	Development and validation of a new 18 X TR typing assay for forensic applications. Electrophoresis, 2021, 42, 766-773.	2.4	3
8	Identification and sequencing of 59 highly polymorphic microhaplotypes for analysis of DNA mixtures. International Journal of Legal Medicine, 2021, 135, 1137-1149.	2.2	15
9	Characterizing stutter variants in forensic STRs with massively parallel sequencing. Forensic Science International: Genetics, 2020, 45, 102225.	3.1	21
10	Identification of sequence polymorphisms at 58 STRs and 94 iiSNPs in a Tibetan population using massively parallel sequencing. Scientific Reports, 2020, 10, 12225.	3.3	20
11	Characterization of genetic polymorphisms in Nigerians residing in Guangzhou using massively parallel sequencing. Forensic Science International: Genetics, 2020, 48, 102323.	3.1	3
12	Genetic analysis of 39 Y-STR loci in a Han population from Henan province, central China. International Journal of Legal Medicine, 2019, 133, 95-97.	2.2	4
13	Genetic polymorphism and population structure of Torghut Mongols and comparison with a Mongolian population 3000 kilometers away. Forensic Science International: Genetics, 2019, 42, 235-243.	3.1	11
14	Identification of the new <i>HLAâ€ÐRB5*01:01:01:02</i> allele in a Chinese individual. Hla, 2019, 93, 56-57.	0.6	3
15	Improved pairwise kinship analysis using massively parallel sequencing. Forensic Science International: Genetics, 2019, 38, 77-85.	3.1	46
16	Revisiting the potential power of human leukocyte antigen (HLA) genes on relationship testing by massively parallel sequencing-based HLA typing in an extended family. Journal of Human Genetics, 2019, 64, 29-38.	2.3	7
17	The novel <i>HLAâ€DQB1*03:01:01:12</i> allele, identified by nextâ€generation sequencing in a Chinese family. Hla, 2018, 91, 221-222.	0.6	7
18	Genome-wide screen for universal individual identification SNPs based on the HapMap and 1000 Genomes databases. Scientific Reports, 2018, 8, 5553.	3.3	9

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19	Integrated massively parallel sequencing of 15 autosomal STRs and Amelogenin using a simplified library preparation approach. Electrophoresis, 2018, 39, 1466-1473.	2.4	5
20	Nextâ€generation sequencingâ€based typing of a new allele <i>HLAâ€DQB1*03:01:01:20</i> in a Chinese family. Hla, 2018, 91, 223-224.	0.6	6
21	<i>HLAâ€A*02:01:01:28</i> , a novel HLA allele identified by nextâ€generation sequencing in a Chinese family. Hla, 2018, 91, 195-196.	0.6	5
22	An SNP panel for the analysis of paternally inherited alleles in maternal plasma using ion Torrent PGM. International Journal of Legal Medicine, 2018, 132, 343-352.	2.2	10
23	SNP typing using the HID-Ion AmpliSeqâ,,¢ Identity Panel in a southern Chinese population. International Journal of Legal Medicine, 2018, 132, 997-1006.	2.2	22
24	The new allele <i>HLAâ€DRB1*14:54:01:04</i> in a Chinese family identified using nextâ€generation sequencing. Hla, 2018, 92, 320-321.	0.6	2
25	Development of New mRNA Markers for the Identification of Menstrual Blood. Annals of Clinical and Laboratory Science, 2018, 48, 55-62.	0.2	3
26	Noninvasive fetal genotyping of paternally inherited alleles using targeted massively parallel sequencing in parentage testing cases. Transfusion, 2017, 57, 1505-1514.	1.6	16
27	Analysis of 24 Y-STR haplotype data in a Chinese Han population from Guangdong Province. International Journal of Legal Medicine, 2016, 130, 689-691.	2.2	7
28	Genetic polymorphisms and mutation rates of 27 Y-chromosomal STRs in a Han population from Guangdong Province, Southern China. Forensic Science International: Genetics, 2016, 21, 5-9.	3.1	76
29	Analysis of 17 Yâ€STR loci haplotype and Yâ€chromosome haplogroup distribution in five Chinese ethnic groups. Electrophoresis, 2015, 36, 2546-2552.	2.4	8
30	Investigation of the Application of miR10b and miR135b in the Identification of Semen Stains. PLoS ONE, 2015, 10, e0137067.	2.5	12
31	Haplotype analysis of the polymorphic 40 Y-STR markers in Chinese populations. Forensic Science International: Genetics, 2015, 19, 255-262.	3.1	34
32	Identification of the sequence variations of 15 autosomal STR loci in a Chinese population. Annals of Human Biology, 2014, 41, 524-530.	1.0	6
33	Future directions of forensic DNA databases. Croatian Medical Journal, 2014, 55, 163-166.	0.7	47
34	Identification of a rare off-ladder allele of the D13S325 locus during paternity testing. Legal Medicine, 2014, 16, 48-51.	1.3	0
35	Epigenomeâ€wide <scp>DNA</scp> methylation assay reveals placental epigenetic markers for noninvasive fetal singleâ€nucleotide polymorphism genotyping in maternal plasma. Transfusion, 2014, 54, 2523-2533.	1.6	23
36	Detection of the deletion on Yp11.2 in a Chinese population. Forensic Science International: Genetics, 2014, 8, 73-79.	3.1	19

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37	Comparison of southern Chinese Han and Brazilian Caucasian mutation rates at autosomal short tandem repeat loci used in human forensic genetics. International Journal of Legal Medicine, 2014, 128, 1-9.	2.2	25
38	Complete paternal uniparental isodisomy for Chromosome $\hat{e}f2$ revealed in a parentage testing case. Transfusion, 2013, 53, 1266-1269.	1.6	18
39	Detecting hypermethylated fetal <i><scp>RASSF</scp>1<scp>A</scp></i> sequences in maternal plasma: implications for noninvasive paternity testing in pregnancy. Transfusion, 2013, 53, 1856-1858.	1.6	7
40	Polymorphism analysis and evaluation of 19 STR loci in the Han population of Southern China. Annals of Human Biology, 2013, 40, 191-196.	1.0	42
41	A paternity case with mutations at three CODIS core STR loci. Forensic Science International: Genetics, 2012, 6, e61-e62.	3.1	9
42	Increased Expression and Altered Methylation of HERVWE1 in the Human Placentas of Smaller Fetuses from Monozygotic, Dichorionic, Discordant Twins. PLoS ONE, 2012, 7, e33503.	2.5	28
43	Null alleles of the X and Y chromosomal amelogenin gene in a Chinese population. International Journal of Legal Medicine, 2012, 126, 513-518.	2.2	33
44	Predicting Human Age with Bloodstains by sjTREC Quantification. PLoS ONE, 2012, 7, e42412.	2.5	42
45	Non-invasive prenatal assessment of trisomy 21 by multiplexed maternal plasma DNA sequencing: large scale validity study. BMJ: British Medical Journal, 2011, 342, c7401-c7401.	2.3	641
46	Genetic polymorphisms of twelve X-chromosomal STR loci in Chinese Han population from Guangdong Province. Forensic Science International: Genetics, 2011, 5, e114-e116.	3.1	26
47	Detection and quantification of the age-related sjTREC decline in human peripheral blood. International Journal of Legal Medicine, 2011, 125, 603-608.	2.2	29
48	Maternal Plasma DNA Sequencing Reveals the Genome-Wide Genetic and Mutational Profile of the Fetus. Science Translational Medicine, 2010, 2, 61ra91.	12.4	878
49	Polymorphism analysis of 15 STR loci in a large sample of the Han population in southern China. Forensic Science International: Genetics, 2009, 4, e27-e29.	3.1	6
50	Development of a five ChX STRs loci typing system. International Journal of Legal Medicine, 2008, 122, 261-265.	2.2	17