Human Platelets Utilize Cyclooxygenase-1 to Generate Dioxolane A₃, a Neutrophil-activating Eicosanoid

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Eicosanoids are important mediators of fever, pain, and inflammation that modulate cell signaling during acute and chronic disease. We show by using lipidomics that thrombin-activated human platelets generate a new type of eicosanoid that both stimulates and primes human neutrophil integrin (Mac-1) expression, in response to formylmethionylleucylphenylalanine. Detailed characterization proposes a dioxolane structure, 8-hydroxy-9,11-dioxolane eicosatetraenoic acid (dioxolane A₃, DXA₃). The lipid is generated in nanogram amounts by platelets from endogenous arachidonate during physiological activation, with inhibition by aspirin in vitro or in vivo, implicating cyclooxygenase-1 (COX). Pharmacological and genetic studies on human/murine platelets revealed that DXA₃ formation requires protease-activated receptors 1 and 4, cytosolic phospholipase A₂ (cPLA₂), Src tyrosine kinases, p38 MAPK, phospholipase C, and intracellular calcium. From data generated by purified COX isoenzymes and chemical oxidation, we propose that DXA₃ is generated by release of an intermediate from the active site followed by oxygenation at C8. In summary, a new neutrophil-activating platelet-derived lipid generated by COX-1 is presented that can activate or prime human neutrophils, suggesting a role in innate immunity and acute inflammation.

Emerging evidence indicates that platelets influence innate immunity during acute infection and injury through their interactions with leukocytes (1–3). Platelets generate soluble lipid-signaling mediators that include eicosanoids such as thromboxane A₂ (TXA₂) and 12-hydroxyeicosatetraenoic acid (12-HETE) and small amounts of the prostaglandins (PGs) PGE₂ and D₂. Currently the effects of platelet-derived lipids on leukocytes are not fully known. In this study, we sought to discover whether platelets release leukocyte-regulating lipids using a lipidomic approach. Analogous methodologies have recently been used for discovery of lipids in diabetes, cardiovascular disease, and hemostasis (4–6).

Herein, we show that thrombin-activated human platelets generate a novel eicosanoid from endogenous substrate, proposed to be a dioxolane (DX), that elevates Mac-1 (CD11b/CD18) on neutrophils at nanomolar concentrations. We also present the detailed cellular and enzymatic mechanisms of formation along with characterization of its proposed covalent structure and established a quantitative assay. These data demonstrate a new platelet-derived leukocyte-activating eicosanoid and the first DX lipid to originate from mammalian cells, suggesting a novel mechanism for promoting neutrophil activities in the early stage of tissue damage/wounding responses.

Experimental Procedures

Materials

Lipids and lipid standards were purchased from Avanti Polar Lipids (Alabaster, AL) or Cayman Chemical (Ann Arbor, MI). Deuterated standards are as follows: arachidonic acid-d₄, 5Z,8Z,11Z,14Z-eicosatetraenoic-5,6,8,6,54 PM 5/12/20169, 11,12,14,15-d₄ acid, ≥99% deuterated forms; PGE₂-d₄, 9-oxo-11α,15S-dihydroxy-prosta-5Z,13E-dien-1-oic-3,3,4,4-d₄ acid, ≥99% deuterated forms; and PGD₂-d₄, 9α,15S-dihydroxy-11-oxo-prosta-5Z,13E-dien-1-oic-3,3,4,4-d₄ acid, ≥99% deuterated forms. HPLC grade solvents were from Thermo Fisher Scientific (Hemel Hempstead, Hertfordshire, UK). PAR-1 and PAR-4 agonists were from Tocris Biosciences (Bristol, UK). COX-1 inhibitor (Sc-560) was from Cayman Chemical. Platelet signaling inhibitors (PO2, oleyloxyethylphosphocholine (OPEC), bromoeno lactone, cytosolic phospholipase A₂α (cPLA₂α) inhibitor (N-[(2S,4R)-4-(biphenyl-2-yl-methyl)-isobuty-l-amino)-1-[2-(2,4-difluorobenzoyl)-benzoyl]-pyrrolidinome-3-nitroso-1-oxyl]-9H-fluoren-9-one (FNTS), NMBHA, N-methyl benzohydroxamic acid; ANOVA, analysis of variance; HPETE, hydroperoxyeicosatetraenoic acid; FTMS, fourier MS; CID, collision-induced dissociation; AA, arachidonic acid; PLC, phospholipase C; LOX, lipoxgenase.
A New Bioactive Eicosanoid Generated by Human Platelets

Human blood donations were approved by the Cardiﬁ University School of Medicine Ethics Committee, with informed consent (SMREC 12/37 and SMREC 12/10), and according to the Declaration of Helsinki. For cPLA₂-deficient samples, samples were approved by St Thomas’s Hospital Research Ethics Committee, reference 07/Q0702/24: patient samples; South East NHS Research Ethics Committee. For studies on isolated platelets, whole blood was collected from healthy volunteers free from non-steroidal anti-inﬂammatory drugs for at least 14 days and added to acid/citrate/dextrose (ACD; 85 mmol/liter trisodium citrate, 65 mmol/liter citric acid, 100 mmol/liter glucose) (blood/ACD, 8:1:19, v/v) then centrifuged at 250 × g for 10 min at room temperature. Platelet-rich plasma was collected and centrifuged at 900 × g for 10 min, and the pellet was resuspended in Tyrode’s buffer (134 mmol/liter NaCl, 12 mmol/liter NaHCO₃, 2.9 mmol/liter KCl, 0.34 mmol/liter Na₂HPO₄, 1.0 mmol/liter MgCl₂,10 mmol/liter Hepes, 5 mmol/liter glucose, pH 7.4) containing ACD (9:1, v/v). Platelets were centrifuged at 800 × g for 10 min and then resuspended in Tyrode’s buffer at 2 × 10⁶/ml. Platelets were activated at 37 °C in the presence of 1 mmol/liter thrombin, as described above, and then lipids were extracted as described below. Exclusion criteria was a known sensitivity to aspirin. For studies on isolated murine platelets, whole blood was collected using cardiac puncture (mice were 28 weeks old) into 150 µl of ACD (85 mmol/liter trisodium citrate, 71 mmol/liter citric acid, 100 mmol/liter glucose). 150 µl of 3.8% sodium citrate and 300 µl of Tyrode’s buffer (145 mmol/liter NaCl, 12 mmol/liter NaHCO₃, 2.95 mmol/liter KCl, 1.0 mmol/liter MgCl₂,10 mmol/liter Hepes, 5 mmol/liter glucose, pH 7.35) were added, and the blood was centrifuged at 150 × g for 5 min at room temperature. Platelet-rich plasma was collected, and 400 µl of Tyrode’s buffer was added to the red cells and centrifuged again at 150 × g for 5 min at room temperature. Platelet-rich plasma was combined and centrifuged at 530 × g for 5 min at room temperature. Platelets were resuspended in Tyrode’s buffer at 2 × 10⁸/ml. All animal experiments were performed in accordance with the 1986 United Kingdom Home Office Animals Act (Scientific Procedures). 12/15-LOX knock-out mice were generated as described previously (10), and wild-type male C57BL/6 mice (25–30 g) from Charles River, UK, were kept in constant temperature cages (20–22 °C) and given free access to water and standard chow.

Isolation and Activation of Human Neutrophils

Human neutrophils were isolated from 20 ml of citrate anti-coagulated whole blood and resuspended in Krebs buffer. Briefly, blood was mixed 1:3 with 2% trisodium citrate (w/v) and HetaSep (Stemcell Technologies) and allowed to sediment for 45 min at 20 °C. The upper plasma layer was recovered and underlaid with ice-cold Lymphoprep™ (2:1 for Plasma/Lymphoprep™) and centrifuged at 800 × g for 20 min at 4 °C. The pellet was resuspended in ice-cold PBS and 0.4% sodium tricinate (w/v) and centrifuged at 400 × g for 5 min at 4 °C. Contaminating erythrocytes were removed using up to three cycles of hypotonic lysis. Finally, cells were resuspended in a small volume of Krebs buffer (100 mmol/liter NaCl, 50 mmol/liter Hepes, 5 mmol/liter KCl, 1 mmol/liter MgCl₂, 1 mmol/liter NaH₂PO₄, 1 mmol/liter CaCl₂, and 2 mmol/liter d-glucose, pH 7.4), counted, and kept on ice. Neutrophils were diluted to 2 × 10⁶ cells/ml and incubated with or without DXA₃ for 10 min at 37 °C. In some experiments, 10 µM fMLP was then added, and neutrophils were incubated for a further 10 min at 37 °C. Cells were blocked using 5% mouse serum in PBS (containing 0.5% BSA, 5 mmol/liter EDTA, and 2 mmol/liter sodium azide) for 1 h on ice and centrifuged at 320 × g for 5 min at 4 °C. Anti-human CD11b-Alexa Fluor 647 (0.0625 µg, eBioscience) or isotype control were added and incubated for 30 min on ice. Neutrophils were washed twice with ice-cold PBS (containing 0.5% BSA, 5 mmol/liter EDTA, and 2 mmol/liter sodium azide) and dissolved in the same buffer for flow cytometric analysis. Neutrophils were analyzed on a cyan ADP flow cytometer (Beckman Instruments) and identified by forward and side scatter and Alexa Fluor 647. DXA₃ used for these experiments was purified from COX-1 incubations. Other lipids were not detectable in these preparations using MS.

Culturing and Activation of RAW 264 Cells

RAW 264 cells were cultured in DMEM (10% FBS, 1 × penicillin/streptomycin) at 37 °C and 5% CO₂.
A New Bioactive Eicosanoid Generated by Human Platelets

To determine PG synthesis, cells were incubated in serum-free DMEM (with 1× penicillin/streptomycin), and the cells were incubated for 1 h at 37 °C, 5% CO₂. Where used, 200 ng/ml LPS was added, and cells were incubated for 24 h. Cells (8 × 10⁶ ml⁻¹) were treated with 10 μM ionophore at 37 °C for 10 min, and lipids were extracted and analyzed as described below.

**Isolation of Human Serum**

Whole blood from healthy volunteers was clotted at 37 °C for 15 min in glass and centrifuged (1500 rpm, 10 min, 4 °C). Serum was re-spun (2900 rpm, 10 min, 4 °C), and 3 volumes of MeOH/water (20:80 v/v) were added. Protein precipitates were spun down (13,000 rpm, 10 min, 4 °C), and supernatants were applied to preconditioned Waters C18 Sep-Pak columns. These were washed with 10 ml of water, 6 ml of hexane, and the eicosanoids were then eluted using 7 ml of methyl formate into tubes containing 6 μl of MeOH/glycerol (70:30 v/v) (11). Lipid were redissolved in methanol, chilled (−80 °C, 60 min), and re-spun (13,000 rpm, 10 min, 4 °C) before LC/MS/MS analysis for DXA₃.

**Lipid Extraction**

Lipids were extracted by adding a solvent mixture (1 mol/liter acetic acid, isopropyl alcohol, hexane (2:20:30, v/v/v)) to the sample at a ratio of 2.5 ml to 1 ml of sample, vortexing, and then adding 2.5 ml of hexane (12). Where quantitation was required, 5–10 ng of PGE₂₇₄₄, PGD₂₇₄₄, and 12-HETE-₇₄₄ were added to the samples before extraction, as internal standards. After vortexing and centrifugation, lipids were recovered in the upper hexane layer. The samples were then re-extracted by addition of an equal volume of hexane. The combined hexane layers were dried and analyzed for DXA₃ using LC/MS/MS as described below.

**Generation of DXA₃ through Oxidation of 11-HPETE or by Purified or Recombinant COX Isoforms**

Arachidonic acid was oxidized using N-methylbenzhydroxamic acid (NMBHA) and 2,2’-azobis(4-methoxy-2,4-dimethylvaleronitrile) (MeOAMVN) as detailed below. To a 6.5 μM arachidonic acid solution in chlorobenzene, 3.5 eq of NMBHA and 0.1 eq of MeOAMVN were added, and the mixture was stirred at 37 °C for 5 h under O₂. After drying under N₂, the sample was dissolved in methanol and stored at −80 °C until purification. Isolation of positional isomers used a Spherisorb ODS2 column (5 μm, 150 × 4.6 mm; Waters) with a gradient of 50–90% solvent B (acetonitrile, 0.1% formic acid) in solvent A (water, 0.1% formic acid) for 60 min, 90% solvent B for 4.5 min, and then re-equilibrating to 50% solvent B over 9.5 min with a flow rate 1 ml/min⁻¹. Elution was monitored at 205 nm (unoxidized lipid) and 235 nm (HPETE). Fractions were collected and positional isomers identified using MS/MS transitions for the free HPETEs as follows: m/z 317.2 → 115.1 (5-HPETE); m/z 317.2 → 155.1 (8-HPETE); m/z 317.2 → 151.1 (9-HPETE); m/z 317.2 → 167.1 (11-HPETE); m/z 317.2 → 179.1 (12-HPETE); and m/z 317.2 → 219.1 (15-HPETE). Next, purified 11-HPETE was oxidized using 0.1 eq of MeOAMVN in 5 ml of chlorobenzene by stirring at 37 °C for 5 h under O₂, and the hydroperoxides were then reduced to corresponding hydroxides using tin chloride (SnCl₂). Lipids were extracted using the hexane/isopropyl alcohol extraction method, as described earlier. Apo-COX-1 was stored in 80 mM Tris, pH 7.8, at −80 °C. In some experiments, a commercial preparation was used (Cayman Chemical). Wild-type murine COX-2 (recombinant) was at 10.61 mg ml⁻¹. For heme reconstitution, apo-COX-1 or -2 (35 μg) was preincubated on ice for 20 min with 2 molar equivalents of hematin in phosphate buffer (100 mM potassium phosphate buffer, pH 7.4). Then, 3.5 μg of the reconstituted enzyme was added to 1 ml of phosphate buffer and 500 μmol/liter phenol and incubated for 3 min at 37 °C in the presence of 150 mM arachidonate (AA or AA-d₃). Where [¹⁴C]AA was used, 9.8 μg of enzyme was incubated with 70 μM AA (259 kBq). In some experiments, wild-type COX-2 was compared with active site mutants (V349A and W387F). The reaction was stopped by using ice-cold lipid extraction solvent and immediate extraction of lipids after addition of 5 ng each of PGE₂₇₄₄ and PGD₂₇₄₄ as internal standards, when required. In some experiments, 10 μM dihydrodiensatinepentaacetic acid was added just before holo-COX-1. DXA₃ was analyzed using reverse phase LC/MS/MS as described below.

In some experiments, AA was replaced with 1-stearoyl-2-arachidonoyl-phosphatidylethanolamine. Free and esterified DXAs were analyzed using reverse phase LC/MS/MS as below.

**Reversed Phase LC/MS/MS and LC/MS³ of DXA₃ and Platelet Eicosanoids**

Several different LC separations were used on a 4000 Q-Trap platform. For high resolution mass analysis and fragmentation of free DXA₃, a reversed-phase UPLC Fourier Transform MS method was used (Thermo Scientific Orbitrap Elite) using a Spherisorb ODS2 column (5 μm, 150 × 4.6 mm; Waters) with a flow rate of 1 ml/min⁻¹. Solvent B was increased from 20 to 42.5% over 50 min, then increased to 90% over 10.5 min, held for 4 min, and then returned to 20% over 1 min. Equilibration time between runs was 4.5 min. Analysis was performed using heated ESI in negative ion mode at sheath, auxiliary and sweep gas flows of 70, 20, and 0, and capillary and source heater temperatures at 300 and 350 °C, respectively. LC/MS of parent ions was monitored using accurate mass in Fourier MS mode. Negative MS/MS spectra were acquired using higher energy collision-induced dissociation. Data-dependent MS³ of m/z 351 was carried out in ion trap-MS mode on the LTQ ion trap.

For MS/MS of m/z 351 or m/z 359, collision-induced dissociation (CID) was used with a resolving power of 30,000 in negative FTMS mode. Data-dependent MS³ of m/z 351 or m/z 359 from DXA₃-d₃ was carried out in negative FTMS mode with a resolving power of 15,000.

**Generation of a Quantitative Assay for DXA₃**

[¹⁴C]AA was oxidized using COX-1 as described for unlabelled AA above. The amount of [¹⁴C]DXA₃ was determined by comparison with a [¹⁴C]AA standard curve analyzed using LC separation with radiochemical detection (Berthold Technologies) using a Spherisorb ODS2 column (5 μm, 150 × 4.6 mm; Waters) with a gradient of 20–42.5% solvent B (acetonitrile, 0.1% formic acid) in solvent A (water, 0.1% formic acid) over 50 min, 42.5–90% solvent B from 50 to 60 min, 90% solvent B from...
Purification and Derivatization of DXA and GC/MS Analysis

DXA was purified from lipid extracts of thrombin-activated platelets or COX-1 reactions using HPLC/UV on a Spherisorb ODS2 column (5 μm, 150 × 4.6 mm; Waters) with a gradient of 20–42.5% solvent B (acetonitrile, 0.1% formic acid) in solvent A (water, 0.1% formic acid) over 50 min, 42.5–90% solvent B from 50 to 60 min, 90% solvent B from 60 to 64.5 min, 90% to 20% from 64.5 to 65.5 min, and 20% solvent B from 65.5 to 75 min with a flow rate of 1 ml min⁻¹, and fractions were collected at 30-s intervals for LC/MS/MS confirmation of [¹⁴C]DXA. The same gradient was also used for LC/MS/MS detection of [¹⁴C]DXA (m/z 353.2 → 165.1).

Acid Hydrolysis of DXA₃

Semi-purified DXA₃ generated by COX-1 was solubilized in acetonitrile (1 ml) before addition of 1% acetic acid (4 ml). Samples were left at room temperature for 30 min before extraction using a C₁₈ solid phase extraction cartridge.

Tin(II) Chloride Reduction of DXA₃

DXA₃ generated via oxidation of 11-HPETE was reduced using 95 μg of SnCl₂ in water for 10 min at room temperature. Lipids were re-extracted as above using hexane/isopropyl/acetate acid.

Statistics

Data on platelets are representative of at least three separate donors, with samples run in triplicate for each experiment. Data are expressed as mean ± S.E. of three separate determinations. Statistical significance was assessed using an unpaired two-tailed Student's t test. Where the differences between more than two sets of data were analyzed, one-way ANOVA was used followed by Bonferroni multiple comparisons test, as indicated in the figure legends. p < 0.05 was considered statistically significant.

Results

Platelets Generate DXA₃—We initially sought to discover esterified eicosanoids by scanning for precursors of m/z 351.2 in negative ion mode and of lipid extracts from thrombin-activated platelets. This work is published as the characterization of phospholipid esterified PGE₂ and is described elsewhere (13). During precursor scanning for 351.2, we uncovered an unknown lipid also attached to phospholipids that was also generated as a free acid. This is visible when analyzing free acid lipids at m/z 351.2, where two lipids are seeded, including PGE₂, and a more prominent ion at 48 min (marked by *, Fig. 1A) (13). MS/MS demonstrated a complex spectrum with major ions at m/z 163.2 and 165.2 that did not match any known eicosanoids in the LipidMaps database (Fig. 1B). However, a number of ions were indicative of prostaglandins, specifically m/z 333, 315, 289, and 271. The daughter ion at m/z 165.2 was then used to selectively detect the lipid in multiple reaction monitoring mode. A single lipid was visible at 48 min (Fig. 1C). The high resolution m/z [M – H]⁻ of 351.2177 suggests an elemental composition C₃₀H₄₃O₅, corresponding to arachidonic acid plus three oxygen atoms, and the presence of 5 rings/double bonds (Fig. 1B). Its elution on reverse phase LC/MS/MS, considerably later than PGE₂, indicated a less polar lipid (Fig. 1A).

Extensive structural characterization was undertaken using GC/MS, derivatization, MS/MS, and MS³ and suggested the lipid as DXA₃. GC/MS presented in the supplemental material and demonstrates one hydroxyl and no carbonyls (supplemental Fig. 1). For further structural confirmation, DXA₃ or DXA₃-d₉ generated by COX-1 was analyzed using high resolution MS² on an Orbitrap Elite, during LC elution. MS³ of daughter ions at m/z 333.1 and 315.2 indicate the origin of m/z 271.2 (Fig. 1, D–F), whereas MS³ of m/z 225.1 and 207.1 shows the origin of m/z 163.1 (Fig. 2). Of note, m/z 155 is a prominent ion generated on CID fragmentation of 8-HETE, thus supporting the position of the –OH group at C8. To further confirm, analogous experiments were undertaken using deuterated AA as substrate for DXA₃ generation. MS³ of DXA₃-d₉ showed the same fragmentation pattern; however, many ions showed an additional ion at 1 atomic mass unit lower (e.g. m/z 340 and 321, as well as the expected 341 and 322), supporting our proposed fragmentation mechanisms as shown (supplemental Figs. 2 and 3). The lower m/z ions represent ring opening with addition of –H and concomitant loss of a single deuterium, as shown. Finally, a UV spectrum was acquired during purification of low nanogram amounts of COX-1-derived DXA₃, showing a λmax at 238 nm, similar to that reported for a similar DX by Teder et al. (14). This confirms the presence of a UV chromophore and is consistent with a conjugated diene at C12–15 of the lipid backbone (Fig. 3, A and B).
DXA3 is named based on the dioxolane structure, “A” for the first member of the class discovered, and 3 for the number of double bonds, as per traditional eicosanoid naming conventions (15). Further work is required to confirm the structure using NMR and to determine enantiomeric/geometric isomer composition, using synthetic standards once these become available.

**DXA3 Activates Neutrophil Surface Integrin Expression**—Neutrophils incubated with purified DXA3 elevated surface Mac-1 (integrin, CD11b/CD18), comparable with fMLP activation (Fig. 3, C and D). Together, fMLP and DXA3 caused an additive effect on Mac-1 expression, suggesting they activate neutrophils by distinct pathways (Fig. 3, C−F). However, at 10 nM, DXA3 effectively primed for fMLP activation after a 10 min pre-incubation (Fig. 3, G and H).

**Human Platelets Acutely Generate DXA3 on Thrombin Activation via COX-1**—As a purified standard is not yet available, we synthesized and purified a biogenic standard by COX-1 oxidation of [14C]AA oxidation in vitro, using COX-1. This was quantified using radiochemical detection and HPLC-purified. The radiolabeled standard was then utilized in LC/MS/MS.
assays, to quantify cold DXA₃ generated by COX-1, which was then used as a primary standard for quantitation, against PGE₂-d₄ as internal standard. Fig. 4, A–D, shows the LC/MS/MS and MS/MS spectrum of purified [¹⁴C]DXA₃ along with standard curves for both radiochemical detection of [¹⁴C]DXA₃ and DXA₃ versus PGE₂-d₄.

DXA₃ was undetectable basally with levels rising by 10 min of thrombin activation (n = 7–10 separate donors, mean ± S.E.). Levels were higher than PGE₂, but lower than TXB₂ or 12-HETE (a representative donor is shown in Fig. 4E, data for all donors as Fig. 4F). Levels varied between genetically unrelated donors for all eicosanoids. DXA₃ formed on activation by collagen, ionophore, or collagen/thrombin of platelets, with high levels already apparent 2 min post-activation (Fig. 5A). Its thrombin-dependent formation was blocked by the selective COX-1 inhibitor, SC560, aspirin, or indomethacin in vitro or in vivo following administration of 75 mg/day aspirin for 7 days in healthy donors (Fig. 5, B–D). DXA₃ was absent in thrombin-activated platelets from a patient with genetic deficiency of cPLA₂ (16) or in the presence of the cPLA₂ inhibitor, cPLA₂i (Fig. 5, E and F). In contrast, neither iPLA₂ nor sPLA₂ appeared significantly involved (data not shown). Pharmacological inhibitors/agonists implicated PAR-1 and -4 receptors, src-tyrosine kinase, p38 MAPK, intracellular calcium, and PLC, but ruled out phosphatidylinositol 3-kinase, although PKC played an inhibitory role (Fig. 6, A–D). Murine platelets also generated

![Diagram of DXA₃ fragmentation](https://example.com/diagram.png)
FIGURE 3. **DXA3 contains a UV chromophore and primes and activates neutrophil integrin expression.** A, DXA3 can be detected at 235 nm, during LC elution. DXA3 was generated using COX-1 as described under “Experimental Procedures” and then purified using LC/UV. B, DXA3 contains a UV chromophore. A UV spectrum was acquired at the apex of the peak at 46.7 and shows a \( \lambda_{\text{max}} \) at 238 nm. C, DXA3 activates neutrophil Mac-1 expression. Neutrophils were incubated with fMLP, DXA3, or both, before addition of anti-human CD11b (Mac-1)-Alexa Fluor 647 and flow cytometry analysis as under “Experimental Procedures.” A representative experiment repeated with three individual donors is shown (\( n=3 \), mean ± S.E.). D–F, representative histograms depicting increased Mac-1 expression following activation with fMLP, DXA3, or fMLP/DXA3. Line represents the Mac-1-positive neutrophil gate, as set using untreated neutrophils. G, DXA3 primes neutrophil responses to fMLP. Bar chart showing activation of Mac-1 expression in three donors by fMLP with/without DXA3 priming for 10 min (\( n=3 \) mean ± S.E.). H, representative histogram showing increased fMLP-stimulated Mac-1 expression following priming by DXA3. Bar represents the Mac-1 positive neutrophil gate, as set using untreated neutrophils. Statistical significance used Mann-Whitney U test, ****, \( p < 0.0001 \); *** \( p < 0.001 \); **, \( p < 0.01 \). n = 3 donors.
DXA₃ at levels similar to human cells; however, levels were significantly higher in platelets genetically deficient in a second arachidonate-oxidizing enzyme, 12-lipoxygenase (LOX) (Fig. 6E). Similarly, other COX-1-derived lipids were elevated in these platelets, and 12-HETE was absent (Fig. 6F–H). Collectively, these data show that thrombin-stimulated DXA₃ generation depends on a highly coordinated signaling pathway, culminating in cPLA₂-dependent hydrolysis of AA from phospholips, prior to its oxygenation by COX-1 (Scheme 1).

**Fig. 4.** Setting up a quantitative assay for DXA₃ and determining its levels in human platelets. [¹⁴C]DXA₃ was generated, using COX-1 and purified using HPLC with radiochemical detection, as described in the supplemental material. [¹⁴C]DXA₃ was quantified by comparing the radiochemical response to [¹⁴C]AA. A, LC/MS/MS analysis of m/z 353.2 → 165.1 showing [¹⁴C]DXA₃ eluting at 49.5 min, undertaken on the 4000 Q-Trap as described under “Experimental Procedures.” B, MS/MS spectrum of [¹⁴C]DXA₃ showing m/z 353.2 as parent ion. Experiment was performed on the 4000 Q-Trap platform in enhanced product ion mode, as described under “Experimental Procedures.” C, standard curve for [¹⁴C]DXA₃ using LC/MS/MS detection used for quantification of unlabeled DXA₃. D, standard curve for quantitation of DXA₃ in biological samples. A standard curve was generated with varying DXA₃, but keeping PGE₂-d₄ levels constant, and responses were plotted as shown. E, time course of eicosanoid generation by thrombin-activated platelets. Washed platelets were activated for varying times, using 0.2 unit/ml thrombin, then lipids were extracted and analyzed using reverse phase LC/MS/MS, monitoring parent m/z 351.2 → 165.1 (DXA₃), 319.2 → 179.1 (12-HETE), m/z 351.2 → 271.1 (PGE₂), and m/z 369.2 → 169.1 (TXB₂), as described under “Experimental Procedures.” Data are representative of experiments repeated three times on different donors (n = 3, mean ± S.E.). F, levels of eicosanoids generated by genetically unrelated volunteers. Data are shown as Tukey boxplots, where whiskers represent 1.5 the lower and upper interquartile range, data not included within the whiskers are displayed as an outlier. Statistical significance used Mann-Whitney U test, ***, p < 0.0001; ***, p < 0.001; **, p < 0.01. n = 7–10 donors.
FIGURE 5. Time course of agonist-stimulated generation and demonstrating the requirement for COX-1 and cPLA₂ in DXA₃ formation. A, generation of DXA₃ by human platelets. Washed platelets were activated for varying times, using 0.2 unit/ml thombin, 10 μg/ml collagen, 10 μmol/liter A23187, and then lipids were extracted and analyzed using reverse phase LC/MS/MS, monitoring parent m/z 351.2 → 165.1 as described under "Experimental Procedures." Levels are expressed as analyte/internal standard. Data are representative of experiments repeated at least three times on different donors (n = 3, mean ± S.E.). B and C, requirement for COX-1 for DXA₃ formation. Platelets were incubated with inhibitors 10 min prior to thombin activation (0.2 units/ml for 30 min at 37 °C). Lipids were extracted and analyzed using LC/MS/MS monitoring m/z 351.2 → 165.1, as described under "Experimental Procedures." Data are representative of experiments repeated at least three times on different donors (n = 3, mean ± S.E.). ***, p < 0.001 versus thombin, using ANOVA and Bonferroni post hoc test. Inhibitors used were aspirin, SC-560 (COX-1 selective), or indomethacin/aspirin (non-selective COX inhibition). D, in vivo aspirin blocks platelet DXA₃ generation. Lipids were analyzed following thombin activation of washed platelets, before or after supplementation with 75 mg/day aspirin for 7 days. Data are representative of five independent donors (n = 5, mean ± S.E.); ***, p < 0.001 versus thombin alone, using ANOVA and Bonferroni post hoc test. E, cPLA₂ is required for DXA₃ formation. Platelets were incubated with 50 nM cPLA₂ inhibitor (cPLA₂i) 10 min prior to thombin activation (0.2 units/ml for 30 min at 37 °C). Lipids were extracted and analyzed using LC/MS/MS monitoring m/z 351.2 → 165.1, as described under "Experimental Procedures." Data are representative of experiments repeated at least three times on different donors (n = 3, mean ± S.E.). ***, p < 0.001 versus thombin, using ANOVA and Bonferroni post hoc test. F, platelets genetically deficient in cPLA₂ do not generate DXA₃. Washed human platelets from a patient genetically deficient in cPLA₂ or a healthy control were activated using thombin (0.2 units/ml for 30 min at 37 °C) before lipid extraction and analysis using reverse phase LC/MS/MS, monitoring m/z 351.2 → 165.1 as described under "Experimental Procedures."
and also synthesized the free acid form of DXA<sub>3</sub>, through oxidation of 11-HPETE, as described by Porter and co-workers for generating cholesteryl-esterified dioxolane lipids (17, 18). In all reactions, an ion with same retention time and MS/MS spectrum as the platelet lipid was formed (Fig. 7, A–H). However, for either COX or 11-HPETE oxidation-generated DXA<sub>3</sub>, two additional ions with m/z 351.2 → 165.1 were seen eluting just before and after DXA<sub>3</sub>, which may represent enantiomers or positional isomers, e.g. at C8, C9, or C11. Monocyclic isomers contain three chiral centers, thus eight possible stereoisomers (or four pairs of enantiomers): RRR, RRS, RSS, SSS, SSR, SRR, SRS, and RSR (19). These additional ions show identical MS/MS spectra to DXA<sub>3</sub> (data not shown) suggesting these peaks to be isomers. With enantiomers not being separated by...
our chromatography, up to four peaks of isomers would be expected. The absence of these isomers in platelet extracts (Fig. 1C) indicates a higher degree of control over the cellular biosynthesis of DXA<sub>3</sub>, preventing generation of stereoisomers. This may indicate that additional unknown enzymatic pathways exert control of DXA<sub>3</sub> generation in platelets and will be subject to further study.

Platelets generate significant amounts of 11-HETE via COX-1 turnover (20, 21). This likely results from 11-hydroperoxyl radical intermediates (11-LOO°) exiting the catalytic site, then being reduced to form 11-HETE. Similarly, we reasoned that DXA<sub>3</sub> could form by COX via rearrangement of an enzyme-generated intermediate exiting the active site early, before full prostanoid ring formation. This could occur either at the 11-LOO° or 9,11-dioxolane radical stage (e.g. just before or after formation of the DX ring). To examine this, we measured DXA<sub>3</sub> formation by two COX-2 mutants that generate more 11-HETE and less PGH<sub>2</sub> than wild-type enzyme (22). Thus, these enzymes favor escape of lipid radicals prior to DX/prostanoid ring formation. Both mutants were found to generate less DXA<sub>3</sub>, indicating that the DX ring forms before DXA<sub>3</sub> leaves the active site (Fig. 8, A–C, and Scheme 2). Following escape of a 9,11-DX radical, oxygen addition at C8 is expected, followed by peroxidase-dependent reduction. This could be mediated by COX-1 peroxidase or GSH peroxidase. In support, DXA<sub>3</sub> formation was inhibited by 1–10 mM iodoacetate, a thiol-alkylating reagent (Fig. 8D).

**Generation of DXA<sub>3</sub> Is Independent of Thromboxane Synthase**—To determine the role of enzymatic activities downstream of COX-1, an inhibitor of thromboxane synthase was added to platelets during activation. Picotamide led to inhibition of TXB<sub>2</sub> generation and a corresponding elevation in PGE<sub>2</sub>, because less PGH<sub>2</sub> was being converted by thromboxane synthase (Fig. 8, E and F). However, DXA<sub>3</sub> formation was unaffected (Fig. 8G). Last, there was no correlation between TXB<sub>2</sub> and DXA<sub>3</sub> levels, further supporting the idea that thromboxane synthase is not involved in DXA<sub>3</sub> generation (data not shown).

**Generation of DXA<sub>3</sub> by RAW 264 Cells and Human Serum**—To determine generation of DXA<sub>3</sub> in other cell types, RAW 264 macrophages were treated using LPS for 24 h, with/without...
ionophore activation. Under basal conditions, these cells express only COX-1, although following LPS treatment, they up-regulate COX-2. We found that cells required ionophore for robust PG generation. DXA₃ formation paralleled that of PGP₂, being present basally, but unaffected by inflammatory activation. Thus, the lipid was most likely generated by COX-1

**FIGURE 7.** DXA₃ is generated by purified COXs and via oxidation of 11-HPETE. A and B, COX isoforms generate DXA₃. 3.5 μg of apo- or holo-COX-1 or COX-2, or hematin (control for reconstituted enzyme), or DMSO (vehicle for hematin) was incubated with 150 μM of AA for 3 min at 37 °C, before lipid extraction and analysis as described under “Experimental Procedures.” Levels are expressed as ratio analyte to internal standard/3.5 μg of enzyme generated over 3 min (n = 3, mean ± S.E.). Data are representative of ≈3 separate experiments. C and D, LC/MS/MS of DXA₃ formed in vitro via COX-1 or -2. Lipid extracts were separated using reverse phase LC/MS/MS, monitoring m/z 351.2 → 165.1, with reactions as described under “Materials and Methods.” E, LC/MS/MS of DXA₃ formed in vitro via 11-HPETE oxidation. Purified 11-HPETE was oxidized as described under “Experimental Procedures” and separated using LC/MS/MS. F and G, MS/MS spectra of DXA₃ formed in vitro via COX-1 or -2. Lipid extracts were separated as in C and D. MS spectra were acquired at the apex of elution of DXA₃. * shows position of additional isomers with identical MS/MS spectra to DXA₃ eluting either before or after lipid.
but not COX-2 in these cells (Fig. 9, A–D). Human blood was harvested and allowed to clot. Analysis of serum demonstrated a significant DXA$_3$ peak, indicating that physiological coagulation forms this lipid (Fig. 8E). In contrast, DXA$_3$ was absent from plasma (data not shown).

**Discussion**

Herein, we used a lipidomic approach to identify and characterize a new neutrophil-activating lipid, proposed to be DXA$_3$, formed endogenously by agonist-activated platelets in a COX-1-dependent manner, by a macrophage cell line, and during blood
At this time, we present a proposed structure based on strong and consistent UV, GC/MS, and LC/MS data. Once the sufficient synthetic standard is available, full NMR analysis will be undertaken. We note that many other biologically relevant lipids, including thromboxane, leukotrienes, protectins, etc., were first published as proposed structures in a similar manner to our study.

SCHEME 2. Proposed mechanism of DXA₃ formation by COX. During COX turnover, a dioxolane ring forms between C9 and C11, prior to prostanoid ring formation, resulting in a carbon-centered radical at C8. Leakage of this lipid intermediate from the active site, then addition of oxygen followed by reduction to LOOH, and then LOH leads to formation of DXA₃.
The mechanism of DXA₃ formation in vitro by COX enzymes is described, as well as its detailed cellular biosynthesis pathway in human platelets. DXA₃ represents the first DX eicosanoid isolated and characterized within cells. To date, these have only been demonstrated to form via chemical oxidation of purified arachidonate esters or ω3 fatty acids or by in vitro lipoxygenase oxidation of epoxides, and neither their generation by cells nor any bioactivities have been described (14, 18, 19, 24, 25, 27, 28). Our study greatly extends these old in vitro observations by demonstrating that DX lipids are not only generated by live primary cells under physiological conditions, but they possess biological activity of relevance to innate immunity. This study places this eicosanoid in a new family of products likely relevant as a lipid mediator as are the prostaglandins, leukotrienes, and P450-derived eicosanoids. Extending these cell biology studies to in vivo measurements of leukocyte function and inflammation will be undertaken as soon as the synthetic standard becomes available.

Eicosanoids are essential lipid signaling mediators involved in diverse biological processes (29–32). Identification of new
bioactive eicosanoids from this pathway could pave the way for additional and more selective therapeutic approaches. Thus, the proposed structure for DXA₃ represents a new member of this family, characterized by a unique five-membered endoperoxide ring, and generated by a COX isoform known to play important roles in vascular disease and, more recently, in cancer.

Mac-1 (CD11b/CD18) is the predominant β2 integrin on neutrophils that mediates adhesion-dependent processes, such as binding to the endothelium or phagocytosis, recruitment, and transendothelial migration (33, 34). Herein, we show that DXA₃ enhances Mac-1 on the cell surface (Fig. 3). The only other known Mac-1-inducing eicosanoids are leukotriene B4 and 5-oxo-ETE, both neutrophil-derived lipids (35, 36). Thus, neutrophil integrin activation by platelet-derived DXs could be of relevance during acute inflammation and infection. DXA₃ was generated by platelets utilizing endogenous substrate in nanogram amounts that are ~10-fold higher than platelet PGE₂ (Fig. 4E). Its formation does not require supply of exogenous substrates and can be triggered directly by pathophysiological agonists in healthy primary cells, both important criteria in establishing that a new lipid mediator is endogenously relevant.

As DXA₃ was generated via COX-1 in platelets, we reasoned that it could form through two potential mechanisms, either (i) rearrangement of 11-LOO⁺, known to be released by the enzyme during turnover, or (ii) that the dioxolane ring could form before the lipid exits from the active site (20, 21, 23, 37). In both cases, attack at C9 by the peroxyl radical would form the C8, and finally peroxidase reduction of the resulting LOOH by COX-1 peroxidase or GSH peroxidase in platelets. Our data using mutant COX-2 enzymes that generate less DXA₃ but more 11-HETE suggest that the DX ring forms before lipid release by the enzyme. Thus, dioxolane ring formation occurs first and before prostanoid ring closure between C8 and C12 (Scheme 2). Finally, given that COX-1 generates 11R-HETE, we postulate that the dioxolane ring will likely be 9S,11R. Our observation of a single DX isomer in platelets but several in purified enzyme reactions indicates that platelets exert additional control over its biosynthesis. This may be at the stage of oxygen insertion into the chiral center at C8.

DXA₃ was generated by platelets via a highly coordinated sequence of signaling events, including PAR-1 and -4, src tyrosine kinases, intracellular Ca²⁺, cPLA₂, PLC, p38, and MAPK. This indicates tight control of its formation, similar to generation of other COX metabolites, such as TXA₂. The signaling pathway is distinct from generation of free and esterified HETE and hydroxydocosahexaenonic acids, which form via 12-LOX, and require extracellular calcium, independent of PLC and MAPK (6, 26).

DXA₃ was also generated by RAW cells as a single isomer, similar to platelets. Our preliminary data suggest that it originates primarily from COX-1 in these cells. In contrast, we found that either isomer could generate the lipid in vitro. In line with our observation that cellular DXA₃ is a single isomer in platelets and RAW cells, although three isomers form via COXs in vitro, this collectively suggests that cellular DXA₃ generation is under enzymatic control downstream of its synthesis by COX-1. Future studies will examine the ability of cellular COX-2 to generate the isoform and under which activation conditions. COX-1 is important not only in acute innate immunity but also in gastric function and development, and thus its generation by this isoform may have wider implications for eicosanoid biology in other organs.

Murine platelets also generated DXA₃, and levels of this were enhanced in cells deficient in 12-lipoxygenase. This may be related to greater availability of substrate, although this has not been explored herein.

Eicosanoids include a large number of related structures formed via oxidation of arachidonate, following its release from intracellular membranes by phospholipases. A rapid burst of eicosanoid generation is a key event during cell activation and is stimulated during innate immunity by bacterial products, growth factors, cytokines, thrombin, and collagen. Most known eicosanoids from COXs were identified and structurally characterized in the 1980–1990s and include platelet-derived lipids, TXA₃ and 12-HETE, as well as the PGs, exemplified by PGE₂, and D₂, well known as mediators of pain, fever, cell proliferation, and innate and adaptive immune responses. Our observation of a cellurally generated DX eicosanoid defines a new class of these lipids formed endogenously by mammalian cells. More members of this class are possible, given recent observations of purified LOXs being able to generate DX isomers via oxidation of epoxides in vitro in acellular experiments (14).


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A New Bioactive Eicosanoid Generated by Human Platelets


Supplementary Data.

Supplementary Results

*Derivatization, acid hydrolysis and catalytic hydrogenation of DXA₃*

DXA₃ generated by COX-1 oxidation of AA was partially purified using reverse phase HPLC, as described in Supplementary Methods, derivatized using PFB, MOX and/or TMS, and analyzed using GC/MS and/or LC/MS. First, LC/MS of MOX-derivatized DXA₃ showed no loss of lipid, indicating that it does not contain any carbonyl groups (Supplementary Figure 1 A,B). In contrast, a small amount of contaminating PGE₂ was lost on derivatization. It was not possible to analyze TMS-ethers using LC/MS/MS due to derivatization of the carboxyl group by the reagent (required for PFB derivatization and negative ionization and detection). Derivatization of DXA₃ using PFB, then MOX and TMS followed by GC/MS analysis, yielded signals at m/z 423 (+TMS), indicating one hydroxyl (Supplementary Figure 1 C). Note that during negative ion chemical ionization (NICI) analysis, the PFB group is lost generating the carboxylate anion, which is detected as a negative signal. Also, several lipids are detected on derivatization since three isomers as generated by COX-1 (Figure 7 C). Ions were not detected for addition of carbonyls or more than one hydroxyl (not shown). The MS spectrum at 11.4 min shows a molecular ion of m/z 423.2 and a fragment at m/z 333.1 (-90 amu), representing the loss of –OTMS (Supplementary Figure 1 D). This further supports the presence of a single hydroxyl group. Since the remaining two oxygens are neither hydroxyl or carbonyl, this is consistent with the proposed dioxolane structure. Epoxides, but not dioxolanes are acid sensitive. Thus, purified DXA₃ was incubated with 1% acetic acid, and analyzed using LC/MS. No loss of m/z 351 was observed (Supplementary
Figure 1 E,F). We note that DXA₃ will be considerably more lipophilic than PGE₂ or D₂, which both contain additional hydroxyl and carbonyl functional groups, consistent with our observation of later elution on reverse phase HPLC (Figure 1 A).

Supplementary Figure Legends.

Supplementary Figure 1. Derivatization and analysis of DXA₃ using LC/MS/MS and GC/MS, shows the presence of one hydroxyl and no carbonyl or epoxide functional groups. Panels A,B. LC/MS shows DXA₃ contains no carbonyl groups. Semi-purified DXA₃ generated using COX-1 was derivatized using methyloxime (MOX) and analyzed using LC/MS/MS, on the 2000 Q-Trap, monitoring m/z 351 in Q1. Panel A, no derivatization. Panel B, after derivatization. Note: loss of small peak corresponding to contaminating PGE₂ in this preparation confirms derivatization has been successful. Panels C,D. GC/MS shows that DXA₃ contains one hydroxyl group and no carbonyls. Purified DXA₃ generated using COX-1 was derivatized using PFB, MOX and TMS, and analyzed using GC/MS as described in Methods. Panel C. m/z of derivative with one hydroxyl, showing a major peak at 11.38 min, and two additional smaller peaks at either side. Panel D. MS spectrum at 11.38 shows ion at m/z 432, with loss of 90 amu at m/z 333 (-OTMS). Panels E,F. DXA₃ is insensitive to acid hydrolysis indicating no epoxide groups. Semi-purified DXA₃ generated by COX-1 was solubilized in acetonitrile before the addition of 1 % acetic acid (1:4). Following 30 min at 22 °C, lipids were extracted using a C₁₈ Bond Elute cartridge and analyzed using LC/MS/MS on the 2000 Q-Trap for m/z 351.2 in full scan Q1 mode. Panel E: no hydrolysis, Panel F: after hydrolysis.
Supplementary Figure 2. Characterization of DXA\textsubscript{3-d8} MS/MS and MS\textsuperscript{3} fragmentation, using high resolution FTMS. Panel A. LC/MS/MS of COX-1 derived DXA\textsubscript{3-d8}, generated using AA-d8 as substrate. Analysis was undertaken on the Orbitrap Elite in FTMS mode, separating using reverse phase LC, isolating m/z 359.27 in the Velos Pro, then fragmenting using CID at 50 V, with resolution 15,000 ppm, as described in Methods. Panel B. MS\textsuperscript{3} of daughter ion at m/z 340.25 (smaller ion adjacent to m/z 341 in Panel a), with CID 30V. Panel C. MS\textsuperscript{3} of daughter ion at m/z 322.24, with CID of daughter ion at 30V. Panel d. Proposed fragmentation pathway for m/z 359.2679 generating m/z 341.2573, which fragments to m/z 278.2507 via via m/z 322.2405. DXA\textsubscript{3} loses H\textsubscript{2}O forming m/z 341.2573. Following ring opening m/z 340.2511, leaving a keto group at C9, H\textsubscript{2}O and CO\textsubscript{2} are lost, generating m/z 278.2507 via a m/z 322.2405 intermediate, as shown.

Supplementary Figure 3. Characterization of DXA\textsubscript{3-d8} MS/MS and MS\textsuperscript{3} fragmentation, using high resolution FTMS. Panel A. MS\textsuperscript{3} of daughter ion at m/z 231.15 with CID 30V. Panel B. MS\textsuperscript{3} of daughter ion at m/z 212.13, with CID 30V. Panel c. Proposed fragmentation pathway for m/z 359.2 generating m/z 358.2, then via fragmentation of 340.2, forming 212.1 and 168.1. Following ring opening, with keto group at C11, H\textsubscript{2}O is lost, followed by two 1[5]-sigmatropic shifts generating m/z 340.2511. Following loss of a conjugated triene, m/z 231.1509 is generated, which then loses H\textsubscript{2}O, and via an intermediate fragments to m/z 212.1341 and last 168.1422.
Supplementary Figure 1

A. underivatised DXA₃
m/z 351 = DXA₃ and PGE₂

B. MOX derivatised DXA₃
m/z 351 = DXA₃ and PGE₂

C. GC/MS
NICI m/z 423
Corresponding to DXA₃+TMS

D. GC/MS
MS spectrum at 11.38 min

E. LC/MS
No acid hydrolysis,
Extracted m/z 351

F. LC/MS
Post acid hydrolysis,
Extracted m/z 351
Supplementary Figure 2

A) MS/MS spectrum of m/z 359.27

B) MS/MS/MS spectrum of m/z 340.25 from MS/MS (a)

C) MS/MS/MS spectrum of m/z 322.24 from MS/MS (a)

Dioxolane ring opening

- 9-keto

Proposed mechanism For loss of CO₂

- H₂O

m/z 341 and 340 both seen in MS³

m/z 322.2405

m/z 342.2511

m/z 359.2679

m/z 341.2573

m/z 278.2507
**Supplementary Figure 3**

**A**

LC/MS/MS/MS spectrum of m/z 231.15 from MS/MS

Dioxolane ring opening

11-keto

$m/z$ 359.2679

**B**

LC/MS/MS/MS spectrum of m/z 212.13 from MS/MS

Two 1[5]-sigmatropic shifts

$\text{H}_2\text{O}$

$m/z$ 358.2616

Loss of neutral conjugated Triene from C13 to C20

$m/z$ 231.1509

$\text{H}_2\text{O}$

$m/z$ 340.2511

Proposed generation of $m/z$ 168.1442 from $m/z$ 212.1341

$m/z$ 168.1442
Human Platelets Utilize Cyclooxygenase-1 to Generate Dioxolane A₃, a Neutrophil-activating Eicosanoid

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