Hyperforin, an anti-inflammatory constituent from St. John’s wort, inhibits microsomal prostaglandin E₂ synthase-1 and suppresses prostaglandin E₂ formation in vivo

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The acylphloroglucinol hyperforin (Hyp) from St. John’s wort possesses anti-inflammatory and anti-carcinogenic properties which were ascribed among others to the inhibition of 5-lipoxygenase. Here, we investigated whether Hyp also interferes with prostanoid generation in biological systems, particularly with key enzymes participating in prostaglandin (PG)E₂ biosynthesis, i.e., cyclooxygenases (COX)-1/2 and microsomal PGE₂ synthase (mPGES)-1 which play key roles in inflammation and tumorigenesis. Similar to the mPGES-1 inhibitors MK-886 and MD-52, Hyp significantly suppressed PGE₂ formation in whole blood assays starting at 0.03−1 μM, whereas the concomitant generation of COX-derived 12(S)-hydroxy-5-cis-8,10-trans-heptadecatrienoic acid, thromboxane B₂, and 6-keto PGF₁α was not significantly suppressed up to 30 μM. In cell-free assays, Hyp efficiently blocked the conversion of PGH₂ to PGE₂ mediated by mPGES-1 (IC₅₀ = 1 μM), and isolated COX enzymes were not (COX-2) or hardly (COX-1) suppressed. Intrapitoneal (i.p.) administration of Hyp (4 mg kg⁻¹) to rats impaired exudate volume and leukocyte numbers in carrageenan-induced pleurisy associated with reduced PGE₂ levels, and Hyp (given i.p.) inhibited carrageenan-induced mouse paw edema formation (ED₅₀ = 1 mg kg⁻¹) being superior over indomethacin (ED₅₀ = 5 mg kg⁻¹). We conclude that the suppression of PGE₂ biosynthesis in vitro and in vivo by acting on mPGES-1 critically contributes to the anti-inflammatory efficiency of Hyp.

Keywords: hyperforin, St. John’s wort, prostaglandin E₂, cyclooxygenase, inflammation

INTRODUCTION

Hyperforin (Hyp), a polyprenylated acylphloroglucinol (Figure 1A), is assumed to be one of the main active components of St. John’s wort, which is frequently used for the treatment of mild to moderate depressions (Muller, 2003). Besides its antidepressant activity, Hyp also exerts potent anti-inflammatory and anti-tumoral effects (Medina et al., 2006). Hyp was shown to reduce tumor cell growth (Schempp et al., 2002; Hostanska et al., 2003), cancer invasion, metastasis (Donà et al., 2004), angiogenesis (Martínez-Poveda et al., 2005), and lymphatic capillary growth (Rothley et al., 2009). On the cellular level, Hyp decreases proliferation rates and causes apoptosis of leukemic cells (Schempp et al., 2002; Quiney et al., 2006) and various cancer cell lines (Schempp et al., 2002; Hostanska et al., 2003; Donà et al., 2004), and also inhibits proliferation of non-transformed T lymphocytes (Schempp et al., 2000). In regard of its anti-inflammatory potential, Hyp blocks several pro-inflammatory functions of leukocytes in vitro such as chemotaxis and chemoinvasion (Dell’Aica et al., 2007; Lorusso et al., 2009), suppresses receptor-mediated Ca²⁺-mobilization and eicosanoid release in leukocytes (Albert et al., 2002; Feisst and Werz, 2004), and down-regulates effector functions of activated T lymphocytes (Cabrelle et al., 2008). In vivo, Hyp reduced croton-oil-induced ear edema in mice after topical application (Sosa et al., 2007), impaired acute neutrophil recruitment and enhanced resolution in a pulmonary bleomycin-induced inflammation model reducing consequent fibrosis (Dell’Aica et al., 2007), and suppressed carrageenan-induced rat pleurisy when given intraperitoneally (i.p.; Feisst et al., 2009). Also for humans, topical treatment of atopic dermatitis with a Hypericum cream, standardized to 1.5% Hyp, revealed significant therapeutic efficacy (Schempp et al., 2003). However, despite these well-recognized anti-carcinogenic and anti-inflammatory effects, the underlying molecular mechanisms and targets of Hyp are incompletely understood.

The lipid mediator prostaglandin (PG)E₂ is a key player in inflammation, pain, fever, and cancer but is also known to regulate physiological functions in the gastrointestinal tract, in the kidney, and in the immune and nervous system (Sugimoto and Narumiya, 2007). PGE₂ is formed from arachidonic acid (AA) by cyclooxygenase (COX)-catalyzed synthesis of PGH₂ and further transformation by PGE₂ synthases (Samuelsson et al., 2007). The microsomal PGE₂ synthase (mPGES)-1 is induced by pro-inflammatory stimuli and is responsible for excessive PGE₂ generation connected to pathologies (Jakobsson et al., 1999). Co-expression studies indicate a preferred functional coupling between mPGES-1 and COX-2 (Murakami et al., 2001).
and Kudo, 2006), and data from mPGES-1 knockout mice and animal studies with selective mPGES-1 inhibitors suggest key roles of mPGES-1 in inflammation, pain, fever, atherosclerosis, and tumorigenesis (Samuelsson et al., 2007; Friesen and Mancini, 2008; Koeberle and Werz, 2009).

We have previously shown that Hyp inhibits leukotriene biosynthesis in neutrophils (IC_{50} = 1–2 μM) by interference with the C2-like domain of 5-lipoxygenase (5-LO), and we observed an inhibitory effect of Hyp also on COX-1 activity in human platelets (IC_{50} = 3 μM; Albert et al., 2002; Feist et al., 2009). However, COX-2 activity (measured as 6-keto PGF_{1α} synthesis) in monocytes Mono Mac 6 cells was not affected up to 30 μM Hyp. This is surprising because Hammer et al. (2007) recently demonstrated that Hyp inhibits the release of PGE, from lipopolysaccharide (LPS)-stimulated murine (RAW264.7) macrophages, a process coupled to COX-2 as well. Here, we identified mPGES-1 as functional target of Hyp and suggest that interference with mPGES-1 suppresses PGE biosynthesis in vivo and contributes to the anti-inflammatory effectiveness in related animal models.

**MATERIALS AND METHODS**

**REAGENTS**

Hyperforin dicyclohexylammonium salt and the Hypericum perforatum extract were obtained from Dr. Willmar Schwabe GmbH & Co (Karlsruhe, Germany), respectively. The compounds and the extract were dissolved in dimethyl sulfoxide (DMSO) and kept in the dark at −20°C, and freezing–thawing cycles were kept to a minimum. For animal studies, Hyp was dissolved in DMSO and diluted with saline achieving a final DMSO concentration of 2–4%. The thromboxane (TX) synthase inhibitor CV4151 (Kato et al., 1985) and the mPGES-1 inhibitor 2-(2-chlorophenyl)-1-phenanthro[9,10-d]-imidazol (MD-52; Côté et al., 2007) were kindly provided by Dr. S. Laufer (University of Tübingen, Germany) and Dr. M. Schubert-Zsilavecz (University of Frankfurt, Germany), respectively. The antibody against human mPGES-1 was from Cayman Chemical (Ann Arbor, MI, USA), antibodies against COX-2 and β-actin were obtained from Sigma-Aldrich (Deisenhofen, Germany) and the antibody against cleaved poly(ADP-ribose)polymerase (PARP) was purchased from Cell Signaling Technology (Danvers, MA, USA). Materials used: DMEM high glucose (4.5 g l^{-1} medium, penicillin, streptomycin, trypsin–EDTA solution, sodium pyruvate, PAA (Coelbe, Germany); PGH_{1}, Larodan (Malmö, Sweden); 11β-PGE_{1}, MK-886, PGB_{1}, human recombinant COX-2, ovine isolated COX-1, Cayman Chemical (Ann Arbor, MI, USA). A-Carrageenan type IV isolated from Gigartina aciculare and G. pistillata was purchased from Sigma-Aldrich (Milan, Italy). H-PGE_{1}, was from PerkinElmer Life Sciences (Milan, Italy) and PGE_{1}, antibody from Sigma-Aldrich (Milan, Italy). All other chemicals were obtained from Sigma-Aldrich (Deisenhofen, Germany) unless stated otherwise.

**CELLS AND CELL VIABILITY ASSAY**

Human lung carcinoma A549 cells were cultured in DMEM high glucose (4.5 g l^{-1}) medium supplemented with heat-inactivated fetal calf serum (FCS, 10%, v/v), penicillin (100 U ml^{-1}), and streptomycin (100 μg ml^{-1}) at 37°C in a 5% CO_{2} incubator. After 3 days, confluent cells were detached using 1× trypsin–EDTA solution and reseeded at 2 × 10^{4} cells in 20 ml medium in 175 cm^{2} flasks. Cell viability was assessed using the colorimetric thiazolyl blue tetrazolium bromide dye reduction assay (MTT assay). A549 cells (4 × 10^{4} cells per 100 μl medium) were plated into a 96-well microplate and incubated at 37°C and 5% CO_{2} for 16 h. Then, Hyp (30 μM) was added, and the samples were incubated for another 5 h. Thiazolyl blue tetrazolium bromide (20 μl, 5 mg ml^{-1}) was added and the incubations were continued for 4 h. The formazan product was solubilized with sodium dodecylsulfate (SDS, 10%, m v^{-1} in 20 mM HCl), and the absorbance was measured at 595 nm relative to the absorbance of vehicle (DMSO)-treated control cells using a multiwell scanning spectrophotometer (Victor plate reader, PerkinElmer, Rodgau-Juegesheim, Germany).

**ANIMALS**

Male adult CD1 mice (25–35 g, Harlan, Milan, Italy) and Wistar Han rats (190–200 g, Harlan, Milan, Italy) were housed in a controlled environment and provided with standard rodent chow and water. Animal care complied with Italian regulations on protection of animals used for experimental and other scientific purpose (Ministerial Decree 116192) as well as with the European Economic Community regulations (Official Journal of E.C. L 358/1 12/18/1986).

**DETERMINATION OF PGE_{2}, 6-KETO PGF_{1α}, AND TXB_{2} FORMATION IN LPS-STIMULATED HUMAN WHOLE BLOOD**

Peripheral blood from healthy adult volunteers, who had not received any medication for at least 2 weeks under informed consent, was obtained by venepuncture and collected in syringes containing heparin (20 U ml^{-1}). For determination of PGE_{2}, and 6-keto PGF_{1α} aliquots of whole blood (0.8 ml) were mixed with the TX synthase inhibitor CV4151 (1 μM) and with aspirin (50 μM) to establish experimental conditions where prostanoids are essentially produced via the COX-2 pathway (Koeberle et al., 2008). For the determination of TXB, whole blood aliquots were not pretreated with aspirin and CV4151. A total volume of 1 ml was adjusted with sample buffer (10 mM potassium phosphate buffer pH 7.4, 3 mM KC1, 140 mM NaCl, and 6 mM d-glucose). After pre-incubation with the indicated compounds for 5 min at room temperature, the samples were stimulated with LPS (10 μg ml^{-1}) for 5 h at 37°C. Prostanoid formation was stopped on ice, the samples were centrifuged (2300 × g, 10 min, 4°C), and 6-keto PGF_{1α} and TXB, were quantified in the supernatant using a 6-keto PGF_{1α} or TXB, high sensitivity EIA kit (Assay Designs, Ann Arbor, MI, USA) according to the manufacturer’s protocol. PGE_{2} was determined after solid phase extraction and HPLC separation using a PGE_{2} high sensitivity EIA kit (Assay Designs, Ann Arbor, MI, USA) as described (Koeberle et al., 2008). The mPGES-1 inhibitors MK-886 (30 μM) and MD52 (6 μM), the COX-1/2 inhibitor indomethacin (50 μM) and the COX-2-selective inhibitor celecoxib (20 μM) were used as control. Care should be taken when interpreting the controls in respect to COX-1/2 isoenzyme specificity which was not further evaluated for our experimental settings.

**DETERMINATION OF PROSTANOID FORMATION FROM EXOGENOUS AA IN HUMAN WHOLE BLOOD**

Heparinized human blood, supplemented with penicillin (100 U ml^{-1}) and streptomycin (100 μg ml^{-1}), was treated with 10 μg ml^{-1} LPS for 16 h at 37°C and 5% CO_{2}. Then, CV4151 (1 μM)
was added, and after pre-incubation with the indicated compounds for 10 min at 37°C, prostanoid formation was initiated by addition of 20 μM AA. In contrast to the LPS-stimulated whole blood assay described above, whole blood aliquots were not pretreated with aspirin (inhibition of COX-1 product formation) because both COX-1 and -2 are described to contribute to mPGES-1-dependent PGE_2 synthesis at high AA concentrations (Murakami et al., 2000). PGE_2, or 6-keto PGF_1α, formation within 10 min was determined as described for LPS-stimulated whole blood. Calculated prostanoid levels were corrected by the amount of PGE_2 formed during pre-incubation with LPS.

For determination of the COX product 12(S)-hydroxy-5-cis,8,10-trans-heptadecatrienoic acid (12-HHT), freshly drawn human blood (2 ml) was pre-incubated with the indicated compounds at 37°C for 10 min, and formation of 12-HHT was initiated by addition of 30 μM Ca^{2+}-ionophore A23187 plus 100 μM AA. After 10 min at 37°C, 12-HHT was extracted from plasma and analyzed by HPLC using PGB_1 as internal standard as described (Capdevila et al., 1995).

**ACTIVITY ASSAYS OF ISOLATED COX-1 AND COX-2**

Inhibition of the activities of isolated ovine COX-1 and human COX-2 was performed as described (Koeberle et al., 2008). Briefly, purified COX-1 (ovine, 50 units) or COX-2 (human recombinant, 20 units) were diluted in 1 ml reaction mixture containing 100 mM Tris buffer pH 8, 5 mM glutathione, 5 μM hemoglobin, and 100 μM EDTA at 4°C and pre-incubated with the test compounds for 5 min. Samples were pre-warmed for 60 s at 37°C, and AA (5 μM for COX-1, 2 μM for COX-2) was added to start the reaction. After 5 min at 37°C, the COX product 12-HHT was extracted and then analyzed by HPLC as described (Albert et al., 2002).

**PREPARATION OF CRUDE mPGES-1 IN MICROSOMES OF A549 CELLS AND DETERMINATION OF PGE_2 SYNTHASE ACTIVITY**

Preparation of A549 cells and determination of mPGES-1 activity was performed as described previously (Koeberle et al., 2008). In brief, cells were incubated for 16 h at 37°C and 5% CO₂, the medium was replaced, 1 ng ml⁻¹ interleukin (IL)-1β was added, and cells were incubated for another 48 h. After sonication, the homogenate was subjected to differential centrifugation at 10,000 x g for 10 min and 174,000 x g for 1 h at 4°C. The pellet (microsomal fraction) was resuspended in 1 ml homogenization buffer (0.1 M potassium phosphate buffer pH 7.4, 1 mM phenylmethylsulfonyl fluoride, 60 μg ml⁻¹ soybean trypsin inhibitor, 1 μg ml⁻¹ leupeptin, 2.5 mM glutathione, and 250 mM sucrose), and the total protein concentration was determined. Microsomal membranes were diluted in potassium phosphate buffer (0.1 M, pH 7.4) containing 2.5 mM glutathione to a final protein concentration of approx. 70 μg ml⁻¹. Test compounds or vehicle were added, and after 15 min at 4°C, the reaction (100 μl total volume) was initiated by addition of PGH_2 (20 μM, final concentration). After 1 min at 4°C, the reaction was terminated using stop solution (100 μl: 40 mM FeCl_3, 80 mM citric acid, and 10 μM of 11β-PGE_2 as internal standard). PGE_2 was separated by solid phase extraction and analyzed by HPLC as described (Koeberle et al., 2008).

**SDS-PAGE AND WESTERN BLOT**

Cells (4 x 10⁶ cells) were resuspended in 50 μl PBS buffer pH 7.2, mixed with the same volume of 2 x SDS/PAGE sample loading buffer (20 mM Tris–HCl, pH 8, 2 mM EDTA, 5% (v/v) SDS, and 10% (v/v) β-mercaptoethanol), and boiled for 5 min at 95°C. Aliquots (20 μl) corresponding to equivalents of 0.8 x 10⁶ cells were mixed with 4 μl glycerol/0.1% bromophenol blue (1:1, v:v), and proteins were separated by SDS-PAGE. After electroblotting to nitrocellulose membrane (GE Healthcare, Munich, Germany) and blocking with 5% BSA for 1 h at room temperature, membranes were washed and incubated with primary antibodies overnight at 4°C. The membranes were washed and incubated with a 1:1000 dilution of alkaline phosphatase-conjugated immunoglobulin G for 3 h at room temperature. After washing, proteins were visualized with nitro blue tetrazolium and 5-bromo-4-chloro-3-indolylphosphate.

**CARRAGEENAN-INDUCED PAW EDEMA IN MICE**

Mice were divided into groups (n = 10 for each group) and lightly anesthetized with enflurane 4% mixed with O_2 0.5 l min⁻¹, N_2 0.5 l min⁻¹. Each group of animals received subplantar administration of saline (0.05 ml) or of α-carrageenan type IV 1% (w/v), 0.05 ml) in saline. The paw was marked in order to immerse it always at the same extent in the measurement chamber. The volume was measured by using a hydrodynamisimeter, specially modified for small volumes (Ugo Basile, Milan, Italy), immediately before subplantar injection and 2, 4, and 6 h thereafter (Posadas et al., 2004). The assessment of paw volume was performed always in double blind and by the same operator. The increase in paw volume was calculated by subtracting the initial paw volume (basal) to the paw volume measured at each time point.

**CARRAGEENAN-INDUCED PLEURISY IN RATS**

Rats were anesthetized with enflurane 4% mixed with O_2 0.5 l min⁻¹, N_2 0.5 l min⁻¹ and submitted to a skin incision at the level of the left sixth intercostal space. The underlying muscle was dissected, and saline (0.2 ml) or λ-carrageenan type IV 1% (w/v), 0.2 ml) was injected into the pleural cavity. The skin incision was closed with a suture, and the animals were allowed to recover. At 4 h after the injection of carrageenan, the animals were killed by inhalation of CO₂. The chest was carefully opened, and the pleural cavity was rinsed with 2 ml saline solution containing heparin (5 U ml⁻¹). The exudate and washing solution were removed by aspiration, and the total volume was measured. Any exudate that was contaminated with blood was discarded. The amount of exudate was calculated by subtracting the volume injected (2 ml) from the total volume recovered. Leukocytes in the exudate were resuspended in phosphate buffered saline (PBS) and counted with an optical light microscope in a Burkner’s chamber after vital trypan blue staining.

The amount of PGE_2 in the supernatant of centrifuged exudate (800 x g for 10 min) was assayed by radioimmunoassay according to manufacturer’s protocol. The results are expressed as nanograms per rat and represent the mean ± SE of 10 rats.

**STATISTICS**

Data are expressed as mean ± SE. IC₅₀ values were calculated by nonlinear regression using SigmaPlot 9.0 (Systat Software Inc., San Jose, USA) one site binding competition. The program Graphpad
Instat (Graphpad Software Inc., San Diego, CA, USA) was used for statistical comparisons. Statistical evaluation of the data was performed by one-way or two-way ANOVAs for independent or correlated samples followed by Tukey HSD post hoc tests. A P value < 0.05 (*) was considered significant.

RESULTS

EFFECTS OF HYPERFORIN ON PROSTANOID FORMATION IN HUMAN WHOLE BLOOD

Hyperforin was recently shown to inhibit the production of PGE$_2$ in LPS-stimulated RAW264.7 mouse macrophages (Hammer et al., 2007). Here, we investigated the effects of Hyp on prostanoid formation in human whole blood pre-incubated with Hyp for 5 min and stimulated with LPS for 5 h. Hyp concentration-dependently suppressed PGE$_2$ formation with an apparent IC$_{50}$ of approx. 3 μM reaching significant inhibition at 1 μM (25 ± 3% inhibition; Figure 1B). Acute (5 h) cytotoxic effects of Hyp (up to 30 μM) can be excluded as determined by an A549 cell viability assay measuring mitochondrial dehydrogenase activity (see Materials and Methods, data not shown). The selective mPGES-1 inhibitor MD-52 (Côté et al., 2007) and the combined 5-LO-activating protein (FLAP), COX-1 and mPGES-1 inhibitor MK-886 (Claveau et al., 2003; Koeberle et al., 2009b) were used as reference (see also Koeberle et al., 2008). PGE$_2$ levels were neither completely suppressed by the mPGES-1 inhibitors nor by Hyp, even not at high Hyp concentrations (30 μM) or prolonged incubation times (24 h, data not shown). In contrast, the COX-1/2 inhibitor indomethacin (50 μM) and the COX-2 selective celecoxib (20 μM) effectively reduced PGE$_2$ (Figure 1B) almost to the level of the unstimulated control (PGE$_2$; 14.2 ± 8.4%). Celecoxib and/or indomethacin also inhibited the formation of 6-keto PGF$_{1\alpha}$ (the stable metabolite of PGI$_2$, Figure 1C) and TXB$_2$ (Figure 1D) as expected, whereas Hyp (up to 30 μM), MD-52, and MK-886 failed to suppress the formation of 6-keto PGF$_{1\alpha}$ (Figure 1C) and TXB$_2$ (Figure 1D) under the

![Figure 1](image)

**FIGURE 1** Effects of hyperforin on prostanoid formation in LPS-stimulated human whole blood. (A) Chemical structure of Hyp. (B,C) Heparinized human whole blood, treated with 1 μM TX synthase inhibitor CV4151 and 50 μM aspirin, was pre-incubated with the indicated agents or vehicle (DMSO) for 5 min at room temperature, and then, prostanoid formation was induced by addition of 10 μg ml$^{-1}$ LPS. After 5 h at 37°C, PGE$_2$ (B) was extracted from plasma by RP-18 solid phase extraction, separated by RP-HPLC, and quantified by ELISA as described. 6-Keto PGF$_{1\alpha}$ (C) and TXB$_2$ (D) were directly determined in the plasma by ELISA. The 100% values correspond to prostanoid levels in the range of 14–27 ng ml$^{-1}$ PGE$_2$, 3–5 ng ml$^{-1}$ 6-keto PGF$_{1\alpha}$, and 47–64 ng ml$^{-1}$ TXB$_2$ in the individual experiments, respectively. MK-886 (30 μM), MD-52 (6 μM), indomethacin (Indo, 50 μM), and celecoxib (Cele, 20 μM) were used as controls. Data are given as mean ± SE, $n$ = 3–4, *p < 0.05, **p < 0.001 vs. vehicle (0.1% DMSO) control, ANOVA + Tukey HSD post hoc tests.
same experimental conditions. Together, unlike COX inhibitors but as like mPGES-1 inhibitors, Hyp selectively suppresses PGE₂ formation but not the synthesis of other prostanoids such as 6-keto PGF₁α or TXB₂.

Hyperforin may suppress PGE₂ formation by interference with LPS-signaling, mPGES-1 expression or liberation of AA as substrate for COX enzymes. Thus, heparinized human whole blood was first incubated with LPS for 16 h to induce the expression of COX-2 and mPGES-1 (in contrast to constitutively expressed COX-1 and cPGES). Then, the blood was pre-incubated with Hyp for 10 min, and prostanooid formation was initiated by addition of 20 μM AA. After 10 min at 37°C, generation of PGE₂ and 6-keto PGF₁α was determined in plasma as described above. PGE₂ formation was potently inhibited by Hyp in a concentration-dependent manner with an IC₅₀ of 0.25 μM reaching significance at 30 nM (Figure 2A), whereas formation of 6-keto PGF₁α was not significantly affected up to 30 μM (Figure 2B), again indicating selective suppression of PGE₂ biosynthesis by Hyp. Note that Hyp (3–30 μM) was more efficient in suppressing PGE₂ synthesis after long-term pre-stimulation with LPS (induction of COX-2 and mPGES-1, Figure 2A) than during short term stimulation (Figure 2B) supporting that Hyp interferes with the inducible COX-2/mPGES-1 pathway.

In order to assess the effects of Hyp on the activity of COX-1 in whole blood (constitutively expressed in platelets), freshly withdrawn human blood (no incubation with LPS) was pre-incubated with Hyp for 10 min at 37°C, treated with Ca²⁺-ionophore and 100 μM AA for another 10 min, and the formation of 12-HHT (mainly derived from platelet COX-1) was determined. Hyp (up to 30 μM) failed to suppress 12-HHT formation under these assay conditions, whereas indomethacin and aspirin almost completely blocked it (Figure 2C). In conclusion, the data suggest that Hyp potently and selectively blocks PGE₂ formation in whole blood without markedly affecting COX-1/2.

**EFFECTS OF HYPERFORIN ON THE ACTIVITY OF ISOLATED COX-1 AND COX-2**

We determined the direct interference of Hyp with the catalytic activity of isolated COX enzymes. Purified ovine COX-1 and human recombinant COX-2 were pre-incubated with Hyp for 5 min prior to addition of AA (5 μM for COX-1, 2 μM for COX-2). Formation of 12-HHT, the major COX-1/2-derived product under these experimental conditions (Capdevila et al., 1995), was analyzed by RP-HPLC. Hyp concentration-dependently reduced 12-HHT formation by COX-1 with an IC₅₀ = 12 μM (Figure 3A), whereas COX-2 was not significantly inhibited up to 30 μM (Figure 3B). The reference compounds indomethacin (10 μM) and celecoxib (5 μM) potently inhibited 12-HHT formation, as expected.

**INHIBITION OF mPGES-1 ACTIVITY BY HYPERFORIN**

Because Hyp potently inhibited PGE₂ formation in whole blood from AA but failed to block COX-2, inhibition of PGE₂ synthesis downstream of COX-2 (i.e., interference with PGE₂ synthases) appeared reasonable. mPGES-1 is the major PGE₂ synthase under pathological conditions related to inflammation and cancer (Samuelsson et al., 2007), and its expression is strongly increased in blood upon LPS treatment, primarily contributing to PGE₂ formation in blood (Mosca et al., 2007). To assess the effects of Hyp on mPGES-1 activity, micro-

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**Figure 2** Effects of hyperforin on arachidonic acid-induced prostanoid formation in human whole blood. (A,B) Heparinized human whole blood was treated with 10 μg ml⁻¹ LPS for 16 h at 37°C and 5% CO₂, supplemented with TX synthase inhibitor CV4151 (1 μM), and pre-incubated with Hyp or vehicle (DMSO) for 10 min at 37°C. (A) Then, PGE₂ formation was initiated with 20 μM AA, and PGE₂ formed within 10 min was separated by RP-HPLC and quantified by ELISA. The 100% value corresponds to PGE₂ levels in the range of 18–31 ng ml⁻¹ in the individual experiments, respectively. (B) 6-keto PGF₁α was directly determined in the plasma by ELISA. The 100% value corresponds to 6-keto PGF₁α levels in the range of 4–7 ng ml⁻¹. Indomethacin (Indo, 50 μM) and celecoxib (Cele, 20 μM) were used as controls. (C) 12-HHT formation in whole blood. Heparinized whole blood was pre-incubated with Hyp or vehicle (DMSO) for 10 min, and AA (100 μM) and Ca²⁺-ionophore (30 μM) were added to induce 12-HHT product formation. After 10 min at 37°C, 12-HHT was extracted form the plasma by RP-18 solid phase extraction and analyzed by RP-HPLC as described. The 100% value corresponds to 1.5–2.4 μg ml⁻¹ 12-HHT. Indomethacin (Indo, 20 μM) and aspirin (ASA, 30 μM) were used as controls. Data are given as mean ± SE, n = 3–5. *p < 0.05, **p < 0.01 or ***p < 0.001 vs. vehicle (0.1% DMSO) control, ANOVA + Tukey HSD post hoc tests.
Cell line to study mPGES-1 functions), and also β-actin levels were slightly reduced at 10 μM (Figure 5). Since previous studies showed that Hyp causes apoptosis of various cancer cell lines (Schempp et al., 2002; Hostanska et al., 2003; Dona et al., 2004), we reasoned that reduced expression of COX-2 and mPGES-1 in A549 cells by Hyp may be related to induction of apoptotic cell death. In fact, the decrease of COX-2 and mPGES-1 protein correlated with the induction of apoptosis as determined by the cleavage of PARP (Figure 5), and the cell viability was strongly reduced after 24 h (EC_{50} = 3 μM, not shown).

**HYPERFORIN SUPPRESSES CARRAGEENAN-INDUCED MOUSE PAW EDEMA**

The anti-inflammatory properties of Hyp were investigated in carrageenan-induced mouse paw edema in which PG_2E_1 (and also other mediators) critically contributes to the inflammatory response (Guay et al., 2004). Hyp (0.25, 1, 4 mg kg^{-1}) was administered i.p. 30 min prior to injection of carrageenan into the mouse paw. The increase in paw volume was time-dependently assessed and reached a maximum at 4-h post-carrageenan treatment (Figure 6A). In fact, in mice treated with 0.25, 1, and 4 mg kg^{-1} of Hyp, the peak of the response to carrageenan at 4 h was reduced by 35, 63, and 78%, respectively, whereas indomethacin (5 mg kg^{-1}), used as reference, caused 57% inhibition. Moreover, comparison of the areas under the curves of each group, between 2 and 6 h after carrageenan injection, yielded similar degrees of inhibition confirming a more potent effect of Hyp (ED_{50} = 1 mg kg^{-1}) over indomethacin (ED_{50} = 5 mg kg^{-1}; Figure 6B).

**HYPERFORIN SUPPRESSES CARRAGEENAN-INDUCED PLEURISY IN RATS**

Hyperforin has previously been shown to suppress carrageenan-induced pleurisy in rats by decreasing pleural LT_E_1 levels (Feist et al., 2009). Here, we further assessed the formation of PG_E_1 during pleurisy in order to get some information about its ability to inhibit PG_E_1 synthesis in vivo. Injection of carrageenan into the pleural cavity of rats (DMSO 4% group) elicited an acute inflammatory response characterized by the accumulation of fluid that contained large numbers of inflammatory cells (Table 1). As observed in carrageenan-induced paw edema, Hyp (4 mg kg^{-1} i.p., 30 min before carrageenan) significantly inhibited the inflammatory response 4 h after carrageenan injection as demonstrated by the significant attenuation of exudate formation (64%) and cell infiltration (50%). Indomethacin (5 mg kg^{-1}) also reduced exudate formation and cell infiltration (75 and 65%, respectively), and its anti-inflammatory action is not significantly different from Hyp (Table 1). In comparison with the corresponding exudates from DMSO-treated rats, exudates of Hyp-treated animals exhibited significantly decreased PG_E_1 levels, and indomethacin almost completely suppressed PG_E_2 formation as expected (Table 1).

**DISCUSSION**

Systematic investigations of Hyp's pharmacological action within the PG_E_2 biosynthetic pathway revealed mPGES-1 as a putative molecular target of Hyp. Thus, Hyp potently and concentration-dependently inhibited the enzymatic conversion of PGH_2 to PG_E_2 catalyzed by mPGES-1 in a defined cell-free assay, diminished the formation of PG_E_2 in human whole blood without affecting the
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5-LO forms the basis of the anti-inflammatory effectiveness of Hyp and St. John’s wort observed in vitro and in vivo (Schempp et al., 2003; Medina et al., 2006; Hammer et al., 2007; Sosa et al., 2007). St. John’s wort is frequently used for the treatment of mild and moderate depressions but also to intervene with inflammatory disorders, peptic ulcers, and skin wounds (Medina et al., 2006). Although the anti-depressive, anti-inflammatory, and anti-carcinogenic effects of St. John’s wort extracts could be ascribed to Hyp (Muller, 2003; Medina et al., 2006), the underlying molecular mechanisms and molecular targets of Hyp are still elusive. Postulated anti-inflammatory mechanisms include down-regulation of the expression of the chemokine receptor CXCR3 on activated T cells (Cabrelle et al., 2008), down-modulation of matrix metalloproteinase 9, seemingly due to inhibition of leukocyte elastase, and reduced expression of the adhesion molecule CD11b in neutrophils.

FIGURE 4 | Effects of hyperforin on the activity of mPGES-1. (A) Concentration–response curves for Hyp, octahydro-hyperforin, the acylphloroglucinol core, and humulone. (B) Chemical structures of octahydro-hyperforin, the acylphloroglucinol core, and humulone. (C) Concentration–response curves for a St. John’s wort extract. (A,C) For studies of mPGES-1 inhibition, microsomal preparations of IL-1β-stimulated A549 cells were pre-incubated with vehicle (DMSO) or the test compounds at the indicated concentrations for 15 min at 4°C, and the reaction was started with 20 μM PGH₂. After 1 min at 4°C, the reaction was terminated using a stop solution containing FeCl₂ and 11β-PGE₂ (1 nmol) as internal standard. The 100% values in the individual experiments are in the range of 3–4 μg ml⁻¹ PGE₂. Data are given as mean ± SE, n = 3–4. (D) Reversibility of mPGES-1 inhibition by Hyp. Microsomal preparations of IL-1β-stimulated A549 cells were pre-incubated with 3 μM inhibitor for 15 min at 4°C. An aliquot was diluted 10-fold to obtain an inhibitor concentration of 0.3 μM. For comparison, microsomal preparations were pre-incubated for 15 min with 0.3 μM Hyp or with vehicle (DMSO), and then, 20 μM PGH₂ was added (no dilution). Then, all samples were incubated for 1 min on ice, and PGE₂ formation was analyzed as described by RP-HPLC. Data are given as mean ± SE, n = 3–4, * p < 0.01 vs. vehicle (0.1% DMSO) control, ANOVA + Tukey HSD post hoc tests.
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(Dell’Aica et al., 2007), as well as suppression of Ca2+ mobilization, leukocyte elastase release, and formation of reactive oxygen species in neutrophils (Feisst and Werz, 2004). In addition, we recently identified Hyp to inhibit 5-LO and COX-1 in pro-inflammatory eicosanoid biosynthesis in vitro (Albert et al., 2002) thereby exhibiting a unique molecular interference of 5-LO at its C2-like domain (Feisst et al., 2009). Nevertheless, most of these postulated anti-inflammatory mechanisms have been essentially identified and characterized based on cell-free assays and assays based on isolated cells, and thus, the pharmacological relevance of these postulates is unclear. These assays lack the influence of a physiologically relevant environment as compared to studies using whole blood or tissues, and important factors (plasma protein binding, cell–cell interactions, etc.) that may potentially modulate the efficacy are therefore neglected. Moreover, the effective concentrations of Hyp for intervention with certain targets or mechanisms are often substantially higher (>1 μM) than Hyp levels reached in vivo after oral administration of St. John’s wort preparations. Thus, daily oral intake of 3 × 300 mg hypericum extract by human volunteers led to Hyp steady-state concentrations of 0.28 μM (Biber et al., 1998). In light of these facts, suppression of PGE, formation by Hyp was evident in human whole blood with significant suppressive effects at 0.03–1 μM, congruent with the ability of Hyp to directly interfere with mPGES-1 activity in the cell-free assay (IC50 = 1 μM).

Hyperforin almost completely blocked mPGES-1 activity in the cell-free assay, but PGE2 formation in LPS-stimulated whole blood was only partially inhibited (residual PGE2 formation approx. 50% of vehicle control) as observed for mPGES-1 inhibitors such as MK-886 (this study and Koeberle et al., 2008), MD-52 (this study and Koeberle et al., 2009a), and others (Koeberle and Werz, 2009). In contrast, COX inhibitors (i.e., indomethacin or celecoxib) were markedly more effective to suppress PGE2 formation in whole blood. Possibly, constitutively expressed homeostatic PGE synthases like cPGES and mPGES-2, Murakami and Kudo, 2006), contributed to PGE2 synthesis under these conditions and are not inhibited by Hyp. In fact, when whole blood was first pre-incubated with LPS to induce COX-2- and mPGES-1-expression (in contrast to the expression of constitutively expressed COX-1 or PGE2 synthases like cPGES and mPGES-2, Murakami and Kudo, 2006), Hyp showed an increased efficiency to reduce PGE2 formation. The contribution of PGE2 synthases others than mPGES-1 may also explain the incomplete suppression of PGE2 formation in the pleurisy model by Hyp as compared to indomethacin. Future studies will address the dissection of the different enzymatic pathways leading to PGE2 biosynthesis in cellular assays and focus on the respective interference with Hyp.

The anti-inflammatory efficiency of Hyp in vivo was confirmed in carrageenan-induced rat pleurisy and further demonstrated for carrageenan-induced mouse paw edema. During paw edema formation, PGE2 levels were significantly elevated (Harada et al., 1982; Guay et al., 2004), and COX inhibitors were shown to prevent the inflammatory response (Gemmell et al., 1979). Also in the pleurisy model, PGE2 essentially contributes to the inflammatory

**Table 1 | Effect of Hyp on rat carrageenan-induced pleurisy 4 h after i.p. injection.**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Exudate volume (ml)</th>
<th>Inflammatory cells (&gt;10⁴)</th>
<th>PGE2 (ng/rat)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vehicle (DMSO)</td>
<td>0.44 ± 0.04</td>
<td>103.0 ± 15.4</td>
<td>1.81 ± 0.16</td>
</tr>
<tr>
<td>Hyperforin (4 mg kg⁻¹)</td>
<td>0.16 ± 0.05***</td>
<td>51.3 ± 72**</td>
<td>0.97 ± 0.15***</td>
</tr>
<tr>
<td>Indomethacin (5 mg kg⁻¹)</td>
<td>0.11 ± 0.04***</td>
<td>35.8 ± 4.2***</td>
<td>0.19 ± 0.05***</td>
</tr>
</tbody>
</table>

Data are given as mean ± SE, n = 10, ***p < 0.01, **p < 0.001 vs. vehicle (DMSO 4%), ANOVA + Tukey HSD post hoc tests.
response following carrageenan stimulation (Harada et al., 1996; Kawamura et al., 2000). Hyp efficiently suppressed paw edema and was even more potent in this respect than indomethacin. Similarly, Hyp impaired PGE, levels in the exudates of rats during pleurisy and markedly attenuated the concomitant exudate formation and inflammatory cell infiltration. However, our data cannot exclude that inhibition of 5-LO (as previously reported in cell-free and cellular studies and for carrageenan-induced rat pleurisy; Albert et al., 2002; Feisst et al., 2009) contributes to the suppression of PGE, formation by reducing leukocyte recruitment. Taken together, both mPGES-1 and 5-LO seem to be relevant targets underlying the therapeutic efficiency of Hyp in eicosanoid-related disorders.

Besides inflammation, excessive PGE, formation is also associated with tumorigenesis, particularly of colon, breast, prostate, and lung carcinomas. Pharmacological intervention with PGE, formation (COX-2 selective inhibitors) has shown a chemopreventive potential (Rao and Reddy, 2004), and in vitro studies suggest induction of apoptosis as critical step (Jana, 2008). mPGES-1 is overexpressed in various cancers such as non-small cell lung cancer, invasive breast cancer, colorectal cancer, and gastric cancer (Samuelsson et al., 2007). Hyp induced apoptosis of various cancer cells such as mammary carcinoma, squamous cell carcinoma, malignant melanoma, and lymphoma cells (Schempp et al., 2002; Hostanska et al., 2003; Dona et al., 2004; Quiney et al., 2006), and we could demonstrate that also human A549 lung carcinoma cells, constitutively expressing mPGES-1 (Jakobsson et al., 1999), undergo apoptosis in response to Hyp (EC50 ≈ 3 μM) as monitored by the cleavage of PARP. It is tempting to speculate whether a functional correlation between the inhibition of PGE, formation and the induction of apoptosis by Hyp exists. Interestingly, also major depression represents an inflammatory process (Leonard, 2007), where PGE, levels are significantly elevated, and suppression of COX-2 by celecoxib showed clinical efficacy (Muller et al., 2006). A recent meta-analysis suggests significant effectiveness of St. John’s wort in major depression (Linde et al., 2008) but whether inhibition of mPGES-1 contributes to the antidepressant activity of Hyp cannot be answered so far.

In conclusion, Hyp potently inhibits cellular and cell-free PGE, formation via interference with mPGES-1 at physiologically achievable plasma concentrations and shows anti-inflammatory effectiveness in vivo. Hyp also suppresses leukotriene formation by inhibition of 5-LO (Albert et al., 2002), which is also involved in inflammation and carcinogenesis (Werz and Steinhilber, 2006) like mPGES-1 (Murakami and Kudo, 2006). Dual inhibition of mPGES-1 and 5-LO might therefore provide a molecular basis for Hyp’s anti-inflammatory and anti-carcinogenic properties. Such a pharmacological profile would be advantageous, particularly in view of the emerging link between chronic inflammation and cancer, where mPGES-1 and 5-LO may act as potential players.

REFERENCES


