

# Niels Morling

## List of Publications by Citations

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137  
papers

3,094  
citations

30  
h-index

50  
g-index

141  
ext. papers

3,650  
ext. citations

3.6  
avg, IF

5.39  
L-index

#	Paper	IF	Citations
137	Next generation sequencing and its applications in forensic genetics. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 18, 78-89	4.3	249
136	A global analysis of Y-chromosomal haplotype diversity for 23 STR loci. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 12, 12-23	4.3	171
135	Massively parallel sequencing of forensic STRs: Considerations of the DNA commission of the International Society for Forensic Genetics (ISFG) on minimal nomenclature requirements. <i>Forensic Science International: Genetics</i> , <b>2016</b> , 22, 54-63	4.3	148
134	Toward male individualization with rapidly mutating y-chromosomal short tandem repeats. <i>Human Mutation</i> , <b>2014</b> , 35, 1021-32	4.7	130
133	Recommendations of the DNA Commission of the International Society for Forensic Genetics (ISFG) on quality control of autosomal Short Tandem Repeat allele frequency databasing (STRidER). <i>Forensic Science International: Genetics</i> , <b>2016</b> , 24, 97-102	4.3	91
132	Second-generation sequencing of forensic STRs using the Ion Torrent® HID STR 10-plex and the Ion PGM®. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 14, 132-40	4.3	88
131	Evaluation of the Ion Torrent® HID SNP 169-plex: A SNP typing assay developed for human identification by second generation sequencing. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 12, 144-54	4.3	79
130	Second generation sequencing of three STRs D3S1358, D12S391 and D21S11 in Danes and a new nomenclature for sequenced STR alleles. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 12, 38-41	4.3	67
129	Paternity Testing Commission of the International Society of Forensic Genetics: recommendations on genetic investigations in paternity cases. <i>Forensic Science International</i> , <b>2002</b> , 129, 148-57	2.6	67
128	Analysis of 12 X-STRs in Greenlanders, Danes and Somalis using Argus X-12. <i>International Journal of Legal Medicine</i> , <b>2012</b> , 126, 121-8	3.1	62
127	Characterization of mutations and sequence variants in the D21S11 locus by next generation sequencing. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 8, 68-72	4.3	60
126	Evaluation of the Precision ID Ancestry Panel for crime case work: A SNP typing assay developed for typing of 165 ancestral informative markers. <i>Forensic Science International: Genetics</i> , <b>2017</b> , 28, 138-145	4.3	58
125	Typing of 30 insertion/deletions in Danes using the first commercial indel kit--Mentype® DIPplex. <i>Forensic Science International: Genetics</i> , <b>2012</b> , 6, e72-4	4.3	58
124	DNA commission of the International society for forensic genetics: Assessing the value of forensic biological evidence - Guidelines highlighting the importance of propositions: Part I: evaluation of DNA profiling comparisons given (sub-) source propositions. <i>Forensic Science International: Genetics</i> , <b>2018</b> , 36, 189-202	4.3	56
123	Genetic determinants of hair and eye colours in the Scottish and Danish populations. <i>BMC Genetics</i> , <b>2009</b> , 10, 88	2.6	51
122	Genetic investigations of sudden unexpected deaths in infancy using next-generation sequencing of 100 genes associated with cardiac diseases. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 817-22	5.3	44
121	Forensic and population genetic analyses of Danes, Greenlanders and Somalis typed with the Yfiler® Plus PCR amplification kit. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 16, 232-236	4.3	43

120	Statistical model for degraded DNA samples and adjusted probabilities for allelic drop-out. <i>Forensic Science International: Genetics</i> , <b>2012</b> , 6, 97-101	4.3	43
119	Genetic investigation of 100 heart genes in sudden unexplained death victims in a forensic setting. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 1797-1802	5.3	41
118	Forensic genetic SNP typing of low-template DNA and highly degraded DNA from crime case samples. <i>Forensic Science International: Genetics</i> , <b>2013</b> , 7, 345-52	4.3	41
117	A method for the analysis of 32 X chromosome insertion deletion polymorphisms in a single PCR. <i>International Journal of Legal Medicine</i> , <b>2012</b> , 126, 97-105	3.1	39
116	The discrete Laplace exponential family and estimation of Y-STR haplotype frequencies. <i>Journal of Theoretical Biology</i> , <b>2013</b> , 329, 39-51	2.3	39
115	Massively parallel pyrosequencing of the mitochondrial genome with the 454 methodology in forensic genetics. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 12, 30-7	4.3	36
114	Allelic drop-out probabilities estimated by logistic regression--further considerations and practical implementation. <i>Forensic Science International: Genetics</i> , <b>2012</b> , 6, 263-7	4.3	36
113	ISO 17025 validation of a next-generation sequencing assay for relationship testing. <i>Electrophoresis</i> , <b>2016</b> , 37, 2822-2831	3.6	35
112	miR-125b induces cellular senescence in malignant melanoma. <i>BMC Dermatology</i> , <b>2014</b> , 14, 8	2.1	34
111	Evaluation of DNA variants associated with androgenetic alopecia and their potential to predict male pattern baldness. <i>PLoS ONE</i> , <b>2015</b> , 10, e0127852	3.7	34
110	MDM2 inhibitor nutlin-3a induces apoptosis and senescence in cutaneous T-cell lymphoma: role of p53. <i>Journal of Investigative Dermatology</i> , <b>2012</b> , 132, 1487-96	4.3	34
109	Increasing the reference populations for the 55 AISNP panel: the need and benefits. <i>International Journal of Legal Medicine</i> , <b>2017</b> , 131, 913-917	3.1	32
108	Analysis of ancestry informative markers in three main ethnic groups from Ecuador supports a trihybrid origin of Ecuadorians. <i>Forensic Science International: Genetics</i> , <b>2017</b> , 31, 29-33	4.3	30
107	Introduction of the Python script STRinNGS for analysis of STR regions in FASTQ or BAM files and expansion of the Danish STR sequence database to 11 STRs. <i>Forensic Science International: Genetics</i> , <b>2016</b> , 21, 68-75	4.3	29
106	Whose DNA is this? How relevant a question? (a note for forensic scientists). <i>Forensic Science International: Genetics</i> , <b>2013</b> , 7, 467-70	4.3	27
105	Multiplex PCR with minisequencing as an effective high-throughput SNP typing method for formalin-fixed tissue. <i>Electrophoresis</i> , <b>2007</b> , 28, 2361-7	3.6	27
104	DNA commission of the International society for forensic genetics: Assessing the value of forensic biological evidence - Guidelines highlighting the importance of propositions. Part II: Evaluation of biological traces considering activity level propositions. <i>Forensic Science International: Genetics</i> , <b>2020</b> , 44, 102186	4.3	27
103	Evaluation of the predictive capacity of DNA variants associated with straight hair in Europeans. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 19, 280-288	4.3	26

102	The half-life of 25(OH)D after UVB exposure depends on gender and vitamin D receptor polymorphism but mainly on the start level. <i>Photochemical and Photobiological Sciences</i> , <b>2017</b> , 16, 985-995	4.2	25
101	Sequencing of 231 forensic genetic markers using the MiSeq FGx forensic genomics system: an evaluation of the assay and software. <i>Forensic Sciences Research</i> , <b>2018</b> , 3, 111-123	3.6	25
100	The effect of gender on eye colour variation in European populations and an evaluation of the IrisPlex prediction model. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 11, 1-6	4.3	25
99	Genetic analyses of the human eye colours using a novel objective method for eye colour classification. <i>Forensic Science International: Genetics</i> , <b>2013</b> , 7, 508-15	4.3	25
98	Performance of two 17 locus forensic identification STR kits-Applied Biosystems's AmpFIBTR and NGMSelect and Promega's PowerPlex ES17 kits. <i>Forensic Science International: Genetics</i> , <b>2012</b> , 6, 523-31	4.3	25
97	Characterization of a new HLA-G allele encoding a nonconservative amino acid substitution in the alpha3 domain (exon 4) and its relevance to certain complications in pregnancy. <i>Immunogenetics</i> , <b>2001</b> , 53, 48-53	3.2	25
96	Towards broadening Forensic DNA Phenotyping beyond pigmentation: Improving the prediction of head hair shape from DNA. <i>Forensic Science International: Genetics</i> , <b>2018</b> , 37, 241-251	4.3	24
95	Kinship Analysis with Diallelic SNPs - Experiences with the SNPforID Multiplex in an ISO17025 Accredited Laboratory. <i>Transfusion Medicine and Hemotherapy</i> , <b>2012</b> , 39, 195-201	4.2	23
94	Evaluation of Forensic DNA Traces When Propositions of Interest Relate to Activities: Analysis and Discussion of Recurrent Concerns. <i>Frontiers in Genetics</i> , <b>2016</b> , 7, 215	4.5	23
93	New clues to the evolutionary history of the main European paternal lineage M269: dissection of the Y-SNP S116 in Atlantic Europe and Iberia. <i>European Journal of Human Genetics</i> , <b>2016</b> , 24, 437-41	5.3	22
92	Frequencies of HID-ion ampliseq ancestry panel markers among greenlanders. <i>Forensic Science International: Genetics</i> , <b>2016</b> , 24, 60-64	4.3	22
91	Importance of nonsynonymous OCA2 variants in human eye color prediction. <i>Molecular Genetics &amp; Genomic Medicine</i> , <b>2016</b> , 4, 420-30	2.3	21
90	Major inter-personal variation in the increase and maximal level of 25-hydroxy vitamin D induced by UVB. <i>Photochemical and Photobiological Sciences</i> , <b>2016</b> , 15, 536-45	4.2	20
89	Technological innovations in forensic genetics: social, legal and ethical aspects. <i>Recent Advances in DNA &amp; Gene Sequences</i> , <b>2014</b> , 8, 98-103		19
88	Evaluation of four automated protocols for extraction of DNA from FTA cards. <i>Journal of the Association for Laboratory Automation</i> , <b>2013</b> , 18, 404-10		19
87	Platelet alloimmunization after transfusion. A prospective study in 117 heart surgery patients. <i>Vox Sanguinis</i> , <b>1997</b> , 72, 238-41	3.1	19
86	Graft-derived anti-HPA-2b production after allogeneic bone-marrow transplantation. <i>British Journal of Haematology</i> , <b>1994</b> , 86, 651-3	4.5	19
85	High-throughput sequencing of forensic genetic samples using punches of FTA cards with buccal swabs. <i>BioTechniques</i> , <b>2016</b> , 61, 149-51	2.5	19

84	PCR in forensic genetics. <i>Biochemical Society Transactions</i> , <b>2009</b> , 37, 438-40	5.1	18
83	Comparison of manual and automated AmpliSeq workflows in the typing of a Somali population with the Precision ID Identity Panel. <i>Forensic Science International: Genetics</i> , <b>2017</b> , 31, 118-125	4.3	17
82	Quantification of massively parallel sequencing libraries - a comparative study of eight methods. <i>Scientific Reports</i> , <b>2018</b> , 8, 1110	4.9	17
81	Stutter analysis of complex STR MPS data. <i>Forensic Science International: Genetics</i> , <b>2018</b> , 35, 107-112	4.3	17
80	Collaborative EDNAP exercise on the IrisPlex system for DNA-based prediction of human eye colour. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 11, 241-51	4.3	17
79	Forensic genetics. <i>Lancet, The</i> , <b>2004</b> , 364 Suppl 1, s10-1	4.0	17
78	A comparison of genetic findings in sudden cardiac death victims and cardiac patients: the importance of phenotypic classification. <i>Europace</i> , <b>2015</b> , 17, 350-7	3.9	16
77	Identifying contributors of DNA mixtures by means of quantitative information of STR typing. <i>Journal of Computational Biology</i> , <b>2012</b> , 19, 887-902	1.7	16
76	Characterization of mutations and sequence variations in complex STR loci by second generation sequencing. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2013</b> , 4, e218-e219	0.5	15
75	Forensic and phylogeographic characterisation of mtDNA lineages from Somalia. <i>International Journal of Legal Medicine</i> , <b>2012</b> , 126, 573-9	3.1	15
74	DNA polymorphism of HLA class II genes in systemic lupus erythematosus. <i>Tissue Antigens</i> , <b>1994</b> , 43, 34-7		15
73	HLA-DPB1 typing with polymerase chain reaction and restriction fragment length polymorphism technique in Danes. <i>Tissue Antigens</i> , <b>1992</b> , 40, 140-4		14
72	Decrease DNA contamination in the laboratories. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2017</b> , 6, e577-e578	0.5	13
71	Weight of the evidence of genetic investigations of ancestry informative markers. <i>Theoretical Population Biology</i> , <b>2018</b> , 120, 1-10	1.2	13
70	Peopling of the North Circumpolar Region--insights from Y chromosome STR and SNP typing of Greenlanders. <i>PLoS ONE</i> , <b>2015</b> , 10, e0116573	3.7	13
69	The peopling of Greenland: further insights from the analysis of genetic diversity using autosomal and X-chromosomal markers. <i>European Journal of Human Genetics</i> , <b>2015</b> , 23, 245-51	5.3	13
68	SPERM HY-LITER for the identification of spermatozoa from sexual assault evidence. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 12, 161-7	4.3	12
67	The role of the glutathione S-transferase genes GSTT1, GSTM1, and GSTP1 in acetaminophen-poisoned patients. <i>Clinical Toxicology</i> , <b>2012</b> , 50, 27-33	2.9	12

66	Reinvestigations of six unusual paternity cases by typing of autosomal single-nucleotide polymorphisms. <i>Transfusion</i> , <b>2012</b> , 52, 425-30	2.9	12
65	Statistical modelling of Ion PGM HID STR 10-plex MPS data. <i>Forensic Science International: Genetics</i> , <b>2017</b> , 28, 82-89	4.3	11
64	Analysis of mainland Japanese and Okinawan Japanese populations using the precision ID Ancestry Panel. <i>Forensic Science International: Genetics</i> , <b>2018</b> , 33, 106-109	4.3	11
63	Whole genome and transcriptome sequencing of post-mortem cardiac tissues from sudden cardiac death victims identifies a gene regulatory variant in NEXN. <i>International Journal of Legal Medicine</i> , <b>2019</b> , 133, 1699-1709	3.1	11
62	Genomic and immunohistochemical characterisation of a lacrimal gland oncocytoma and review of literature. <i>Oncology Letters</i> , <b>2017</b> , 14, 4176-4182	2.6	11
61	Identifying the most likely contributors to a Y-STR mixture using the discrete Laplace method. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 15, 76-83	4.3	11
60	Pigment genes not skin pigmentation affect UVB-induced vitamin D. <i>Photochemical and Photobiological Sciences</i> , <b>2019</b> , 18, 448-458	4.2	10
59	A study of the peopling of Greenland using next generation sequencing of complete mitochondrial genomes. <i>American Journal of Physical Anthropology</i> , <b>2016</b> , 161, 698-704	2.5	10
58	A report of the 2009-2011 paternity and relationship testing workshops of the English Speaking Working Group of the International Society For Forensic Genetics. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 9, e1-2	4.3	10
57	Ancestry prediction efficiency of the software GenoGeographer using a z-score method and the ancestry informative markers in the Precision ID Ancestry Panel. <i>Forensic Science International: Genetics</i> , <b>2020</b> , 44, 102154	4.3	10
56	Evaluation of the iPLEX <sup>®</sup> ADME PGx Pro Panel and allele frequencies of pharmacogenetic markers in Danes. <i>Clinical Biochemistry</i> , <b>2016</b> , 49, 1299-1301	3.5	8
55	Cluster analysis of European Y-chromosomal STR haplotypes using the discrete Laplace method. <i>Forensic Science International: Genetics</i> , <b>2014</b> , 11, 182-94	4.3	8
54	On the Bayesian approach to forensic age estimation of living individuals. <i>Forensic Science International</i> , <b>2017</b> , 281, e24-e29	2.6	8
53	Detection of fetal-specific DNA after enrichment for trophoblasts using the monoclonal antibody LK26 in model systems but failure to demonstrate fetal DNA in maternal peripheral blood. <i>Prenatal Diagnosis</i> , <b>1999</b> , 19, 271-8	3.2	8
52	eDNA <sup>™</sup> An expert software system for comparison and evaluation of DNA profiles in forensic casework. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2015</b> , 5, e400-e402	0.5	7
51	Estimating Y-STR allelic drop-out rates and adjusting for interlocus balances. <i>Forensic Science International: Genetics</i> , <b>2013</b> , 7, 327-36	4.3	7
50	Targeted molecular genetic testing in young sudden cardiac death victims from Western Denmark. <i>International Journal of Legal Medicine</i> , <b>2020</b> , 134, 111-121	3.1	7
49	Analysis of 49 autosomal SNPs in an Iraqi population. <i>Forensic Science International: Genetics</i> , <b>2013</b> , 7, 198-9	4.3	6

48	Sequence variants of allele 22 and 23 of DYS635 causing different stutter rates. <i>Forensic Science International: Genetics</i> , <b>2012</b> , 6, e161-2	4.3	6
47	Refining the genetic portrait of Portuguese Roma through X-chromosomal markers. <i>American Journal of Physical Anthropology</i> , <b>2012</b> , 148, 389-94	2.5	6
46	Characterization of sequence variations in the D21S11 locus in Danes, Somalis and Greenlanders by second generation sequencing. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2013</b> , 4, e302-e303	0.5	5
45	Estimating drop-out probabilities of STR alleles accounting for stutters, detection threshold truncation and degradation. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2013</b> , 4, e51-e52	0.5	5
44	Next-generation sequencing of multiple individuals per barcoded library by deconvolution of sequenced amplicons using endonuclease fragment analysis. <i>BioTechniques</i> , <b>2014</b> , 57, 91-4	2.5	5
43	Typing of 49 autosomal SNPs by single base extension and capillary electrophoresis for forensic genetic testing. <i>Methods in Molecular Biology</i> , <b>2012</b> , 830, 87-107	1.4	5
42	Pigmentary Markers in Danes--Associations with Quantitative Skin Colour, Nevi Count, Familial Atypical Multiple-Mole, and Melanoma Syndrome. <i>PLoS ONE</i> , <b>2016</b> , 11, e0150381	3.7	5
41	Association between brown eye colour in rs12913832:GG individuals and SNPs in TYR, TYRP1, and SLC24A4. <i>PLoS ONE</i> , <b>2020</b> , 15, e0239131	3.7	5
40	GenoGeographer DA tool for genogeographic inference. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2017</b> , 6, e463-e465	0.5	4
39	Non-invasive prenatal paternity testing using a standard forensic genetic massively parallel sequencing assay for amplification of human identification SNPs. <i>International Journal of Legal Medicine</i> , <b>2019</b> , 133, 1361-1368	3.1	4
38	Body fluid identification of blood, saliva and semen using second generation sequencing of micro-RNA. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2013</b> , 4, e204-e205	0.5	4
37	Biomek 3000: the workhorse in an automated accredited forensic genetic laboratory. <i>Journal of the Association for Laboratory Automation</i> , <b>2012</b> , 17, 378-86		4
36	Sequences of microvariant/bff-ladder/STR alleles. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2011</b> , 3, e204-e205	0.5	4
35	Repeated extraction of DNA from FTA cards. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2011</b> , 3, e345-e346	0.5	4
34	Prediction of Eye Colour in Scandinavians Using the EyeColour 11 (EC11) SNP Set. <i>Genes</i> , <b>2021</b> , 12,	4.2	4
33	Yfiler Plus population samples and dilution series: stutters, analytic thresholds, and drop-out probabilities. <i>International Journal of Legal Medicine</i> , <b>2017</b> , 131, 1503-1511	3.1	3
32	The multivariate Dirichlet-multinomial distribution and its application in forensic genetics to adjust for subpopulation effects using the Ecorrection. <i>Theoretical Population Biology</i> , <b>2015</b> , 105, 24-32	1.2	3
31	Monozygotic twins discordant for narcolepsy type 1 and multiple sclerosis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , <b>2016</b> , 3, e249	9.1	3

30	Modelling noise in second generation sequencing forensic genetics STR data using a one-inflated (zero-truncated) negative binomial model. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2015</b> , 5, e416-e417	0.5	3
29	Evaluation of Y-STR analyses of sperm cell negative vaginal samples. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2011</b> , 3, e141-e142	0.5	3
28	A comparative study of single nucleotide variant detection performance using three massively parallel sequencing methods. <i>PLoS ONE</i> , <b>2020</b> , 15, e0239850	3.7	3
27	Associations between second to fourth digit ratio, cortisol, vitamin D, and body composition among Polish children. <i>Scientific Reports</i> , <b>2021</b> , 11, 7029	4.9	3
26	Sequence variants in muscle tissue-related genes may determine the severity of muscle contractures in cerebral palsy. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , <b>2019</b> , 180, 12-24	3.5	3
25	The stratum corneum transcriptome in atopic dermatitis can be assessed by tape stripping. <i>Journal of Dermatological Science</i> , <b>2021</b> , 101, 14-21	4.3	3
24	Modelling allelic drop-outs in STR sequencing data generated by MPS. <i>Forensic Science International: Genetics</i> , <b>2018</b> , 37, 6-12	4.3	3
23	Serum 25(OH)D levels after oral vitamin D supplementation and UVB exposure correlate. <i>Photodermatology Photoimmunology and Photomedicine</i> , <b>2019</b> , 35, 344-353	2.4	2
22	Identical twins in forensic genetics - Epidemiology and risk based estimation of weight of evidence. <i>Science and Justice - Journal of the Forensic Science Society</i> , <b>2015</b> , 55, 408-14	2	2
21	Skin pigmentation and genetic variants in an admixed Brazilian population of primarily European ancestry. <i>International Journal of Legal Medicine</i> , <b>2020</b> , 134, 1569-1579	3.1	2
20	Targeted exon sequencing in deceased schizophrenia patients in Denmark. <i>International Journal of Legal Medicine</i> , <b>2020</b> , 134, 135-147	3.1	2
19	Forensic application and genetic diversity of 21 autosomal STR loci in five major population groups of Pakistan. <i>International Journal of Legal Medicine</i> , <b>2021</b> , 135, 775-777	3.1	2
18	Evaluation of a custom GeneRead <sup>™</sup> massively parallel sequencing assay with 210 ancestry informative SNPs using the Ion S5 <sup>™</sup> and MiSeq platforms. <i>Forensic Science International: Genetics</i> , <b>2021</b> , 50, 102411	4.3	2
17	Cortisol concentration affects fat and muscle mass among Polish children aged 6-13 years. <i>BMC Pediatrics</i> , <b>2021</b> , 21, 365	2.6	2
16	Association of saliva 25(OH)D concentration with body composition and proportion among pre-pubertal and pubertal Polish children. <i>American Journal of Human Biology</i> , <b>2020</b> , 32, e23397	2.7	2
15	Reproducibility of the Infinium methylationEPIC BeadChip assay using low DNA amounts.. <i>Epigenetics</i> , <b>2022</b> , 1-10	5.7	2
14	Results of the 2015 Relationship Testing Workshop of the English Speaking Working Group. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2015</b> , 5, e320-e321	0.5	1
13	Study of 25 X-chromosome Single Nucleotide Polymorphisms in African and Asian populations. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2011</b> , 3, e139-e140	0.5	1



12	Survival of fetuses and viruses: universal mechanisms of co-existence with an immunological potent host. <i>American Journal of Reproductive Immunology</i> , <b>1999</b> , 41, 353-5	3.8	1
11	The transcriptome of hand eczema assessed by tape stripping. <i>Contact Dermatitis</i> , <b>2021</b> ,	2.7	1
10	Sequencing of human identification markers in an Uyghur population using the MiSeq FGxTM Forensic Genomics System. <i>Forensic Sciences Research</i> , <b>2020</b> , 1-9	3.6	1
9	Evaluation of the Precision of Ancestry Inferences in South American Admixed Populations. <i>Frontiers in Genetics</i> , <b>2020</b> , 11, 966	4.5	1
8	Differential Methylation in the Regulatory Region in Sudden Unexplained Death and Sudden Unexpected Death in Epilepsy. <i>International Journal of Molecular Sciences</i> , <b>2021</b> , 22,	6.3	1
7	Genetic investigations of 100 inherited cardiac disease-related genes in deceased individuals with schizophrenia. <i>International Journal of Legal Medicine</i> , <b>2021</b> , 135, 1395-1405	3.1	0
6	Use of Next-Generation Sequencing in Forensic Genetics <b>2017</b> , 1-9		
5	The effect of wild card designations and rare alleles in forensic DNA database searches. <i>Forensic Science International: Genetics</i> , <b>2015</b> , 16, 98-104	4.3	
4	Group specific component in serum and otosclerosis: no association. <i>Acta Oto-Laryngologica</i> , <b>1994</b> , 114, 303-4	1.6	
3	GENETIC PORTRAIT OF THE PUNJABI POPULATION FROM PAKISTAN USING THE PRECISION ID ANCESTRY PANEL. <i>Forensic Science International: Genetics Supplement Series</i> , <b>2019</b> , 7, 87-89	0.5	
2	A 48-plex autosomal SNP GenPlex assay for human individualization and relationship testing. <i>Methods in Molecular Biology</i> , <b>2012</b> , 830, 73-85	1.4	
1	Anthropological analyses of 30 insertion/deletion autosomal markers in five major ethnic groups of Pakistan. <i>Forensic Sciences Research</i> , 1-5	3.6	