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Discovery of an AHR pelargonidin agonist that counter-regulates Ace2 expression and attenuates ACE2-SARS-CoV-2 interaction

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ABSTRACT

The severe acute respiratory syndrome (SARS)-CoV-2 is the pathogenetic agent of Corona Virus Induced Disease (COVID-19). The virus enters the human cells after binding to the angiotensin converting enzyme (ACE)2 receptor in target tissues. ACE2 expression is induced in response to inflammation. The colon expression of ACE2 is upregulated in patients with inflammatory bowel disease (IBD), highlighting a potential risk of intestinal inflammation in promoting viral entry in the human body. Because mechanisms that regulate ACE2 expression in the intestine are poorly understood and there is a need of anti-SARS-CoV-2 therapies, we have settled to investigate whether natural flavonoids might regulate the expression of Ace2 in intestinal models of inflammation. The results of these studies demonstrated that pelargonidin activates the Aryl hydrocarbon Receptor (AHR) in vitro and reverses intestinal inflammation caused by chronic exposure to high fat diet or to the intestinal braking-barrier agent TNBS in a AhR-dependent manner. In these two models, development of colon inflammation associated with upregulation of Ace2 mRNA expression. Colon levels of Ace2 mRNA were directly correlated with Tnf-α mRNA levels. Molecular docking studies suggested that pelargonidin binds a fatty acid binding pocket on the receptor binding domain of SARS-CoV-2 Spike protein. In vitro studies demonstrated that pelargonidin significantly reduces the binding of SARS-CoV-2 Spike protein to ACE2 and reduces the SARS-CoV-2 replication in a concentration-dependent manner. In summary, we have provided evidence that a natural flavonoid might hold potential in reducing intestinal inflammation and ACE2 induction in the inflamed colon in a AhR-dependent manner.

1. Introduction

The coronavirus disease 2019 (COVID-19) is a respiratory tract infection caused severe acute respiratory syndrome (SARS)-CoV-2, a newly emerged coronavirus first identified in the city of Wuhan in China in December 2019 [1]. The SARS-CoV2 virus entry into host cells is mediated by interaction between the spike (S) glycoprotein, that assembly into an homotrimeric complex protruding from the viral surface, with the angiotensin-converting enzyme 2 (ACE2) [2]. The spike glycoprotein comprises two functional subunits: the S1 subunit contains a specific motif, the receptor-binding domain (RBD), that is responsible for binding to ACE2, while the S2 subunit is essential for fusion of the viral and cellular membranes [2] Thus, targeting the interaction of spike’s RBD with ACE2 might hold potential for the treatment of COVID 19. Accordingly, several agents that target the SARS-CoV2/ACE2 interaction, including several monoclonal antibodies, have been identified and tested clinically. Following a similar approach, it has been suggested that modulation of ACE2 expression could also be of interest in reducing SARS-CoV2 entry in target cells [2–7].

The expression of ACE2 vary greatly among human tissues. In the normal lung, ACE2 mRNA is mainly expressed by type I and II alveolar (AT1 and AT2) epithelial cells and endothelial cells [8,9]. However, it
has been shown that the expression of ACE2 in the lung increases in response to inflammation or exposure to interferon IFN-γ [10]. In normal individual, ACE2 mRNA and protein are highly expressed in the gastrointestinal system, in particular in the duodenum, jejunum, ileum, and colon, although the functional role of this peptidase in intestinal epithelial cells remains poorly elucidated [11]. Furthermore, similarly to the lung, the intestinal expression of ACE2 is modulated in response to intestinal inflammation in inflammatory bowel disease (IBD) in a phenotype dependent manner [12]. More specifically, while expression of ACE2 is reduced in the ileum of patients with active Crohn disease, an elevated colonic expression of ACE2 mRNA has been reported in ulcerative colitis patients with active inflammation and severe disease [13–15]. Since the colonic ACE2 levels normalize after anti-cytokine therapy and the anti-cytokine therapy associates with reduced morbidity of IBD patients from COVID19, these data establish a relation between the levels of colonic ACE2 expression and beneficial effects exerts by anti-cytokine therapies. Moreover, the evaluation of SECURE-IBD registry (available http://www.covidibd.org) suggest that patients with IBD are under-represented in those diagnosed with COVID-19 compared to the general populations [16]. Despite these data raise interest on ACE2 as potential target in IBD in the frame of COVID19 pandemic, the mechanisms involved in ACE2 regulation in the intestine are only partially understood. Further on, it is unclear whether ACE2 expression could be modulated with dietary factors that might have also utility in reducing inflammation.

The aryl hydrocarbon receptor (AhR) is a ligand-activated transcription factor that belongs to the basic helix-loop-helix (bHLH)/Per-ARNT-Sim (PAS) superfamily. Although AhR was originally described as a xenobiotics sensor, it is now well established that dietary/intestinal ARNT-Sim (PAS) superfamily. Although AHR was originally described a xenobiotics sensor, it is now well established that dietary/intestinal AhR ligands bind to the AhR, leading to the activation of the AhR transcription factor. The results of these studies show that not only a dietary agent might regulate the expression of the Ace2 through its interaction with IBD and hence the potential of the natural pelargonidin in regulating orange colours of berries such as raspberries and strawberries, as well as expressed at high levels by intestinal epithelial cells where it is essential as a xenobiotics sensor, it is now well established that dietary/intestinal ARNT-Sim (PAS) superfamily. Although AHR was originally described

### 2. Material and methods

#### 2.1. Transactivation assay

For AhR-mediated transactivation, HepG2 cells were plated at 7.5 × 10⁴ cells/well in a 24-well plate. Cells were transiently transfected with 200 ng of the reporter vector pLightAhRE, 100 ng of pCMVXL4Ahr, and 100 ng of pGL4.70 (Promega, Madison, WI, USA), a vector encoding the human RENILLA luciferase gene. At 24 h post-transfection, cells were stimulated 18 h with Pelargonidin 10 and 50 µM (Merk Group, Darmstadt, GE). Dose-response curves were performed in HepG2 cells transfected with AHR, as described above, and then treated with increasing concentrations of Pelargonidin (1, 2.5 5, 10, 12.5, 25, 50, 75, 100 µM).

At 18 h post-stimulation, cellular lysates were assayed for luciferase and RENILLA activities using the Dual-Luciferase Reporter assay system (Promega, Madison, WI, USA). Luminescence was measured using GloMax 20/20 luminometer (Promega, Madison, WI, USA). LUCIFERASE activities (RLU) were normalized with RENILLA activities (RRU).

#### 2.2. Mouse spleen macrophages purification

Spleens were collected from Ahr−/− and Ahr+/+ mice. Spleens were homogenized and then red blood cell lysis was performed using a hypotonic solution. After cell count, the cell pellet was resuspended at a concentration of 10⁶/mL, to proceed to the positive selection of CD11b+ cells. In particularly, macrophages were purified using CD11b microbeads (Miltenyi Biotec, Bergisch Gladbach, GE), according to the manufacturer’s instructions. Subsequently the selected macrophages have been plated at a concentration of 500,000/mL and were activated with LPS (5 ng/mL, Merk Group, Darmstadt, GE) in combination with IFN-γ (20 ng/mL, ebioscience, San Diego, CA, USA) for 16 h alone or in addition to Pelargonidin (20 µM) for gene expression profile analysis.

#### 2.3. Animals

All animal care and experimental procedures complied with the guidelines of Animal Care and were approved by Use Committee of the University of Perugia and by the Italian Minister of Health and Istituto Superiore di Sanità (Italy) and was in agreement with the European guidelines for use of experimental animals (permissions n. 583/2017-PR and 1126/2016-PR). The general health conditions of the animals were monitored daily by the veterinarian in the animal facility. The study protocol caused minimal suffering. However, mice with an high grade of pain were eutanized. AhR knock out mice (Ahr−/−) on C57BL/6 back-ground and their C57BL/6 congenic littermates wild type (Ahr+/+) were originally supplied by Charles River (Wilmington, MA, USA). Colonies were housed under controlled temperature (22 °C) and photoperiods of 12:12 h light/dark cycle in restricted access area and abled to acclimate to these conditions for at least 7 days before inclusion in an experiment.

To reproduce a mouse model of Acute Colitis a 2,4,6-trinitrobenzenesulfonic acid (Sigma Chemical Co, St Louis, MO) (TNBS)-colitis model is used as previously described [27]. Briefly, according to this 8–10 weeks old male mice C57BL/6 wild-type and Ahr−/− and Ahr+/+ on C57BL/6 genetic background were administrated. Briefly, mice were fasted for 12 h overnight (Day −1). The next day (Day 0), mice were sedated by administration of Zoletil at a dose of 50 mg/Kg of body weight. Then a 3.5F = 11.55 mm catheter was inserted into the colon such that it was up to 4 cm from the anus and 1 mg of TNBS in 50% ethanol administered via the catheter into the colon lumen using a 1 mL syringe (injection volume of 100 µL). Control mice received 50% ethanol alone. Pelargonidin was administrated by oral gavage (1, 5 or 10 mg/kg) daily from day 0 to the 4 day, that is the day of sacrifice. The severity of colitis was scored daily for each mouse by assessing body weight, the fecal occult blood, and stool consistency. Each parameter was scored from 0 to 4 and the sum represents the Colitis Disease Activity Index (CDAI). The scoring system was as follows: Percent of body weight loss: None = 0; 1–5% = 1; 5–10% = 2; 10–20% = 3; >20% = 4. Stool consistency: Normal = 0; soft but still formed = 1; very soft = 2; diarrhea = 3; liquid stools that stick to the anus or anal occlusion = 4. Fecal blood: None = 0; visible in the stool = 2; severe bleeding with fresh blood around the anus and very present in the stool = 4. On day 4, surviving mice were sacrificed, blood samples collected by facial vein, and the colon excised, measured length and weight, and evaluated for macroscopic damage.

To investigate the effects of Pelargonidins in a mouse model of NASH, 10–12 weeks old C57BL/6 male mice, Ahr wild type (Ahr+/+) and their congenic littermates Ahr null mice (Ahr−/−) were fed a high fat diet containing 59 KJ% fat plus 1% cholesterol, w/o sugar (sniff, Soest, GE) and fructose in drinking water (42 g/L) for 8 weeks as previously
described [28]. Food intake was valued as the difference of weight between the provided and the remnant amount of food at 2-day intervals. The food was provided as pressed pellets and the residual spillage was not considered. After 8 days, Ahr−/− and Ahr+/+ HFD-F mice were randomized to receive HFD-F alone or plus Pelargonidin (5 mg/Kg/die) by oral gavage for remaining 7 weeks. On the sacrifice day, fed mice were deeply anesthetized with Zoletil at a dose of 50 mg/Kg and sacrificed before 12 AM. Thus, blood, liver, small intestine, colon, spleen, epididymal adipose tissue (eWAT), brown adipose tissue (BAT) and gastrocnemius were collected. The abdominal circumference (AC) (immediately anterior to the forefoot), body weight, and body length (nose-to-anus or nose-anus length), were measured in anaesthetized mice at the day of sacrifice. The body weight and body length were used to calculate the Body mass index (BMI) (body weight (g)/length² (cm²)).

2.4. Intestinal permeability

Intestinal permeability test was performed using fluorescein isothiocyanate conjugated dextran (FITC-dextran) (Sigma-Aldrich, St. Louis, MO, USA). FITC-dextran was dissolved in PBS at a concentration of 100 mg/mL. The evening before the test day mice were fasted overnight. In the next morning, after weighing each mouse, FITC was administered to each one at a dose of 44 mg/100 g of body weight by oral gavage. After 4 h, the mice were anesthetized by inhalation of isoflurane and 300–400 μL of blood collected from the facial vein in microtubes containing EDTA (Invitrogen, Carlsbad, CA, USA). Blood was stored in the dark. Once blood has been collected from all the mice, microtubes were processed to separate the serum. For analysis, serum was diluted with an equal volume of PBS (Euroclone S.p.a Milan, IT). The concentration of FITC in serum was determined by spectrophotometry with an excitation of 485 nm and an emission wavelength of 528 nm using as standard serially diluted FITC-dextran (0, 125, 250, 500, 1000, 2000, 4000, 6000, 8000 ng/mL). Serum from mice not administered with FITC-dextran was used to determine the background.

2.5. Histology

Colon sample (2–3 cm up anus) were first fixed in 10% Formalin (Bio-Optica Milano S.p.A. Milan, IT), embedded in Paraffin (Bio-Optica Milano S.p.A. Milan, IT), cut into 5-μm-thick sections and then stained with Hematoxylin/Eosin (H&E) (Bio-Optica Milano S.p.A. Milan, IT) for histopathological analysis. The histological score of the colon was assessed as previously described by U. Erben et al. [29]. This score evaluated the level of tissue inflammation in relation to the extension of the inflammatory cell infiltrate (mild severity if the cellular infiltrate is present in the mucosa, moderate if they are involved mucosa and submucosa, marked if the infiltration is transmural), and epithelial changes of the intestinal mucosal architecture like erosions or ulcerations.

Immunohistochemistry was performed on paraffin embedded colon. In brief, Ag retrieval was achieved by incubation of the slides for 90 min in the hot (95 °C) sodium citrate buffer (pH 6.0) and 30 min of cooling at room temperature. Immunostaining technique was performed using the commercial kit Elabscience®/2-step plus Poly-HRP Anti Rabbit/Mouse IgG Detection System (with DAB Solution) (Houston, Texas 77079, USA.) ACE2 Recombinant Rabbit Monoclonal Antibody (SN0754A) Invitrogen, Thermofisher scientific (Waltham, Massachusetts, USA) was incubated overnight at 4°C. Subsequently, sections were incubated with Polyperoxidase-anti-Mouse/Rabbit IgG and then with DAB Working Solution, both supplied by the kit. Slides were counterstained with hematoxylin, dehydrated through ethanol and xylen, and coverslipped using a xylene-based mounting medium.

2.6. Quantitative Real-Time PCR analysis

RNA was extracted from mouse colon and spleen macrophages using respectively TRIzol reagent (Invitrogen, Carlsbad, CA, USA) and Directzol™ RNA MiniPrep w/ZymoSpin™ IIC Columns (Zymo Research, Irvine, CA, USA). 1 μg of RNA from each sample was reverse transcribed using FastGene Scriptase Basic Kit (Nippon Genetics Europe, Düren, GE) in a 20-μL reaction volume; 50 ng of cDNA was amplified in a 20-μL solution containing 200 nM of each primer and 10 μL of SYBR Select Master Mix (Thermo Fisher Scientific, Waltham, MA, USA). All reactions were performed in triplicate using the following thermal cycling conditions: 3 min at 95 °C, followed by 40 cycles of 95 °C for 15 s, 66 °C for 20 s, and 72 °C for 30 s, using a StepOnePlus system (Applied Biosystems, Foster City, CA, USA). The relative mRNA expression was calculated according to the ΔCt method. Primers were designed using the software PRIMER3 (http://frodo.wi.mit.edu/primer3/), using data published in the NCBI database. The primer used were as following (forward and reverse): mGapdh (for CTGAGTATGCTGGATGCTCT; rev GTTGTTGTGGTCCAGGATGCTTG), mTnfr1 (for CCA-CACGCTCTTCTGCTTA; rev AGGGCTGCGGCAATAAGA), ml16 (for CTCACAAGTCCGGAGCTTA; rev TTCTGCAAGTGCATCATG), mTfpi (for TTGCTTCAGGCCTCAAGAGA; rev TGGTGTGAGGGGAGCAAGAC), mAce2 (for AGATGGCGCGAAGATTGTCT; rev GGGGCTGTCGAAGAAGTTGCTC), mMas (for CTGCTGACAGCAGCAGTGT; rev ACAGAAGGGCCAGCAGGAAT), ml11 (for GCTGAAAGCTTCCACACCTCA; rev AGGCCCAGATTATTTTGGTCG), Cyp1a1 (for ACTCTCTTCCCTGAGTCCCT; rev GTCTGTGATGCTCCCCAGATG) (Thermo Fisher Scientific, Waltham, MA, USA).

2.7. Cell culture

Caco-2 cells, a human intestinal epithelial cell line (Sigma-Aldrich St. Louis, MO, USA) was grown at 37 °C in D-MEM (Euroclone S.p.A Milan, IT) containing 10% FBS (Euroclone S.p.A Milan, IT), 1% l-glutamine (Euroclone S.p.A Milan, IT), and 1% penicillin/streptomycin (Euroclone S.p.A Milan, IT). Cells were regularly passaged to maintain exponential growth. Caco-2 cells were classically activated with hTNF-α (Sino Biological, Beijing, CN) 100 ng/ml for 24 h alone or in combination with Pelargonidin (5, 10, and 20 μM) for the analysis of expression of several genes.

2.8. Mouse colon epithelial cells (IECs) purification and culture

Colon were collected from 6 to 10 weeks Ahr+/+ and Ahr−/− mice. Organs were cleaned of feces by flushing with PBS 2% Fetal bovine serum (FBS) using a syringe and were cut first longitudinally and then laterally into 0.5 cm length pieces. Each tissue pieces were incubated for 30 min at 37 °C in 10 mL digestion solution containing DMEM with 5% FBS, 10 mM HEPES (Euroclone S.p.A Milan, IT), 5 mM EDTA, 1 mM DTT (Sigma-Aldrich, St. Louis, MO, USA). After incubation, samples were mixed for 30 s using a vortex mixer. Cell suspensions were stored on ice and supernatant were separate from colon tissue pieces and transferred into new tube after were applied into a MACS SmartStrainer 100 μm (Miltenyi Biotec, Bergisch Gladbach, GE) placed on a 15 mL tube. After centrifugation cell suspension were applied onto a MACS SmartStrainer 15 mL reaction volume; 50 ng of cDNA was amplified in a 20-μL solution containing 200 nM of each primer and 10 μL of SYBR Select Master Mix (Thermo Fisher Scientific, Waltham, MA, USA). The relative mRNA expression was calculated according to the ΔCt method. Primers were designed using the software PRIMER3 (http://frodo.wi.mit.edu/primer3/), using data published in the NCBI database. The primer used were as following (forward and reverse): mGapdh (for CTGAGTATGCTGGATGCTCT; rev GTTGTTGTGGTCCAGGATGCTTG), mTnfr1 (for CCA-CACGCTCTTCTGCTTA; rev AGGGCTGCGGCAATAAGA), ml16 (for CTCACAAGTCCGGAGCTTA; rev TTCTGCAAGTGCATCATG), mTfpi (for TTGCTTCAGGCCTCAAGAGA; rev TGGTGTGAGGGGAGCAAGAC), mAce2 (for AGATGGCGCGAAGATTGTCT; rev GGGGCTGTCGAAGAAGTTGCTC), mMas (for CTGCTGACAGCAGCAGTGT; rev ACAGAAGGGCCAGCAGGAAT), ml11 (for GCTGAAAGCTTCCACACCTCA; rev AGGCCCAGATTATTTTGGTCG), Cyp1a1 (for ACTCTCTTCCCTGAGTCCCT; rev GTCTGTGATGCTCCCCAGATG) (Thermo Fisher Scientific, Waltham, MA, USA).
2.9. ACE2/SARS-CoV-2 Spike inhibitor Screening assay Kit

Pelargonidin at different concentrations (1–10–20–50 μM) was tested using the ACE2: SARS-CoV-2 Spike Inhibitor Screening Assay Kit (BPS Bioscience San Diego, CA, USA) according to the manufacturer’s instructions. Briefly, we have thawed ACE2 protein on ice and diluted to 1 μg/mL in PBS. We have used 50 μL of ACE solution to coat a 96-well nickel-coated plate (1 h). The plate was washed 3 times and incubated for 10 min with a Blocking Buffer. Next, 10 μL of inhibitor solution containing the Pelargonidin was added to wells and incubated for 1 h at room temperature with slow shaking. For the “Positive Control” and “Blank,” 10 μL of inhibitor buffer (5% DMSO, Sigma-Aldrich, St. Louis, MO, USA, solution) were used. After the incubation (1 h), SARS-CoV-2 SPIKE (RBD)-Fc was thawed on ice and diluted to 0.25 ng/μL (≈5 nM) in Assay Buffer 1; 5 nM SPIKE protein was added to each well, except to the blank. The reaction was incubated for 1 h at room temperature, with slow shaking. After 3 washes and incubation with a Blocking Buffer (10 min), we have treated the plate with an Anti-mouse-Fc-HRP and the plate was incubated for 1 h at room temperature with slow shaking. Finally, HRP substrate was added to the plate to produce chemiluminescence, which was measured using FluoroStar Omega microplate reader (BMG LABTECH Ottenberg, GE).

To confirm the validity of the assay used in this study, remnants of plasma samples used to test levels of anti-SARS-CoV2 IgG in post COVID-19 patients were used. The original samples were collected at the blood bank of Azienda Ospedaliera of Perugia from post COVID-19 donors who participate to a program of plasma biobanking. The program’s protocol included the quantitative analysis of the anti-SARS-CoV-2 IgG antibodies directed against the subunits (S1) and (S2) of the virus SPIKE protein. An informed consent was obtained by each donor for the use of the plasma sample remnants and the protocol was approved by the Ethical Committee of the University of Perugia: authorization n. 61,843 (July 13, 2020).

2.10. Virus isolation and cell cultures

The experiments were performed in Biological Safety Level 3 (BSL-3) virology laboratory of “Santa Maria della Misericordia” Hospital, Perugia, Italy. The SARS-CoV-2 strain was isolated as previously described [30]. For viral propagation, Vero E6 cell line was maintained in Eagle minimal essential medium (MEM) supplemented with 10% foetal bovine serum (FBS) and 100 U/mL penicillin–streptomycin solution. After isolation, SARS-CoV-2 titer was determined by Median Tissue Culture Infectious Dose (TCID50) endpoint dilution and stock aliquots were stored at −80 °C. The stock virus titer was 4.22 × 10^7 TCID50/mL.

2.11. SARS-CoV-2 yield reduction assay

Vero E6 cells (20,000 cells/well) were seeded in 96-well clear flat-bottom plates and incubated at 37 °C with 5% CO2 for 24 h. After incubation, Pelargonidin at concentration of 20, 50 and 100 μM, and Remdesivir 10 μM (Veklury, Gilead, US) all diluted in complete medium were added to each well and then, cells were infected with SARS-CoV-2 (50 or 100 TCID50/well). Negative controls (pelargonidin or Remdesivir alone), infected positive controls (SARS-CoV-2 alone) and mock-infected cells were included in each plate. Plates were incubated at 37 °C with 5% CO2 for 48 h. After the incubation, supernatants of 5 technical replicates were pooled and stored at −80 °C for further analysis.

2.12. Plaque-reduction assays

 Supernatants viral titer was determined by plaque assay as previously described [31], with some modifications. Vero E6 cells (600,000 cells/well) were seeded in a 6-well plate and incubated at 37 °C with 5% CO2 for 24 h. After incubation, the medium was removed and cells were infected with 500 μL of ten-fold serial dilution of supernatant previously obtained, rocking the plates every fifteen minutes. In the meanwhile, the overlay medium (complete medium with Agar 0.1%) was prepared and maintained in a 50 °C water bath. After 1 h of infection, the overlay medium (2 mL) was poured into each well and the plates incubated for 3 days. Finally, the overlay was discarded, cells were fixed for 30 min with 4% formalin and stained with 0.5% crystal violet. Viral titer was determined as plaque-forming units per mL, considering wells with plaques ranging from 2 to 100. For each pool of supernatants, plaque-reduction assay was performed in duplicate.

2.13. Computational studies

Two cryo-electron microscopy structures of SARS-CoV2 were employed for docking calculations, in the central b-sheet core and in the flavonoids binding pocket (PDB ID: 6VSB) [32], while for dockings in the fatty acids (FA) pocket we used the (PDB ID 6ZBS) [33]. The receptor was treated with the Protein Preparation [34] tool implemented in Maestro ver. 11.8. (Schrödinger Release 2019–1: Maestro, New York). The 3D structure of pelargonidin was built using the Graphical User Interface (GUI) of Maestro ver. 11.8. (Schrödinger Release 2019–1: Maestro, New York). The protonation state of Pelargonidin at pH 7.4 in water has been calculated using the Epik module [35]. Finally, the compound was then minimized using the OPLS 2005 force field through 2500 iteration steps of the Polak-Ribiere Conjugate Gradient (PRCG) [36] algorithm.

The docking procedure was realized with the Glide software package (Glide, version 7.1. New York, NY: Schrödinger, LLC, 2019) using the Standard Precision (SP) algorithm of the GlideScore function [37] and the OPLS 2005 force field [38]. A grid box of 20 × 20 × 20 Å for SARS-CoV2 receptor centered on the putative binding pocket was created. A total amount of 100 poses was generated. Docking conformations of pelargonidin were then clustered based on their atomic RMSD and five clusters were obtained. Among them, only the conformation included in the most populated cluster with both the Glide Emodel and GlideScore docking lowest-energy value was considered.

2.14. Statistical analysis

The two-tailed unpaired Student’s t test was used for statistical comparisons (*P < 0.05) using the Prism 6.0 software (GraphPad). The calculation of Pearson r, P value and R squared was performed in the correlation graphs using the Prism 6.0 software (GraphPad).

3. Results

3.1. Activity of the natural compounds toward AHR

We have first investigated whether the natural pelargonidin exerted agonistic effects on AHR. The activity of pelargonidin was evaluated in a Luciferase reporter assay using HepG2 cells, transiently transfected with an AHR reporter gene cloned upstream to the LUCIFERASE. HepG2 were incubated with 5 nM TCDD, an AHR agonist as a control, or vehicle (0.1% v/v DMSO) in the presence of the Pelargonidin (10–50 μM) for 18 h. As shown in Fig. 1A, Pelargonidin was found effective in trans-activating the AHR; furthermore, in a concentration–response curve analysis, we found that pelargonidin transactivated AHR with a relative potency, expressed as EC50, of 12 μM (Fig. 1B).

3.2. Pelargonidin rescues from intestinal inflammation in TNBS-induced colitis in mice and restore Ace2 expression

We have investigated whether pelargonidin administration attenuates intestinal inflammation in a mouse model of colitis. We first performed a dose finding experiment by administering Pelargonidin at a dose of 1, 5 or 10 mg/kg in a mouse model of TNBS-induced colitis (Fig. 2). Clinical data and analysis of the macroscopic and microscopic...
features of the colon showed for doses of 1 and 5 mg/kg of pelargonidin a dose-dependent effect with the lowest dose exerting only mild beneficial effects. However, the 10 mg/kg dose, contrary to expectations, showed intermediate effects between the 1 and 5 mg/kg dose. Because these data suggest that 5 mg/kg of Pelargonidin was fully effective in reversing signs of intestinal inflammation we used this dose to further investigate the mechanism of action of pelargonidin on intestinal immunity. For this purpose, we induced a colon inflammation by administering TNBS to Ahr\(^{-/-}\) and Ahr\(^{-/-}\)mice. As shown previously by us and others, Ahr\(^{-/-}\) mice are very susceptible to the TNBS colitis with high mortality rate [26]. In our experimental set we found a mortality rate of 80% in the Ahr knock-out group and therefore it was not possible to further investigate in this strain (data not shown). In wild-type mice, however, the severity of wasting disease and intestinal inflammation induced by TNBS was reversed by treatment with pelargonidin as measured by lower body weight lost, lower CDAI and by assessing the macroscopic and microscopic feature of the colon (Fig. 3A–D). We analyzed the colon expression of Ace2 protein and mRNA (Fig. 3E, F). Immunohistochemistry and qPCR confirmed that induction of colitis by administration of TNBS increased the expression of Ace2 both mRNA and protein as found in patients with IBD. In contrast, the administration of pelargonidin was able to counteract the increase in Ace2 expression (Fig. 3E, F) [12]. To further confirm that pelargonidin administration in vivo activates the AHR receptor, we have evaluated the expression of one of the main target genes, Cyp1a1. The data showed that pelargonidin administration up-regulates Cyp1a1 expression in the colon (Fig. 3G). We then analyzed the colon expression of Mas and cytokines (Fig. 3H–J).

In this mouse model of colitis, we found exposure to TNBS promoted a strong upregulation of pro-inflammatory cytokines (Fig. 3I–J); Il-1β was up-regulated \( \approx 20 \) times in the colon of TNBS-treated mice compared to naïve mice (Fig. 3I). Furthermore, a five fold increase in Tnf-a mRNA (Fig. 3J) was detected in mice exposed to TNBS. In contrast, Mas mRNA expression was decreased (Fig. 3H). Administering TNBS mice with pelargonidin reversed this pattern while induced the expression of Tgf-β (Fig. 3I–K). The anti-inflammatory effect exerted by pelargonidin also resulted in a strong reduction of Ace2 mRNA expression (Fig. 3F). The correlation analysis between Il and I/\( \beta \)/Ace2 and Tnf-a/Ace2 confirmed a statistically significant correlation between the expression of the two pro-inflammatory cytokines and Ace2 (Il-I/\( \beta \)/Ace2 \( P \) value = 0.0244; Tnf-a/Ace2 \( P \) value < 0.0001) (Fig. 3L, M). Again, there was a closer correlation between the expression of Ace2 and that of Tnf-a (Il-I/\( \beta \)/Ace2 \( R \) squared = 0.1867; Tnf-a/Ace2 \( R \) squared = 0.7695) (Fig. 3L, M).

Due to the inability to analyze the effects of pelargonidin treatment in the colitis model in Ahr\(^{-/-}\) mice for the high mortality, to investigate whether the immunomodulatory effect exerted by Pelargonidin was mediated by binding on AHR, we tested the compound on murine macrophages purified from the spleen of Ahr\(^{-/-}\) and Ahr\(^{-/-}\) mice. Macrophages were subjected to pro-inflammatory stimuli (LPS + IFN-\( \gamma \)) and treated with pelargonidin for 16 h. Stimulation with LPS + IFN-\( \gamma \) induced an increase in the production of pro-inflammatory cytokines (Fig. 4A, B) and a reduction in the production of Tgf-β (Fig. 4C) in macrophages deriving from both genotypes. Il-6 up-regulation was much higher in Ahr\(^{-/-}\) macrophages compared to wild-type (Fig. 4E). Treatment with Pelargonidin reverted the pro-inflammatory polarization but only in wild-type macrophages, while it had no effect on Ahr\(^{-/-}\) macrophages, underlining the role of the AHR receptor in the mechanism of action of Pelargonidin.

3.3. Pelargonidin protects against colon inflammation induced by HFD-F and down-regulates Ace2 expression

Taking into consideration obesity pandemia and that exposure to high caloric intake promotes a state of intestinal sub-inflammation, and that obesity is a well-defined risk factor for development of severe COVID19, we have tested the effects of Pelargonidin also in a mouse model of a high-fat/high-sugar diet [39–41].

In this model, Ahr\(^{-/-}\) and Ahr\(^{-/-}\) mice were treated with exposed a chronic high caloric [26] intake by feeding them a diet enriched cholesterol and fructose (HFD-F) for 8 weeks. Starting from the second week (day 8) an experimental group of mice for each genotype was treated with pelargonidin daily. As shown in Fig. 5, mice exposed to chronic high caloric intake gained significantly more body weight than mice feed a normal chow diet, and they had significantly higher BMI at the end of the study without significant differences between the two genotypes (Fig. 5A–C). Treating mice with 5 mg/kg/day Pelargonidin for 7 weeks protected against body weight gain and resulted in a lower BMI in comparison to mice feed an HFD-F alone (Fig. 5A–C). Instead, the beneficial effects exerted by Pelargonidin were completely abrogated in Ahr\(^{-/-}\) mice (Fig. 5A–C). We have therefore focused our attention on intestinal permeability. The histological analysis of the colon showed no major morphological abnormalities, although further functional assessment revealed a robust increase of intestinal permeability in the mice exposed to HFD-F (Fig. 5D, E). Treatment mice with pelargonidin restored the normal colon permeability in wild-type mice, but had no beneficial effects in Ahr\(^{-/-}\) mice which exhibited much more severe changes in permeability than in Ahr\(^{-/-}\) mice (Fig. 5D, E).
Elevated colonic ACE2 levels have been detected in the colon of IBD patients and are associated with inflammation and severe disease, suggesting a possible correlation between ACE2 expression and colon inflammation in IBD patients [12]. We have therefore investigated the expression of Ace2 and Mas in the colon and compared them with colon profiles of various cytokines (Fig. 5 F–J). HFD-F increased Ace2 expression and reduced Mas expression in both Ahr<sup>+</sup>/ and Ahr<sup>−/−</sup> mice, while increased the colon expression of Il-1β and Tnf-α and downregulated the Tgf-β mRNA. Pelargonidin treatment reversed the inflammatory pattern by down-regulating the expression of both Il-1β and Tnf-α in Ahr<sup>−/−</sup> mice. In wild-type mice exposure to the anthocyanin also reduced the expression of Ace2, bringing it back to the value measured in untreated wild-type mice, while exerted only a slight effect on the expression of Mas mRNA (Fig. 2 F–J). To investigate whether there was also in this mouse models with a mild inflammation in the colon, a direct correlation between the expression of Ace2 and pro-inflammatory cytokines, we performed correlation analyses between Il and 1β and Ace2 and between Tnf-α and Ace2. As shown in Fig. 5 K and L, the colon levels of the two cytokines were positively correlated with Ace2 expression in both Ahr<sup>+/+</sup> and Ahr<sup>−/−</sup> mice. The fact that the R squared value was greater in the Tnf-α/Ace2 correlation than Il-1β/Ace2 confirmed a stronger correlation between the expression of Ace2 and Tnf-α.

3.4. In vitro characterization of pelargonidin effects on intestinal epithelial cells

Because, ACE2 expression in the intestine is highly restricted to epithelial cells, we have then examined whether exposure of Caco-2 cells, a human intestinal epithelial cell line, to TNF-α modulates the expression of ACE2. The results of these experiments demonstrated that exposure to TNF-α promotes the expression of pro-inflammatory genes Il-8, Il-6 and Il-1β by Caco-2 cells (Fig. 6B–D) and increased the expression of Ace2 (~2.5 times) confirming the close correlation between TNF-α and ACE2 (Fig. 6A). Administration of Pelargonidin reversed this pattern in a concentration-dependent manner (Fig. 6A–D).

To investigate the molecular mechanism behind the correlation between TNF-α and ACE2 and behind the beneficial effect exerted by Pelargonidin, we purified intestinal epithelial cells (IECs) from the colon
Fig. 3. Pelargonidin effect on acute colitis. Colitis was induced by TNBS. After induction of colitis the mice were treated daily with Pelargonidin (5 mg/Kg) or vehicle. Disease was monitored by daily evaluation of (A) changes in body weight (%), (B) colitis disease activity index (CDAI), (C) colon length (cm) and ratio of colon weight/colon length (g/cm). (D) H&E staining of colon sections (10× magnification) and Histological Score from each experimental group. (E) The figure shows immunohistochemistry representative images of the colon of one mouse for each experimental group stained with anti-ACE2 Ab (20x magnification). RNA extracted from colon was used to evaluate, by quantitative real-time PCR, the relative mRNA expression of (F) Ace2, (G) Cyp1a1, (H) Mas, (I) Il-1β, (J) Tnf-α and (K) Tgf-β. Values are normalized relative to Gapdh mRNA. The values are expressed relative to those of control group (NT) which are arbitrarily set to one. Correlation
of Ahr\(^{+/+}\) and Ahr\(^{-/-}\) mice and stimulated them with TNF-\(\alpha\). By analyzing the expression of the Ace2 and Il-6 genes we demonstrated that also in these primary murine intestinal epithelial cells TNF-\(\alpha\) induces inflammation by up-regulating both genes (Fig. 7A and B). Furthermore, the absence of the AHR receptor exacerbated the inflammation induced by TNF-\(\alpha\). Pelargonidin reverted the upregulation of both genes in a concentration-dependent manner but only in cells purified from wild-type mice (Fig. 7A and B). In \(\text{Ahr}^{-/-}\) mice, the absence of the receptor abrogated the beneficial effect exerted by Pelargonidin. It is known that TNF-\(\alpha\), through its receptor, induces the transcription of several pro-inflammatory genes by activating the NF-kB pathway [42,43]. For this reason, we have investigated the effect of NF-kB pathway inhibition on Ace2 expression. Administration of an NF-kB inhibitor was able to down-regulate the expression of Ace2 and Il-6 in both wild-type and knock-out cells confirming that the mechanism by which TNF-\(\alpha\) up-regulate Ace2 and Il-6 involves the activation of NF-kB.

Since ACE2 represents the host receptor used by the SARS-Cov2 virus to invade cells, and reducing the expression of \(\text{ACE2}\) might represent an interesting host defense mechanism against virus invasion. In this context we also investigated whether Pelargonidin was able to reduce the binding of the viral Spike protein on ACE2 (Fig. 8).

We therefore investigated, through docking calculations, the possibility of Pelargonidin to bind several pockets suggested in previous studies: i) the hydrophobic pockets on the b-sheet core of the RBD [44], ii) the fatty acid (FA) pocket identified by Toelzer et al. [33,45], and finally, iii) the flavonoids binding pocket suggested by Allam et al. [46]. All docking calculations were performed using Glide software package. The scores of the best poses highlighted a marked preference of the results obtained in the FA pocket (best docking score \(7.7\) kcal/mol), with respect to the results obtained in the hydrophobic pockets on the b-sheet core of the RBD (best docking score \(5.2\) kcal/mol) or in the pocket proposed by Allam (best docking score \(5.8\) kcal/mol), thus suggesting that Pelargonidin binds the FA pocket on the RBD of Spike, as already reported for other highly unsaturated naturally occurring compounds (e.g. Vitamin k, retinoids) [45]. Analysis of the best scored and clustered docking pose in the FA pocket (Fig. 8A–C) showed the Pelargonidin polyphenolic ring contacting with Leu368, Leu387, Phe388, Phe342 and Ile434. Furthermore, Phe377 makes a p-cation interaction with the oxygen of the ring C, whereas Tyr365 and Tyr369 are engaged in a p-p stacking with the biphenyl ring and with the ring B, respectively. Finally, the binding mode is further stabilized by the H-bond formed between the ring B hydroxyl group and the backbone of Ala372.
Given the results of the docking calculations, we have then investigated whether the Pelargonidin impact on the binding of S protein to the ACE2 receptor. For this purpose, a Spike/ACE2 Inhibitor Screening Assay Kit was used. The results obtained by us showed that pelargonidin was able to reduce binding in a concentration-dependent manner. At the highest tested concentration, equal to 50 μM, we measured a reduction in Spike binding on ACE2 of about 40% (Fig. 8D). To confirm this data, we also performed a SARS-CoV-2 virus infection test on Vero E6 cell line, kidney epithelial cells extracted from monkey. In this assay, pelargonidin showed an ability to reduce virus entry, measured by plaque assay, of approximately 70% at the highest concentration of 100 μM (Fig. 8E).

The data obtained from docking and in vitro tests associated with the down-regulation of ACE2 expression exerted by Pelargonidin and the anti-inflammatory activity make pelargonidin very interesting in the prevention and treatment of SARS-CoV-2 infection.

4. Discussion

ACE2, the SARS CoV2 receptor, in widely expressed in the...
gastrointestinal tract, although the terminal ileum is one of the human body tissues with the highest expression of the receptor [21]. In the ileum, ACE2 regulates amino acids uptake and is essential for nutrients absorption. The receptor is also expressed in normal human colon, thought at significantly lower levels than the ileum, but its functional role in this portion of the intestine is less defined. Of relevance, however, the expression of ACE2 in the two tissues undergoes an opposite regulation in response to inflammation. Thus, while the ileal expression of ACE2 is reduced in patients with active Crohn’s disease of the ileum, the colonic expression of the receptor is robustly induced in patients with ulcerative colitis or Crohn’s disease showing a colon involvement [47]. Further on, a linear correlation has been detected linking the severity of colon inflammation, colon expression of pro-inflammatory cytokines including IFN-$\gamma$, TNF-$\alpha$, IL-1$\beta$, and IL-6 and the expression of ACE2 mRNA [12,13,48,49]. This inflammation-modulated expression of Ace2 in the colon show similarity with the receptor regulation in the

Fig. 7. TNF-$\alpha$ up-regulates Ace2 expression by NF-$\kappa$B on intestinal epithelial cells and pelargonidin inhibits this pathway by activating AHR. Intestinal epithelial cells were purified from the colon of Ahr$^{+/+}$ and Ahr$^{-/-}$ mice. Intestinal epithelial cells were cultured for 24 h with TNF-$\alpha$ 100 ng/ml and treated with Pelargonidin (5, 10, and 20 $\mu$M) or with the NF-$\kappa$B inhibitor (INF-$\kappa$B 100 nM). At the end of stimulation, the relative mRNA expression of (A) Ace2 and (B) IL-6 was evaluated by Real-Time PCR. Values are normalized relative to Gapdh mRNA and the values are expressed relative to those of control group (Ahr$^{+/+}$ NT) which are arbitrarily set to one. are expressed as mean $\pm$ SEM ($n$ = 5); * $p$ < 0.05.
lung, the target tissue of COVID19 [10,50]. It is now well established that the expression of the receptor in alveolar cells increases in a cytokine dependent manner in response to SARS-CoV2 infection, with a direct correlation with the tissue expression of IFN-γ. While these data highlight a potential role for an inflammatory network in promoting intestinal ACE2 expression, the molecular mechanisms involved in its counter-regulation remains elusive. To answer this question, we have settled up a study to investigate the role of the AHR in regulating Ace2 expression in mouse models of IBD. Additionally, since obesity and diabetes promote a state of sub-inflammation in the gastrointestinal tract [51,52] and it are two well-defined risk factors for severe COVID19, we have investigated whether this regulation is maintained in mice exposed to a high caloric intake.

Gastrointestinal symptoms, including vomiting, diarrhea, or abdominal pain of various severity, occurs frequently in the early stage of COVID19, highlighting the role of intestinal ACCE in promoting SARS CoV2 infection. This view is of specific relevance in patients with IBD. Two recent studies [53,54] have shown that SARS CoV2 virus might directly infect the human gut, and therefore the modulation of ACE2 expression in the gastrointestinal tract might have therapeutic relevance in reducing the virus load. The data we have gathered from the two mouse models of IBD strongly support the notion that upregulation of Ace2 in the colon is due to a state of inflammation marked by strong induction of pro-inflammatory cytokines [47,55]. Importantly, in contrast to Ace2 the expression of the Ang 1–7 receptor Mas, was markedly downregulated in both models (Figs. 3 and 5). The correlation...
of Ace2 mRNA expression with the levels of inflammatory biomarkers was very stringent. Indeed, while in the TNBS model the levels of various cytokines were increased by 5–20 folds and the expression of Ace2 mRNA was induced by ≈7-fold in comparison to naive mice (Fig. 3E), in the HFD-F model only a mild inflammation was detected and the levels of IL-1β and Tnf-α and Ace2 mRNA levels were increased by approximately two-folds (Fig. 5H–J). Moreover, in the colon of animals subjected to a high-calorie intake, the expression of Mas was halved in comparison to that in mice fed a chow-diet (Fig. 5F, G). These data establish, across the various models, a direct correlation between the severity of colonic inflammation and Ace2 gene transcription.

In both mouse models we have found that values of R squared indicated a close correlation between the expression of Ace2 and Tnf-α. These data are deeply consistent with the finding that in IBD patients, anti-TNF-α therapies negatively regulate ACE2 expression in the colon [12,56]. Confirmation of the direct regulation of ACE2 expression by TNF-α was given obtained also by results of in vitro experiment, since exposure of human intestinal epithelial cells, Caco 2, to TNF-α directly up-regulates the expression of ACE2 (Fig. 6A).

Similarly, to other flavonoids, pelargonidin, a common constituent of many red plant foods, including berries, grapes or cabbage, have been used in the traditional medicine in the treatment of a number of human alignments [57,58]. Pelargonidin display a several biological activities, the most prominent being those related to their anti-oxidant effects. We have confirmed in this study that pelargonidin exerts anti-inflammatory effects in intestine in mouse models of IBD and obesity through the interaction with the AHR. Treatment with pelargonidin reduced the expression of pro-inflammatory cytokines in the colon in both models of intestinal inflammation, alleviating the disease (Figs. 2, 3 and 5). Furthermore, treatment with Pelargonidin restored Ace2 expression to the level measured in the colon of naive mice (Figs. 3E and 5F). The beneficial effects exerted by the compound were abrogated in the absence of AHR confirming the functional relevance of the receptor in mediating the anti-inflammatory activities of this flavonoid (Figs. 4, 5 and 7). In vitro studies also confirmed that pelargonidin might directly inhibit ACE2 induction caused by exposure of human intestinal epithelial cell line to the TNF-α (Fig. 6). Moreover, through the use of an NF-kB inhibitor, we have shown that the up-regulation of ACE2 by TNF-α is mediated by the activation of NF-kB (Fig. 7). Using intestinal epithelial cells purified from wild-type and Ahr−/− mice we showed that the effect of down-regulating of Ace2 exerted by Pelargonidin requires the presence of AHR (Fig. 7). As demonstrated in other studies, the activated AHR receptor can exert anti-inflammatory effects by inhibiting the NF-kB pathway [59,60].

The regulatory effects of pelargonidin on the intestinal ACE2 expression might be of relevance in the context of SARS-CoV-2 infection, since high levels of ACE2 are compatible with an increased risk of SARS-CoV-2 uptake by the inflamed colon [12]. Importantly, in addition to ACE2 regulation, we have shown that beneficial effects of Pelargonidin might extend to a direct regulation of SARS-CoV-2/ACE2 interaction. Indeed, we found that Pelargonidin not only reduces the interaction between ACE2 and the Spike protein of the SARS-CoV-2 but also reduces the ability of the SARS-CoV-2 to infect Vero E6 cell line, a monkey kidney epithelial cell line used to investigate the SARS-CoV-2 replication (Fig. 8). We have investigated the molecular mechanisms involved in preventing the binding of SARS-CoV2 virus to its receptor identifying the fatty acid binding pocket in the RBD as the putative binding site of pelargonidin.

In summary we shown that the natural flavonoid, Pelargonidin, reverses Ace2 induction in rodent models of inflammation and this effect is mediated by AHR. In addition, the Pelargonidin might reduce SARS-CoV-2/ACE2 interaction affecting the virus uptake and replication. These data highlight the potential of natural agents in reducing inflammation and invasion by SARS-CoV-2.

CRediT authorship contribution statement

Michele Biagioli: Conceptualization, Investigation, Data curation, Writing - original draft. Silvia Marchiano: Investigation, Data curation. Rosalinda Roselli: Investigation, Data curation. Cristina Di Giorgio: Investigation, Data curation. Rachele Bellini: Investigation, Data curation. Martina Bordoni: Investigation, Data curation. Anna Gidari: Methodology, Data curation. Samuele Sabbatini: Methodology, Data curation. Daniela Francisc: Conceptualization, Supervision. Bianca Fiorillo: Investigation, Data curation. Bruno Catalonotti: Conceptualization, Writing - review & editing. Eleonora Distrutti: Conceptualization, Writing - review & editing. Adriana Carino: Investigation, Data curation. Angela Zampella: Conceptualization, Writing - review & editing. Stefano Fiorucci: Conceptualization, Writing - original draft, Funding acquisition.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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