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Neutrophil Transmigration Mediated by the Neutrophil-Specific Antigen CD177 Is Influenced by the Endothelial S_{536}-N Dimorphism of Platelet Endothelial Cell Adhesion Molecule-1


The human neutrophil-specific adhesion molecule CD177 (also known as the NB1 alloantigen) becomes upregulated on the cell surface in a number of inflammatory settings. We recently showed that CD177 functions as a novel heterophilic counterreceptor for the endothelial junctional protein PECAM-1 (CD31), an interaction that is mediated by membrane-proximal PECAM-1 IgD6, which is known to harbor an S_{536}-N single nucleotide polymorphism of two major isoforms V_{98}N_{536}G_{643} and L_{98}S_{536}R_{643} and a yet-to-be-determined region on CD177. In vitro transendothelial migration experiments revealed that CD177+ neutrophils migrated significantly faster through HUVECs expressing the LSR, compared with the VNG, allelic variant of PECAM-1 and that this correlated with the decreased ability of anti-PECAM-1 Ab of ITIM tyrosine phosphorylation in HUVECs expressing the LSR allelic variant relative to the VNG allelic variant. Moreover, engagement of CD177 with rCD177-Fc (to mimic heterophilic CD177 binding) suppressed Ab-induced tyrosine phosphorylation to a greater extent in cells expressing the LSR isoform compared with the VNG isoform, with a corresponding increased higher level of β-catenin phosphorylation. These data suggest that heterophilic PECAM-1/CD177 interactions affect the phosphorylation state of PECAM-1 and endothelial cell junctional integrity in such a way as to facilitate neutrophil transmigration in a previously unrecognized allele-specific manner. The Journal of Immunology, 2010, 184: 000–000.

Neutrophils are the most abundant leukocytes in blood and function as the first line of defense in the innate immune response. Neutrophils detect bacterial components, such as LPS and iMLP, via Toll-like or G-protein coupled receptors (1, 2), resulting in upregulation of migratory activities and neutrophil accumulation at sites of acute inflammation, vascular injury, or infection.

Neutrophil recruitment is a tightly regulated process and involves a multistep cascade of adhesive and migratory events that are mediated by three classes of adhesion receptors: selectins, integrins, and adhesion receptors of the Ig superfamily (3–5). One important Ig superfamily member is PECAM-1, a cell adhesion and signaling receptor that is expressed on platelets, monocytes, neutrophils and some T cells, as well as abundantly at endothelial cell–cell junctions (5, 6). PECAM-1 is composed of six extracellular IgDs (IgD1–IgD6), and IgD1 is known to mediate cation-independent homophilic interactions that play an important role during monocyte and neutrophil transendothelial migration (7, 8).

In addition to PECAM-1 IgD1-mediated homophilic interactions, we recently reported that CD177 (also known as NB1 Ag), a neutrophil-specific 58–64-kDa GPI-anchored member of cysteine-rich Ly-6 family, functions as a novel heterophilic binding partner that engages PECAM-1 in membrane-proximal IgD6 (9). A characteristic feature of CD177 is its variable expression on the surface of neutrophil subpopulations; CD177+ neutrophils in individuals can vary from 0 to 100% (10). However, the molecular basis of heterogeneous CD177 expression is not completely understood. In different clinical conditions, such as myeloproliferative disorder, essential thrombocythemia, and after G-CSF administration, CD177 becomes significantly upregulated on the neutrophil surface (10, 11).

Recently, three linked single nucleotide polymorphisms (SNPs) within the PECAM-1 gene have been identified that encode amino acid substitutions within IgD1 (exon 3; L_{98}V), IgD6 (exon 8; S_{536}N), and the cytoplasmic domain (exon 12; R_{643}G), resulting in the expression of two major PECAM-1 isoforms within the human population, termed LSR and VNG (frequency 0.42 versus 0.58) (12). Because the S_{536}N dimorphism is proximal to the CD177 binding site within IgD6 of PECAM-1, the purpose of the present investigation was to further examine the effect of the S_{536}N dimorphism on CD177-dependent neutrophil migration.

**Material and Methods**

**Abs and reagents**

mAbs PECAM-1.1 (specific for IgD5), PECAM-1.3 (against IgD1), and PECAM-1.2 (against IgD6) were produced and characterized as described (7). MAb Gi18 specific for IgD1 of PECAM-1 and Gi11 (against JAM-C) were generated in our laboratory (13). mAb CD62e for detection of E-selectin...
was purchased from Serotec (Düsseldorf, Germany); mAbs specific for phosphorylated epidermal growth factor receptor (Y1173), ERK (T202/Y204), and β-catenin (T41/S45) and mAb against ICAM-1 were purchased from Santa Cruz Biotechnology (Heidelberg, Germany). Hyridoma cells producing mAb 7D8 specific for CD177 was a gift from Dr. D. Stroence (National Institutes of Health, Bethesda, MD). A polyclonal anti-peptide Ab specific for the phosphorylated form of tyrosine 686 (anti-pY686) was produced in rabbits and used to detect PECAM-1 ITM phosphorylation. Unlabeled and labeled secondary Abs were obtained from Dako (Cytomation, Glostrup, Denmark). Protease inhibitor mixture and chemolattractants (MLP, TNF-α, leukotriene B4 (LTB4), and IL-8) were from Sigma-Aldrich (Taufkirchen, Germany). FITC-labeled albumin, calcine, and 2’-7’-bis-(2-carboxyethyl)-5-(and-6)-carboxyfluorescein acetoxymethyl ester were from Invitrogen (Karlsruhe, Germany). Pneumolysin (PLY) was a generous gift from Dr. T. Mitchell, Glasgow, U.K.

Genotyping of HUVECs and neutrophils for PECAM-1 polymorphisms

Total RNA was extracted using thepeqGOLD RNA Pure kit, according to the manufacturer’s instructions (peqLAB, Erlangen, Germany). RNA (1 μg) was reverse transcribed using a Ready-To-Go kit (GE Healthcare, Munich, Germany) with random hexamer primers, as recommended by the manufacturer (Invitrogen). Specific primer pairs encompassing nucleotides 188–673 (5’-TGGGCTGACTAGGCTTCATC-3’) and 5’-CAGAAATGTGGACGCTAGC-3’) and 1795–2287 (5’-GACTGTCGACTCATCCT-3’) and 5’-CCGTGCTGACTGAGTCCAG-3’) were designed according to the database (accession number: NM_000442.3, www.ncbi.nlm.nih.gov/nucleotide) to amplify PECAM-1 polymorphic regions containing the L98S, V356N, and R44G dimorphisms. An aliquot of 2 μl cDNA was amplified with 5 μl each primer (5 μM), 8 μl 2’-deoxynucleoside 5’-triphosphate (1.25 mM each 2’-deoxynucleoside 5’-triphosphate), and 1 μl AmpliTaq Gold Polymerase (5 U/μl; Applied Biosystems, Weiterstadt, Germany) in total volume of 50 μl for 30 cycles in a Thermal Cycler (Thermo Electron, Dreieich, Germany) under the following conditions: denaturation 95°C, 1 min; annealing 56°C, 1 min; and extension 72°C, 1 min. PCR products were sequenced with PCR primers (see above) using a Dye Terminator Cycle sequencing kit on an ABI 3100 DNA sequencer (Applied Biosystems).

Maintenance of HUVECs

Primary HUVECs derived from umbilical cords of single individuals were purchased from Lonza (Basel, Switzerland) and were cultured in endothelial cell basal medium (Lonza) in T-75 flasks, as recommended by the manufacturer. Cells were grown (~2) d up to 70–80% confluence and were subsequently used for experiments or were split by direct Accutase (PAAs Laboratory, Colbe, Germany). Only HUVECs maintained for less than four passages were used in this study. Aliquots of cells were frozen in liquid nitrogen.

Flow cytometric analysis of neutrophils and endothelial cells

Neutrophils were isolated from EDTA-anticoagulated whole blood from healthy donors by dextran sedimentation and gradient centrifugation. Aliquots of 107 washed fresh neutrophils were incubated with 20 μg FITC-labeled Gi18 or TD8 against PECAM-1 or CD177, respectively, for 30 min at 4°C. After washing with PBS, fluorescence-labeled neutrophils were analyzed on a FACS Calibur flow cytometer (BD Biosciences, Heidelberg, Germany). In some cases, neutrophils were stimulated with MLP (10-7 M) or PLY (25 ng/ml) for 30 min at room temperature prior to labeling with mAbs. HUVECs were isolated as described above and incubated with 20 μg FITC-labeled mAbs Gi18, Gi11, ICAM-1, or CD62e for 30 min at 4°C. Cells were washed with PBS and analyzed by flow cytometry, as above. For ICAM-1 and CD62e expression analysis, cells were stimulated with TNF-α (20 ng/ml), Immunol Tools, Friesoythe, Germany) for 2 h at 37°C prior to incubation with mAbs.

Permeability assays

The permeability of HUVECs was assessed by the passage of FITC-albumin (M, 66,000; Sigma-Aldrich), as previously described (14). Briefly, 3 × 105 HUVECs with known PECAM-1 phenotypes (LSR or VNG) were plated onto 48 Transwells fibronectin-coated 3-μm-pore-size polycarbonate membrane inserts (Costar, Cambridge, MA) and left for 2 d to form confluent monolayers. Confluency of cells was monitored by microscopy. Aliquots of 200 μl RPMI 1640 media (Life Technologies, Carlsbad, CA) containing FITC-albumin (40 μg/ml) were applied in the upper chamber. Every 15 min for 1 h, a sample from the bottom chamber was read in triplicate in a fluorescent microtiter plate reader (BioTek, Bad Friedrichshall, Germany). After incubation with 200 μl RPMI 1640 media containing 0.2 U/ml thrombin (Siemens Healthcare Diagnostic, Marburg, Germany) or histamine (3.2 × 10-5 mol/l; Sigma-Aldrich) at 37°C for 15 min, the passage of FITC-albumin was measured as described above. Additional experiments were conducted with HUVECs treated with 2.5 μg/ml rCD177/Fc or JAM-C/Fc for 30 min. In some experiments, cells were pretreated with 2.5 μg/ml PECAM-1 mAb against domain 5 (PECAM-1.1) at 37°C for 15 min prior to performing permeability assays. All data were from at least four independent experiments, and statistical analysis was performed using SPSS software (IBM, Munich, Germany).

Neutrophil transmigration assay

Neutrophil transmigration through HUVECs was performed as previously described (15). In brief, PECAM-1–phosphoryted HUVECs (1 × 105) were cultured on 6.5-mm Transwells with fibronectin-coated 3-μm-pore-size polycarbonate membrane inserts (Costar), as described above. Human neutrophils were obtained from CD177-phosphoryted blood donors and were isolated by Ficolli (Sigma-Aldrich) gradient centrifugation followed by hypotonic red cell lysis. Phosphotyrosine neutrophils (7.5 × 105 cells/ml) in RPMI 1640 medium (Life Technologies) were labeled with 30 μl 2’,7’-bis-(2-carboxyethyl)-5-(and-6)-carboxyfluorescein acetoxymethyl ester (2.5 μg/ml). Aliquots of 200 μl labeled neutrophils (5 × 105) were placed in the upper chamber, and cells were allowed to migrate at 37°C to the lower chamber containing MLP (10-8 M), IL-8 (50 ng/ml), or LTB4 (10-7 M) as chemolattractants. After 90 min, neutrophils found in the lower chamber were measured in triplicate in a fluorescent microparticle reader. For inhibition experiments, HUVECs were treated with PECAM-1.3 or -1.2 at a concentration of 2.5 μg/ml for 20 min at room temperature prior to migration assay. In some experiments, neutrophils were labeled with calcine (10 μg/ml) in RPMI 1640 at room temperature for 30 min prior stimulation with FMLP (10-7 M) or PLY (25 ng/ml).

Stimulation of HUVECs with rCD177/Fc, PECAM-1 mAbs, and H2O2

HUVECs were cultured on six-well plates (Greiner, Frickenhausen, Germany) for 48 h until confluent. The media were changed to endothelial basal medium (Life Technologies) before the experiments. HUVECs were incubated with rCD177/Fc (2.5 μg/ml), or mAb PECAM-1.1, PECAM-1.2 (2.5 μg/ml), PECAM-1.3 (2.5 μg/ml) for 60 min at 37°C, or with H2O2 (0.9 μM, Merck, Darmstadt, Germany) for 10 min at 37°C. Stimulation was followed by further incubation (30 min) with a secondary Ab to achieve cross-linking (final concentration 2.5 μg/ml). In some experiments, stimulation with PECAM-1.1 or H2O2 was followed by treatment with rCD177/Fc for 30 min at 37°C. Cells were then lysed with 100 μl 20 mM TBS, 1% Triton X-100, 7.5 μl Protease Inhibitor Cocktail, and 10 μl 5% EDTA for 30 min at 4°C. After centrifugation at 10,000 × g for 30 min at 4°C, supernatants were collected, and the protein concentration was measured using a bicinchoninic acid assay (Pierce, Rockford, IL). Cell lysates (10 μg) were analyzed by SDS-PAGE/Western blot analysis using mAb Gi18 against PECAM-1 (2.5 μg/ml), a rabbit polyclonal Ab specific for phosphorylated PECAM-1 (1 μg/ml) against phosphorylated tyrosine (pY) antibodies (1:1000 dilution). Proteins were visualized using HRP-labeled donkey anti-mouse IgG or goat anti-rabbit IgG followed by chemiluminescence detection (ECL plus; GE Healthcare, Piscataway, NJ).

Production of L98S536 and V98N536 PECAM-1 proteins

A construct encoding the extracellular domain of L98S536 PECAM-1 (residues 1–574) in the expression vector pcDNA3 was used as a template for the preparation of the V98N536 PECAM-1 using the QuikChange Site-Directed Mutagenesis Kit (Stratagene, La Jolla, CA). The vector was transfected into CHO cells, and high-expression clones were selected. The clones were grown in CELLline factory flasks as per the manufacturer’s instructions (Wilson Wolf Manufacturing, New Brighton, MN), and culture supernatant was collected. The PECAM-1 isoforms were purified using PECAM-1.3 affinity columns prepared with the Amino Link Plus Immobilization Kit (Pierce, Rockford, IL) and analyzed on SDS-PAGE gels for purity.

Production of CD177 protein

Full-length CD177 cDNA containing the human Ig Fc domain in the plg plus vector were cloned into pBluescript II (Stratagene, La Jolla, CA) containing PECAM-1.1 affinity columns prepared with the Amino Link Plus Immobilization Kit (Pierce, Rockford, IL) and analyzed on SDS-PAGE gels for purity.
the presence of blasticidin (20 mg/ml; Invitrogen, Karlsruhe, Germany) for stable expression. Soluble rCD177 was collected from cell culture supernatants and purified using mAb 7D8 affinity column, as previously described (9). Native CD177 protein was purified from human granulocytes, as described previously (16).

Surface plasmon resonance analysis
Surface plasmon resonance (SPR) analysis was performed on a ProteOn XPR36 system (Bio-Rad, Hercules, CA), as previously described (9). The purified soluble PECAM-1 proteins (VN or LS allelic form), at a concentration of 100 μM/ml (in 10 mM sodium acetate buffer [pH 4.5]), were immobilized on a general layer medium sensor chip by amine coupling using standard procedures. CD177 (1 μM in PBS) or mAb PECAM-1.2 (5–80 nM in PBS) were injected as analyte over the chip at a flow rate of 20 and 100 μl/min, respectively, in a total volume of 250 μl at 25°C. The sensorsgrams were evaluated using the ProteOn evaluation software package.

Results
Genotyping HUVECs and neutrophils
To identify the allelic isoforms of PECAM-1 present in neutrophils and HUVECs, mRNA was isolated from HUVECs (n = 15) and neutrophils (n = 10) derived from different donors and amplified by RT-PCR. Two regions encompassing the three SNPs (L98V, S536N, and R643G) were analyzed by direct DNA sequencing (Fig. 1). These results indicate that HUVECs used in this study expressed equal levels of adhesion molecules, such as JAM-C, E-selectin, and ICAM-1. These results were detected. Similar observations were noted with other adhesion molecules bearing NN or SS allelic variants of PECAM-1 by flow cytometry. As compared in a time-dependent manner (Fig. 4A), neutrophils derived from six blood donors were surface labeled with FITC-conjugated anti-CD177 and anti–PECAM-1 and analyzed by flow cytometry. Neutrophils expressing high or low levels of CD177, but having similar PECAM-1 expression, were selected for further studies (Fig. 3). As shown in Fig. 4A, CD177High neutrophils transmigrated to a significantly greater extent through the homozygous SS, compared with heterozygous SN or homozygous NN, HUVEC monolayer toward fMLP. These differences were not detectable when CD177Null neutrophils were applied (Fig. 4A). These results indicate that the PECAM-1 S536N dimorphism within IgD6 has functional consequences for neutrophil transendothelial migration. To exclude the possibility that the genetically linked V98L dimorphism within IgD1, which mediates homophilic adhesion, might contribute to the observed difference in neutrophil transmigration, these experiments were repeated in the presence of mAbs PECAM-1.3 (specific for IgD1) or PECAM-1.2 (specific for IgD6). As shown in Fig. 4B, the addition of mAb PECAM-1.2 eliminated the phenotype-specific quantitative difference in transmigration efficiency, while suppressing the total degree of transmigration of CD177+ neutrophils across HUVEC monolayers. In contrast, the addition of PECAM-1.3 (specific for IgD1), which blocks homophilic interactions (7) and loosens endothelial junctions, overshadows the impact of heterophilic interaction and, thereby, eliminates the PECAM-1 allele-dependent effects on neutrophil migration. Thus, IgD6 plays a more significant role in the CD177-facilitated component of this process than does IgD1.
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migrated faster through HUVECs phenotyped for SS in comparison with NS and NN in response to all three chemoattractants, indicating that PECAM-1 allelic-dependent neutrophil migration is not chemoattractant specific.

To further characterize the influence of the S536N polymorphism on neutrophil transendothelial migration, we activated neutrophils with the bacterial toxin PLY prior to reperforming the transendothelial cell migration experiments. As shown in Fig. 5A, PLY treatment resulted in a significant upregulation of CD177 in the CD177+ but not CD177- neutrophil subpopulation, while depressing PECAM-1 expression. Despite the lower levels of PECAM-1, PLY-treated neutrophils exhibited slightly increased transendothelial migration across all HUVEC phenotypes (Fig. 5B), suggesting that neutrophil CD177, and not neutrophil PECAM-1, plays a dominant role in this process.

Allele-specific differences in PECAM-1 ITIM phosphorylation

Previous studies showed that the level of PECAM-1 ITIM phosphorylation correlates with endothelial cell barrier integrity (18, 19). To determine whether binding of domain-specific anti–PECAM-1 mAbs or CD177 might differentially stimulate phosphorylation of PECAM-1 expressed in SS versus NN homozygous HUVEC lines, these reagents were added to the cells, and coimmunoprecipitation/phosphotyrosine immunoblot analysis was performed. As shown in Fig. 6A, although CD177 by itself was unable to induce significant tyrosine phosphorylation of PECAM-1 ITIMs, engagement with IgD5-specific mAb PECAM-1.1 or IgD6-specific mAb PECAM-1.2, but not with IgD1-specific mAb PECAM-1.3, induced significant PECAM-1 tyrosine phosphorylation. Of note, engagement by mAb PECAM-1.2 induced less tyrosine phosphorylation in HUVECs expressing the SS allelic isoform of PECAM-1 than it did in NN-bearing endothelial cells. Because mAb PECAM-1.1 caused

FIGURE 2. A, FACS analysis of adhesion molecules on SS (white) and NN (black) phenotyped HUVECs. HUVECs in resting (−TNF-α) or stimulated conditions (+TNF-α) were labeled with mAbs against PECAM-1, JAM-C, ICAM-1, E-Selectin, and mIgG (as control). Bound Abs were detected by FITC-labeled secondary Ab and were analyzed by flow cytometry. B, Permeability of three phenotyped HUVEC lines. HUVECs were cultured on fibronectin-coated polycarbonate filter chambers for 48 h. The passage of FITC-albumin through a confluent monolayer of cells at different time periods (5–60 min) was measured in the presence and absence of 0.2 U/ml thrombin or 3.2 × 10−5 mol/l histamine (upper panel). The lower panel shows permeability after treatment with mAb PECAM-1.1 in the presence and absence of rCD177/Fc or rJAM-C/Fc (as control). The intensity of migrated FITC-albumin in the lower chamber was measured by fluorescence microtiter plate reader. *p < 0.01; FITC-albumin passage between homozygous SS and NN (n = 4). AU, arbitrary unit.

FIGURE 3. Analysis of CD177 and PECAM-1 surface expression in neutrophils by FACS. Neutrophils of CD177− (left panel) and CD177+ (right panel) individuals were labeled with mAb 7D8 or mAb Gi18. After washings, bound Abs were detected by FITC-labeled secondary Ab and were analyzed by flow cytometry.
equal PECAM-1 phosphorylation in both isoforms, we sought to investigate the influence of CD177 on PECAM-1-pretreated HUVECs. Interestingly, rCD177/Fc suppressed PECAM-1 phosphorylation significantly in SS HUVECs compared with NN HUVECs. To further demonstrate the specificity of this phenomenon, we examined the phosphorylation of epidermal growth factor receptor and ERK, but we did not observe any PECAM-1 allele-specific differences (Supplemental Fig. 1). These effects were also not due to the differential affinity of mAb PECAM-1.2 or CD177 for the SS versus NN form of PECAM-1. SPR analysis showed that mAb PECAM-1.2 and CD177 bound equally to SS and NN allelic forms of PECAM-1 protein with similar binding kinetics (Fig. 7). Interestingly, postengagement of IgD6 by CD177 selectively diminished the degree of PECAM-1 tyrosine phosphorylation induced by the addition of H₂O₂ (Fig. 6B) in SS but not NN-bearing HUVECs.

Several lines of evidence suggest that PECAM-1 ITIM phosphorylation might regulate endothelial permeability by phosphorylation/degradation of the adherent junction protein β-catenin (18, 19). Our permeability experiments in which we add the IgD5-specific mAb PECAM-1.1 to prestimulated HUVECs treated with rCD177/FC showed increased permeability in HUVECs expressing the SS versus NN isoform of PECAM-1 (Fig. 3B), and this was correlated with suppression of PECAM-1 phosphorylation (Fig. 6B). Thus, the ability of CD177 to suppress PECAM-1 phosphorylation in SS homozygous HUVECs might account for the different degree of β-catenin phosphorylation. To prove this hypothesis, we analyzed β-catenin phosphorylation in NN- and SS-bearing...
between homozygous SS and two NS and NN phenotypes (rCD177/Fc. Indeed, treatment of these cells with H₂O₂ and b-catenin phosphorylation was observed in both cells treated with PLY. Neutrophils were treated with 25 ng/ml PLY for 30 min and labeled with mAbs 7D8 (anti-CD177) or Gi18 (anti–PECAM-1). After washing, bound mAb was detected with FITC-labeled secondary Ab and analyzed by FACS. B, PLY-untreated (−) or -treated (+) fluorescence-labeled neutrophils were allowed to migrate through HUVECs toward fMLP and were measured as above. *p < 0.05; **p < 0.01; neutrophil migration between homozygous SS and two NS and NN phenotypes (n = 4).

HUVECs after stimulation with H₂O₂ in the presence or absence of rCD177/Fc. Indeed, treatment of these cells with H₂O₂ and rCD177/Fc induced significantly greater β-catenin phosphorylation in SS- versus NN-bearing HUVECs (Fig. 6B). In contrast, equal β-catenin phosphorylation was observed in both cells treated with H₂O₂ alone. Taken together, our results suggest that PECAM-1 allele-dependent phosphorylation of β-catenin following CD177 engagement may be involved in the difference in endothelial cell permeability that we observed.

Discussion

Previous studies showed that the neutrophil-specific Ag CD177 supports transendothelial migration by interacting with membrane-proximal IgD6 of endothelial PECAM-1 (9). In the present investigation, we found that the S₃₅₆N dimorphism located in IgD6 of endothelial PECAM-1 has a significant influence on the transendothelial migration of neutrophils (Fig. 4A). The rate of neutrophil transmigration through SS homozygous HUVECs was faster than through NN HUVECs, an effect that was dependent on the expression of neutrophil CD177 but independent of neutrophil PECAM-1 (Fig. 3A). Moreover, engagement of PECAM-1 by CD177 selectively suppressed PECAM-1 ITIM phosphorylation in SS but not NN homozygous HUVECs (Fig. 6B). Taken together, these data suggest that heterophilic PECAM-1/CD177 interactions that occur during neutrophil diapedesis reduce the overall PECAM-1 phosphorylation state, leading to weakened junctional stability that is conducive to neutrophil transmigration in a previously unrecognized allele-specific manner.

The S₃₅₆N dimorphism within PECAM-1 IgD6 is in strong linkage disequilibrium with two other SNPs that occur within the coding region of the molecule, resulting in two common PECAM-1 haplotypes that have been termed LSR and VNG (12). Goodman et al. (20) were the first to describe an influence of these isofoms on leukocyte/endothelial cell interaction: heterozygous (LSR/VNG) monocytes were reported to adhere better to endothelium under conditions of flow than were LSR/VNG homozygous cells. This influence was attributed to PECAM-1 expressed on monocytes, rather than to endothelial cell PECAM-1. These findings are difficult to compare with our study because monocytes do not express CD177 (10), and adhesion under flow requires different molecular interactions than transmigration through static endothelial cells.

It is unlikely that any of the three amino acid polymorphisms of PECAM-1 by itself is solely responsible for the allele-specific effects that we observed downstream of PECAM-1/CD177 interaction. The L₉₈V dimorphism located in IgD1 is known to mediate cation-independent homophilic interaction between PECAM-1 molecules on adjacent cells (21). Inhibition of this interaction affects allele-specific neutrophil transmigration (Fig. 4B), indicating that homophilic interactions between neutrophil and endothelial cell PECAM-1 do not play a major role in the allele-specific differences that were observed. The S₃₅₆N dimorphism within IgD6 did not affect its binding affinity for CD177 (Fig. 7), and blocking IgD6 eliminated the differences in neutrophil transmigration rates between the two haplotypes. Finally, the R₆₄₃G dimorphism is located within the cytoplasmic domain; however, its physical relationship to the ITIM phosphorylation sites has not been determined.

PECAM-1 is an adhesion and signaling receptor that is expressed abundantly at endothelial intracellular borders (22, 23), and it becomes phosphorylated when endothelial cell junctions are perturbed (18, 19). Elrayess et al. (24) reported higher tyrosine phosphorylation levels in VNG compared with LSR HUVECs, a finding that is consistent with the results in our study (Fig. 6A). Interestingly, although engaging endothelial cell PECAM-1 with a soluble dimeric form of CD177 did not by itself induce PECAM-1 phosphorylation, it was able to suppress PECAM-1 phosphorylation induced by Ab-mediated cross-linking or H₂O₂ treatment. Biswas et al. (19) reported that phosphorylated PECAM-1 ITIMs serve as a molecular scaffold that simultaneously recruits tyrosine-phosphorylated β-catenin and the protein-tyrosine phosphatase Src homology region 2 domain-containing phosphatase 1 (SHP-2). Dephosphorylation of phosphorylated β-catenin by PECAM-1-bound SHP-2, in turn, allows β-catenin to rebind VE-cadherin, thereby supporting reassembly of the adherens junctional complex. Our finding that the binding of CD177 selectively suppresses PECAM-1 ITIM phosphorylation in SS (but not NN) homozygous HUVECs is consistent with the observed increase in FITC-albumin leakage as well as neutrophil transendothelial migration across endothelial cell monolayers expressing this allelic isofom of PECAM-1. Because our results showed that CD177-mediated suppression of PECAM-1 ITIM tyrosine phosphorylation leads to greater β-catenin phosphorylation in the PECAM-1 SS phenotype, it would be expected to delay the recruitment of SHP-2 and β-catenin and postpone restoration of endothelial cell junctional integrity in these cells. The R₆₄₃G dimorphism is situated closed to
the ITIM phosphorylation sites, but it seems more likely that two or all three dimorphisms together induce molecular changes that allow PECAM-1 to respond differently toward CD177 binding.

CD177 becomes significantly upregulated on the surface of circulating neutrophils in response to bacterial infections (11). PLY is a potent bacterial toxin produced during infections by *Streptococcus*...
The authors have no financial conflicts of interest.

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Disclosures
The authors have no financial conflicts of interest.

References
Supplemental figure 1:

Immunoblot analysis of EGFR and ERK phosphorylation in PECAM-1 phenotyped HUVEC. Homozygous PECAM-1 NN or SS HUVEC was stimulated with H$_2$O$_2$. After lysis, aliquots of protein (5 µg) were run on 10% SDS-PAGE and were analyzed by immunoblotting using mAb against phosphorylated EGFR pY1173 (2.5 µg/ml) or ERK pT202/Y204 (2.5 µg/ml), or against GAPDH (2.5 µg/ml).
Supplemental figure 1