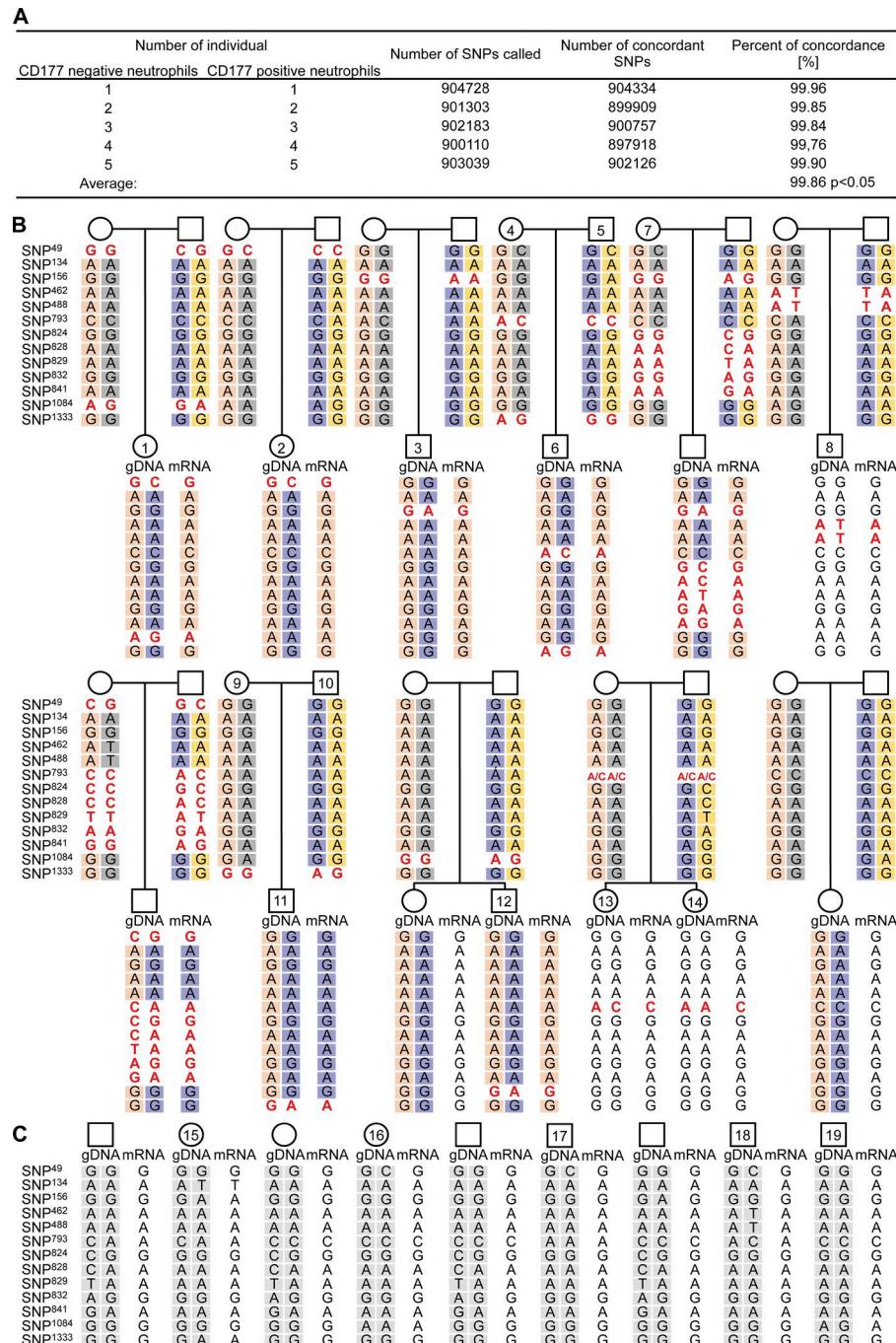
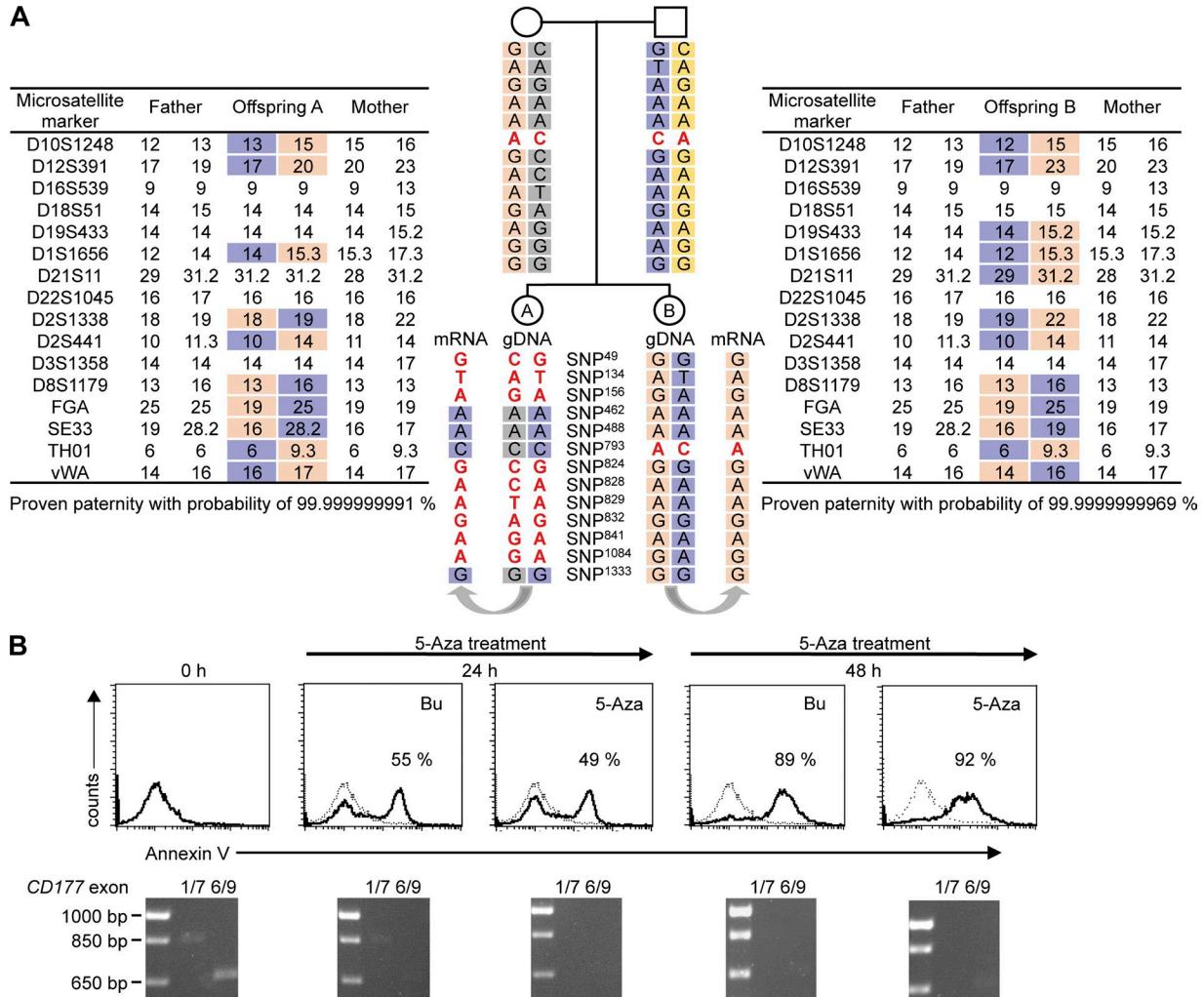


SUPPLEMENTAL MATERIAL

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**Figure S1. Genome-wide SNPs analysis of CD177<sup>neg</sup> and CD177<sup>pos</sup> neutrophil subsets and haplotype analysis of the CD177 gene.** (A) Genome-wide SNP analysis using an Affymetrix SNP array 6.0 was performed in the two sorted CD177 neutrophil subsets from five different individuals. Shown are the number of SNPs called and the number of concordant SNPs between the two subsets in a given individual (five biological replicates, 99.9% concordance by Fisher's exact test). (B) Haplotype analysis of the CD177 gene. Analyses were performed in 12 families with 17 offspring, and (C) 9 additional individuals. 13 SNPs are depicted in the haplotype analysis that were selected because they were informative with respect to the allelic choice in at least one trio. The informative parental haplotype that is inherited by the offspring is boxed in light orange (maternal) and blue (paternal), and the SNP positions are indicated. SNPs that were informative for the allelic choice in a given trio are marked in red. Individuals numbered 1–19 are homozygous for g7497A (SNP<sup>829</sup>). Note that these individuals had several informative SNPs indicating monoallelic CD177 expression (see Discussion).



**Figure S2. Haplotype analysis, sibling testing of an informative family with respect to the allelic choice of the CD177 MAE, and 5-Aza-2-deoxycytidin (5-Aza) treatment of primary human neutrophils.** (A) Haplotype analysis of a family with two siblings. The informative parental haplotype that is inherited by the offspring is boxed in light orange (maternal) and blue (paternal). SNPs that were informative for the allelic choice are marked in red. One offspring showed MAE of the paternal (left) and one of the maternal allele (right). The corresponding tables show the microsatellite markers obtained by PowerPlex ESX-17 for both offspring establishing full-sibling status. Red and blue color-coding as described above. (B) Primary human neutrophils were treated with buffer (Bu) or 10  $\mu$ M 5-Aza-2-deoxycytidin (5-Aza) for 24 h and 48 h, respectively. The percentage of Annexin V-positive apoptotic neutrophils was assessed by flow cytometry, and histograms are depicted. mRNA was purified and subjected to CD177 PCR. Freshly isolated cells, but not 5-Aza-treated cells, showed a PCR product. One of two representative experiments is given.

Table S1. SNPs identified in the *CD177* exons by sequencing

SNP position	Allele	SNP-ID no.	Location in the <i>CD177</i> gene	Genomic position on chromosome 19 (hg19)
SNP <sup>49</sup>	G/C	rs45441892	exon 1	<b>43.857.873-43.857.873</b>
SNP <sup>134</sup>	A/T	rs45553433	exon 2	<b>43.858.044-43.858.044</b>
SNP <sup>156</sup>	G/A	rs45571738	exon 2	<b>43.858.066-43.858.066</b>
SNP <sup>387</sup>	C/T	no	exon 3	43.858.510-43.858.510
SNP <sup>462</sup>	A/T	no	exon 4	<b>43.859.853-43.859.853</b>
SNP <sup>488</sup>	A/T	no	exon 4	<b>43.859.879-43.859.879</b>
SNP <sup>593</sup>	T/G	rs71337594	exon 5	43.860.192-43.860.192
SNP <sup>652</sup>	G/A	rs199668750	exon 5	43.860.251-43.860.251
SNP <sup>656</sup>	T/G	rs200662237	exon 5	43.860.255-43.860.255
SNP <sup>664</sup>	T/G	rs73559882	exon 6	43.864.419-43.864.419
SNP <sup>667</sup>	C/T	no	exon 6	43.864.422-43.864.422
SNP <sup>752</sup>	T/C	rs57802244	exon 6	43.864.507-43.864.507
SNP <sup>793</sup>	C/A	<b>rs10425835</b>	<b>exon 6</b>	<b>43.864.548-43.864.548</b>
SNP <sup>798</sup>	T/C	no	exon 6	43.864.553-43.864.553
SNP <sup>824</sup>	C/G	rs200660811	exon 7	43.865.316-43.865.316
SNP <sup>828</sup>	C/A	rs587670082	exon 7	43.865.320-43.865.320
SNP <sup>829</sup>	T/A	rs201821720	exon 7	43.865.321-43.865.321
SNP <sup>832</sup>	A/G	rs200145410	exon 7	43.865.324-43.865.324
SNP <sup>841</sup>	G/A	rs201266439	exon 7	43.865.333-43.865.333
SNP <sup>1084</sup>	G/A	rs61625631	exon 8	43.865.692-43.865.692
SNP <sup>1114</sup>	C/T	rs61002457	exon 8	43.865.722-43.865.722
SNP <sup>1333</sup>	A/C/G/T	rs78718189	exon 9	43.866.449-43.866.449

SNP-ID is SNP identification number. SNP used for haplotype analysis are shown in bold.

Table S2. Probes in the *CD177* promoter and gene body that are covered by the Illumina Infinium HumanMethylation450 BeadChips used for genome-wide DNA methylation

Probe name	P-value	Student's t test	Mean (N)	Mean (P)	Mean ratio (N/P)	Mean diff (N-P)	UCSC ref gene	hg19 (bps) chr.19
cg22537604	7.10E-07	30.51	0.94	-0.80	3.3	1.74	<i>CD177</i>	43857074
cg15742245	2.98E-05	14.33	2.80	1.07	3.3	1.73	<i>CD177</i>	43857717
cg12032655	3.64E-05	13.76	-0.49	-1.58	2.1	1.09	<i>CD177</i>	43856746
cg05275595	4.92E-04	8.00	0.38	-0.21	1.5	0.59	<i>CD177</i>	43857874
cg11906444	4.44E-01	-0.83	1.57	1.64	1.0	-0.07	<i>CD177</i>	43858174

cg means CpG cluster number.

Table S3. Oligonucleotides for PCR assays

Name	5'-3' sequence
RACE Abridged Anchor Primer	GGCCACCGCGTCGACTAGTACGGGIIGGGIIGGGI
Abridged Universal Primer	GGCCACCGCGTCGACTAGTAC
Universal Amplification Primer	CUACUACUACUAGGCCACCGCGTCGACTAGTAC
5' RACE CD177-102R	CACTCTGGCAGTGGGAGGGATGAA
5' RACE CD177-265R	CCTTGGAGAGCACCAAGGCTCACT
5' RACE CD177-616R	GTGTCCCATTGAGCAGGTGCAAAC
qRT-PCR human CD177-F	TTGATGCTCATTGAGAGCGG
qRT-PCR human CD177-R	GCCCTCGTGAGCCCT
qRT-PCR human CD177-probe	FAM-CCCCAAGTGAGCTGGTGCCTCC-TAMRA
qRT-PCR human 18S-F	ACATCCAAGGAAGGCAGCAG
qRT-PCR human 18S-R	TTTCGTCACACCTCCCG
qRT-PCR human 18S-probe	FAM-CGCGCAATTACCCACTCCGAC-TAMRA
CD177-43.857.788-F	GCTGGCTGCTTAAGGGCTGGTATAA
CD177-43.870.898-R	CATCCGTACGGGAAATCAGACTTG
CD177-43.857.825-F	CTGCTGAAAAAGCAGAAAGAGA
CD177-43.858.437-R	ATCCGGTGCAGTGACG
CD177-43.858.316-F	AGGCCTGACCTCCATCCT
CD177-43.859.013-R	CTGTGTTACAAGCTGTTC
CD177-43.859.487-F	AACTCCGTACCTCGTGATCC
CD177-43.860.178-R	GCATCCCTGAGCTCTCAGAT
CD177-43.860.080-F	CTCTGAGGGTTGGTGGTC
CD177-43.860.770-R	CCACCATGGCTCTGT
CD177-43.864.250-F	TGGGATTTGACTCCCAAGT
CD177-43.864.943-R	TGTTTGTTCTGTTCTCAAGG
CD177-43.864.829-F	TCAGTGGCTTCATACATTCTGG
CD177-43.865.520-R	GTGTCCTGGCCTCATTTCTC
CD177-43.865.418-F	CCTCGGACCTGTGCAATAGT
CD177-43.866.126-R	AGTCCAGCTCTCAGACC
CD177-43.866.021-F	CTCTGGACTCCTGGGTCTGA
CD177-43.866.676-R	CCTCTCCCAGGGCAAGTC
CD177-43.857.862-F	GGGTCATGAGCGGGTATTAC
CD177-43.864.453-F	CACACGGAAACTGGCTCAAGAAC
CD177-43.865.382-R	GGAGGGGCTGAGTGGATGGTGG
CD177-43.865.688-F	GGGCGCCACTCATGTTATG
CD177-43.866.478-R	ATAGAGTTAGCAGGAAGGGAAAC
Pyro-SNP49-134-156-F	AGCCACAGACGGGTATGAG
Biotin-Pyro-SNP49-134-156-R	Biotin-CTGGTGTCTTAGGGGTCATTG
PyroSeq-SNP49-134-156-F	AGACGGGTATGAGC
Pyro-SNP462-488-F	AACTCCCTCCGCTTGG
Biotin-Pyro-SNP462-488-R	Biotin-GGTCCCCTGGGGCAGAT
PyroSeq-SNP462-488-F	CCAGTCCTGCTCATG
Pyro-SNP793-F	GGCAGGTGTCAAGGAGAC
Biotin-Pyro-SNP793-R	Biotin-CCAGGGTTGATGTGAGTCTAC
PyroSeq-SNP793-F	GAGACGCTGCTGTC
Biotin-Pyro-SNP1084-F	Biotin-GCTCCCCCGAATGACCT
Pyro-SNP1084-R	AGCCCACCTCTGAGAGATGA
PyroSeq-SNP1084-F	CCATCATAACAATGAGTGG
Luciferase-P1-F	TGCGTGGGAATCCGCCCTCCCTCTTGCTT
Luciferase-P2-F	TGCGTGGGATCCCTCTGGTGGTAGGAAACTGG
Luciferase-P3-F	TGCGTGGGATCCGGGTCAGATGCCACAGTT
Luciferase-P4-F	TGCGTGGGATCCAAGAAAGGGGGTCCCTGATAG
Luciferase-P5-F	TGCGTGGGATCCGGCTGCTTAAGGGCTGGTAT
Luciferase-P6-F	TGCGTGGGATCCGTATGAGCGGGTATTACT
Luciferase-R	CACGCAGGATCCACAGGGATCAAGGGACTGA
M13-For	GTAAAACGACGGCCAG
M13-Rev	CAGGAAACAGCTATGAC
ChIP1-43.856.451-F	TTGCAAGTCACATTCAATTCTCT
ChIP1-43.856.582-R	AGGAAGGAACCTGGGGTATCT
ChIP2-43.857.044-F	GGGAAGACAGTGGAGGGTAGA
ChIP2-43.857.159-R	CCACCTCTGACCCAGATG
ChIP3-43.857.186-F	ACTGCCTTCCCAAACACTG
ChIP3-43.857.307-R	AGAAACTGTGGCATCTTGACC
ChIP4-43.857.630-F	GTCCCTGGACCCCTCATCTG

Table S3. Oligonucleotides for PCR assays (*Continued*)

Name	5'-3' sequence
ChIP4-43.857.720-R	GCCGCAGTGGCCTATCAG
ChIP5-43.857.792-F	GCTGCTTAAGGGCTGGTATAAA
ChIP5-43.857.896-R	CAGGAGGGCCAGCAGTAATA
ChIP6-43.857.862-F	GGGTCAATGAGCGCGGTATTAC
ChIP6-43.858.070-R	GGGCAGGTGGACACACCTT
Ctrl-43.870.329-F	CTCATCCAGGAGTTATCGAGAATA
Ctrl-43.870.406-R	GAAATGAACCACAGCACTTCC
hACTgProm-H3K4-F	ACCGGCAGAGAACCGCGA
hACTgProm-H3K4-R	CGGAAAGATGCCATATATGGAC
hALDOA-H3K4-F	TCCTGGCAAGATAAGGAGTTGAC
hALDOA-H3K4-R	ACACACGATAGCCCTAGCAGTTC

ACT is actin and ALDOA aldolase A.