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Dipeptidyl Peptidase IV Is a Human and Murine Neutrophil Chemorepellent

Sarah E. Herlihy, Darrell Pilling, Anu S. Maharjan, and Richard H. Gomer

In Dictyostelium discoideum, AprA is a secreted protein that inhibits proliferation and causes chemorepulsion of Dictyostelium cells, yet AprA has little sequence similarity to any human proteins. We found that a predicted structure of AprA has similarity to human dipeptidyl peptidase IV (DPPIV). DPPIV is a serine protease present in extracellular fluids that cleaves peptides with a proline or alanine in the second position. In Insall chambers, DPPIV gradients below, similar to, and above the human serum concentration difference between the front and back of the cell is sufficient to cause chemorepulsion. Neutrophil speed and viability are unaffected by DPPIV. DPPIV inhibitors block DPPIV-mediated chemorepulsion. In a murine model of acute respiratory distress syndrome, aspirated bleomycin induces a significant increase in the number of neutrophils in the lungs after 3 d. Oropharyngeal aspiration of DPPIV inhibits the bleomycin-induced accumulation of mouse neutrophils. These results indicate that DPPIV functions as a chemorepellent of human and mouse neutrophils, and they suggest new mechanisms to inhibit neutrophil accumulation in acute respiratory distress syndrome. The Journal of Immunology, 2013, 190: 000–000.

The online version of this article contains supplemental material.
CD244+ cells with pan Mouse IgG Dynabeads (Invitrogen). Cells were resuspended to a final volume of 1 ml in 2% human albumin-RPMI 1640.

To determine the purity of CD244-depleted neutrophil preparations, cells were resuspended in PBS containing 2% BSA and cell smears were prepared as described previously (31). Cells smears were air-dried overnight, fixed in methanol, stained with H&E, and cell morphology was determined by microscopy, as described previously (30).

Human albumin was isolated from human serum (Lonza) using Anti-Flag blue gel beads (Bio-Rad). After washing beads three times with PBS, beads were incubated with serum at room temperature for 2 h. Beads were collected by centrifugation at 300 × g for 2 min and washed with buffer (20 mM Tris, 140 mM NaCl, 2 mM calcium) and the albumin was eluted off overnight with 500 mM NaCl. Albumin was then concentrated and buffer exchanged to Earl’s balanced salt solution (Sigma-Aldrich) using a 10-kDa centrifugal filter (Millipore) and stored at 4 °C.

**Insall chamber assays**

Human soluble rDPPIV was purchased from Enzo Life Sciences. To measure the effect of DPPIV on cell dispersion, we used Insall chambers, which allows direct visualization of cell movement, as described previously for melanoma cells, Dictyostelium, and neutrophils (13, 32, 33). Briefly, 22 × 22-mm glass coverslips were etched with 1 M HCl, rinsed with deionized water, and coated with 20 μg/ml bovine plasma fibronectin (Sigma-Aldrich) for 30 min at 37°C. Coverslips were then washed twice with PBS, and 300 μl neutrophils at 5 × 10^6 cells/ml was allowed to adhere to the coverslip for 15 min at 37°C. We then used an Insall chamber slide, a gift from Robert Insall (32). Two concentric depressions and the separating bridge were filled with 2% BSA-RPMI 1640. The media was then removed from the outer chamber and was separated bridge were filled with 2% BSA-RPMI 1640. The media was then removed from the coverslips, which were then placed face down on the slide, a gift from Robert Insall (32). Two concentric depressions and the separating bridge were filled with 2% BSA-RPMI 1640. The media was then removed from the outer chamber and was replaced by DPPIV alone, DPPIV inhibitor alone (diprotin A from Enzo Life Sciences or DPP1 1c hydrochloride from Tocris Bioscience), DPPIV plus DPPIV inhibitor (all in 2% BSA-RPMI 1640), or 2% BSA-RPMI 1640. Cells located on the bridge between the square depressions were then filmed as previously described (34), using a ×10 objective for 1 h at 37°C in a humidified 5% CO2 incubator. Displacement of at least 10 random chosen cells per experiment was measured during periods of 10 min. Cell tracking and track analysis was done as previously described (13) with the exception that videos of an hour in length were processed (36). The primary BAL cells were collected by centrifugation at 500 × g for 5 min. The cells were resuspended in 500 μl 4% BSA-PBS and counted with a hemocytometer. Diluted cells (100 μl) were then aliquoted into cytospin funnels and spun onto glass slides (Superfrost Plus white slides; VWR Scientific, West Chester, PA) at 400 rpm for 5 min using a cytospin centrifuge (Shandon, Cheshire, U.K.). These cells were then air-dried and stained with Gill’s hematoxylin. More than 200 cells were counted for total cell number, and neutrophils, macrophages, and lymphocytes were identified by morphology. The percentage of neutrophils, macrophages, or lymphocytes was then multiplied by the total number of cells recovered from the BAL to obtain the number of each cell type in the BAL. The mice were used in accordance with guidelines published by the National Institutes of Health, and the protocol was approved by the Texas A&M University Animal Use and Care Committee.

**Immunohistochemistry**

After BAL, lungs were inflated with prewarmed OCT (VWR Scientific) and then embedded in OCT, frozen on dry ice, and stored at −80 °C as described previously (37). Lung tissue sections (5 μm) were prepared and immunohistochemistry was done as described previously (37) except that slides were incubated with 5 μg/ml primary Abs in 4% BSA-PBS for 60 min. Active caspase-3 staining was done overnight at 4°C. The lung sections were stained for Ly6G (BD Biosciences) to detect neutrophils, CD11b (BioLegend) to detect infiltrating macrophages and neutrophils, CD11c (MBL International) to detect lung-resident macrophages and dendritic cells, CD206 (BioLegend) to detect the mannose receptor on macrophages and dendritic cells, and CD107b (Mac3) (BioLegend) to detect alveolar and tissue macrophages and granulocytes, cleaved caspase-3 (Cell Signaling Technologies) to detect activated caspase-3, and isotype-matched mouse or rabbit irrelevant Abs as controls. Slides were then washed three times with PBS for 30 min and incubated with 5 μg/ml biotinylated mouse F(ab’2) anti-rat IgG or 5 μg/ml biotinylated donkey F(ab’2) anti-rabbit IgG in 4% BSA-PBS for 30 min. Slides were then washed three times in PBS for 30 min and incubated with a 1:500 dilution of streptavidin-alkaline phosphatase (Vector Laboratories) in 4% BSA-PBS for 30 min. Staining was then developed with a VectorRed alkaline phosphatase kit (Vector Laboratories) for 10 min. Slides were then mounted as described previously (37). Five or ten 450-μm fields of view were counted for Ly6G-stained or active caspase-3–stained positive cells, respectively. Areas containing large blood vessels and bronchioles were excluded from analysis.

**Effect of DPPIV peptidase activity on albumin stability**

BSA-RPMI 1640 (2%) and 2% human albumin-RPMI 1640 were incubated with 300 ng/ml DPP1V or an equivalent volume of buffer for 30 min at 37°C in tubes that were precoated with 4% BSA-RPMI 1640. Where indicated, the DPP1V inhibitor DPP1 1c hydrochloride was added to the conditioned medium to a final concentration of 200 nM. Cells were collected by centrifugation at 500 × g for 5 min and the supernatant was added to the outside chamber of an Insall chamber. Cells were then washed twice with PBS for 30 min and incubated with 5 μg/ml biotinylated mouse F(ab’2) anti-rat IgG or 5 μg/ml biotinylated donkey F(ab’2) anti-rabbit IgG in 4% BSA-PBS for 30 min. Slides were then washed three times in PBS for 30 min and incubated with a 1:500 dilution of streptavidin-alkaline phosphatase (Vector Laboratories) in 4% BSA-PBS for 30 min. Staining was then developed with a VectorRed alkaline phosphatase kit (Vector Laboratories) for 10 min. Slides were then mounted as described previously (37). Five or ten 450-μm fields of view were counted for Ly6G-stained or active caspase-3–stained positive cells, respectively. Areas containing large blood vessels and bronchioles were excluded from analysis.

**Neutrophil influx in mice**

Four-week-old C57BL/6 male mice (The Jackson Laboratory, Bar Harbor, ME) were treated with an oropharyngeal aspiration of 50 μl saline or 3 μg/ml bleomycine (Calbiochem) (35). The successful aspiration of bleomycine into the lungs was confirmed by listening for the crackling noise heard after the aspiration. Twenty-four hours following bleomycin aspiration (day 1), mice were treated with an oropharyngeal aspiration of 50 μl 0.9% saline with 0.9 μg rDPPIV (Enzo Life Sciences) or an equal volume of 0.9% saline. Mice were weighed daily and euthanized at day 3 after bleomycin aspiration. Blood was collected by cardiac puncture from the euthanized mice and blood glucose was measured using Accu-Check Advantage (Roche). The lungs were perfused with 300 μl PBS three times to collect cells by bronchoalveolar lavage (BAL) as described previously (36). The primary BAL cells were collected by centrifugation at 500 × g for 10 min. Primary BAL pellets were resuspended in the secondary and tertiary BAL fluid and the combined cells were collected by centrifugation at 500 × g for 5 min. The cells were resuspended in 500 μl 4% BSA-PBS and counted with a hemocytometer. Diluted cells (100 μl) were then aliquoted into cytospin funnels and spun onto glass slides (Superfrost Plus white slides; VWR Scientific, West Chester, PA) at 400 rpm for 5 min using a cytospin centrifuge (Shandon, Cheshire, U.K.). These cells were then air-dried and stained with Gill’s hematoxylin. More than 200 cells were counted for total cell number, and neutrophils, macrophages, and lymphocytes were identified by morphology. The percentage of neutrophils, macrophages, or lymphocytes was then multiplied by the total number of cells recovered from the BAL to obtain the number of each cell type in the BAL. The mice were used in accordance with guidelines published by the National Institutes of Health, and the protocol was approved by the Texas A&M University Animal Use and Care Committee.

**FIGURE 1.** Superimposition of the predicted structure of AprA with the structure of the α/β hydrolase domain of human DPPIV. The catalytic domains of DPPIV (the α/β hydrolase domain of Protein Data Bank ID 1J2E) and AprA (predicted structure) were superimposed and the catalytic triads highlighted. The α/β hydrolase domain of DPPIV is shown in green and its catalytic triad (Asp108, His740, and Ser636) is orange. The predicted structure of AprA is shown in cyan and its potential catalytic triad (Asp288, His110, and Ser155) is blue. The β-propeller domain of DPPIV was removed for simplicity because the predicted structure of AprA had no overlap with this domain.
Neutrophils show biased movement away from a source of DPPIV

Isolated neutrophils were cultured in RPMI 1640 containing 10% (v/v) FBS and 2 mM glutamine in 96-well tissue culture plates at 37°C for 20 h, as described previously (38). Neutrophils were cultured in the presence or absence of 25 ng/ml human IL-8, human TNF-α (both from PeproTech), or 400 ng/ml DPPIV. After 20 h, neutrophils were labeled with annexin V (BioLegend) and propidium iodide (Sigma-Aldrich) according to the manufacturers’ instructions to identify early and late stages of cell death. Additionally, cytospin preparations were used to assess for morphological changes associated with apoptosis, as described previously (38–40).

Statistical analysis

Statistics were done using Prism (GraphPad Software, San Diego, CA). One-way ANOVA was used to compare between multiple groups and a Student t test was used to compare between two groups.

Results

AprA has structural similarity to the human protein DPPIV

AprA has little sequence similarity to any mammalian protein (11). Using I-TASSER, we generated a predicted structure for AprA (Fig. 1). The Dali server, used to compare protein structures, composes a list of the proteins with the most structural similarity to the query protein. The proteins with the top structural similarities to the predicted structure of AprA were cephalosporin C deacetylase (a β-lactamase from the fungus Acremonium), acetyl xylan esterase (from the bacteria Bacillis pumilus, which cleaves carboxyl-ester bonds), acylaminic acid–releasing enzyme (from at least three different volunteers were used. DPPIV (human soluble form, which cleaves terminal amino acids with proline or alanine in the second position), AprA has an 11% structural identity to the entire structure of soluble human DPPIV (Fig. 1, the αβ hydrolase domain of DPPIV is shown). Because DPPIV, similar to AprA, can be found as an extracellular protein, these results suggested that AprA might have functional similarity to DPPIV or vice versa.

Neutrophils show biased movement away from a source of DPPIV

AprA functions as a chemorepellent of Dicystostelium cells (13). Dicystostelium and neutrophils share many properties of chemotaxis (41, 42). Therefore, we hypothesized that DPPIV may regulate human neutrophil motility owing to its structural similarity to AprA. To examine neutrophil chemotaxis in gradients of DPPIV, we used an Insall chamber. When neutrophil movement was tracked for 10-min periods, there was no bias of movement in the media control (Fig. 2B, Supplemental Fig. 1). A biased movement away from a source of DPPIV was observed (Fig. 2A, Supplemental Fig. 1). Cells were tracked, and the average center of mass observed for the endpoints of cells in each population was determined. The center of mass of cell endpoints showed displacement away from the source of DPPIV (Fig. 2, Supplemental Fig. 1). The concentration of DPPIV in human blood ranges from 400 to 800 ng/ml, or 4–7–9.4 nM (43, 44). Therefore, we tested the ability of DPPIV to affect neutrophil migration above, below, and within this concentration range. Neutrophils showed biased movement away from higher concentrations of rDPPIV in a variety of DPPIV concentration gradients (Table I). An equal concentration of DPPIV in both

Table I. The effect of DPPIV on forward migration and directness of neutrophil movement

<table>
<thead>
<tr>
<th>rDPPIV Gradient (nM)</th>
<th>Forward Migration Index</th>
<th>Directionality</th>
<th>No. of Cells Moving toward rDPPIV</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Media Control (0 nM rDPPIV)</td>
<td>rDPPIV</td>
<td>Media Control (0 nM rDPPIV)</td>
</tr>
<tr>
<td>0–1.2</td>
<td>0.00 ± 0.03</td>
<td>0.21 ± 0.03***</td>
<td>0.35 ± 0.02</td>
</tr>
<tr>
<td>0–3.5</td>
<td>−0.04 ± 0.04</td>
<td>0.21 ± 0.06***</td>
<td>0.26 ± 0.02</td>
</tr>
<tr>
<td>0–11.7</td>
<td>0.01 ± 0.06</td>
<td>0.15 ± 0.05**</td>
<td>0.47 ± 0.03</td>
</tr>
<tr>
<td>4.7–4.7</td>
<td>0.01 ± 0.06</td>
<td>−0.01 ± 0.04</td>
<td>0.47 ± 0.03</td>
</tr>
<tr>
<td>4.7–11.7</td>
<td>−0.02 ± 0.06</td>
<td>0.22 ± 0.06**</td>
<td>0.50 ± 0.03</td>
</tr>
<tr>
<td>9.4–23</td>
<td>−0.01 ± 0.05</td>
<td>0.13 ± 0.05**</td>
<td>0.36 ± 0.03</td>
</tr>
<tr>
<td>0–3.5*TNF</td>
<td>−0.02 ± 0.03</td>
<td>0.10 ± 0.03**</td>
<td>0.32 ± 0.02</td>
</tr>
</tbody>
</table>

The data from at least three independent sets of cell population tracks (see Fig. 2 and Supplemental Fig. 1 as examples) were analyzed to determine the forward migration index (FMI) and directionality. FMI is a measure of migration of cells along the gradient, where 0 equals no movement, a positive number equals movement away from the source, and a negative number indicates movement toward the source. Directionality is the ratio of Euclidean distance to accumulated distance. For each gradient, neutrophils from at least three different volunteers were used.

*p < 0.05, **p < 0.01, ***p < 0.001 compared with the media control (t test).

N/A, not applicable. TNF, gradient of DPPIV with TNF-α-stimulated neutrophils.
Neutrophil chemorepulsion is sensitive to DPPIV enzyme inhibitors

DPPIV enzyme activity has been implicated in the chemotaxis of several types of immune cells through cleavage of human chemokines (25, 47–49). To determine whether DPPIV enzyme activity affects DPPIV-induced neutrophil chemorepulsion, we used two enzyme inhibitors of DPPIV, diprotin A and DPPIV 1c hydrochloride (50, 51). The inhibitors alone caused no attraction or repulsion of human neutrophils (Fig. 3). When either of the inhibitors was added with DPPIV, the chemorepulsion of neutrophils away from the source of DPPIV was significantly reduced compared with DPPIV alone (Fig. 3). This suggests that DPPIV inhibitors block the ability of DPPIV to induce neutrophil chemorepulsion.

DPPIV does not appear to cleave albumin

Several possibilities exist to explain why DPPIV enzymatic activity seems to be required for neutrophil chemorepulsion. In addition to DPPIV directly regulating neutrophil chemorepulsion, DPPIV could cleave a component of the media, creating a breakdown component of BSA that acts as a chemorepellent. To determine whether a component of the media was broken down by rDPPIV, BSA-RPMI 1640 or human albumin-RPMI 1640 was incubated with rDPPIV for 3 h. Although BSA is not pure, the human albumin media appear to contain only human albumin and DPPIV as protein components (Fig. 4A, 4B). In both cases, there was no indication of cleavage products accumulating over time or when time points are compared in the presence and absence of rDPPIV (Fig. 4A, 4B).

Purified neutrophils are chemorepelled by DPPIV

A protein or peptide released by contaminating cells into the media and then cleaved by DPPIV could also act as the chemorepellent. Neutrophils were further purified after isolation, resuspended in pure human albumin, and assayed for chemorepulsion from DPPIV. Both the number of contaminating eosinophils and monocytes in the neutrophil preparation decreased significantly following depletion (Fig. 4C). Neutrophils were 97% pure following CD244 depletion, making a gradient of attractant or repellent from remaining contaminating cells unlikely (Fig. 4C). Pure neutrophils showed a biased movement away from rDPPIV (Fig. 4D). These data suggest that it is unlikely that DPPIV cleaves a component of the media or from contaminating cells to induce chemorepulsion.

Conditioned medium from neutrophils does not affect chemorepulsion

Proteases on the neutrophil surface could create a neutrophil chemorepellent or cleave a breakdown component of the media that

### Table II. Effect of DPPIV on the average cell speed (μm/min) of neutrophils

<table>
<thead>
<tr>
<th>rDPPIV gradient (nM)</th>
<th>Media control (0 nM rDPPIV)</th>
<th>rDPPIV</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-1.2</td>
<td>27 ± 1</td>
<td>26 ± 1</td>
</tr>
<tr>
<td>0-3.5</td>
<td>23 ± 1</td>
<td>23 ± 1</td>
</tr>
<tr>
<td>0-11.7</td>
<td>24 ± 2</td>
<td>25 ± 2</td>
</tr>
<tr>
<td>4.7-4.7</td>
<td>17 ± 1</td>
<td>16 ± 1</td>
</tr>
<tr>
<td>4.7-11.7</td>
<td>17 ± 1</td>
<td>18 ± 2</td>
</tr>
<tr>
<td>9.4-23</td>
<td>17 ± 1</td>
<td>17 ± 1</td>
</tr>
</tbody>
</table>

The data from at least three independent sets of cell population tracks (see Fig. 2 and Supplemental Fig. 1 for examples) were used to determine the average speed of neutrophils. Values are means ± SEM (n = 3 or more).

### Table III. Percentage of cells in the population moving in a biased direction over 10 min

<table>
<thead>
<tr>
<th>rDPPIV Concentration Gradient (nM)</th>
<th>P₀ (Control)</th>
<th>P₀ (rDPPIV)</th>
<th>P₁ (Control)</th>
<th>P₁ (rDPPIV)</th>
<th>P₂ (Control)</th>
<th>P₂ (rDPPIV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-1.2</td>
<td>0.39 ± 0.14</td>
<td>0.54 ± 0.02</td>
<td>0.32 ± 0.08</td>
<td>0.16 ± 0.01*</td>
<td>0.29 ± 0.06</td>
<td>0.31 ± 0.02</td>
</tr>
<tr>
<td>0-3.5</td>
<td>0.27 ± 0.13</td>
<td>0.28 ± 0.08</td>
<td>0.39 ± 0.08</td>
<td>0.27 ± 0.04</td>
<td>0.34 ± 0.06</td>
<td>0.45 ± 0.06*</td>
</tr>
<tr>
<td>0-11.7</td>
<td>0.10 ± 0.03</td>
<td>0.13 ± 0.06</td>
<td>0.46 ± 0.01</td>
<td>0.35 ± 0.02*</td>
<td>0.44 ± 0.03</td>
<td>0.52 ± 0.05</td>
</tr>
<tr>
<td>4.7-4.7</td>
<td>0.17 ± 0.07</td>
<td>0.20 ± 0.05</td>
<td>0.44 ± 0.03</td>
<td>0.43 ± 0.03</td>
<td>0.39 ± 0.04</td>
<td>0.37 ± 0.03</td>
</tr>
<tr>
<td>4.7-11.7</td>
<td>0.19 ± 0.06</td>
<td>0.22 ± 0.06</td>
<td>0.44 ± 0.03</td>
<td>0.26 ± 0.05*</td>
<td>0.38 ± 0.04</td>
<td>0.53 ± 0.02*</td>
</tr>
<tr>
<td>9.4-23</td>
<td>0.29 ± 0.02</td>
<td>0.22 ± 0.06</td>
<td>0.39 ± 0.03</td>
<td>0.38 ± 0.03</td>
<td>0.33 ± 0.02</td>
<td>0.40 ± 0.08</td>
</tr>
<tr>
<td>0-3.5TNF</td>
<td>0.26 ± 0.04</td>
<td>0.26 ± 0.03</td>
<td>0.38 ± 0.04</td>
<td>0.31 ± 0.03</td>
<td>0.36 ± 0.01</td>
<td>0.43 ± 0.02*</td>
</tr>
</tbody>
</table>

The data from at least three independent sets of cell population tracks (see Fig. 2 and Supplemental Fig. 1 for examples) were used to determine the probability of cell movement toward or away from the source of DPPIV. P₀ and P₁ are the probabilities that a cell would move away from or toward the source of DPPIV, respectively, in a 13-s interval. P₀ is the probability that a cell will not move. P₂ and P₁ are statistically significantly different in all DPPIV gradients at p < 0.05 (t test), except for 0-11.7 nM DPPIV, which is significant at p < 0.01. In the controls, P₂ and P₁ are statistically significant only in the 4.7-11.7 and 9.4-23 nM controls (p < 0.05).

TNF, Gradient of DPPIV with TNF-α-stimulated neutrophils.
acts as a chemorepellent. Additionally, neutrophils could be releasing a chemorepellent or chemoattractant that is then affected by DPPIV. To determine whether neutrophil proteins are responsible for the chemorepulsion effect, neutrophils were incubated with DPPIV or an equivalent volume of buffer for 30 min at 37°C. Neutrophils were then collected by centrifugation, and media from the outer chamber of the Insall chamber were replaced with these conditioned media. Half of the conditioned medium from cells preincubated with DPPIV was mixed with the DPPIV inhibitor DPPIV 1c hydrochloride immediately before addition to the Insall chamber. Conditioned media from buffer-treated cells or conditioned media from DPPIV-treated cells that were mixed with the inhibitor failed to promote chemorepulsion of neutrophils (Fig. 5A). Conditioned media from DPPIV-treated cells did promote chemorepulsion of neutrophils (Fig. 5A). As determined by Western blots stained with anti-DPPIV Abs, there was no detectable DPPIV present in conditioned media from buffer-treated cells, whereas there was DPPIV present in the conditioned media from DPPIV-treated cells (Fig. 5B). Taken together, these data suggest that DPPIV promotes neutrophil chemorepulsion, rather than cleaving a component of the media or a substrate secreted from cells.

**DPPIV reduces the number of neutrophils in lungs of mice treated with bleomycin**

Oropharyngeal aspiration of bleomycin in mice causes neutrophils to accumulate in the lungs within 24 h of bleomycin administration (52). If DPPIV functions as a neutrophil chemorepellent, then administering DPPIV to a tissue could drive neutrophils out of the tissue or prevent their entry. We examined the ability of DPPIV to affect neutrophil accumulation in the lungs of bleomycin-treated mice. Assuming the volume of liquid in the airways of mouse lungs is not 0.1 ml, administering 0.9 mg DPPIV creates a DPPIV concentration considerably greater than the DPPIV serum concentration of 0.4 µg/ml (53). Mice were treated with bleomycin on day 0 and were treated with aspirated rDPPIV or an equivalent volume of saline on day 1. Bleomycin is metabolized and excreted within 24 h, so the DPPIV treatment at 24 h does not block the effect of bleomycin (54, 55). Three days after bleomycin treatment, mice were euthanized and weakly adhered cells from the airways were collected by BAL. Although the BAL from DPPIV-treated mice appeared to have fewer total number of cells than did that from mice given bleomycin alone, this difference was not significant (Fig. 6A). BALs from mice given rDPPIV following saline had significantly fewer cells than mice given saline or bleomycin alone (Fig. 6A). Cell morphology was used to determine the total number of neutrophils, macrophages, and lymphocytes in the BAL. Significantly fewer neutrophils were present in the BALs from bleomycin plus rDPPIV– and saline plus rDPPIV–treated mice compared with mice treated with bleomycin alone (Fig. 6B). There was no difference in total macrophage or lymphocyte numbers in the BAL between DPPIV-treated mice and their untreated counterparts (Fig. 6B). Following BAL, lungs were sectioned and stained with anti-mouse Ly6G to detect neutrophils.
There were significantly fewer Ly6G+ cells in the post-BAL lungs of mice treated with bleomycin and then treated with DPPIV on day 1 than the post-BAL lungs of mice treated with bleomycin alone (Fig. 6C, 6D). There was no difference in the numbers of CD11b, CD11c, CD206, or CD107b (Mac3)-positive cells in the lungs of mice treated with bleomycin compared with those treated with bleomycin and then treated with DPPIV (Fig. 6E), suggesting that the effect of DPPIV is specific to neutrophils.
DPPIV regulates glucagon-like peptide-1, which regulates glucose levels in the blood (56). Inhibiting DPPIV affects glucose levels in mice (57). To determine whether oropharyngeal aspiration of DPPIV at day 1 affected weight gain or serum glucose levels, we measured weight change for 3 d and measured glucose levels in the blood of mice euthanized on day 3. Although not statistically significant, DPPIV administration appeared to rescue the weight loss caused by bleomycin and had no significant impact on weight gain when administered on day 1 to mice given saline on day 0 (Supplemental Fig. 2A). Additionally, DPPIV administration had no significant effect on serum glucose levels at day 3 (Supplemental Fig. 2B). Taken together, our results indicate that DPPIV reduces the accumulation of neutrophils in the lungs of mice but does not cause significant changes in weight or serum glucose levels 2 d after administration.

**DPPIV does not promote or inhibit neutrophil survival**

The number of cells in any tissue is dependent on four factors: proliferation, death, recruitment, and emigration (58). Because neutrophils are terminally differentiated cells, proliferation is not likely to regulate neutrophil numbers in an inflammatory environment. To determine whether the reduction in the number of neutrophils in DPPIV-treated lungs was due to altered cell survival, we cultured human neutrophils in the presence or absence of DPPIV for 20 h. In the absence of exogenous proteins, most neutrophils were apoptotic, as determined by annexin V staining and morphological changes (Fig. 7A, 7B). The addition of DPPIV had no significant effect on neutrophil survival (Fig. 7A–C). To determine whether DPPIV was regulating neutrophil survival in vivo, we stained lung sections with Abs against active caspase-3. Similar to previous observations in the presence or absence of bleomycin, there was no difference in the number of cells labeled with active caspase-3 Abs with the addition of DPPIV (Fig. 7D) (59). These data indicate that the reduction in the number of neutrophils in the lungs of mice following DPPIV administration is unlikely to be due to increased neutrophil cell death.

**Discussion**

We previously characterized AprA as an endogenous chemo-repellent of Dictyostelium cells (13). In this study, we identified structural similarities between AprA and human DPPIV and found that DPPIV appears to act as a neutrophil chemorepellent at physiological concentrations.

In the Insall chambers, we observed a strong DPPIV chemorepellent effect using a 0–100 ng/ml (0–1.2 nM) gradient. The gradient in the Insall chamber forms over a 970-μm gap. Making the first-order assumption that the gradient is linear, DPPIV is effective at 0.10 ng/ml/μm, or 0.0012 nM/μm. With an observed average neutrophil length of 10.5 μm (consistent with previous observations) (60), a cell in the middle of the gradient would correspond to a difference of 1.1 ng over the length of the cell at a place where the average DPPIV concentration is 50 ng/ml. This would then represent a 2.2% difference in the DPPIV concentration between the front and the back of the cell. For the 4.7–11.7 nM and the 9.4–23 nM gradients, the difference is 0.9%. This is similar to the 1.25% concentration difference of cAMP that induces chemotaxis for aggregating Dictyostelium cells (45), or the 1% fMLP concentration gradient that induces neutrophil chemotaxis (61).

For unknown reasons, we observed that for all of the DPPIV gradients in the Insall chambers, ∼17% of the neutrophils showed movement toward the source of DPPIV during 10 min. There was no obvious difference in the percentage of cells moving toward DPPIV as a function of the DPPIV concentration gradient. This effect is strikingly similar to the observation that 17% of Dictyostelium cells also move toward a source of AprA in the Insall chambers (13). At the lowest gradient concentration used, the
DPPIV concentration at the middle of the gradient was 0.6 nM, or ~110 DPPIV molecules per cell volume in the extracellular environment. By Poisson statistics, there could be momentary conditions where the cell does not detect the gradient or detects an “opposite” gradient. However, during 10 min all cells should detect the gradient. As the percentage of non-neutrophils in both isolated and purified neutrophils is <17% of the total population, some portion of the backward-moving cells must be neutrophils. The existence of cells moving backward thus indicates an unknown heterogeneity in both Dictyostelium cells and human neutrophils.

Inhibitors that block the active site of DPPIV also blocked the ability of DPPIV to act as a chemorepellent. It is possible that the inhibitors disrupt the DPPIV structure and therefore disturb its ability to act as a ligand for a chemorepellent receptor. However, the crystal structure of human DPPIV in a complex with diprotin A shows no significant structural difference compared with DPPIV alone (62). This suggests that the inhibitors block chemorepulsion in some other manner, such as blocking the ability to cleave something in the assay medium. In the Insall chamber assay, neutrophils were in RPMI 1640 containing BSA or human albumin on a fibronectin matrix. The only known enzymatic activity of DPPIV is cleavage of two amino acids from the N terminus of a protein if the second amino acid is a proline or alanine (26). RPMI 1640 does not contain proteins, and BSA and human albumin do not have a proline or an alanine as the second amino acid. Additionally, no cleavage products of BSA or human albumin were seen following incubation with rDPPIV for 3 h. DPPIV does not cleave fibronectin (63). Following CD244 depletion, neutrophils were 97% pure and rDPPIV was still able to cause repulsion, indicating that a factor secreted by cells contaminating the neutrophil preparation is not likely responsible for the effect. There are no known neutrophil-secreted neutrophil chemottractants that are, or could be, affected by DPPIV (25, 64–68). Conditioned media from neutrophils incubated with buffer alone did not cause attraction or repulsion of neutrophils, supporting this idea. Conditioned media from neutrophils incubated with DPPIV caused chemorepulsion of neutrophils. The chemorepulsive activity of the conditioned media was lost with the addition of a DPPIV inhibitor, suggesting that the chemorepulsive activity in the conditioned media is due specifically to the presence of DPPIV activity. Although DPPIV-induced neutrophil chemorepulsion appears to require the DPPIV active site, the chemorepulsion does not appear to be caused by DPPIV’s enzymatic activity on material in the medium, on chemottractants or repellents, or by components from cells.

DPPIV knockout mice have been generated (53) and rat strains lacking DPPIV have been identified (69, 70). DPPIV-deficient animals have normal blood levels of most leukocytes, including neutrophils (71–73). Arthritic joints in mammals contain abnormally high numbers of neutrophils (74). DPPIV knockout mice have increased severity of experimentally induced arthritis, with a 2.4-fold increase in the number of cells in the joint (75). A 2-fold increase in neutrophils occurs in OVA-induced lung inflammation in DPPIV-deficient rats compared with normal rats (76). Reduced levels of DPPIV correlate with increased inflammation in the joints of rheumatoid arthritis patients (77). The increased inflammation observed in DPPIV-deficient situations has been assumed to be due to a persistent chemokine presence, as DPPIV was not present to cleave those chemokines that have an alanine or proline as the second amino acid (78).

We found that oropharyngeal aspiration of DPPIV caused a reduction in the accumulation of neutrophils, but not other immune cells, in the lungs of mice treated with blemcin, suggesting that the chemorepulsive effect of DPPIV is specific to neutrophils. Additionally, the reduction in neutrophils was not due to increased neutrophil death. Our data suggests an alternative hypothesis to the persistence of chemokines in a tissue. We hypothesize that DPPIV acts directly as a neutrophil chemorepellent. Following neutrophil influx, activated T cells enter a site of inflammation (79). Activated T cells highly express membrane DPPIV, and extracellular DPPIV is thought to be cleaved from the membrane of T cells (22). An increase in extracellular DPPIV could promote neutrophil egress from the inflammatory site. In the resolution of zebrafish wounds, neutrophils are able to move out of the tissue and back into the vasculature (80). An intriguing possibility is that T cell–released DPPIV may help cause the egress of neutrophils out of the lung tissue to resolve lung inflammation. Some lung diseases such as acute respiratory distress syndrome involve an excess number of neutrophils in the lungs (81). The ability of DPPIV to induce neutrophil chemorepulsion suggests the existence of new mechanisms that may be used to treat these diseases.

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Disclosures
The authors have, through Texas A&M University, applied for a patent on the use of DPPIV as a neutrophil chemorepellent.

References


