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Effects of intranasal TNF α on granulocyte recruitment and activity in healthy subjects and patients with allergic rhinitis

Henrik Widegren*¹, Jonas Erjefält², Magnus Korsgren³, Morgan Andersson¹ and Lennart Greiff¹

Address: ¹Department of Otorhinolaryngology, Head & Neck Surgery, Lund University Hospital, Lund, Sweden, ²Department of Experimental Medical Sciences, Lund University, Lund, Sweden and ³Department of Clinical Pharmacology, Lund University Hospital, Lund, Sweden

Email: Henrik Widegren* - henrik.widegren@skane.se; Jonas Erjefält - jonas.erjefalt@med.lu.se; Magnus Korsgren - magnus.korsgren@astrazeneca.com; Morgan Andersson - morgan.andersson@skane.se; Lennart Greiff - lennart.greiff@skane.se

* Corresponding author

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Abstract

Background: TNF α may contribute to the pathophysiology of airway inflammation. For example, we have recently shown that nasal administration of TNF α produces late phase co-appearance of granulocyte and plasma exudation markers on the mucosal surface. The objective of the present study was to examine indices of granulocyte presence and activity in response to intranasal TNF α challenge.

Methods: Healthy subjects and patients with allergic rhinitis (examined out of season) were subjected to nasal challenge with TNF α (10 μ g) in a sham-controlled and crossover design. Nasal lavages were carried out prior to and 24 hours post challenge. Nasal biopsies were obtained post challenge. Nasal lavage fluid levels of myeloperoxidase (MPO) and eosinophil cationic protein (ECP) were analyzed as indices of neutrophil and eosinophil activity. Moreover, IL-8 and α_2 -macroglobulin were analyzed as markers of pro-inflammatory cytokine production and plasma exudation. Nasal biopsy numbers of neutrophils and eosinophils were monitored.

Results: Nasal lavage fluid levels of MPO recorded 24 hours post TNF α challenge were increased in healthy subjects ($p = 0.0081$) and in patients with allergic rhinitis ($p = 0.0081$) (c.f. sham challenge). Similarly, α_2 -macroglobulin was increased in healthy subjects ($p = 0.014$) and in patients with allergic rhinitis ($p = 0.0034$). Lavage fluid levels of ECP and IL-8 were not affected by TNF α challenge. TNF α increased the numbers of subepithelial neutrophils ($p = 0.0021$), but not the numbers of eosinophils.

Conclusion: TNF α produces a nasal inflammatory response in humans that is characterised by late phase (i.e., 24 hours post challenge) neutrophil activity and plasma exudation.

Background

Tumour necrosis factor- α (TNF α) is an important pro-inflammatory cytokine of the immune system. Dysregu-

lated TNF α responses are implicated in several inflammatory diseases including rheumatoid arthritis. In this condition, TNF α antagonism, either with recombinant

soluble receptors or neutralising antibodies, improves disease activity scores [1].

In human bronchial airways, TNF α is produced by several cell types including macrophages [2], eosinophils [3,4], epithelial cells [5,6], and mast cells [7,8]. Inhalation of TNF α produces a response that is characterized by increased sputum numbers of granulocytes and increased airway responsiveness [9,10]. Furthermore, in refractory asthma, treatment with etanercept, an anti-TNF α measure, is associated with improvement in asthma symptoms, lung function, and airway hyperresponsiveness [11,12]. While these observations suggest an association between TNF α and bronchial airway inflammation, little is known about TNF α -induced nasal airway responses in man.

Focusing on human nasal airways, patients suffering from allergic rhinitis have been shown to feature increased tissue expression of TNF α mRNA and increased nasal mucosal output of TNF α [13-15]. Furthermore, in a preliminary study, we have shown that while nasal challenge with TNF α fails to produce acute effects, it produces late-phase plasma exudation (i.e., extravasation and luminal entry of plasma) [16], a response characteristic of ongoing inflammation [17]. We have also observed that this response may be associated with increased nasal mucosal output of granulocyte mediators [16]. While our observations suggest a role for TNF α in upper airway conditions characterised by inflammation, information on the nature of this response is scarce. Further information in this field is warranted for several reasons: it may be used to determine whether or not anti-TNF measures should be evaluated for upper respiratory tract conditions. Furthermore, studies in allergic rhinitis may elucidate specific roles for TNF α in type-1 allergic inflammation, which may not be done in allergic asthma where type-1 features may be blurred by chronic inflammatory features.

The aim of the present study was to examine characteristics of the inflammatory process evoked by human nasal administration of TNF α . Nasal challenges with TNF α were carried out and the response was monitored by nasal lavages and biopsies and by analysis of granulocyte activation and plasma exudation markers as well as tissue numbers of granulocytes. Based on our previous study [16], we focused the evaluation on an observation point 24 hours post challenge. Healthy subjects and patients with allergic rhinitis (examined out of season) were included in order to get a preliminary estimation regarding whether or not these group of subjects differ with regard to TNF α -induced effects.

Methods

Study Design

The present study involved nasal administration of a single dose of TNF α (10 μ g) in a single blind, sham-controlled (isotonic saline), randomized, and crossover design with a two-week washout period. Nasal lavages were carried out prior to and 24 hours post challenge. Furthermore, nasal mucosal biopsies were obtained 24 hours post-challenge. Lavage fluid levels of myeloperoxidase (MPO), eosinophil cationic protein (ECP), α_2 -macroglobulin, and IL-8 were measured as indices of neutrophil activity, eosinophil activity, plasma exudation, and pro-inflammatory cytokine production, respectively. Numbers of neutrophils and eosinophils in the biopsy material were determined. Nasal symptoms were scored at strategic time-points (below).

Subjects

Thirteen healthy subjects (5 females, 8 males, aged 20–28 years) and 14 patients with seasonal allergic rhinitis (8 females, 6 males, aged 23–45 years) participated in the study. The protocol was approved by the Regional Ethics Committee and written informed consent was obtained. The study was carried out in compliance with the Helsinki Declaration. For healthy subjects, inclusion criteria were: negative skin prick-test, normal nasal examination, and use of contraceptives (for females). Exclusion criteria were: history of respiratory tract infection within seven days of the start of the study, allergic and non-allergic rhinitis, other nasal disease (structural abnormalities, rhinosinusitis, and polyposis), use of decongestants within seven days of the start of the study, and use of other treatments (except occasional analgesics) within one month of the start of the study. For patients with allergic rhinitis, inclusion and exclusion criteria were the same as described above except that these subjects presented a history of at least two years of seasonal allergic rhinitis and a positive skin-prick test to relevant allergens (patients with positive tests for perennial allergens were excluded). Accordingly, eleven of the allergic subjects were skin-prick positive for both birch and timothy grass allergen, whereas one was sensitized to birch alone and one to timothy grass alone. None of the individuals had positive tests for house dust mite. Eight subjects had positive skin-prick tests for either cat or dog dander, but none of them were exposed to these animals on a regular basis.

TNF α challenge

Recombinant human TNF α (210-TA/CF, R&D Systems, Abingdon, UK) was diluted with isotonic saline to a 100 μ g/ml concentration. The solution was prepared less than one hour before the challenge and it was administered using a spray-device delivering 100 μ l per actuation. The dose of 10 μ g of TNF α was based on experience from our previous study [16]. Isotonic saline was used for the sham

challenge. The challenges were given in a randomized order. The right hand side nasal cavity was challenged at the first visit and the left hand side at the second visit. The reason for using both nasal cavities was to avoid any influence of the first biopsy on the second challenge and sampling procedure. A washout period of two weeks was instituted between the challenges to make the repeated biopsy procedure more tolerable.

Symptom scores

Nasal symptoms were scored by the subjects prior to TNF α challenge and 24 hours thereafter. Sneezes, secretion, and blockage were each scored on a four-graded scale: 0 for no symptoms, 1 for mild symptoms, 2 for moderate symptoms, and 3 for severe symptoms. The scores were added to a total nasal symptom score (range 0–9).

Nasal lavages

Nasal lavages with isotonic saline were carried out using a pool-device [18]. The volume of the pool-fluid was 15 ml and it was kept in the nasal cavity for 5 min at each occasion. The lavages were performed immediately after the symptom scoring, before the challenge and before the nasal biopsy at 24 hours. The recovered fluid was centrifuged and the supernatant was homogenized, prepared in aliquots, and frozen (-30°C) for later analysis. The results were expressed as weight of analyte per volume without correction for any dilution of the lavage fluid: correction for dilution may be of little importance when high volume lavages are employed.

Nasal biopsies

The biopsy was obtained immediately after the second nasal lavage. Topical anaesthesia was applied using a solution of lidocaine hydrochloride (34.0 mg/ml) and nafazoline (0.17 mg/ml), delivered first by a spray-device and thereafter by a cotton swab. The biopsies were taken from the inferior aspect of the inferior turbinate about 0.5 cm from its anterior margin using a cutting-punch forceps with a drilled out punch (2 mm in diameter). Haemostasis was achieved using diathermy.

Analysis

α_2 -Macroglobulin was measured using a RIA sensitive to 7.8 ng/ml. Rabbit anti-human α_2 -macroglobulin (Dako, Copenhagen, Denmark) was used as anti-serum and human serum (Behringwerke, Marburg, Germany) as standard. Human α_2 -macroglobulin (Capell-Organon, Turnhout, Belgium) was iodinated. Tracer and standard or sample was mixed with antiserum before adding goat anti-rabbit anti-serum (AstraZeneca, Lund, Sweden). The bound fraction was measured using a gamma counter. The intra- and inter-assay coefficients of variation are between 3.8–6.0% and 3.1–7.2%, respectively. IL-8 was

measured using a commercially available enzyme-linked immunosorbent assay (R&D Systems, Abingdon, UK). ECP and MPO were also measured using commercially available assays (Phadia, Uppsala, Sweden and Calbiochem, Darmstadt, Germany, respectively). The limits of detection for IL-8, ECP and MPO were 31.2 pg/ml, 2.0 and 1.6 ng/ml, respectively.

The biopsies were immersed overnight in buffered 4% paraformaldehyde (pH 7.2), dehydrated in increasing concentrations of ethanol, embedded in paraffin, and sliced in 3 μ m sections. After rinsing in TRIS-buffered saline, the sections were incubated with primary monoclonal antibodies overnight at 4°C. The monoclonal antibody EG2 (Phadia, dilution 1:200), which detects the eosinophil granule protein ECP, was used to identify eosinophils. Neutrophils were detected by a monoclonal antibody against the neutrophil granule protein MPO (Dako, A390, dilution 1:8000). After incubation with primary antibodies specimens were rinsed in TBS buffer and incubated with a secondary antibody (rabbit anti-mouse, Dako Z0109, dilution 1:80) for 40 min at room temperature. After washing in TBS the sections were incubated in mouse APAAP (Dako D0651, dilution 1:50 in 20% NHS) for 30 min, rinsed, and developed using New Fuchsin (Dako) as substrate. Endogenous alkaline phosphatase activity was inhibited by Levamisol (Dako). All sections were counter-stained with Harris's haematoxylin, coated with Aqua Perm mountant (484975 Life Sci. International), dried overnight, and mounted in Kaiser's glycerol gelatin (Merck, Darmstadt, Germany). The sections were blinded and the numbers of ECP-positive (eosinophils) and MPO-positive (neutrophils) subepithelial cells were counted down to a depth of 210 μ m below the epithelial basement membrane. The numbers of eosinophils and neutrophils were expressed as cells/mm² of tissue compartment.

Statistics

Differences in nasal lavage fluid levels of MPO, ECP, IL-8, and α_2 -macroglobulin as well as biopsy numbers of neutrophils and eosinophils, between observations following TNF α and sham challenge, were analyzed using the Wilcoxon signed rank test. A p-value less than 0.05 was considered significant. Data is presented as mean \pm S.E.M.

Results

One patient with allergic rhinitis was excluded from the study because of airway infection. Otherwise, all individuals complied with the protocol. Nasal administration of TNF α did not produce any nasal symptoms: score 0.8 \pm 0.3 post sham challenge and score 0.8 \pm 0.2 post TNF α (range 0–9).

Nasal lavage fluid levels of MPO recorded 24 hrs post TNF α challenge were increased in healthy subjects ($p = 0.0081$) and in patients with allergic rhinitis ($p = 0.0081$) (c.f. sham challenge) (Fig. 1). Furthermore, α_2 -macroglobulin was increased in both these study groups ($p = 0.014$ and $p = 0.0034$, respectively) (Fig. 2), indicating plasma exudation. ECP and IL-8 were not affected by TNF α challenge (Figs. 3 and 4).

TNF α increased the numbers of subepithelial neutrophils in the study material as a whole ($p = 0.0021$) (Fig. 5): a subgroup analysis revealed that this change reflected a statistically significant increase in the healthy group ($p = 0.0015$), but not in the allergic rhinitis group ($p = 0.204$) (Fig. 6). Nasal biopsy numbers of eosinophils were unaffected by the TNF α challenge (Fig. 7). (Note, a non-significant difference 24 hours post sham challenge between healthy subjects and patients with allergic rhinitis also for eosinophils ($p = 0.06$, Mann Whitney).) Fig. 8 shows micrographs of nasal mucosal neutrophils and eosinophils 24 hours post TNF α and sham challenge, respectively.

Discussion

The present study, involving healthy subjects and patients with allergic rhinitis, demonstrates that intranasal TNF α produces a local inflammatory response. Our observation, together with our previous findings [16], suggests that this effect predominantly involves recruitment and activation of neutrophils and a low-grade exudative response 24 hours post challenge. The observation is of interest in terms of nasal airway defence/physiology and in terms of whether or not TNF α should be viewed as a pro-inflammatory factor in respiratory conditions characterised by inflammation.

The dose of TNF α and the time-point for monitoring of the present response was chosen based on our previous observation: the 10 μg dose may be low, but has been shown to produce inflammatory effects 24 hours post challenge [16]. In the present study, the plasma exudation producing effect of TNF α (10 μg) at this time-point was confirmed. Taken together, our observations suggest that TNF α may not *per se* produce acute plasma exudation in human nasal airways (like, e.g., histamine would), but can induce an inflammatory condition that is characterised by "late-phase" plasma exudation. Plasma exudation is a specific feature of inflammation and increased levels of plasma proteins in airway mucosal surface liquids characterize allergic rhinitis and upper respiratory tract infections [19,20]. Accordingly, and supported by observations on increased levels of TNF α in allergic rhinitis [13,14] as well as common cold [21,22], TNF α may be a pro-inflammatory factor in inflammatory conditions of the nasal airway.

Nasal lavage fluid levels of MPO were increased 24 hours post TNF α administration in healthy subjects and in patients with allergic rhinitis (this study). The finding is in agreement with our previous observation in allergic rhinitis, although that particular increase in MPO only reached borderline statistical significance [16]. Moreover, in the present study, nasal biopsy numbers of neutrophils, in the study group as a whole, were significantly increased post TNF α challenge further indicating a neutrophil-active effect of this cytokine. Our finding is in agreement with observations by Thomas *et al.* [9,10], who reported sputum neutrophilia in healthy subjects as well as in patients with mild asthma following TNF α inhalation. Also in agreement, observations *in vitro* have identified TNF α as a factor that attracts [23] and, potentially, activates [24] neutrophil granulocytes. Taken together, avail-

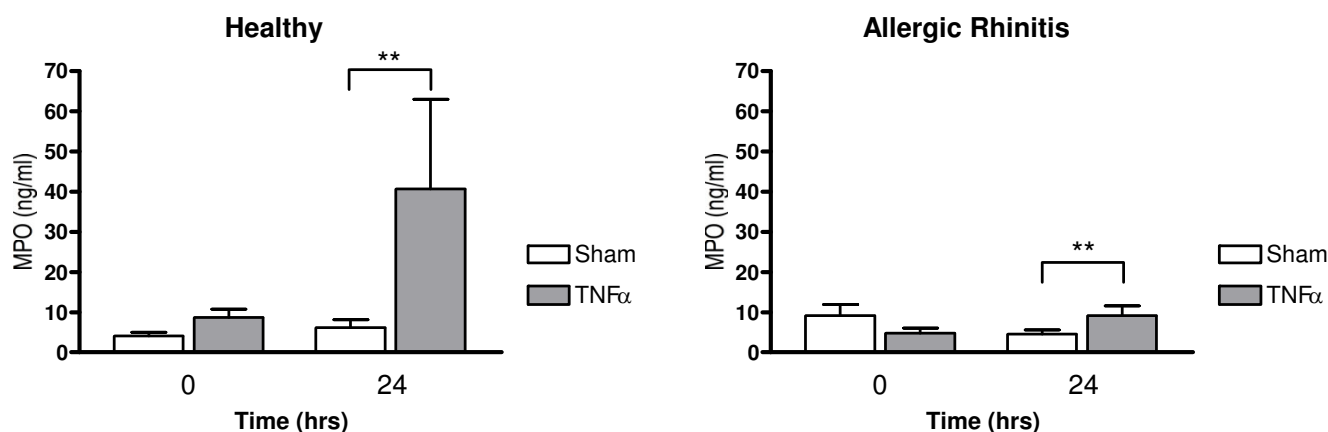


Figure 1

Nasal lavage fluid levels of MPO prior to and 24 hours post TNF α and sham challenge. (** $p < 0.01$).

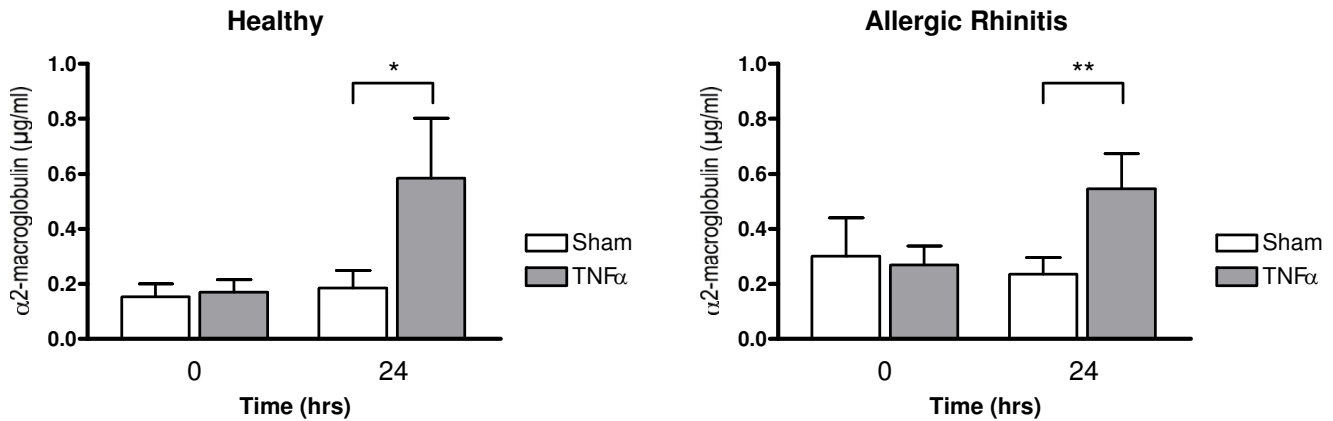


Figure 2
Nasal lavage fluid levels of α_2 -macroglobulin prior to and 24 hours post TNF α and sham challenge. (*p < 0.05, **p < 0.01).

able information now indicates that TNF α can be involved in neutrophil recruitment and activity in human airways. We have no specific explanation for the present numerically higher levels of MPO observed in healthy subjects (*c.f.* patients with allergic rhinitis) following TNF α challenge. However, this difference is not statistically significant ($p = 0.5$, Mann Whitney).

In our previous study, involving patients with allergic rhinitis examined out of season, we observed moderately increased lavage fluid levels of ECP 24 hours after nasal TNF α challenge [16]. In contrast, in the present study, this observation could not be confirmed. We have no specific explanation for the discrepant findings, but a variable eosinophil activity at baseline may be one option: Thomas *et al.* observed that inhalation of TNF α to mild asthmatics increased eosinophil infiltration [9] whereas this

was not the case in healthy subjects [10]. In the present study, the biopsy analysis, which provides an additional aspect of cellular activity, showed that TNF α did not produce recruitment of eosinophils. Accordingly, our overall conclusion is that TNF α administered to patients with allergic rhinitis out of season, when there may be some degree of nasal eosinophilia but not at all as marked as during season allergen exposure, does not produce any marked pro-eosinophil effects. Further studies are warranted to examine effects of TNF α on eosinophil activities, tentatively involving patients with on-going allergic rhinitis.

The present study was not powered to discriminate between healthy subjects and patients with allergic rhinitis, but may allow for a preliminary comparison. The only difference in TNF α induced effects between the groups

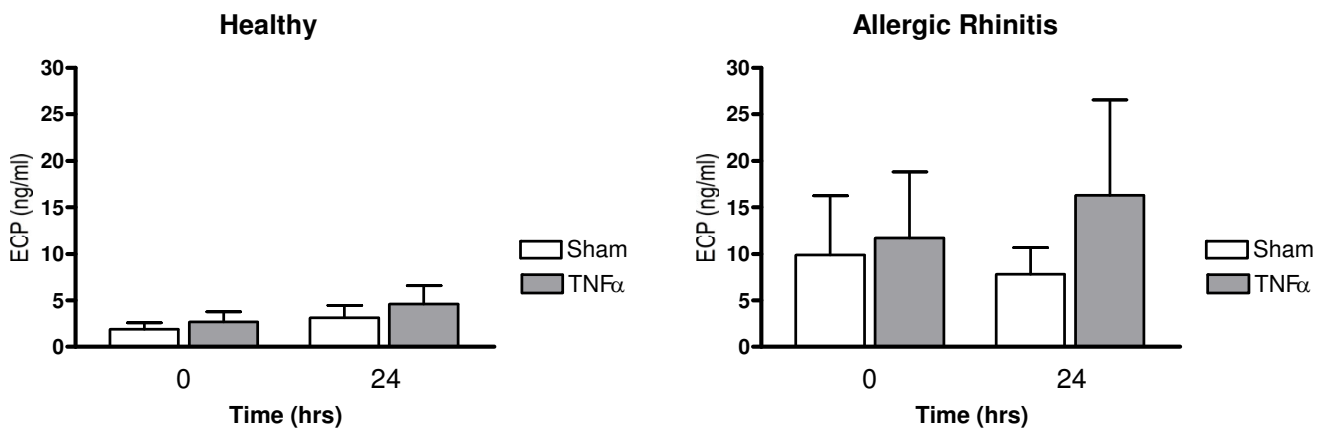


Figure 3
Nasal lavage fluid levels of ECP prior to and 24 hours post TNF α and sham challenge.

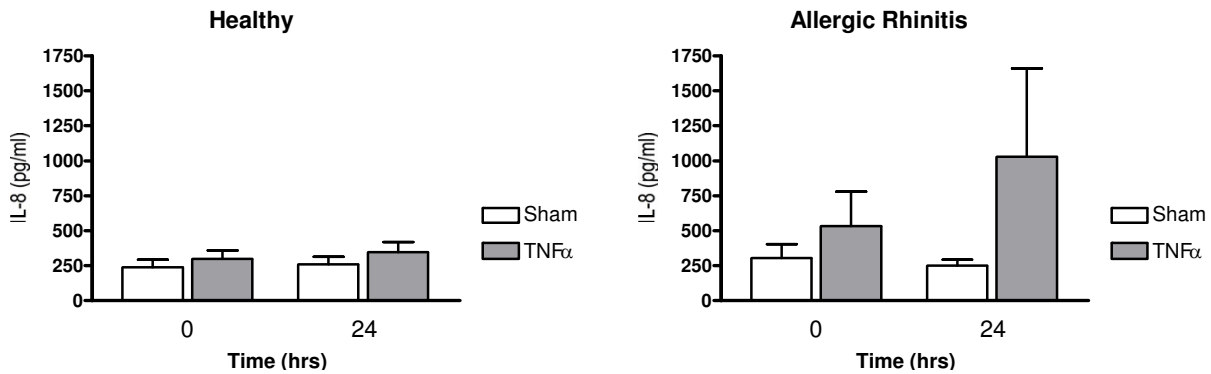


Figure 4
Nasal lavage fluid levels of IL-8 prior to and 24 hours post TNF α and sham challenge.

was that the nasal mucosal numbers of neutrophils were increased in healthy subjects following TNF α challenge but not in patients with allergic rhinitis. However, TNF α did produce a numerical increase in neutrophils also in allergic rhinitis, which did not reach statistical significance (possibly due to a somewhat higher baseline). Furthermore, MPO, as marker of neutrophil activity, was significantly increased in both study groups. Taken together, no marked differences emerged between healthy subjects and patients with allergic rhinitis regarding TNF α -induced effects. However, further studies may be warranted in order to elucidate whether TNF α has a specific role in allergic airway conditions: such studies are warranted based on observations that patients with allergic rhinitis feature increased tissue expression of TNF α mRNA and TNF α protein [13-15]. Moreover, a basis for further exploration may be the recent observation that mice sensitized to ovalbumin and subjected to gene ther-

apy (soluble TNF α receptor IgGfc fusion gene) present decreased symptoms and attenuated aggregation of mast cells, eosinophils, and IL-5 positive cells following nasal ovalbumin challenge [25]. Tentatively, further human *in vivo* studies may involve intervention with anti-TNF α measures in allergic rhinitis at seasonal allergen exposure.

Anti-TNF α drugs, including etanercept, have emerged as a possible treatment option for refractory asthma [11,12]: reduction of symptoms, improvement in lung function, and reduction of bronchial hyperresponsiveness as a result of anti-TNF α intervention. A similar use of anti-TNF α drugs in allergic rhinitis is in our opinion not likely, in part reflecting that existing treatment options are very good and in part inherited difficulties with current anti-TNF α approaches. However, the human nasal airway and allergic rhinitis can be used to monitor effects of TNF α (this study) and anti-TNF α drugs. This may be particularly valid since there are key similarities between inflammatory processes of the upper and lower airways, since observations made in upper airway conditions frequently can

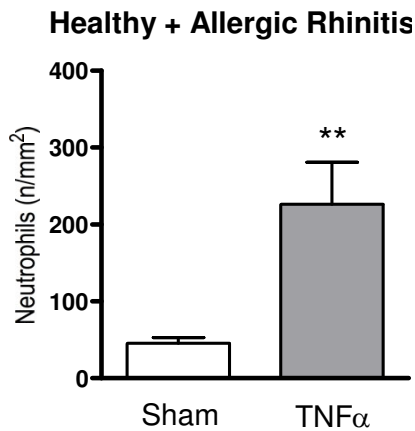


Figure 5
Numbers of subepithelial neutrophils in nasal biopsies obtained 24 hours post TNF α and sham challenge in the study material as a whole. (**p < 0.01).

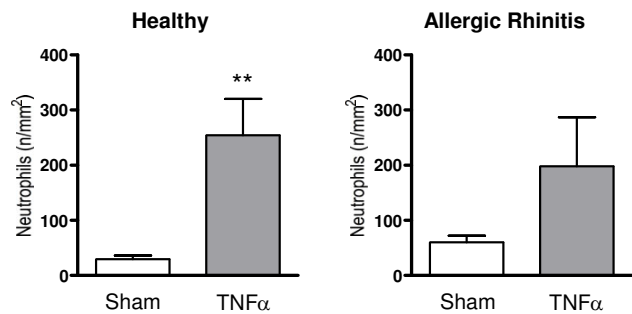


Figure 6
Numbers of subepithelial neutrophils in nasal biopsies obtained 24 hours post TNF α and sham challenge in healthy subjects and patients with allergic rhinitis. (**p < 0.01).

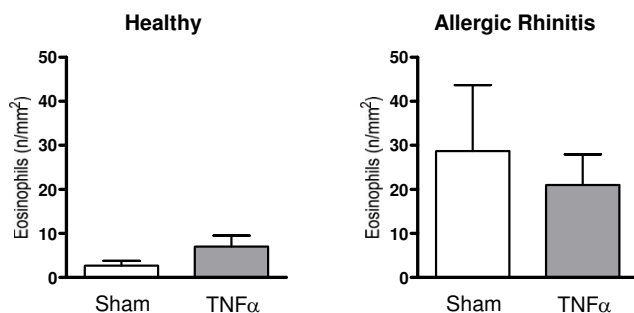


Figure 7
Numbers of subepithelial eosinophils in nasal biopsies obtained 24 hours post TNF α and sham challenge in healthy subjects and patients with allergic rhinitis.

be translated to the bronchial airway diseases, and since nasal airway studies are easier to perform. In this context, the present findings suggest, indirectly, that beneficial effects of anti-TNF α drugs in refractory asthma may reflect anti-neutrophil rather than anti-eosinophil effects.

Conclusion

Topical TNF α produces a nasal inflammatory response in humans that potentially is mediated through increased neutrophil activity.

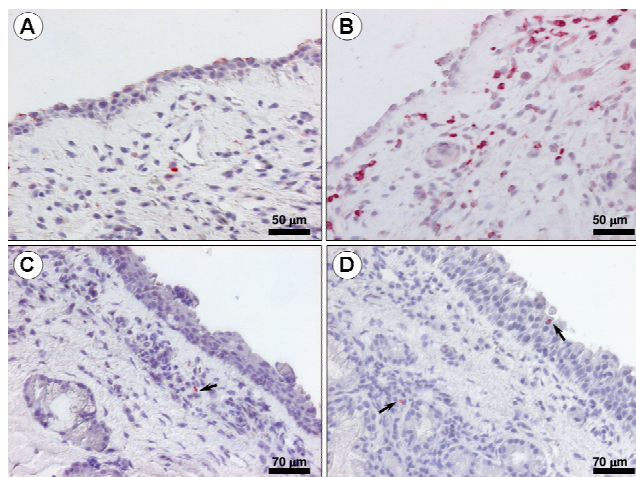


Figure 8
Micrographs showing nasal mucosal neutrophils and eosinophils 24 hours post TNF α and sham challenge. A: Sham-challenged healthy subject: biopsy stained for the neutrophil granule protein MPO. B: TNF α challenged patient with allergic rhinitis: biopsy stained for MPO. C: Sham-challenged healthy subject: biopsy stained for the eosinophil granule protein ECP. D: TNF α challenged healthy subject: biopsy stained for ECP. Nasal biopsy numbers of neutrophils were increased 24 hours after TNF α challenge.

Competing interests

The author(s) declare that they have no competing interests.

Authors' contributions

All authors participated in the design process of the study. HW, MK, MA, and LG carried out the challenge, lavage/biopsy, and lavage analysis procedures. HW and JE performed the immunohistochemistry. HW and LG drafted the manuscript. All authors read and approved the final manuscript.

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References

- Olsen NJ, Stein CM: **New drugs for rheumatoid arthritis.** *N Engl J Med* 2004, **350(21)**:2167-2179.
- Gosset P, Tsicopoulos A, Wallaert B, Joseph M, Capron A, Tonnel AB: **Tumor necrosis factor alpha and interleukin-6 production by human mononuclear phagocytes from allergic asthmatics after IgE-dependent stimulation.** *Am Rev Respir Dis* 1992, **146(3)**:768-774.
- Costa JJ, Matossian K, Resnick MB, Beil WJ, Wong DT, Gordon JR, Dvorak AM, Weller PF, Galli SJ: **Human eosinophils can express the cytokines tumor necrosis factor-alpha and macrophage inflammatory protein-1 alpha.** *J Clin Invest* 1993, **91(6)**:2673-2684.
- Finotto S, Ohno I, Marshall JS, Gauldie J, Denburg JA, Dolovich J, Clark DA, Jordana M: **TNF-alpha production by eosinophils in upper airways inflammation (nasal polyposis).** *J Immunol* 1994, **153(5)**:2278-2289.
- Ackerman V, Marini M, Vittori E, Bellini A, Vassalli G, Mattoli S: **Detection of cytokines and their cell sources in bronchial biopsy specimens from asthmatic patients. Relationship to atopic status, symptoms, and level of airway hyperresponsiveness.** *Chest* 1994, **105(3)**:687-696.
- Devalia JL, Campbell AM, Sapsford RJ, Rusznak C, Quint D, Godard P, Bousquet J, Davies RJ: **Effect of nitrogen dioxide on synthesis of inflammatory cytokines expressed by human bronchial epithelial cells in vitro.** *Am J Respir Cell Mol Biol* 1993, **9(3)**:271-278.
- Bradding P, Roberts JA, Britten KM, Montefort S, Djukanovic R, Mueller R, Heusser CH, Howarth PH, Holgate ST: **Interleukin-4, -5, and -6 and tumor necrosis factor-alpha in normal and asthmatic airways: evidence for the human mast cell as a source of these cytokines.** *Am J Respir Cell Mol Biol* 1994, **10(5)**:471-480.
- Gordon JR, Galli SJ: **Mast cells as a source of both preformed and immunologically inducible TNF-alpha/cachectin.** *Nature* 1990, **346(6281)**:274-276.
- Thomas PS, Heywood G: **Effects of inhaled tumour necrosis factor alpha in subjects with mild asthma.** *Thorax* 2002, **57(9)**:774-778.
- Thomas PS, Yates DH, Barnes PJ: **Tumor necrosis factor-alpha increases airway responsiveness and sputum neutrophilia in normal human subjects.** *Am J Respir Crit Care Med* 1995, **152(1)**:76-80.
- Howarth PH, Babu KS, Arshad HS, Lau LC, Buckley MG, McConnell W, Beckett P, Ali MA, Chauhan A, Wilson SJ, et al.: **Tumour Necrosis Factor (TNF{alpha}) as a novel therapeutic target in symptomatic corticosteroid-dependent asthma.** *Thorax* 2005.
- Berry MA, Hargadon B, Shelley M, Parker D, Shaw DE, Green RH, Bradding P, Brightling CE, Wardlaw AJ, Pavord ID: **Evidence of a role of tumor necrosis factor alpha in refractory asthma.** *N Engl J Med* 2006, **354(7)**:697-708.
- Bensch GW, Nelson HS, Borish LC: **Evaluation of cytokines in nasal secretions after nasal antigen challenge: lack of influ-**

- ence of antihistamines. *Ann Allergy Asthma Immunol* 2002, **88(5)**:457-462.
14. Bradding P, Mediawake R, Feather IH, Madden J, Church MK, Holgate ST, Howarth PH: **TNF alpha is localized to nasal mucosal mast cells and is released in acute allergic rhinitis.** *Clin Exp Allergy* 1995, **25(5)**:406-415.
 15. Ying S, Robinson DS, Varney V, Meng Q, Tsicopoulos A, Moqbel R, Durham SR, Kay AB, Hamid Q: **TNF alpha mRNA expression in allergic inflammation.** *Clin Exp Allergy* 1991, **21(6)**:745-750.
 16. Widegren H, Korsgren M, Andersson M, Greiff L: **Effects of TNFalpha on the human nasal mucosa in vivo.** *Respir Med* 2007, **101(9)**:1982-1987.
 17. Persson CG, Erjefalt JS, Greiff L, Andersson M, Erjefalt I, Godfrey RW, Korsgren M, Linden M, Sundler F, Svensson C: **Plasma-derived proteins in airway defence, disease and repair of epithelial injury.** *Eur Respir J* 1998, **11(4)**:958-970.
 18. Greiff L, Pipkorn U, Alkner U, Persson CG: **The 'nasal pool' device applies controlled concentrations of solutes on human nasal airway mucosa and samples its surface exudations/secretions.** *Clin Exp Allergy* 1990, **20(3)**:253-259.
 19. Svensson C, Andersson M, Persson CG, Venge P, Alkner U, Pipkorn U: **Albumin, bradykinins, and eosinophil cationic protein on the nasal mucosal surface in patients with hay fever during natural allergen exposure.** *J Allergy Clin Immunol* 1990, **85(5)**:828-833.
 20. Akerlund A, Greiff L, Andersson M, Bende M, Alkner U, Persson CG: **Mucosal exudation of fibrinogen in coronavirus-induced common colds.** *Acta Otolaryngol* 1993, **113(5)**:642-648.
 21. Corne JM, Lau L, Scott SJ, Davies R, Johnston SL, Howarth PH: **The relationship between atopic status and IL-10 nasal lavage levels in the acute and persistent inflammatory response to upper respiratory tract infection.** *Am J Respir Crit Care Med* 2001, **163(5)**:1101-1107.
 22. Klemens C, Rasp G, Jund F, Hilgert E, Devens C, Pfrogner E, Kramer MF: **Mediators and cytokines in allergic and viral-triggered rhinitis.** *Allergy Asthma Proc* 2007, **28(4)**:434-441.
 23. Kyan-Aung U, Haskard DO, Poston RN, Thornhill MH, Lee TH: **Endothelial leukocyte adhesion molecule-1 and intercellular adhesion molecule-1 mediate the adhesion of eosinophils to endothelial cells in vitro and are expressed by endothelium in allergic cutaneous inflammation in vivo.** *J Immunol* 1991, **146(2)**:521-528.
 24. Shalaby MR, Aggarwal BB, Rinderknecht E, Svedersky LP, Finkle BS, Palladino MA Jr: **Activation of human polymorphonuclear neutrophil functions by interferon-gamma and tumor necrosis factors.** *J Immunol* 1985, **135(3)**:2069-2073.
 25. He J, Wang T, Yao L, Chen A, Zhou B, Yu H, Jia R, Cheng C, Huan L, Zhang Z: **Construction and delivery of gene therapy vector containing soluble TNFalpha receptor-IgGFc fusion gene for the treatment of allergic rhinitis.** *Cytokine* 2006, **36(5-6)**:296-304.

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