Neutrophilia, gelatinase release and microvascular leakage induced by human mast cell tryptase in a mouse model: Lack of a role of protease activated receptor 2 (PAR2)

Running title: Pro-inflammatory actions of tryptase and the role of PAR2

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Key words: Transgenic/Knockout Mice, Mast Cells/Basophils, Neutrophils, Inflammation, Allergy.

Abbreviations used in this article: MMP, Matrix metalloproteinase; PAR2, Protease-activated receptor 2.

This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1111/cea.13108

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Abstract

Background: Tryptase, the most abundant protease of the human mast cell, has been implicated as a key mediator of allergic inflammation that acts through activation of PAR2.

Objectives: To investigate the contribution of PAR2 in the pro-inflammatory actions mediated by tryptase in a mice model.

Methods: We have injected recombinant human βII-tryptase into the peritoneum of PAR2-deficient and wild-type C57BL/6 mice. After 6, 12 and 24 hours mice were euthanized, peritoneal lavage performed and inflammatory changes investigated.

Results: Tryptase stimulated an increase in neutrophil numbers in the peritoneum, but responses did not differ between PAR2-deficient and wild-type mice. Heat-inactivation of tryptase or pre-incubation with a selective tryptase inhibitor reduced neutrophilia, but neutrophil accumulation was not elicited with a peptide agonist of PAR2 (SLIGRL-NH2). Zymography indicated that tryptase stimulated the release of matrix metalloproteinases (MMP) 2 and 9 in the peritoneum of both mouse strains. Studies involving immunomagnetic isolation of neutrophils suggested that neutrophils represent the major cellular source of tryptase-induced MMP2 and MMP9. At 24 h after tryptase injection there was increased microvascular leakage as indicated by high levels of albumin in peritoneal lavage fluid, and this appeared to be partially abolished by heat-inactivating tryptase or addition of a protease inhibitor. There was no corresponding increase in levels of histamine or total protein. The extent of tryptase-induced microvascular leakage or gelatinase release into the peritoneum did not differ between PAR2-deficient and wild-type mice.

Conclusions: Our findings indicate that tryptase is a potent stimulus for neutrophil accumulation, MMP release and microvascular leakage. Though these actions required an intact catalytic site, the primary mechanism of tryptase in vivo would appear to involve processes independent of PAR2.

Introduction

Tryptase, the most abundant product of human mast cells is emerging as a key mediator of inflammation, as well as being an important marker for mast cell activation in allergic disease. Though relatively few naturally occurring substrates have been identified for this tetrameric serine protease, a range of potent pro-inflammatory actions have been described for tryptase following its administration into animal models, or when it is transferred to cells or tissues. Reports that tryptase can activate protease activated receptor 2 (PAR2) [1-3] have led to suggestions that this G-protein coupled receptor may represent a key cellular target [4-7]. Tryptase can like certain other trypsin proteases cleave PAR2 to expose a ‘tethered ligand’ leading to signal transduction. The idea that PAR2 has a pivotal role in mediating the actions of tryptase has received support from comparisons of the actions of PAR2 agonists with those of tryptase. However, some discrepancies have been reported, raising questions over the degree to which PAR2 activation may be involved.

Transfer of purified human tryptase into the peritoneum [8] or trachea of mice [9], or the skin of guinea pigs [10] or sheep [11] has been found to stimulate massive accumulation of neutrophils or eosinophils, and in some cases prolonged microvascular leakage. Peritoneal injection of the mouse tryptase mMCP-6, is also associated with intraperitoneal neutrophilia [12], while injection with

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mMCP-7, another mouse tryptase, has induced eosinophil accumulation in the peritoneal cavity [9].

In all in vivo models studied, the actions of tryptase have been dependent on an intact catalytic site, and inhibitors of tryptase have shown efficacy in models of asthma in sheep [13], mice [14] and humans [15, 16], as well as in a rat model of colitis [17]. Transfer of peptide agonists of PAR2 such as SLIGRL-NH₂ in vivo has also been associated with induction of microvascular leakage, and neutrophilia [18-20], though paradoxically there have been reports also of anti-inflammatory actions in vivo [21].

Tryptase has been demonstrated to induce profound changes in the behaviour of various cell types including mast cells, neutrophils, eosinophils, endothelial cells, airway smooth muscles, and several cell lines. Tryptase can induce the degranulation of human mast cells [10] and eosinophils [22], induce mitogenic responses and cytokine release from endothelial cells [23], epithelial cells [24], and airway smooth muscle cells [25, 26]. These cell types express PAR2, and in most cases effects similar to those for tryptase have been found to be elicited by PAR2 agonists [22, 27-30]; and cell signalling responses described in some cell models have been similar with both tryptase and PAR2 agonists [31, 32]. A lack of effective PAR2 antagonists has hindered their application in such studies, though the peptide antagonist FSLLRY-NH₂ has been reported to reverse some of the cell signalling responses of tryptase in cells [27, 33]. There have been relatively few direct comparisons between tryptase and other potential PAR2 agonists, but peptide agonists of PAR2 have failed to elicit certain actions of tryptase such as stimulation of IL-8 release from airway smooth muscle cells [34], or the degranulation of lung mast cells [35], and eosinophils [22].

The advent of PAR2 knockout mice with C57BL/6J background (which are apparently phenotypically normal) has provided evidence for a key role for this receptor in models for studying the spread of melanoma metastases [36], and joint swelling in arthritis [37, 38]. Intra-articular injection of tryptase has been reported to stimulate joint swelling and hyperaemia in wild-type, but not in PAR2 knockout mice suggesting that there is a dependence on PAR2 activation, but detailed investigation of cellular changes or other features of inflammation were not examined.

In the present study we have demonstrated that tryptase is able to stimulate inflammatory cell accumulation, microvascular leakage and gelatinase release in the peritoneum of both wild-type and PAR2 deficient mice models. Our findings cast doubt on PAR2 having a key role in mediating these changes.

Materials and Methods

Preparation and purification of tryptase

Tryptase was isolated from a Pichia pastoris expression system for βII-tryptase following protocols similar to those described by Niles et al. [39], with sequential purification by hydrophobic interaction chromatography followed by heparin affinity chromatography. The initial step involved passing the tryptase-rich supernatant down a 25 ml butyl Sepharose (GE Healthcare, Amersham) column at 22°C, washing with Buffer A (10 mM MES, 1 M (NH₄)₂SO₄, 0.5 M NaCl, 10% (v/v glycerol), pH 6.1) and

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eluting with Buffer B (10 mM MES, 0.2 M NaCl, 10% (v/v) glycerol, pH 5.5). Fractions of 5 ml were collected and tryptase activity determined using the chromogenic substrate N-α-benzoyl-DL-arginine p-nitroanilide hydrochloride (BApNA; *vide infra*). Tryptase-rich fractions were passed down a 25 ml heparin–agarose column (Sigma, Gillingham, UK), washing with buffer B and eluting with a 0.2 to 2 M NaCl gradient mixing buffer B with 0 to 75% Buffer C (10 mM MES, 2 M NaCl, 10% (v/v) glycerol, pH 6.1). Fractions (5 ml) with high tryptase activity were eluted between 1.04 to 1.29 M NaCl, and these were concentrated using centrifugal concentrators with 30 kDa cut-off (Merck Millipore, Watford, UK), and injected into a BioSep-Sec-S-3000 size exclusion column employing an HPLC ICS 3000 pump (Dionex /Thermo Fisher, Sunnyvale, CA). Fractions of 0.5 ml were collected and analysed for tryptase activity.

**Characterization of purified tryptase**

SDS-PAGE analysis was performed with a NuPAGE BisTris 4-12% gradient gel (Invitrogen/Thermo Fisher, Inchinnan, UK) under reducing conditions, a single band was observed with a molecular weight of 35 kDa consistent with that of the monomeric form of tryptase. The identity as tryptase was confirmed by western blotting with the tryptase-specific monoclonal antibody AA5. Endotoxin levels as assayed by the chromogenic Limulus Amoebocyte Lysate (LAL) endotoxin assay kit, Toxin Sensor™ from GenScript (Piscataway, NJ) were less than 0.08 EU/1U tryptase in all preparations used in the study.

Tryptase activity was measured by determining cleavage of BApNA spectrophotometrically at 410 nm for 10 minutes at 25°C in a Thermomax microplate reader (Molecular Devices, Wokingham, UK) according a procedure described previously (39). The extinction value (ε) of BApNA was taken as 8800 M⁻¹ cm⁻¹. A colorimetric protein assay using bicinchoninic acid was employed in accordance with the manufacturer’s instructions (Sigma) using bovine serum albumin as standard. The specific activity of the tryptase preparations employed ranged from 9-12.2 U mg⁻¹, where 1 unit was taken as the amount of tryptase that can cleave BApNA to release 1 µmol p-nitroanilide per min at 25°C.

**Animals**

Colonies of mice lacking the PAR2 gene (PAR2⁻/⁻) and the corresponding wild type (PAR⁺/+⁺) were kind gifts from Kowa Company Ltd (Tokyo, Japan). Both colonies were C57BL/6 genetic background and generated as described by Ferrell et al. [40]. The animals were housed and maintained at the University of Strathclyde (courtesy of Professor Robin Plevin) before being transferred to and maintained at the University of Southampton. All of the animals employed weighed between 25 and 35 g, and were housed in standard cages in a temperature-controlled room (from 19 to 24°C) in an SPF environment, with a light cycle of 10 h off and 14 h on. Food and tap water was available *ad libitum*, with stock mice fed RM1 rodent maintenance pellets and breeding mice RM3 rodent breeding pellets (till weaning). Bedding material was sawdust with shredded cardboard. There were no more than five animals per cage. Mice were randomly assigned to each treatment group. Mice from different treatment groups were maintained in the same cage. The studies were conducted in accordance with the Animals (Scientific Procedures) Act 1986 (UK), under project

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Three mice tail biopsy samples from each genotype were subjected to genotyping. Tissues was homogenized using a RiboLyser™ homogenizer (Hybaid Ltd., Ashford, UK) in 1.5 ml Lysing Matrix D tubes (MP Biomedicals, Solon, OH) with TRI Reagent (Sigma). Confirmation of presence of the gene for PAR2 was confirmed by performing the PCR using the following primers: 5′-ATGCGAAGTCTCAGCCTGGCG-3′ and 5′-GAGAGGAGGTCGGCCAAGGCC-3′ to yield a 380-bp PCR product. The PCR was generally performed for 35 cycles with initial denaturation at 95°C for 5 min, annealing at 51°C then for 2 cycle/min and extension at 72°C/cycle (3 min), and a final extension (last cycle) at 72°C for 10 min. Absence of the PAR2 gene was tested by detection of neomycin gene using the following primers; NeoFwr 5’GAGGAAGCGGTCAGCCCATT3’ and NeoRev 3’TCTTCCTATTGACTAAACGG5’ with amplicon size of 281bp. The PCR was generally for 33 cycles with initial denaturation 95°C for 10 min, annealing at 68°C for 1 min and 30 s extension at 72°C, and a final extension (last cycle) at 72°C for 10 min. The PCR was concluded by cooling down to 4°C. PCR products were analysed immediately or stored at 4°C until further analysis. PCR products were separated on 2% agarose gel and visualized with ethidium bromide staining.

**Injection of mice**

Animals were injected i.p. as described previously with 0.5 µg tryptase in 0.5 ml 0.9% saline, a quantity selected as it had provoked extensive cell accumulation in this model [8]. The vehicle control was 0.9% saline. Other experimental controls consisted of tryptase that had been heat-inactivated for 20 minutes at 94°C, or incubated with 50 µg ml⁻¹ with selective tryptase inhibitor A (Sanofi-Aventis, Bridgewater, NJ; Ki 39 Nm) [41] for 60 min and the degree of inhibition was assessed by chromogenic substrate BApNA as described above. Also injected were inhibitor A alone, or PAR peptide agonist SLIGRL-NH₂ (1 µg ml⁻¹) or a peptide with the same amino acids but with a ‘scrambled’ sequence (LSIGRL-NH₂; 1 µg ml⁻¹). Tryptase was handled with care to avoid loss of enzymatic activity and was kept on ice and diluted with saline immediately before injection of the animal. The degree of enzyme inhibition was assessed spectrophotometrically using the chromogenic substrate BApNA. Where inadvertent injection or damage to the gut was suspected, such animals were excluded from the study.

**Peritoneal lavage**

At 6, 12 and 24 hours following injection, mice were sacrificed by cervical dislocation. After swabbing the abdominal skin with 70% ethanol, a midline incision of the abdomen was made and 5 ml saline injected into the peritoneum. Abdominal massage was performed for 30 seconds then saline was collected into tubes maintained on ice. Total numbers of nucleated cells were determined using an improved Neubauer haemocytometer with 0.4% trypan blue. The peritoneal lavage fluid was centrifuged at 300g for 10 minutes at 4°C. The supernatant was aspirated and stored at -20°C.

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Cell pellets were re-suspended in MEM with 5% FCS and cell number was adjusted to $10^6$ cell per ml. Cytocentrifuge preparations were made using a Cytospin 3 (Shandon Southern, Runcorn, UK) with 100 µl aliquots of cell suspension and slides left to air dry overnight. One slide for each sample was stained with eosin/methylene blue stain (Rapid Romanowsky, Raymond A Lamb/Thermo Fisher, Loughborough, UK) according to the manufacturer’s instructions. Differential cell counts were performed, counting a minimum of 500 cells per slide in at least 5 fields at a x1000 magnification using a Standard 20 microscope (Carl Zeiss, Göttingen, Germany). Two researchers were involved in collection of samples to facilitate blinding. The initial code for each was randomly assigned a new code prior to analysis of data, and the code broken on statistical analysis.

**Gelatin zymography**

Supernatant from peritoneal lavage fluid was added to a non-reducing sample buffer (4% SDS, 0.125 M Tris-HCl pH 6.8, 0.003% bromophenol blue and 20% glycerol) and applied to an 8% polyacrylamide gel containing 1 mg ml$^{-1}$ gelatin. After electrophoresis, gels were incubated overnight at 37ºC in MMP proteolysis buffer (50 mM Tris-HCl pH 7.8, 0.5 mM NaCl, 50 mM CaCl$_2$), then stained with Coomassie blue dye. Gels were photographed using a Molecular Imager GS-800 calibrated densitometer (Bio-Rad Laboratories, Hemel Hempstead, UK), and intensity and size of the bands determined relative to a positive control for MMP2 and MMP9 activity, a supernatant from the human fibrosarcoma cell line (HT1080) was employed [42].

**Immuno-magnetic purification of neutrophils**

In order to examine the potential contribution of neutrophils to the response of mice to tryptase, peritoneal cells were collected from naïve mice and from mice which had been injected intraperitoneally with tryptase (0.5 µg/mouse) or casein, an established stimulus for neutrophil accumulation, (0.5 ml of 9 % solution/mouse) [43]. The casein for injection was prepared as described previously [44] with 9% casein hydrolysate in PBS pH 7.2 containing 0.9 mM CaCl$_2$ and 0.5 mM MgCl$_2$. After 24 h, peritoneal lavage was performed as described above.

Cells were passed through nylon mesh (30 µm, BD Falcon, Erembodegem, Belgium), cell numbers determined, and the cell suspension centrifuged at 300g for 10 minutes at 4ºC. Cells were re-suspended in 200 µl buffer (0.5% BSA, 2 mM EDTA in PBS, pH 7.2). Biotinylated anti-lymphocyte antigen 6G (LY-6G; Miltenyi Biotec, Bisley, UK) was added, mixed well and incubated for 10 minutes at 4ºC. A 150 µl aliquot of the buffer was mixed well with anti-biotin MicroBeads (Miltenyi Biotec) and incubated for 15 minutes at 4ºC. Cells were washed and re-suspended in the buffer. Labelled cells were collected using a ferromagnetic column (LS column, Miltenyi Biotec), cell number determined and the cell suspension re-suspended in RPMI 1640 conditioned medium and seeded at a density of $10^6$ cells/well in 24 well tissue culture plates (Greiner Bio-One; Stonehouse, UK). Sorted cell populations were treated with tryptase or other compounds, and at 1, 6 or 24 hours, supernatants were collected for gelatin zymography.

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Assays for markers in peritoneal lavage fluid

Quantitative colorimetric albumin bromocresol green (BCG) assay kits (QuantiChrom, BioAssay Systems, Hayward, CA) with a detection range from 0.1 mg ml\(^{-1}\) to 50 mg ml\(^{-1}\) were employed measuring the optical density over the range 570 to 670 nm in a Thermomax plate reader (Molecular Devices). Total protein concentrations were determined using the bicinchoninic acid method referred to above. An automated glass fibre-based technique was employed to determine levels of histamine in peritoneal lavage fluid where samples loaded to pre-coated FiberPlate assay plates (RefLab, Copenhagen, Denmark). Piperazine-1, 4-bis (2-ethanesulphonic acid) (PIPES) buffer was added, plates incubated for 60 minutes at 37°C, and after washing shipped to RefLab, Denmark for analysis. Elastin zymography was carried out according to the procedure described by Forough et al. [45] with samples electrophoresed on a 10-12% SDS-polyacrylamide gel with 1 mg ml\(^{-1}\) soluble κ–elastin under non-reducing conditions, and porcine pancreatic elastase type II-A as standard (both Sigma). After electrophoresis, the gels were washed in 2.5% Triton X-100 for 30 min, and then incubated at 37°C in 50 mM Tris buffer, pH 7.8, containing 10 mM CaCl\(_2\) for 20 h prior to staining with 0.002% Coomassie brilliant blue (G250).

Statistics

The Kruskal-Wallis non-parametric test was used to investigate differences between groups, and if significant, then the Mann-Whitney U test was employed for pre-planned comparisons between two groups of mice. The degree of association between variables was analysed by calculation of Spearman’s coefficient of rank correlation (\(r_s\)). SPSS version 18 software and Graphpad Prism version 6 were used for analysis of data and preparation of graphs. P<0.05 was taken as significant.

Results

Tryptase-induced cell accumulation

Cells in peritoneal lavage fluid that were recovered from saline-injected mice were predominantly macrophages, with smaller numbers of lymphocytes, and very much smaller numbers of other cell types (Table 1; Fig 1). Injection of tryptase into the peritoneum of wild type and PAR2 deficient mice had little effect on the total number of nucleated cells recovered up to 24 h afterwards in peritoneal lavage fluid. Tryptase injection was associated with a substantial increase in neutrophil numbers in wild type mice at 24 h (Table 1; Fig. 1b). A similar degree of neutrophilia was seen in the PAR2 deficient mice. Relative numbers of eosinophils, lymphocytes, macrophages and mast cells appeared to be little affected by injection of tryptase. There were no significant differences between these two mouse strains in the numbers of other cells types in the peritoneum.

As the pattern of cell accumulation in mice of the C57BL/6 strain differed from that reported previously following injection of human lung tryptase in BALB/c mice [8], comparison was made between the responses of C57BL/6 and BALB/c mice injected with recombinant tryptase under the same conditions. Injection of tryptase in the peritoneum of BALB/c mice was associated with

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increases in numbers of neutrophils, eosinophils, lymphocytes, macrophages and mast cells as well as in the total number of cells, compared with those in the saline-injected group (Fig. 1c-g). This contrasted with the findings in the C57BL/6 mice, in which there were no significant increases in numbers of cells other than neutrophils. Total cell numbers as well as numbers of macrophages, mast cells and eosinophils were all higher in the BALB/c mice when compared with those in the C57BL/6 mice. Although the baseline lymphocyte numbers were lower in BALB/c mice, the degree of tryptase-induced accumulation of lymphocytes was greater. On the other hand, tryptase did not induce lymphocyte accumulation in C57BL/6.

Heat-inactivation of tryptase was associated with lower numbers of neutrophils compared with the tryptase-injected groups particularly in the PAR2 deficient mice (Fig. 1i). Tryptase-induced neutrophilia was much less when administered with the selective tryptase inhibitor and a similar trend was seen with leupeptin (data not shown). The peptide agonist of PAR2 SLIGRL-NH$_2$ failed to replicate the actions of tryptase, and the control peptide employed (LSIGRL-NH$_2$) actually stimulated a small increase in numbers of certain cell (Table 1).

**Effect of tryptase on peritoneal gelatinase and elastolytic activity**

Peritoneal lavage fluid from saline-injected mice exhibited activity on gelatin zymography, with bands at positions corresponding to those for MMP2 but not for MMP9 (Fig. 2a). Tryptase injection, on the other hand, was associated with the presence of bands in peritoneal lavage fluid for both MMP2 and MMP9. Levels of MMP2 were higher in mice 24 h following tryptase injection than at 6 and 12 h, and at this time point were significantly greater than in the saline-injected mice (Fig. 2b). At all of these time points MMP9 activity was substantially greater in the tryptase-injected than the saline-injected mice, though levels at 24 h were greater than at 6 or 12 h (Fig. 2c).

Peritoneal lavage fluid MMP2 activity was higher in tryptase-injected mice deficient in PAR2 than in the corresponding wild-type mice at 12 h (p < 0.0001) and 24 h (p < 0.005), but not at 6 h (Fig 2b). For the saline-injected mice, there were no apparent strain-related difference in MMP2 levels. For MMP9 activity there was greater activity in the PAR2 knockout mice than in the wild-type at 12 h (p < 0.005), but not at 6 or 24 h. Mice injected with heat-inactivated tryptase did not show a marked tryptase-induced increase in MMP2 and MMP9 activity in peritoneal lavage fluid (Table 2). Similarly incubation of tryptase with the selective inhibitor A appeared to abolish the ability of tryptase to stimulate an increase in MMP2 and MMP9 activity. Injecting the PAR2 agonist SLIGRL-NH$_2$ and the non-PAR2 activating control peptide had little effect on levels of MMP2 or MMP9 in this model (data not shown).

In peritoneal lavage fluid levels of MMP2 and MMP9 activity were closely correlated in both tryptase-injected wild type mice ($r_s$ = 0.882, n = 33, p<0.005) and PAR2 deficient mice ($r_s$ = 0.975, n = 30 p<0.005). The activity of MMP2 was associated with the numbers of neutrophils recovered from the peritoneum of wild-type ($r_s$ = 0.335, n = 33, p<0.005) and knockout mice ($r_s$ = 0.328, n = 30, p<0.005). Similarly the activity of MMP9 was correlated with the number of neutrophils from both
wild-type ($r_s = 0.452, n = 33, p<0.005$) and knockout mice ($r_s = 0.360, n = 30, p<0.005$). Associations were not observed between MMP2 or MMP9 levels and the numbers of any of the other cell types enumerated. With elastase as a positive control in elastin zymography at quantities as low as 4.5 µg, there were bands at 28 and 22 kDa (data not shown). However, supernatants from peritoneal lavage fluid from mice of both the saline and tryptase-treated groups failed to show any activity.

**Identification of neutrophils as source of gelatinase activity**

Supernatant from a neutrophil-rich cell population (88% neutrophils) obtained from the peritoneum of 10 tryptase-injected PAR2 wild-type mice (by positive selection with magnetic beads) had substantially more MMP9 than that from a neutrophil-depleted cell population (12% neutrophils; Fig. 3). This was observed when cells were incubated *in vitro* for 1, 6 and 24 h. There was little MMP2 activity in supernatants of peritoneal cells at any of these time points. The supernatant of a mixed cell population from casein-injected mice (n = 5) also showed MMP9 activity, but that from naïve mice (n = 3) had negligible quantities (data not shown).

Treatment of either neutrophil-rich or neutrophil-depleted cell populations with 6.5 or 13 µg ml$^{-1}$ (20 or 40 mU ml$^{-1}$) tryptase *in vitro* was not associated with alterations in levels of MMP9 in cell culture supernatants (Fig 3b). The release of MMP9 into cell supernatants thus appeared to be constitutive. As found for peritoneal lavage fluid, no elastase activity was detected by elastin zymography in supernatants from cultured neutrophil-rich or neutrophil-depleted cell populations.

**Albumin, total protein and histamine levels in peritoneal lavage fluid**

The albumin concentration in peritoneal lavage fluid was greater in tryptase-injected mice at 24 h compared with those injected with saline (Table 2). Apparent increase in albumin levels was induced with inactivated tryptase by heating or the selective inhibitor. Differences were not observed between wild-type and PAR2 knockout mice in the concentrations of albumin in lavage fluid. At 6 h or 12 h following injection of tryptase, the albumin levels in the peritoneal lavage fluid were not significantly different from those in the saline-injected mice (data not shown). There was no apparent association between levels of albumin with either histamine concentrations in peritoneal lavage or with numbers of mast cells recovered.

The total protein concentration in peritoneal lavage fluid as detected by BCA binding did not differ between mice injected with tryptase, PAR2 agonist at 24 h, or at 6 h or 12 h (data not shown). There was no association between concentrations of albumin and total protein levels in mouse peritoneal lavage fluid. For both wild-type and PAR2-deficient mice injected with tryptase, at 24 h peritoneal lavage fluid histamine levels were lower than in the saline-injected mice. This effect was not seen with heated tryptase, or at the earlier timepoints with catalytically active tryptase.

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Discussion

The present study confirms and extends the idea that mast cell tryptase has potent pro-inflammatory actions, but suggests that the effects are not mediated through activation of PAR2. Tryptase was a potent stimulus for the accumulation of neutrophils in vivo, and induced MMP release and prolonged microvascular leakage. Despite several studies suggesting that the actions of tryptase are mediated by cleavage of PAR2, at least in the present model, the role tryptase as a mediator of inflammation appears to be independent of PAR2.

Care was taken to ensure that the tryptase employed in this study was of high purity and activity, and on SDS-PAGE recombinant tryptase appeared as a single band whose identity was confirmed by western blotting. The endotoxin levels were extremely low, and our finding that heating the tryptase at 90ºC for 20 min abolished its proteolytic activity, would argue against a role for endotoxin in the inflammatory changes provoked. The finding that the β-tryptase is a potent stimulus for recruitment of neutrophils in vivo, provided confirmatory evidence that tryptase can induce massive neutrophilia in the mouse peritoneum [8]. It is possible, therefore, that β-tryptase represents the variant of tryptase primarily responsible for this action.

This observation of tryptase-induced increases in the numbers of eosinophils, macrophages and lymphocytes as well as of neutrophils (at both 6 h and 16 h) in the peritoneum of BALB/c mice [8] was differ from the present study where we have used C57BL/6 mice (selected on account of the background of the PAR2 deficient mice). However, the recruitment of eosinophils and macrophages, and even of mast cells (a cell type whose presence was not investigated previously) was induced in the present studies by intraperitoneal injection of BALB/c mice with β-tryptase. The discrepancy between studies is thus likely to be related to the mouse strain rather than to the type of tryptase injected. HayGlass et al. have reported a little difference between BALB/c and C57BL/6 in the degree of neutrophilia or eosinophilia when these mice were immunized with Der p1, ovalbumin or human serum albumin (46). On the other hand, Th1 responses have been reported to be more prominent in C57BL/6 mice, and Th2 responses to be stronger in BALB/c mice infected with leishmania [46]. Strain-related differences in the present study are likely to be related to the nature of the inflammatory infiltrate, as adoptive transfer of Th2 cells has been reported to stimulate eosinophilia in a mouse model of ovalbumin challenge, but neutrophilia in mice reconstituted with Th1 cells [47].

Injection of tryptase into the mouse peritoneum was associated with high levels of albumin in peritoneal lavage fluid 24 h after of injection, indicating increased microvascular permeability. Intradermal injection of tryptase has been reported previously to induce microvascular leakage and oedema in the skin of guinea pigs or sheep 20 minutes following injection [10, 11]. The skin reactions were prolonged and still apparent 60 to 80 min following injection, but at least in the guinea pig model had resolved by 6 h. Microvascular leakage was attributed to the ability of tryptase to stimulate release of histamine through activation of mast cells, and this has been demonstrated experimentally in vitro with guinea pig tissue, and subsequently also in human mast cells [35]. A role for mast cells in the microvascular leakage provoked as late as 24 h after tryptase injection in the present study cannot be excluded, but albumin concentrations were not associated with numbers of

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PAR2 [38, 53]. The tetrameric structure of trypase with the catalytic sites positioned within a central pore [54], is likely to restrict access of the extracellular domain of PAR2 as a substrate; and heavy glycosylation of the N-terminal sequence of PAR2 has been reported to prevent trypase-induced receptor activation [55]. Moreover, trypase can cleave PAR2 not just at a site that would result in exposure of the tethered ligand, but also at a point that could ‘disarm’ the receptor [2].

The non-PAR2 mediated processes involved in the present model remain open to conjecture. Certain other proteases have been shown to alter cell behaviour through regulation of growth factor receptors. Thus, hormone-like cellular signaling has been described through the actions of tryptic proteases on receptors themselves (as with insulin receptors or insulin like growth factor-1 receptors) [56], and the release of a membrane tethered agonist e.g. heparin-binding epidermal growth factor (EGF) by metalloproteinases [57]. In theory also a growth factor agonist could be generated from precursors through protolytic activity. The ability of trypase to control the bioavailability of cytokines has been suggested on the basis of the finding that this protease can activate TGFβ, and at least in mice, to inactivate IL-6 from mast cells [26, 58].

In conclusion our findings indicate that trypase is a potent stimulus of inflammation in vivo, and as such deserves attention as a target for therapeutic intervention. At quite low concentrations this protease can stimulate neutrophilia, microvascular leakage and the generation of MMP9 and MMP2. The studies with PAR2 knockout mice challenge the assumption that activation PAR2 represents the primary mechanism by which trypase can mediate inflammatory changes. The actions of trypase were dependent on an intact catalytic site, but in the model employed, the pro-inflammatory actions of trypase appear to be largely independent of PAR2.

Acknowledgement

We are grateful to Kowa Company Ltd (Tokyo, Japan) for providing the PAR2+/+ and PAR2−/− mice and to Professor Robin Plevin, University of Strathclyde for making available these mice from a colony maintained by him, and to Dr Anne Minnich (formerly of Sanofi, Bridgewater, NJ) and Dr Yong-Mi Choi-Sledeski (Sanofi, Cambridge, MA) for allowing use of the selective trypase inhibitor A in this study. The technical assistance provided by the Biomedical Research Facility, Southampton is gratefully acknowledged.

Funding

This work was supported by grants from the Egyptian Ministry of Higher Education, Asthma UK and the Medical Research Council, UK.

Conflict of interest

The authors declare no conflict of interest.

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Author contributions

MMS Khedr and AM Abdelmotelb were responsible for data acquisition and analysis. AF Walls, X Zhou and SLF Pender contributed to the conception and design of the work or of parts of it, and to its interpretation. MMS Khedr and AF Walls drafted and revised the manuscript, and AM Abdelmotelb, SLF Pender and X Zhou revised it critically for intellectual content.

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**Figures’ legend**

**Figure 1** Eosin/methylene blue stained cytocentrifuge preparations of cells recovered by peritoneal lavage from C57BL/6 mice 24 h following injection with (a) tryptase or (b) saline. Examples of neutrophils (N), eosinophils (E), macrophages (M), lymphocytes (L) and mast cells (MC) are indicated. Numbers are indicated of (c) nucleated cells, (d) neutrophils, (e) eosinophils, (f) macrophages, (g) lymphocytes and (h) mast cells recovered from the peritoneum of BALB/c, C57BL/6 PAR2<sup>−/−</sup> and C57BL/6 PAR2<sup>+/+</sup> mice 24 h following injection of tryptase (Tryp; 0.5 µg/mouse). Saline-injected mice (Sal). Also shown are (i) neutrophil numbers in peritoneal lavage fluid from PAR2<sup>−/−</sup> and PAR2<sup>+/+</sup> mice 24 h following injection of tryptase, heated tryptase, tryptase pre-incubated with the selective inhibitor, the inhibitor alone and the saline vehicle. N = 5-13 mice per group. Median values and inter-quartile range are shown.* P<0.05, ** P<0.005, (Mann-Whitney U test). n.s., not significant.
Figure 2 (a) Gelatin zymography showing the presence of MMP2 and 9 in a standard (Std) supernatant from the HT1080 fibrosarcoma cell line, and supernatants from peritoneal lavage fluid from mice 24 h after injection of saline alone, or tryptase (mice 1 to 3). (b) MMP2 and (c) MMP9 activity in peritoneal lavage fluid from PAR2\(-/-\) and PAR2\(+/+\) mice 6, 12 and 24 h following intraperitoneal injection of tryptase (Tryp; 0.5 µg/mouse). Median values and inter-quartile range are shown. * P<0.05 ** P<0.005 *** P<0.0001, (Mann-Whitney U test). n.s., not significant.

Figure 3 (a) Gelatin zymography illustrated for neutrophil-rich and neutrophil-depleted peritoneal lavage cell populations isolated from 8 mice 24 h following tryptase injection. Culture supernatants were incubated for 1 h with tryptase at 13 µg ml\(^{-1}\) (40 mU ml\(^{-1}\)), heat-inactivated tryptase (of the same original concentration), or medium alone. Bands for molecular weight (MW) markers are also shown. (b) Relative MMP9 activity in culture supernatants of neutrophil-rich and neutrophil-depleted cell populations maintained in culture for 1, 6 or 24 h (n = 6 for each group). Median and inter-quartile range are indicated. * P<0.05, ** P<0.005, (Mann-Whitney U test) compared with activity in the neutrophil-depleted cell population.
Table 1: Total and differential cell counts in peritoneal lavage fluid recovered from mice 24 h following injection of saline vehicle, 0.5 µg tryptase, 0.5 µg SLIGRL-NH₂, or 0.5 µg LSIGRL-NH₂ in wild-type and PAR2 deficient mice.

<table>
<thead>
<tr>
<th>Compounds injected</th>
<th>Mouse strain</th>
<th>Total cells</th>
<th>Neutrophils</th>
<th>Eosinophils</th>
<th>Lymphocytes</th>
<th>Macrophages</th>
<th>Mast cells</th>
</tr>
</thead>
<tbody>
<tr>
<td>Saline</td>
<td>PAR2+/+</td>
<td>3.60 (3.07 - 4.59)</td>
<td>0.02 (0 - 0.06)</td>
<td>0.07 (0.03 - 0.13)</td>
<td>0.69 (0.50 - 1.01)</td>
<td>2.41 (2.24 - 3.33)</td>
<td>0.03 (0.02 - 0.06)</td>
</tr>
<tr>
<td></td>
<td>PAR2−/−</td>
<td>4.81 (3.01 - 5.70)</td>
<td>0.01 (0 - 0.11)</td>
<td>0.07 (0.05 - 0.28)</td>
<td>0.92 (0.12 - 1.14)</td>
<td>3.23 (1.71 - 4.80)</td>
<td>0.06 (0.02 - 0.07)</td>
</tr>
<tr>
<td>Tryptase</td>
<td>PAR2+/+</td>
<td>4.11 (2.45 - 7.32)</td>
<td>0.45** (0.10 - 1.95)</td>
<td>0.08 (0.04 - 0.33)</td>
<td>0.70 (0.37 - 0.92)</td>
<td>2.45 (1.67 - 4.49)</td>
<td>0.03 (0.01 - 0.05)</td>
</tr>
<tr>
<td></td>
<td>PAR2−/−</td>
<td>3.96 (3.21 - 5.52)</td>
<td>0.80** (0.49 - 0.92)</td>
<td>0.06 (0.03 - 0.18)</td>
<td>0.63 (0.29 - 0.86)</td>
<td>3.02 (2.21 - 3.77)</td>
<td>0.01 (0.01 - 0.03)</td>
</tr>
<tr>
<td>SLIGRL-NH₂</td>
<td>PAR2+/+</td>
<td>4.02 (3.33 - 4.88)</td>
<td>0.03* (0.01 - 0.06)</td>
<td>0.18* †† (0.09 - 0.43)</td>
<td>0.36 (0.28 - 1.09)</td>
<td>2.80 (2.20 - 3.85)</td>
<td>0.03 (0.0 - 0.06)</td>
</tr>
<tr>
<td></td>
<td>PAR2−/−</td>
<td>4.22 (3.22 - 5.67)</td>
<td>0.07†† (0.01 - 0.28)</td>
<td>0.17 (0.09 - 0.32)</td>
<td>0.45 (0.17 - 1.99)</td>
<td>3.47 (1.10 - 4.83)</td>
<td>0.03† (0.02 - 0.10)</td>
</tr>
<tr>
<td>LSIGRL-NH₂</td>
<td>PAR2+/+</td>
<td>3.57 (2.95 - 5.18)</td>
<td>0.28** (0.12 - 0.48)</td>
<td>0.35** †† (0.19 - 0.66)</td>
<td>0.19** †† (0.14 - 0.31)</td>
<td>2.96 (2.19 - 3.63)</td>
<td>0.02* (0.01 - 0.03)</td>
</tr>
<tr>
<td></td>
<td>PAR2−/−</td>
<td>6.31** † (5.39 - 10.1)</td>
<td>0.02†† (0.01 - 0.07)</td>
<td>0.03 (0.02 - 0.12)</td>
<td>0.30 (0.20 - 0.51)</td>
<td>5.78** †† (4.85 - 9.17)</td>
<td>0.20** †† (0.16 - 0.21)</td>
</tr>
</tbody>
</table>

Median values (x10⁶) are shown for 10-12 mice in each group. *P<0.05 ** P<0.005, compared with responses in the saline-injected group; and †P<0.05, ††P<0.005, compared with the responses in the tryptase-injected group.
Table 2  MMP2 and MMP9 activity (Arbitrary units), albumin levels (mg ml\(^{-1}\)), total protein concentration (µg ml\(^{-1}\)) and histamine levels (ng ml\(^{-1}\)) in peritoneal lavage fluid from mice 24 h following injection of the saline vehicle, tryptase, heated tryptase, tryptase pre-incubated with the selective inhibitor and the inhibitor alone in PAR2\(^{+/+}\) and PAR2\(^{-/-}\) mice.

<table>
<thead>
<tr>
<th>Mouse strain</th>
<th>Saline</th>
<th>Tryptase alone</th>
<th>Heated tryptase</th>
<th>Inhibitor alone</th>
<th>Tryptase + inhibitor</th>
</tr>
</thead>
<tbody>
<tr>
<td>MMP2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PAR2 (^{+/+})</td>
<td>2.2 (2.0 - 2.4)</td>
<td>3.3** (2.4 - 4.8)</td>
<td>2.4†† (2.2 - 2.5)</td>
<td>1.9†† (1.8 - 2.2)</td>
<td>1.9†† (1.7 - 2.5)</td>
</tr>
<tr>
<td>PAR2 (^{-/-})</td>
<td>2.0 (1.9 - 2.0)</td>
<td>5.3** (5.0 - 6.4)</td>
<td>2.3†† (2.1 - 2.3)</td>
<td>2.1†† (1.9 - 2.3)</td>
<td>1.7†† (1.7 - 1.8)</td>
</tr>
<tr>
<td>MMP9</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PAR2 (^{+/+})</td>
<td>ND</td>
<td>4.9** (3.3 - 5.6)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>PAR2 (^{-/-})</td>
<td>ND</td>
<td>4.7** (4.3 - 5.1)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Albumin</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PAR2 (^{+/+})</td>
<td>2.5 (2.0 - 2.8)</td>
<td>8.0** (5.4 - 11)</td>
<td>8.9** (8.2 - 9.0)</td>
<td>12†† (11 - 13)</td>
<td>8.0* (6.1 - 8.6)</td>
</tr>
<tr>
<td>PAR2 (^{-/-})</td>
<td>2.1 (1.8 - 2.7)</td>
<td>11** (8.6 - 12)</td>
<td>8.0†† (8.2 - 8.4)</td>
<td>11 (3.7 - 12)</td>
<td>8.5†† (9.0 - 11)</td>
</tr>
<tr>
<td>Total protein</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PAR2 (^{+/+})</td>
<td>510 (395 - 547)</td>
<td>200** (54.0 - 368)</td>
<td>319* (193 - 520)</td>
<td>393†† (361 - 442)</td>
<td>524†† (439 - 611)</td>
</tr>
<tr>
<td>PAR2 (^{-/-})</td>
<td>389 (315 - 460)</td>
<td>541 (244 - 906)</td>
<td>528* (455 - 628)</td>
<td>484 (368 - 580)</td>
<td>283† (242 - 322)</td>
</tr>
<tr>
<td>Histamine</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PAR2 (^{+/+})</td>
<td>133 (79 - 159)</td>
<td>54.5** (44.5 - 99.0)</td>
<td>115 (49.2 - 253)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>PAR2 (^{-/-})</td>
<td>254 (126 - 268)</td>
<td>47.8** (16.8 - 80.9)</td>
<td>265†† (233 - 269)</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Median and inter-quartile range values are shown for 10-12 mice each group. * P<0.05 ** P<0.005 *** P<0.0001, compared with response in the saline-injected group. † P<0.05 †† P<0.005, compared with response in the tryptase-injected group. ND, none detected; -, not tested
Figure 1

(a) (b) (c) (d) (e) (f) (g) (h) (i)
Figure 2

(a) 

(b) 

(c) 

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Figure 3

(a) Gelatin zymography showing MMP9 and MMP2 activity in neutrophil-rich and neutrophil-depleted cell populations. (b) Table summarizing MMP9 activity (arbitrary units) in different conditions.

<table>
<thead>
<tr>
<th>Cell population</th>
<th>MMP9 activity (arbitrary units)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1h</td>
</tr>
<tr>
<td>Neutrophil-rich</td>
<td>5.04** (4.34 - 7.51)</td>
</tr>
<tr>
<td>Neutrophil-depleted</td>
<td>0.86 (0.34 - 1.42)</td>
</tr>
</tbody>
</table>