Gelatin tannate reduces the proinflammatory effects of lipopolysaccharide in human intestinal epithelial cells

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Background: Gelatin tannate is a mixture of tannic acid and gelatin. Tannic acid has astringent properties, due to its capacity to form protein–macromolecular complexes, as well as antibacterial and antioxidant properties. However, little is known about its anti-inflammatory properties.

Purpose: To evaluate the anti-inflammatory activity of gelatin tannate by quantifying the suppression of key molecules produced during inflammatory events in lipopolysaccharide (LPS)-stimulated human intestinal cells.

Methods: Intercellular adhesion molecule-1 (ICAM-1) expression was determined by Western blot analysis; interleukin-8 (IL-8) and tumor necrosis factor-α (TNF-α) concentrations were measured by enzyme-linked immunosorbent assays in Caco-2 cells 24 hours after treatment with LPS (1 µg/mL) in presence of different concentrations of gelatin tannate.

Results: ICAM-1 is induced on a wide variety of cells by inflammatory stimuli such as LPS. Our results have shown gelatin tannate as a potent inhibitor of ICAM-1 expression in LPS-stimulated Caco-2 cells. IL-8 and TNF-α are important inflammatory mediators, recruiting neutrophils and T-lymphocytes. Together with LPS, adding gelatin tannate at different concentrations induced a dose-dependent inhibition of IL-8 and TNF-α released by Caco-2 cells.

Conclusion: These results suggest that gelatin tannate exerts anti-inflammatory effects by inhibiting the specific cytokines and adhesion molecules involved in several inflammatory disorders.

Keywords: Caco-2, ICAM-1, IL-8, TNF-α.
Whilst they are only secondary metabolites of plants, tannins are mostly water-soluble phenolic compounds, and are well known antioxidants sourced from medicinal plants, foods, and edible fruits. Vegetable tannins have attracted a lot of attention in recent years because their multifunctional properties are beneficial to human health. Their diverse biological properties and potential for disease prevention have been demonstrated through various in vitro and in vivo assays.6

According to their structure, tannins are categorized as falling into three categories: hydrolyzable tannins, condensed tannins, or complex tannins.7 Hydrolyzable tannins are the esters of 3,4,5-trihydroxybenzoic acid (gallic acid). Gallic acid molecules are esterified to a core polyol, and the galloyl groups may be further esterified or oxidatively cross-linked to form more complex structures. Gallotannins, the polygalloyl esters of glucose, are the simplest hydrolyzable tannins.1,2,3,4,6-penta-O-galloyl-β-D-glucose (PGG) is a prototypical gallotannin, and the central compound in the biosynthetic pathway of hydrolyzable tannins.8 Though PGG itself is thought to stimulate the secretion of tumor necrosis factor-α (TNF-α) and interleukin-1β (IL-1β), which are both proinflammatory cytokines, it can also attenuate the stimulating effect of lipopolysaccharide (LPS) when treated together; this effect has been tested in both in vitro and in vivo models.9 Additionally, PGG is believed to suppress the expression of adhesion molecules, including intercellular adhesion molecule-1 (ICAM-1) and vascular cell adhesion molecule-1 (VCAM-1), induced by TNF-α. The expression of TNF-α-induced monocyte chemoattractant protein-1 (MCP-1) has also been shown to be attenuated by PGG.10

Tannic acid is a polymer of gallic acid molecules and glucose, and is hydrolyzed into glucose and gallic or ellagic acid units. The natural antioxidant, antimicrobial, and antiviral activity of hydrolyzable tannins such as tannic acid and epigallocatechin gallate have been reported.11–13

Tannic acid is present in many different plant species and foods, and is considered a generally recognized safe (GRAS) food substance, as well as an official drug in Europe and North America.11,14 As an antioxidant compound, tannic acid was shown to inhibit lipid oxidation and radical-mediated DNA cleavage by scavenging oxygen and oxygen-derived radicals.15,16 In addition, numerous studies have suggested that tannic acid has antimicrobial activity against foodborne pathogens such as *Escherichia coli* and Listeria monocytogenes.17–20 Tannic acid has astringent properties due to its ability to form macromolecular complexes with the proteins to which it binds by means of hydrogen links. Some controlled studies have shown tannins to be effective in the treatment of diarrhea, with significantly better results than placebo, shortening the duration of the disorder with no undesirable effects.21,22 Tannic acids do have several undesirable gastrointestinal effects: they induce digestive symptoms such as nausea and vomiting whilst inhibiting the absorption of iron and other metals. The use of tannin complexes such as gelatin tannate (TAN), which is hydrolyzed to gelatin and tannic acid upon reaching the intestine, prevents gastric lesions caused by tannic acid.

It has been demonstrated recently that TAN, a mixture of tannic acid and gelatin, exerts an antidiarrhoeal effect.23 The efficacy and safety of a similar compound in the treatment of acute diarrhea in pediatric patients were studied by Loeb et al where infants aged 3–21 months with acute diarrhea of bacterial and viral origin randomly received either a tannin-rich carob pod powder or an equivalent placebo for up to 6 days. Participants given the tannate-containing compound showed normalization of defecation, body temperature, and weight, and ceased vomiting much earlier than those who received placebo.21

The aim of the present study was to evaluate the potential anti-inflammatory effect of TAN, through an in vitro model to study the immune response of intestinal epithelial cells. The expression of some inflammatory mediators such as ICAM-1, IL-8 and TNF-α was evaluated in human epithelial colorectal adenocarcinoma (Caco-2) cells stimulated with LPS, a major structural component of Gram-negative bacteria, known as one of the most potent inflammatory mediators in intestinal epithelium.

**Materials and methods**

**Reagents**

Gelatin tannate (Tasectan®; Novintethical Pharma Sagl, Lugano, Switzerland) is composed of tannic acid and gelatin. The gelatin used in the manufacturing of TAN is a type A or acid gelatin with a dissolution pH of 5.5 and a Bloom gel strength of 40 to 60 g (viscosity of 27 mPs).

Gelatin is a pure natural protein, easily digested, with no cholesterol, odor or flavor. Gelatin contains 84%–90% collagen protein, 1%–2% mineral salts, and the rest is water. The protein is composed of 18 amino acids (glycine, 25.5%; alanine, 8.7%; valine, 2.5%; leucine, 3.2%; isoleucine, 1.4%; cysteine, 0.1%; methionine, 1%; phenylalanine, 2.2%; proline, 18%; hydroxyproline, 14%; serine, 0.4%; threonine, 1.9%; tyrosine, 0.5%; aspartic acid, 6.6%; glutamic acid, 11.4%; arginine, 8.1%, lysine, 4.1%; histidine, 0.8%).
Cell culture and treatments
Caco-2 cells were maintained in minimum essential medium (MEM; Sigma-Aldrich, Milan, Italy), supplemented with 10% fetal calf serum, 100 U/mL penicillin, and 100 µg/mL streptomycin, incubated at 37°C within a humidified 95% air and 5% CO₂ atmosphere. The medium was changed every 2 to 3 days.

Caco-2 cells usually reached confluence 3 days after seeding, and differentiated into enterocyte-like cells 18–20 days postconfluence. All experiments were carried out in these fully differentiated cells.

For experiments, 24 hours before they were required, the cells were trypsinized and plated in either 96-well plates (for enzyme-linked immunosorbent assay [ELISA] tests) or 100 mm Petri dishes (for Western blot). Caco-2 cells were stimulated or not (untreated controls) with LPS at a concentration of 1 µg/mL, used to reproduce the mechanisms involved in the pathogenesis of inflammatory processes, in the absence or presence of different concentrations of TAN (1, 10, and 100 µg/mL). After 24-hours’ incubation, each sample was tested for ICAM-1 expression, and for IL-8 and TNF-α release.

Western blot analysis
The expression of ICAM-1 was evaluated by Western blot analysis. Briefly, both the untreated and treated Caco-2 cells were washed twice with icecold phosphate-buffered saline (PBS) and collected with lysing buffer (10 mM Tris-hydrochloride plus 10 mM potassium chloride, 2 mM magnesium chloride, 0.6 mM phenylmethylsulfonyl fluoride, and 1% sodium dodecyl sulfate, pH 7.4). After cooling for 30 minutes at 0°C, the cells were sonicated. A total protein of 20 µg, present in the supernatant, was loaded onto each lane and separated by 4% to 12% Novex Bis-Tris gel electrophoresis (NuPAGE; Invitrogen, Milan, Italy). Proteins were then transferred into nitrocellulose membranes (Invitrogen) in a wet system. Membranes were blocked with Tris-buffered saline containing 0.1% Tween-20 (TBST) and 5% nonfat dry milk at 4°C overnight. Mouse monoclonal anti-ICAM-1 (1:200; Santa Cruz Biotechnologies, Santa Cruz, CA), and mouse monoclonal anti-α-tubulin antibodies (1:5000; Sigma-Aldrich) were diluted in TBST, and the membranes incubated for 24 hours at room temperature. Antibodies were detected with horseradish peroxidase-conjugated secondary antibody, using the enhanced SuperSignal® West Pico Chemiluminescent Substrate (Pierce Chemical Co, Rockford, IL). Bands were measured densitometrically, and their relative density calculated based on the density of the α-tubulin bands in each sample. Values were expressed as arbitrary densitometric units corresponding to signal intensity.

ELISA assay
IL-8 and TNF-α were measured on cell-free supernatants collected 24 hours after treatment with specific sandwich ELISA (Biotrak, EIA System; Amersham Biosciences, Piscataway, NJ). Cell cultures were carried in triplicate for each condition. All assays were performed as specified by the kit manufacturer. A standard curve was created for each assay using known concentrations of IL-8 or TNF-α as provided in the kit. Plates were read at 450 nm using a microplate reader. The amount of cytokines was quantified within each supernatant in duplicate. Results were given as mean (expressed in pg/mL) and standard error of the mean (SEM).

MTT assay
The cytotoxic effect of the experimental substance was evaluated using a cell viability test based on the cleavage of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) by mitochondrial dehydrogenases from metabolically active cells. Cellular growth was determined using the MTT assay on 96-well microplates. The optical density of each well sample was measured with a microplate spectrophotometer reader at 550 nm.

Statistical analysis
All statistical analyses were performed with the statistical software package SYSTAT (v 9.0; Systat Inc, Evanston, IL). Each result was calculated as the mean value and SEM. Statistical significances between paired and unpaired values were performed using the Student’s t-test. \( P < 0.05 \) were considered statistically significant.

Results
ICAM-1 expression was analyzed at 24 hours in Caco-2 cells treated with LPS (1 µg/mL). Western blot analysis revealed no constitutive expression of ICAM-1 protein in Caco-2 cells, whereas incubation of cultures with LPS for 24 hours induced a strong expression of ICAM-1. Addition of TAN at different concentrations together with LPS induced a dose-dependent inhibition of ICAM-1 expression (Figure 1). At the highest dose, TAN blocked ICAM-1 expression by 69%. TAN did not induce ICAM-1 expression as noted by the lack of bands in the lane corresponding to the treatment of cells with the test product.

The expression of both TNF-α and IL-8 was analyzed in Caco-2 cells treated with LPS (1 µg/mL) by ELISA.
assay at 24 hours. Only a negligible amount of TNF-α was produced by unstimulated Caco-2 cells, but TNF-α was markedly induced by LPS (Figure 2). TAN markedly inhibited LPS-induced release of TNF-α: at a lower concentration (1 µg/mL) there was a very significant inhibition (~35%), that further increased at higher (10 and 100 µg/mL) concentrations of TAN (~43% and ~63% respectively) (Figure 2). The effects of TAN on the release of IL-8 were also investigated. Very low amounts of IL-8 were produced by unstimulated cells. By contrast, LPS potently induced IL-8, which was markedly decreased by TAN in a dose-dependent manner (Figure 3). Inhibition of LPS-induced inflammatory mediator expression in Caco-2 cells by TAN was not due to TAN cytotoxicity, as assessed by MTT assay (Figure 4).

**Discussion**

Our results suggest that TAN has a remarkable anti-inflammatory potential due to its strong ability to inhibit inflammatory biomarkers, such as LPS-induced ICAM-1, IL-8, and TNF-α in Caco-2 cells.

The intestinal epithelium is able to secrete cytokines and other mediators of inflammation, to respond to growth factors and cytokines released by lymphoid cells, and to express integrins and HLA class II molecules, which permits direct cell-to-cell communication with inflammatory and immune cells. Evidence indicates that a dysregulation of intestinal mucosal immunity, as often seen in inflammatory bowel conditions, causes an overproduction of inflammatory cytokines and the trafficking of catalyst leukocytes into the bowel, thus leading to an uncontrolled intestinal inflammation.
Proinflammatory cytokines/chemokines produced in the intestine play an important role in the pathogenesis of complications of critical illness. One of these, IL-8, has been shown to be a particularly potent chemoattractant for leukocytes and subsequent inflammation.25 IL-8 initiates the acute inflammatory cascade, and is also an early marker of the inflammatory process; its expression must be controlled to prevent both excessive tissue injury, and damage to local and distal organs.26

ICAM-1 is a cytokine-inducible glycoprotein belonging to the immunoglobulin supergene family. ICAM-1 is constitutively expressed at low levels on a limited number of cell types, including endothelial cells,28,29 but its expression is markedly upregulated on a wide variety of cells by inflammatory cytokines such as IL-1β, TNF-α, and interferon-γ (IFN-γ).30,31 ICAM-1 participates in leucocyte adhesion to activated endothelial cells, T cell/antigen-presenting cells, T cell-to-T cell, and T cell/B cell interactions.32 Several epithelial cell types express ICAM-1, including human colonic adenocarcinoma cell lines and colonic cancer cells in vivo.33 By downregulating the expression of these molecules on the endothelium, their interaction with immune cells will be reduced and inflammatory activity diminished. This mechanism of action may be of importance to induce remission in inflammatory bowel conditions.

TAN is a mixture of tannic acid and gelatin with antidiarrheal effects. TAN also has attributed antibacterial and antioxidant properties, representing an advantage in the empiric use of conventional antibiotics that can contribute to the development of resistance and dysbacteriosis.33,34 Additionally, unlike other antidiarrheal agents such as loperamide,35 it has no effect upon the central nervous system, and no other undesirable effects such as reactive constipation. More recently, a study reported no TAN-related undesirable effects either during or after treatment of acute diarrhea in children with a mean age of 2.5 years (standard deviation, 2.4; median, 1.7 years); the product was also well tolerated.22 This study demonstrates that the mixture of tannic acid and gelatin acts as a potent inhibitor of ICAM-1 (Figure 1)
gene expression whilst also suppressing the production of IL-8 (Figure 2) and TNF-α (Figure 3) in human enterocytes stimulated with LPS. Previous research has shown that these are key mediators in the pathogenesis of inflammatory bowel conditions, and that the reduction of these mediators has been associated with the amelioration of inflammatory conditions.36,37 The present in vitro study suggests that TAN contributes to decrease the expression and production of such inflammatory mediators in Caco-2 cells. It found, in fact, that TAN inhibits ICAM-1 expression in LPS-stimulated cells, and that it probably acts at the transcriptional level, as evidenced by dose-dependent reductions in its levels of expression.

TAN was also found to decrease the release of IL-8 and TNF-α. IL-8 is known to be an important cytokine for controlling leukocyte trafficking, and the recruitment of both neutrophils and T lymphocytes. TNF-α is a cytokine involved in systemic inflammation, and belongs to a group of cytokines that stimulate the acute phase reaction. Dysregulation of TNF-α production has been implicated in a variety of human diseases, including inflammatory bowel disease (IBD).38 The treatment of IBD has advanced in parallel to the improvement of the knowledge of its physiopathology, leading to the development of biological therapies. An example of this kind of treatment is the use of substances that antagonize TNF-α, such as monoclonal antibodies; several studies have demonstrated that inhibition of TNF-α is useful in the treatment of IBD.39,40

It has been reported recently that a member of the tannin family, known as PGG, is an effective immunoregulator of the LPS-mediated TNF-α response in both human peripheral blood mononuclear cells, and in live rats. PGG was shown to possess anti-inflammatory properties, even if PGG itself stimulated the production of TNF-α and IL-1β.41 Here, we demonstrate that TAN exerts its beneficial properties without inducing the production of inflammatory mediators.

Conclusion

In summary, the present results suggest that TAN could be used not only for its antidiarrheal effect, due to astringent properties, but also for the treatment of intestinal disorders such as inflammatory bowel conditions where inflammatory mediators may play a key role. Our data indicates that TAN may be effective, with minimal or no toxicity, in the treatment of intestinal inflammatory conditions.

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Disclosure

The authors declare no conflict of interest in this work.

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