Sex differences in prostaglandin biosynthesis in neutrophils during acute inflammation

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The severity and course of inflammatory processes differ between women and men, but the biochemical mechanisms underlying these sex differences are elusive. Prostaglandins (PG) and leukotrienes (LT) are lipid mediators linked to inflammation. We demonstrated superior LT biosynthesis in human neutrophils and monocytes, and in mouse macrophages from females, and we confirmed these sex differences in vivo where female mice produced more LTs during zymosan-induced peritonitis versus males. Here, we report sex differences in PG production in neutrophils during acute inflammation. In the late phase (4–8 hrs) of mouse zymosan-induced peritonitis and rat carrageenan-induced pleurisy, PG levels in males were higher versus females, seemingly due to higher PG production in infiltrated neutrophils. Accordingly, human neutrophils from males produced more PGE2 than cells from females. Increased PG biosynthesis in males was accompanied by elevated cyclooxygenase (COX)-2 expression connected to increased nuclear factor-kappa B activation, and was abolished when LT synthesis was pharmacologically blocked, suggesting that elevated PG production in males might be caused by increased COX-2 expression and by shunting phenomena due to suppressed LT formation. Conclusively, our data reveal that the biosynthesis of pro-inflammatory PGs and LTs is conversely regulated by sex with consequences for the inflammatory response.

Sex has emerged as contributing factor in the incidence and progression of diseases associated with the immune system, in particular inflammation, with implications for outcomes and therapies. Most striking sex differences have been observed in asthma and autoimmune diseases (autoimmune thyroid diseases, scleroderma, rheumatoid arthritis etc)1, a spectrum of pathologies in which the patient population is prevalently female. On the other hand, other innate immune disorders such as sepsis2 and post-surgery infections as well as gout1 display a higher incidence and severity in males. The molecular and cellular basis underlying this sex dimorphism is still not completely elucidated and could present important implications for a sex-specific pharmacotherapy.

Sex differences in eicosanoid (leukotriene (LT) and prostaglandin (PG)) production may be responsible, at least in part, for the sex-dependent incidence of many diseases related to inflammation. In fact, we recently reported about a sex-dimorphism in LT biosynthesis (higher in female)4–6 which is of relevance in the light of the well-known sex-biased incidence of several immune diseases, providing a link to asthma pathology in humans7. In particular, androgens exert inhibitory effects on LT formation in human innate immune cells (isolated neutrophils or monocytes, and human whole blood) resulting in a substantial lower LT formation in male cells compared to females4,5. We recently confirmed these sex differences also in vivo, making use of a well-established model of acute inflammation, the zymosan-induced peritonitis in mice6. In fact, LT biosynthesis as well as the inflammatory response were significantly greater in the inflamed peritoneum of female versus male mice. On the cellular level, differential 5-lipoxygenase (LO) subcellular compartmentalization in human leukocytes and in murine peritoneal macrophages (PMs) from males and females might be the basis for the observed differences4,6.

LTs and PGs are locally acting bioactive lipid mediators derived from arachidonic acid (AA) produced by 5-LO and cyclooxygenase (COX) as key enzymes, respectively. They are markedly biosynthesized by monocytes/macrophages, neutrophils, and mast cells and regulate a diverse set of homeostatic and inflammatory processes8.

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linked to numerous sex-dependent diseases. PGs are formed by the action of COX (COX-1 and COX-2) enzymes in a two-step conversion of AA. First, COXs convert AA to a cyclic endoperoxide (PGG₂) and incorporate a 15-hydroperoxy group. This hydroperoxy group of PGG₂ is reduced to a hydroxy moiety yielding PGH₂ that is subsequently converted to the corresponding PGs by specific PG synthases, the nature of which is determined by the enzyme content of the respective cell. COX-1 is constitutively expressed in most cells, thus regarded as a housekeeping protein. On the other hand, the expression of COX-2 is inducible and remains undetectable in most mammalian tissues under basal conditions. Exposure of several types of cells including fibroblasts, neutrophils and monocytes to bacterial endotoxins, cytokines, and hormones induces activation of mitogen-activated protein kinases (MAPK) and nuclear factor-kappa B (NF-κB) which, in turn, induce COX-2 expression and PG production. Inhibiting the formation of PGs by aspirin and other non-steroidal anti-inflammatory drugs during inflammation remains a classic and prevailing strategy to alleviate pain, swelling and fever. A sex-dependent efficacy of aspirin has been underlined in several randomized trials, suggesting differential production or signalling of PGs in males and females. Moreover, as interrelations and crosstalk between the two branches of AA transformation (COX/PGs and 5-LO/LTs) exist, imbalances in LT biosynthesis between genders may translate into differential PG formation.

Here, we show sex differences in PG production in neutrophils during acute inflammation using two different in vivo experimental models, that are, mouse zymosan-induced peritonitis and rat carrageenan-induced pleurisy, as well as in human neutrophils in vitro. Higher PG levels in males are seemingly caused by (i) increased COX-2 expression connected to elevated NF-κB activation versus females and (ii) by AA shunting phenomena due to lower LT production in males.

**Results**

**Sex differences in PG biosynthesis during zymosan-induced peritonitis in mice.** We have previously demonstrated that LT formation in zymosan-induced peritonitis in mice is higher in females compared to males, seemingly due to a divergent subcellular localization of 5-LO in LT-producing peritoneal macrophages. Here, we investigated the temporal PG biosynthesis after intraperitoneal zymosan injection in male and female mice. Both sexes showed a similar time-course in the production of PGE₂ with a peak after 2 hrs and continuous decrease until 8 hrs. Intriguingly, in male mice significant higher levels of PGE₂ in the later phase of inflammation (p < 0.05 at 4 hrs; Fig. 1A) were evident in the peritoneal exudates after zymosan challenge, where the inflammatory cell population was mostly composed of neutrophils. Moreover, in the exudates of male mice after 4 hrs, also higher levels of other PGs, such as 6-keto-PGF₁α, the stable metabolite of PGI₂, were evident as compared with females (p < 0.05, Fig. 1B). In contrast, the levels of LTC₄ that peak after 15 min upon zymosan injection were higher in exudates from female versus male mice (Fig. 1C), which is in agreement with our previous report.

**Sex differences in eicosanoid biosynthesis in carrageenan-induced pleurisy in rats.** Carrageenan-induced pleurisy in rats was chosen as another model of acute inflammation to investigate the sex-related regulation of eicosanoid biosynthesis. Two hrs after pleurisy induction, a peak of LTΒ₄ was observed in both sexes, with a decrease at the later time points (4–8 hrs). In agreement with the data from the peritonitis model in mice, higher levels of LTΒ₄ were found in exudates of female versus male rats (p < 0.001, Fig. 2A) at the peak time (i.e., 2 hrs). The opposite was evident for PGE₂ levels. Thus, a peak of PGE₂ production was reached in both sexes 4 hrs after carrageenan injection and higher PGE₂ levels were obvious in exudates of...
Neutrophils are immune cells involved in the inflammatory reaction, recruited by chemoattractants (i.e. LTB4) that are produced by resident cells17,18. To investigate if the sex difference in PGE2 biosynthesis exists also in humans, neutrophils from blood of female and male donors were freshly isolated and immediately stimulated for PGE2 production by lipopolysaccharide (LPS), a receptor-coupled stimulus, or by A23187 that causes cell activation by substantial amounts of COX enzymes expressed. Based on the sex difference in PG biosynthesis 4 hrs after pleurisy induction in rats, we analyzed the expression of COX-1 and -2 in the infiltrated cells by Western Blot (WB). No significant sex difference was observed for COX-1 expression, whereas COX-2 protein levels were higher in male cells (Fig. 3B,D), with respect to female counterparts. Interestingly, the sex difference in COX-2 protein expression was coupled to a significant higher activation status of NF-κB in male cells, visualized by elevated phospho-NF-κB p65 levels (Fig. 3E and F). Moreover, activation of p38 MAPK was more pronounced in activated phospho-NF-κB in male cells, visualized by elevated phospho-NF-κB p65 levels (Fig. 3E and F). Moreover, activation of p38 MAPK was more pronounced in male cells from carrageenan-treated male rats as compared to cells from female animals (Fig. 3G and H), although the differences did not reach statistical significance.

Inhibition of 5-LO product formation abolishes the sex differences in PG formation in the thoracic exudates of carrageenan-treated rats. Based on the fact that LTB4 levels in the early phase (2 hrs) of the inflammatory response were lower in male versus female rats, we hypothesized that shunting phenomena from LTs to PGs might contribute to the opposite PG levels. Thus, we attempted to block 5-LO product formation by a pharmacological inhibitor (i.e. MK886) in order to investigate if the sex difference in PG biosynthesis at 4 hrs could be abolished. Rats were pre-treated with MK886 (1.5 mg/kg, i.p.) 30 min prior pleurisy induction, sacrificed after 4 hrs, and PGE2 levels were analyzed in the exudates. MK886 administration significantly (p < 0.05) abolished the sex bias in PGE2 (Fig. 4A) as well as in 6-keto-PGF1α production (Fig. 4B). Furthermore, pre-treatment of rats with 1.5 mg/kg MK886 efficiently suppressed LTB4 levels by 75 and 74% in exudates of male and female animals, respectively (Fig. 4C). In agreement with this finding, MK886 treatment reduced the number of infiltrated cells into the thoracic cavity without significant differences between male and female animals (Fig. 4D).

The capacity for PGE2 production in human neutrophils is sex-dependent. Neutrophils are immune cells involved in the inflammatory reaction, recruited by chemoattractants (i.e. LTB4) that are produced by resident cells17,18. To investigate if the sex difference in PGE2 biosynthesis exists also in humans, neutrophils from blood of female and male donors were freshly isolated and immediately stimulated for PGE2 production by lipopolysaccharide (LPS), a receptor-coupled stimulus, or by A23187 that causes cell activation by substantial mobilization of intracellular Ca2+ in a receptor-independent manner. Formation of PGE2 upon stimulation with LPS (1 μg/ml), which is strongly upregulated over the time course of 20 hrs, was much more pronounced for neutrophils derived from male versus cells from female donors (Fig. 5A). Also in response to A23187 (0.5 μM), a trend of higher PGE2 biosynthesis from male cells was evident starting at 30 min post stimulation and reaching significance (p < 0.05) at 4 hrs (Fig. 5B). Increased availability of AA as substrate in male cells may account for higher PGE2 synthesis. However, in agreement with our previous findings4, we observed no differences in the
release of AA upon neutrophil stimulation with 0.5 µM A23187 between cells from male and female donors (Fig. 5C).

**Cyclooxygenase-2 expression is higher in male human neutrophils.** Next, we addressed if COX expression in human neutrophils is affected by the sex. Analysis of COX protein levels (normalized to β-actin) in freshly isolated neutrophils from male and female donors by Western blot revealed no sex difference in the expression of COX-1 (Fig. 6A,B), while a significant (p < 0.05) higher expression of COX-2 in male cells was observed (Fig. 6A,C). This suggests that elevated PGE2 formation in male neutrophils might be connected to higher amounts of COX-2.

The sex difference in PGE2 production in human neutrophils is abolished by inhibition of 5-LO product synthesis. Human neutrophils from females produce significantly higher levels of LTs from AA than cells from males4, implying that in male neutrophils more AA might be available as substrate for COX to produce PGs. In line with the data obtained from the carrageenan-induced pleurisy in rats, pre-treatment of neutrophils with MK886 (30 nM, 15 min) to block 5-LO product formation did not affect PGE2 formation in A23187-stimulated male neutrophils, while it significantly (p < 0.01) increased PGE2 synthesis in female cells. Thus, suppression of 5-LO product biosynthesis abolishes the sex difference in PGE2 formation also in human neutrophils (Fig. 6D).

**Discussion**

A sex bias in inflammatory and immune diseases is clearly evident1, but the underlying biochemical or molecular mechanisms remain unclear. Here, we report about sex differences in PG production in human neutrophils in vitro and in rats and mice during acute inflammation in vivo. Our data suggest that the higher PG levels in neutrophils of males is due to AA substrate shunting phenomena because of lower LT production, but is also governed by higher amounts of COX-2 protein, as compared to neutrophils from females. The two major branches of biosynthetic pathways of pro-inflammatory lipid mediators produced from AA are the COX-mediated cascade...
leading to PGs and the 5-LO-mediated cascade yielding LTs. Sex differences in the regulation of 5-LO in human leukocytes with consequences for LT formation were reported by us before and we recently confirmed sex-biased LT biosynthesis in murine zymosan-induced peritonitis in vivo, a well-recognized model of acute inflammation. Thus, we observed more pronounced LT formation and related inflammatory reactions (i.e., neutrophil infiltration and vascular permeability) in female mice as compared to male animals. The cells responsible for the observed sex difference were resident PMs, occupying the peritoneal cavity under normal physiological conditions, which are the first cells to respond to inflammatory stimuli. In analogy to human neutrophils, the subcellular localization of 5-LO in female and male murine PMs differs, and the limited amount of mobile 5-LO in male PMs is seemingly responsible for lower LT biosynthesis.

The results of the present report confirm higher LT formation in females but also reveal a sex-biased production of PG, in particular by neutrophils. However, while LT formation was higher in female cells and animals, PG formation was higher in males. Our data are consistent with several observations made in vitro and in vivo studies observing a sex dimorphism in PG production, with higher levels in males or in ovariectomized female subjects.

A time-course study of PGE₂ biosynthesis in zymosan-induced peritonitis revealed higher PGE₂ levels in exudates from male mice compared to female animals, 4 hrs or later after zymosan injection. Although PGE₂ levels peaked at 2 hrs, the sex difference was significant only at 4 and 8 hrs, but not earlier. It should be noted that 4 hrs after peritonitis induction the cellular population of the peritoneal cavity is mostly composed of neutrophils as shown previously by us and others. Thus, we suggest that the sex difference is attributable to neutrophils, rather than to PMs that seem to equally generate PGs in the early phase (0–2 hrs) independently of

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**Figure 4.** Inhibition of 5-LO product formation by MK886 abolishes the sex differences in PG formation in the thoracic exudates of carrageenan-treated rats. Rats were pre-treated with MK886 (1.5 mg/kg, i.p.) or vehicle (DMSO 2%, i.p.) 30 min prior to carrageenan intrathoracic injection and sacrificed after 4 hrs. (A) PGE₂, (B) 6-keto-PGF₁α, and (C) LTB₄ levels in the exudates. (D) Number of infiltrated cells. Values represent means ± S.E.M; n = 6 rats; *p < 0.05, male vs female; °p < 0.05, female control rats vs female rats treated with MK886; two-tailed Student’s t test.
the sex. Surprisingly, despite the higher PGE₂ production in males, the number of infiltrating cells were lower at 4 and 8 hrs as compared to female animals. This might be due to reduced levels of LTB₄ in males that acts as potent chemotactic factor for neutrophils, and based on its greater abundance in female mice may recruit neutrophils more efficiently. To confirm the general validity of sex differences in PG biosynthesis, we chose a different animal model of acute inflammation (i.e. pleurisy) with a different stimulus (i.e. carrageenan) and different species (i.e. rat) to support sex-dependent production of eicosanoids as general and model-independent phenomenon. Carrageenan-induced pleurisy in rats represents one of the most commonly used models to investigate eicosanoid biosynthesis and signaling during acute inflammation. In agreement with the results from the zymosan-induced peritonitis, significant higher LTB₄ levels in thoracic exudates of female rats were evident 2 hrs after carrageenan-injection, while superior PGE₂ levels were found in thoracic exudates of male animals at later time points (4–8 hrs). These data highlight the converse transformation of AA to PGs (higher in males) and LTs (higher in females), and suggest that sex differences in PGE₂ production in these experimental models are most likely related to neutrophils. During acute inflammation, the mobilization and recruitment of blood leukocytes into the tissue are mediated by several factors. Among them, LTB⁴ through the BLT1 receptor might be the major chemoattractant molecule responsible for neutrophil infiltration. Since the LTB₄ levels were higher in the thoracic exudates of female rats, we hypothesized that this would cause an increased neutrophil infiltration in the

Figure 5. Capacity of human neutrophils for PGE₂ production is sex-dependent. PGE₂ production in human neutrophils from male and female donors after LPS or A23187 stimulation. Neutrophils from male and female donors were incubated with (A) LPS (1 µg/ml) for 0, 0.5, 3 and 20 hrs, or (B) with 0.5 µM A23187 for 0, 5, 15, 30, 60, 120, and 240 min. The reactions were stopped on ice, and PGE₂ levels were measured by ELISA. (C) After selected time points (0, 5, 15, 30, 60, 120 min) upon stimulation with A23187 (0.5 µM), free ³H-AA content in the supernatant of ³H-AA-pre-labelled neutrophils was evaluated by scintillation counting. Values represents means ± S.E.M of n = 3–6 experiments each in duplicate. *p < 0.05, **p < 0.01, male vs female cells; two-way ANOVA plus Bonferroni.

Figure 6. COX-2 expression is higher in neutrophils from male versus female donors; effects of 5-LO pathway inhibition on PGE₂ production. (A–C) Western blot analysis of total cell lysates of neutrophils from male and female donors. (A) COX-1 and -2 protein expression. Densitometric analysis of (B) COX-1 and (C) COX-2 protein normalized to β-actin expression. Western blots are representatives of 11 independent experiments. Values represents means ± S.E.M of densitometric analyses of COX-1/-2 protein bands, normalized to β-actin. *p < 0.05, male vs female cells. (D) COX-2, COX-1 and -2 protein expression. Densitometric analysis of (B) COX-1 and (C) COX-2 protein normalized to β-actin expression. Western blots are representatives of 11 independent experiments. Values represents means ± S.E.M of densitometric analyses of COX-1/-2 protein bands, normalized to β-actin. *p < 0.05, male vs female cells.
cavity of female animals. Notably, however, the number of infiltrated neutrophils did not differ between males and females at 4 hrs after carrageenan, and MK886 at 1.5 mg/kg that strongly repressed LTB4 formation and caused significant inhibition of cell infiltration in both sexes.

For PG production, AA is converted in two steps by the action of COX enzymes that catalyze the transformation of AA into the endoperoxide PGG2, containing a 15-hydroperoxy moiety. Reduction of the hydroperoxy group to a hydroxyl function then leads to PGH2 that is further converted by specific PG synthases to the respective bioactive PGs, including PGE2, PGF2α, PGD2, PGI2 and thromboxane(s), depending on the tissue-selective expression of the PG synthases28. Since besides PGE2 also the 6-keto-PGF1α, (a stable metabolite and marker of instable PGI2) was consistently higher in males, these data support that upstream COX enzymes might be affected by the sex rather than mPGES-1, the terminal enzyme in pro-inflammatory PG biosynthesis29. Nevertheless, sex differences related to mPGES-1 were found in spontaneously hypertensive rats, where female rats had enhanced mPGES-1 protein expression in the renal inner medulla and greater COX-2 expression in the outer medulla versus males32. In our hands, COX-2 protein, but not COX-1, was more abundant in cells from thoracic exudates of male rats and in human neutrophils from males, and such dominance of COX-2 in males is in agreement with observations by others. Thus, lower COX-2 expression and activity has been noted in the macula densa of female rats compared to males, contributing to the major protection of female to the blood pressure increment and renal damage30. Moreover, the inferior susceptibility to traumatic brain injury of male versus female rats has been related to a robust higher expression of COX-2 in the brain of male rats31. In addition, long-term testosterone treatment augmented COX-2 levels in male rat brain blood vessels, whereas treatment of male rats with 17β-estradiol significantly impaired cerebrovascular COX-2 levels after an inflammatory stimulus21.

Expression of COX-2 is regulated by several transcription factors including NF-κB9, whose activation in thoracic cells from carrageenan-treated rats was sex dependent in our present study. In fact, phospho-NF-κB p65 levels in male rats were significantly higher with respect to female animals. Our data are in line with previously reported sex differences in cerebrovascular pathophysiology that were due to activation of the NF-κB-mediated COX-2 pathway by the androgen 5α-DHT that results in a state of vascular inflammation32.

Our data reveal that activation of human neutrophils by LPS or by A23187 leads to significantly higher PGE2 levels in cells from males versus female counterparts. We showed before that stimulation of human neutrophils with LPS plus FMLP or with A23187 caused higher LTB4 production in female cells as compared to males4. Although A23187 preferentially activates the 5-LO pathway via receptor-independent, massive elevation of intracellular Ca2+, we believe that it represents a suitable stimulus to investigate the sex-regulation of eicosanoid biosynthesis, considering the fact that other stimuli (e.g., LPS) act through receptors that are strongly modulated by sex as well33, 34. We hypothesized that the blockade of 5-LO product formation by using MK886 would redirect AA conversion by COX enzymes. In fact, the sex difference in PGE2 formation in human neutrophils in vitro as well as in carrageenan-treated rats in vivo was abolished by interruption of 5-LO product formation using MK886 that significantly increased PGE2 in females without any alterations in males. In parallel, MK886 strongly reduced LTB4 levels in both test systems. On the other hand, blockade of PGE2 production may have the converse phenotype, however, the COX inhibitor indomethacin did not increase LTB4 levels in male rats during carrageenan-induced pleurisy35.

Taken together, we showed that male mice and rats produce higher levels of PGs in various acute models of inflammation in vivo under conditions where LT production is elevated in female animals at the sites of injury. Neutrophils are abundant innate immune cells in the human body taking part of the first line of defense against host injury, and are considered to be a major source of LTs33. Our findings imply that neutrophils from male subjects have higher capacities to produce PG seemingly due to elevated COX-2 expression and AA substrate availability. These sex differences are of relevance for PG-related functions and pathophysiology, supported also by experimental observations reported by others, and might help to explain, at least in part, the sex dimorphism in innate immune disorders such as sepsis8 and post-surgery infections as well as gout3.

Material and Methods

Materials. Enzyme immunoassay (EIA) kits were from Cayman Chemical Company (BertinPharma, Montigny Le Bretonneux, France) or from Biotrend (Cologne, Germany). 3H-labelled PGE2 and 3H-labelled AA were from PerkinElmer Life Sciences (Milan, Italy and Germany). Unless otherwise stated, all other reagents and compounds were obtained from Sigma-Aldrich (Milan, Italy).

Animals. The animal studies are reported in accordance with the ARRIVE guidelines for reporting animal research36. Age-matched male and female CD-1 mice (8–9 weeks old, 26–40 g Charles River, Calco, Italy) and Wistar male and female rats (200–300 g, Harlan, Milan, Italy) were housed in a controlled environment (21 ± 2°C) and provided with standard rodent chow and water. All animals were allowed to acclimate for four days prior to experiments and were subjected to 12 h light–12 h dark schedule. Experiments were conducted during the light phase. The experimental protocols were approved by the Animal Care Committee of the University of Naples Federico II, in compliance with Italian regulations on protection of animals used for experimental and other scientific purpose (Ministerial Decree 26/2014) as well as with the European Economic Community regulations (Official Journal of E.C. L358/1 12/18/1986).

Induction of peritonitis in mice. Peritonitis was induced in mice as previously described4. In brief, a solution of 2 mg/ml of zymosan A (boiled and washed) was injected intraperitoneally (i.p., 0.5 ml) and at selected time points (0–2–4–8 hrs), mice were sacrificed in a saturated atmosphere with CO2. Peritoneal exudates were collected by washing the cavity with 2 ml of phosphate-buffered saline (PBS) and then centrifuged at 20,000 × g for 20 min at 4°C and supernatants frozen at −80°C for measurements of eicosanoids. PGE2 was evaluated by...
radioimmunoassay (RIA), 6-keto-PGF_1α, and LTC_4 by EIA (Cayman chemicals BertinPharma, Montigny Le Bretonneux, France), according to manufacturer’s protocol. Results are expressed as ng/ml.

**Induction of pleurisy in rats.** Rats were anesthetized with 4% enflurane mixed with 0.51/min O_2, 0.5 l/min N_2O and submitted to a skin incision at the level of the left sixth intercostal space. The underlying muscle was dissected and 1% (w/v) λ-carrageenan type IV (0.2 ml) was injected into the thoracic cavity. The skin incision was closed with a suture and the animals were allowed to recover. At selected time points (0–2–4–8 hrs) after carrageenan injection, animals were sacrificed by CO_2 inhalation. Thoracic exudate was collected by lavage of the cavity with 2 ml of saline solution, after centrifugation (800 × g for 10 min), the leukocyte number was determined by light microscopy using a Bürker chamber. The cells as well as the supernatants were frozen at −80°C for WB analysis and eicosanoid measurement, respectively. In one set of experiments, rats were pre-treated with 1.5 mg/kg MK886 (Cayman Chemical, Bertin Pharma, Montigny Le Bretonneux, France) or vehicle (2% DMSO in saline) 30 min prior to pleurisy induction. Animals were then sacrificed 4 hrs after carrageenan injection. The amount of PGE_2, LTB_4 and 6-keto-PGF_1α in the supernatant of centrifuged exudates was measured by RIA and by EIA, respectively. Results are expressed as the total amount of eicosanoid measured in the thoracic exudate of one rat (nanograms per rat).

**Isolation and stimulation of human neutrophils.** Leukocyte concentrates, prepared from freshly withdrawn peripheral blood of healthy adult human donors who had not taken any anti-inflammatory drugs for the last 10 days were obtained from the Institute of Transfusion Medicine at the University Hospital Jena, Germany. Informed consent was obtained from all subjects. The experimental protocol was approved by the local ethical committee at the University Hospital Jena. All methods were performed in accordance with the relevant guidelines and regulations. Neutrophils were isolated as previously described4, 37. In brief, neutrophils were obtained from leukocyte concentrates by a multi-step procedure: (1) dextran sedimentation; (2) centrifugation on Nycocrep (872 × g, 10 min); (3) hypotonic lysis of erythrocytes. Finally, cells were suspended in ice-cold PBS containing 0.1% glucose (PG buffer) and counted by Vi-CELL™ XR. For PG production, 5 × 10^6 neutrophils, from female and male donors, were resuspended in PG buffer containing 1 mM CaCl_2 (PGc buffer) and stimulated with 1 µg/ml LPS for 0, 0.5, 3 and 20 hrs and with 0.5 µM A23187 for 0, 5, 15, 30, 60, 120, or 240 min. In one set of experiments, freshly isolated neutrophils from male and female donors were pre-treated with 30 nM MK886 or DMSO as vehicle (15 min., 37°C), and then stimulated with 0.5 µM A23187 for 4 hrs. The reaction was stopped on ice and samples were centrifuged (12,000 × g, 5 min, 4°C), PGE_2 levels in the supernatants were measured by ELISA kit (Biotrend, Cologne, Germany).

**Total protein extraction and Western blot analysis.** Protein analysis of COX-1/2, phospho-NF-κB p65, phospho-p38 MAPK and β-actin by Western blot was performed in whole cell lysates. Cells in the thoracic exudates were collected 4 hrs after carrageenan administration and then immediately lysed in a buffer for protein extraction, mixed with sodium dodecyl sulphate (SDS) loading gel buffer and analysed by Western blot according to ref. 6 on a 10% SDS–polyacrylamide gel. The membranes were incubated overnight with rabbit monoclonal antibody anti-COX-2 (1:500, BD Transduction Laboratories, Aurogene, Rome, Italy), mouse monoclonal antibody anti-COX-1 (1:1000, Cell Signaling, Aurogene, Rome, Italy), rabbit monoclonal antibody anti-phospho-NF-κB p65 (Ser536) (1:1000, Cell Signalling Technology, Inc., Germany), rabbit polyclonal antibody anti-phospho-p38 MAPK (Thr180/Thr182) (1:1000, Santa Cruz Biotechnology, Aurogene, Rome, Italy) and β-actin (1:2000, Santa Cruz Biotechnology, Aurogene, Rome, Italy). Membranes were washed six times with 0.1% PBS-Tween and were incubated for 1.5 hrs at room temperature with horseradish peroxidase-conjugated anti-rabbit and anti-mouse secondary antibodies diluted 1:10,000 in 0.1% PBS-Tween containing 0.1% non-fat dry milk. Protein bands were detected by an enhanced chemiluminescence system (Amersham Pharmacia, Aurogene, Rome, Italy). Densitometric analysis was performed by using the Fluor S quantitative imaging system.

**Arachidonic acid release in neutrophils from male and female donors was evaluated as previously reported.** Briefly, 3 × 10^6 cells/ml PBS plus 1 mM EDTA were sonicated (5 × 5 sec, 4°C) and centrifuged (12,000 × g, 15 min, 4°C). Aliquots of the resulting supernatants were mixed 1:1 with ice-cold 2× SDS/PAGE sample loading buffer (SDS-b) and heated for 5 min at 95°C. Samples were loaded (15 µl) and electrophoresed on a 10% SDS–polyacrylamide gel, and transferred to nitrocellulose membranes. After the membranes were incubated with primary antibodies (rabbit anti-COX-1, 1:1000, Cell signalling; rabbit anti-COX-2, 1:1000, Santa Cruz; mouse anti-β-actin, 1:1000; Santa Cruz) they were subsequently detected using IRDye 800CW-labeled anti-rabbit and/or anti-mouse antibodies (1:10,000 each). The immunoreactive bands were visualized using an Odyssey infrared imager (Li-Cor Biosciences, Lincoln, NE).

**Arachidonic acid release in human neutrophils.** Arachidonic acid release in neutrophils from male and female donors was evaluated as reported previously. Briefly, freshly isolated neutrophils (2 × 10^6 cells/ml) were re-suspended in RPMI 1640 without additives, 0.5 µCi Ci-3H-labelled AA/ml were added to the cell suspension and incubated for 2 hrs at 37°C. Cells were washed twice (320 × g, 10 min, 4°C) with incubation buffer (PBS, containing 0.1% glucose and 2 mg/ml fatty acid free BSA). Cells were adjusted to a cell number of 1 × 10^7/0.5 ml and 1 mM CaCl_2, was added to the incubation buffer. Cells were pre-treated with Ca^{2+}-ionophore A23187 (0.5 µM) for 5 to 120 min, as indicated, at 37°C. The reaction was stopped on ice and samples were centrifuged (500 × g, 10 min, 4°C). Aliquots (300 µl) of the supernatants were combined with 2 ml Rotiszint® eco plus and assayed for radioactivity by scintillation counting (Micro Beta Trilux, Perkin Elmer, Waltham, MA).

**Statistical analysis.** Data are expressed as mean ± standard error of the mean (S.E.M.) of n observations, n represents the number of animals, or the number of experiments (in vitro) performed with cells from different donors in duplicates. Statistical evaluation was performed by two-tailed Student t-test for single
comparisons or by two-way ANOVA using GraphPad InStat (Graphpad Software Inc., San Diego, CA) followed by a Bonferroni post-hoc test for multiple comparisons, respectively. P-values < 0.05 were considered as significant.

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Author Contributions
S.P., A.R., O.W. and L.S. designed research; S.P., A.R., V.K., F.D., F.T., R.B., C.W. and S.R. performed research; S.P., A.R., O.W. and L.S. analyzed data; and S.P., A.R., O.W. and L.S. wrote the paper.
Additional Information

Competing Interests: The authors declare that they have no competing interests.

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