The IL-4/IL-13 signaling axis promotes prostatic fibrosis

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Abstract

Background

Lower urinary tract symptoms (LUTS) are a costly and pervasive medical problem for millions of aging men. Recent studies have showed that peri-urethral tissue fibrosis is an untreated pathobiology contributing to LUTS. Fibrosis results from excessive extracellular matrix deposition which increases transition zone and peri-urethral tissue stiffness and compromises prostatic urethral flexibility and compliance, producing urinary obstructive symptoms. Inflammatory cells, including neutrophils, macrophages, and T-lymphocytes, secrete a medley of pro-fibrotic proteins into the prostatic microenvironment, including IFN<sub>γ</sub>, TNF<sub>α</sub>, CXC-type chemokines, and interleukins, all of which have been implicated in inflammation-mediated fibrosis. Among these, IL-4 and IL-13 are of particular interest because they share a common signaling axis that, as shown here for the first time, promotes the expression and maintenance of IL-4, IL-13, their cognate receptors, and ECM components by prostate fibroblasts, even in the absence of immune cells. Based on studies presented here, we hypothesize that the IL-4/IL-13 axis promotes prostate fibroblast activation to ECM-secreting cells.

Methods

N1 or SFT1 immortalized prostate stromal fibroblasts were cultured and treated, short- or long-term, with pro-fibrotic proteins including IL-4, IL-13, TGF-β, TNF-α, IFN<sub>γ</sub>, with or without prior pre-treatment with antagonists or inhibitors. Protein expression was assessed by immunohistochemistry, immunofluorescence, ELISA, immunoblot, or Sircoll assays. Transcript expression levels were determined by qRT-PCR. Intact cells were counted using WST assays.

Results

IL-4Rα, IL-13Rα1, and collagen are concurrently up-regulated in human peri-urethral prostate tissues from men with LUTS. IL-4 and IL-13 induce their own expression as well as that
of their cognate receptors, IL-4Rα and IL-13Rα1. Low concentrations of IL-4 or IL-13 act as cytokines to promote prostate fibroblast proliferation, but higher (>40ng/ml) concentrations repress cellular proliferation. Both IL-4 and IL-13 robustly and specifically promote collagen transcript and protein expression by prostate stromal fibroblasts in a JAK/STAT-dependent manner. Moreover, IL-4 and IL-13-mediated JAK/STAT signaling is coupled to activation of the IL-4Rα receptor.

Conclusions

Taken together, these studies show that IL-4 and IL-13 signal through the IL-4Rα receptor to activate JAK/STAT signaling, thereby promoting their own expression, that of their cognate receptors, and collagen. These findings suggest that the IL-4/IL-13 signaling axis is a powerful, but therapeutically targetable, pro-fibrotic mechanism in the lower urinary tract.

Introduction

Lower urinary tract symptoms (LUTS) are a costly and potentially critical medical problem for millions of aging men. This spectrum disorder encompasses symptoms such as weak stream, nocturia, incomplete emptying and intermittent urination, all of which are indicative of lower urinary tract dysfunction (LUTD). If left untreated or treated ineffectively, LUTD can progress to bladder dysfunction, which can lead to urinary retention, recurrent UTI, bladder calculi, and, eventually, renal impairment [1–4]. LUTD is often, although not always, concomitant with BPH—a proliferative but nonmalignant enlargement of the prostate. Surgical ablation of prostate tissue and medical approaches including 5α-reductase inhibitors, α-adrenergic receptor antagonists, and PDE5 inhibitors improve urinary flow, but such treatments are not effective for all men, can produce adverse effects that result in discontinuation of the therapeutic regimen, and do not abrogate the risk for disease progression [1, 5]. The latter point bears attention, as recent work evaluating Medical Therapy of Prostatic Symptoms (MTOPS) transition zone biopsies showed that high collagen levels consistent with fibrotic changes in tissue architecture were significantly higher in peri-urethral tissues from men whose LUTS progressed under 5α-reductase inhibitor and α-adrenergic receptor antagonist combination therapy.

Previous work by the Marberger group reported that inflammatory infiltrate in transurethral (TUR) prostate tissues from men demonstrating clinical progression of LUTS comprised ~70% CD4+ (helper) T cells and ~15% macrophages [6]. They also reported that the secretory pattern of these CD4+ T-helper cells was consistent with that of both T-helper Type 1 (Th1) cells, which secrete IFNγ and IL-2, and Type 2 (Th2) cells, which secrete IL-4 and IL-13 [6, 7]. All of these inflammatory mediators are pro-fibrotic proteins [6–8]. Torrko et al. reported that high levels of infiltrating CD4+ T-helper cells (p = .002) and CD68+ macrophages (p = .01) in transition zone prostate biopsies sampled as part of the MTOPS study were significantly associated with clinical BPH progression in that study [9]. Taken together, these studies show that Th2 cells and macrophages are highly prevalent in peri-urethral prostate tissues of men who demonstrate progressive LUTS.

Other studies have shown that peri-urethral tissue fibrosis is an untreated pathobiology contributing to LUTD [10–15]. Fibrosis results from excessive extracellular matrix (ECM) deposition which increases transition zone and peri-urethral tissue stiffness and compromises prostatic urethral flexibility and compliance, producing urinary obstructive symptoms and
Fibrosis is consequent to chronic or unresolved inflammation associated with aging, metabolic syndrome, or urinary tract infection. Inflammatory cells, including neutrophils, macrophages, and Th2 cells, secrete a medley of pro-fibrotic proteins into the prostatic micro-environment, including some of those described above: IFN\(\gamma\), TNF\(\alpha\), CXC-type chemokines, and interleukins [6–8]. Together, these studies suggest that high levels of T-cell and macrophage inflammatory infiltrate are coincident with elevated prostate stromal collagen secretion in the periurethral region of the prostate in men suffering from progressive LUTS.

Among the inflammatory mediators secreted by Th2 cells, IL-4 and IL-13 are of particular interest because they share a common JAK/STAT6 signaling axis that promotes fibrillar collagen secretion and accumulation in multiple organ systems (reviewed in [16], most notably in the lung [17, 18], the kidney [19, 20], and the liver [17, 21]. Based on the studies described above, we hypothesized that IL-4 and IL-13 may contribute to peri-urethral prostate ECM accumulation and voiding dysfunction in the lower urinary tract. The studies reported here show that IL-4 and IL-13 promote their own expression, that of their cognate receptors, and ECM components, even in the absence of immune cells. These studies suggest that the IL-4/IL-13 signaling axis is a powerful, but therapeutically targetable, pro-fibrotic mechanism in the lower urinary tract.

Results

IL-4R\(\alpha\) and IL-13R\(\alpha\)1 are up-regulated in peri-urethral prostate with high collagen content

We have previously shown that high levels of collagen accumulation are associated with peri-urethral prostatic tissue fibrosis and lower urinary tracts symptoms (LUTS) [10, 22]. To determine whether collagen levels correlated with IL-4 and/or IL-13 receptor levels, peri-urethral tissue sections from 6 patients, 3 previously characterized with low 4.4–6.5% collagen content (patient specimens 1, 2, 3, Fig 1) and 3 with high 10.4–17.5% collagen content (patient specimens 4, 5, 6, Fig 1) [10] were subjected to immunohistochemistry to examine IL-4R\(\alpha\), IL-13R\(\alpha\)1 and IL-13R\(\alpha\)2 expression levels (S1 Fig). IL-4R\(\alpha\) (\(p<.01\)) (Fig 1A–1D) and IL-13R\(\alpha\)1 (\(p<.01\)) (Fig 1E–1H) expression levels were significantly higher in tissues with high compared to low collagen content. Expression levels for the decoy receptor, IL-13Ra2, were heterogeneous across the 6 patient sample set. These data show that IL-4R\(\alpha\) and IL-13R\(\alpha\)1 are significantly up-regulated concurrent with high collagen content in peri-urethral prostate tissues.

Low doses of IL-4 and IL-13 enhance cellular proliferation

IL-4 and IL-13 can act as cytokines to promote the proliferation of T-lymphocytes and multiple other cell types [23]. To determine whether these interleukins promoted fibroblast proliferation, N1 and SFT1 prostate fibroblasts and primary lung fibroblasts (HLF) (for comparison) were cultured in serum-free defined media supplemented with vehicle or increasing doses of human recombinant IL-4 and/or IL-13 (1–100 ng/ml) for 24 hours. These studies showed that both IL-4 (Fig 2A) and IL-13 (Fig 2B) significantly promoted prostate fibroblast proliferation at low concentrations, with maximal proliferation of 1.3–1.5X over untreated cells observed at 20 ng/ml (S2 Fig). Higher interleukin doses repressed proliferation. Interestingly, primary human lung fibroblasts achieved maximal proliferation of 2X at higher levels of IL-4 or IL-13 (40ng/ml, Fig 2A and 2B) (S2 Fig). Moreover, primary lung fibroblasts promoted maximal levels of cellular proliferation over a wide IL-13 concentration range spanning 40–80ng/ml (Fig 2B) (S2 Fig). These studies show that low, potentially physiological, levels of IL-4 or IL-13 promoted low-level proliferation of prostate fibroblasts, and that prostate fibroblasts were less
responsive to the stimulatory effect of IL-4 and IL-13 compared to primary human lung fibroblasts.

To test the specificity of the IL-4/IL-13 induced proliferative response, cells were pre-treated with increasing doses of antibody against the IL-4Rα or IL-13Rα1 receptor, then stimulated with the concentration of IL-4 or IL-13 that produced the highest proliferative response. IL-4 failure to induce proliferation above basal levels (1-fold proliferation) was observed using 400 ng anti-IL-4Rα (Fig 2C) (S2 Fig), whereas IL-13 failure to induce proliferation above basal levels (1-fold proliferation) was observed using 40 ng anti-IL-13Rα1 (Fig 2D) (S2 Fig). These data suggest that IL-4 solicits a stronger, or perhaps more sustained, proliferative response.

Fig 1. IL-4Ra and IL-13Ra1 are up-regulated in peri-urethral prostate with high collagen content. Tissue sections of peri-urethral prostate tissues from 6 patients were deparaffinized and assessed for collagen content using Masson’s Trichrome Stain [10] or for IL-4R, IL-13Ra1, and IL-13Ra2 protein expression by immunohistochemistry. Tissues from patients 4, 5 and 6 demonstrated significantly higher collagen contents than those from patients 1, 2 and 3 (p < .05) and significantly higher IL13Ra1 (p < .01) and IL4Rα (p < .01) protein expression levels. IL-4Ra staining is shown for patients 2 (A, B) and 4 (C, D), and IL-13Ra1 staining also for patients 2 (E, F) and 4 (G, H). Panels B, D, F and H show the Immunoratio analysis of panels A, C, E and G, respectively, where green indicates incomplete/weak (0, 1+) and red indicates complete/strong (2+, 3+) staining (S1 Fig).
than IL-13, as 10X higher IL-4Rα than IL-13Rα1 antibody levels were required to repress this response. Alternatively, this result may be due to differences in ligand/receptor affinities.

**IL-4 and IL-13 induce self-expression in prostate fibroblasts**

The IL-4/IL-13 signaling axis up-regulates and activates the STAT6 and GATA-3 transcription factors, which then promote their own expression as well as that of the IL-4 and IL-13 interleukins and their cognate receptors, IL-4Rα and IL-13Rα1 in immune cells (reviewed in [24]). Therefore, we investigated whether IL-4 and/or IL-13 promoted their own secretion by prostate fibroblasts. For these experiments, N1 or SFT1 prostate fibroblasts were cultured in serum-free media supplemented with vehicle or 20ng/ml IL-4, IL-13, IFN-γ, TNFα, or 4ng/ml...
TGF-β, for 24 hours with or without 2 hr pre-treatment with 400 ng/ml IL-4Rα or 40 ng/ml IL-13Rα1 antibodies, then switched to fresh serum-free media and incubated for another 24 hrs. Conditioned media was collected and interrogated by ELISA for IL-4 or IL-13 protein secretion. These studies showed that IL-4 induced its own secretion in both N1 and SFT1 cells, and that this effect was ablated upon pretreatment with IL-4Rα antibody (Fig 3A and 3C). Parallel studies showed that IL-13 also induced its own secretion in N1 and SFT1 cells, which was ablated upon pre-treatment with IL-13Rα1 antibody (Fig 3B and 3D). IL-4 and IL-13 induced their own expression at levels significantly higher than that achieved by IFN-γ, TNFα, or TGFβ (Fig 3) (S3 Fig). Taken together, these data show that IL-4 and IL-13 can induce their own production by prostate fibroblasts. Moreover, whereas IL-4 induction was fairly specific, IL-13 induction was promoted by itself as well as other pro-fibrotics.

IL-4 and IL-13 up-regulate collagen expression

IL-4 and IL-13 are pro-fibrotic proteins that can promote the secretion of ECM proteins [25]. We therefore investigated whether IL-4 and/or IL-13 could induce the expression of the COL1A1 gene, which encodes 2 of the 3 strands of the collagen 1 protein trimer, in prostate fibroblasts. As seen in Fig 4A, IL-4 and IL-13 significantly increased COL1A1 gene expression to levels 3-4x higher than vehicle-treated N1 and SFT1 cells (p < .0001). TGF-β induced higher COL1A1 expression than IL-4 or IL-13, to levels 4-5x higher than vehicle-treated cells (p < .0001). Pre-treatment with antibodies against IL-4Rα or IL-13Rα1 ablated the observed IL-4 or IL-13-mediated COL1A1 transcriptional response, respectively, demonstrating the specificity of these responses (S4 Fig). Subsequent studies examined whether increases in COL1A1 transcript expression were mirrored by collagen protein expression. Sircol assays, which detect total soluble collagens I-V, demonstrated significantly higher levels of soluble collagens I-V produced by N1 and SFT-1 cells treated with IL-4 (p < .001) (B) or IL-13 (p < .0001) (C) compared to vehicle-treated cells. Pre-treatment with antibodies against IL-4Rα or IL-13Rα1 ablated the observed IL-4 or IL-13-mediated collagen protein expression to levels similar to those of vehicle-treated cells (p < .05). As expected, TGF-β induced higher levels of soluble collagens I-V than either IL-4 or IL-13. Co-treatment with IL-4 + IL-13 significantly increased production of soluble collagens I-V above those observed for IL-14 alone by N1 (p < .0001) and SFT-1 (p < .01) cells (Fig 4B). These data show that both IL-4 and IL-13 robustly and specifically promote collagen transcript and protein expression by prostate stromal fibroblasts to levels almost equivalent to those induced by the strong pro-fibrotic protein, TGF-β (S4 Fig).

IL-4 and IL-3 do not promote αSMA expression

N1 cells treated with TGF-β1, IL-4, or IL-13 demonstrated significantly increased expression of COL1A1 protein in vitro compared to vehicle-treated cells. (Fig 5A–5D). Pre-treatment with antibodies against the cognate receptors of IL-4 and IL-13, respectively, repressed their induction of COL1A1 protein expression (Fig 5A–5D), demonstrating the specificity of the cellular response to interleukin treatment. αSMA expression, however, was significantly up-regulated by treatment with TGF-β1, but not with IL-4 or IL-13 (Fig 5A–5C and 5E) (S5 Fig). Moreover, N1 cells treated with TGF-β1 exhibited a morphological change from a spindle-like shape to a more stellate shape consistent with the myofibroblast phenotype (Fig 5C) [11], whereas cells treated with IL-4 or IL-13 retained a spindle-like morphology (Fig 5A–5C), suggesting that IL-4 and IL-13 induced fibroblast activation, but not complete differentiation to a myofibroblast phenotype.
Fig 3. IL-4 and IL-13 induce self expression in prostate fibroblasts. N1 or SFT1 prostate fibroblasts were cultured in serum-free media supplemented with vehicle or 20ng/ml IL-4, IL-13, IFN-γ, TNFα, or 4ng/ml TGF-β, for 24 hours with or without 2 hr pre-treatment with 400 ng/ml IL-4R α or 40 ng/ml IL-13R α antibody. The cells were then washed, switched to fresh serum-free media, and incubated for another 24 hrs. Conditioned media was collected and interrogated by ELISA for IL-4 or IL-13 protein secretion (N1 cells, A and B; SFT1 cells, C and D) (S3 Fig), respectively. Statistically significant differences between IL-4 or IL-13 secretion by vehicle-treated compared to other treatments is indicated as * p < .05; ** p < .01; *** p < .001; **** p < .0001.

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IL-4 and IL-13 induce expression of their cognate receptors in vitro

As noted above, the IL-4/IL-13 signaling axis up-regulates and activates the STAT6 and GATA-3 transcription factors, which then promote their own expression as well as that of the IL-4 and IL-13 interleukins and their cognate receptors, IL-4Rα and IL-13Rα1 in immune cells (reviewed in [24]). To begin to examine this in prostate fibroblasts, we first determined whether IL-4 and/or IL-13 induced the expression of their cognate receptors, IL-4Rα and IL-13Rα1. As shown in Fig 6A and quantitated in Fig 6B, N1 cells endogenously expressed both IL-4Rα and IL-13Rα1 under basal conditions. IL-4 significantly (p < .0001) induced expression of IL-4Rα to levels ~3X higher than basal levels or those observed after IL-13 treatment. Conversely, treatment with IL-13 did not induce or repress IL-4Rα expression levels compared to basal expression (Fig 6B). Treatment with IL-13 was associated with a significant (p < .0001) increase in IL-13Rα1 expression levels above basal level, whereas IL-4 treatment had no effect on in IL-13Rα1 expression levels (Fig 6B). Neither IL-4 nor IL-13 induced or repressed expression of the IL-13Rα2 decoy receptor, which was unchanged from basal levels (Fig 6B) (S6 Fig).
IL-4Ra drives IL-4- and IL-13-mediated signal transduction

IL-4 and IL-13 share a common signaling axis that requires the IL-4Rα receptor. IL-4 signals through a IL-4Rα/γc (Type 1) or IL-4Rα/IL-13Rα1 (Type 2) heterodimer, while IL-13 signals solely through the IL-4Rα/IL-13Rα1 Type 2 heterodimer. IL-13Rα2 is thought to act as a decoy receptor to homeostatically sequester IL-13 in normal cells and does not transduce intracellular signaling [26–28]. Signal Transducer and Activator of Transcription (STAT) proteins comprise a family of six members (STATS 1–6) that are phosphorylated primarily by Janus Kinases (JAKs), in response to IL-4 or IL-13 binding to the heterodimeric receptors described above [29]. In order to determine the relative contribution of IL-4 and IL-13 stimulation to STAT6 phosphorylation in human prostate fibroblasts, N1 immortalized human prostate fibroblasts were treated with vehicle, IL-4, or IL-13 with or without pre-treatment with antibodies against their cognate receptors, IL4Rα or IL-13Rα1, respectively. These studies showed that IL-4 stimulation of STAT6 phosphorylation was repressed upon pre-treatment with IL4Rα antibodies (Fig 7A and 7C) but not IL-13Rα1 (Fig 7B and 7C), whereas IL-13 stimulation of STAT6 phosphorylation was repressed upon pre-treatment with either IL-13Rα1 (Fig 7D and 7F) or IL4Rα (Fig 7E and 7F) antibodies (S7 Fig). These data suggest

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downstream signaling mediated by IL-4 may be less suppressible, hence, more robust, than that mediated by IL-13. This is consistent with the ability of IL-4 to signal through both Type 1 and Type 2 heterodimeric receptors.
IL-4 and IL-13 signal through JAK/STAT to induce collagen 1 and collagen 3 expression

Upon activation by ligand binding, the IL-4Rα/IL13Rα1 signaling axis activates the transcription of genes encoding pro-fibrotic collagens, which are key components of the ECM and are heavily upregulated in fibrosis (reviewed in [24]). To directly investigate the potential role(s) of JAK/STAT signaling in IL-4 induction of collagen expression, N1 cells were treated with IL-4 with or without pretreatment with 5μM tofacitinib (CP-690550; Xeljanz), a small molecule JAK2/JAK3 inhibitor with low cytotoxicity shown to repress IL-4-mediated downstream signaling [30–32]. N1 fibroblasts grown in complete media will express high basal collagen levels likely due to the presence of pro-fibrotic proteins in the serum component of the media. If the complete media is removed, the cells washed, and media replaced with serum-free media, the cells will gradually cease producing collagen (Fig 8A). However, cells grown for 24–48 hr in serum-free media, then treated with IL-4 or IL-13, significantly up-regulate collagen 1 and 3 production over basal levels (Fig 8A–8C). Therefore, collagen 1 and 3 accumulation was concordant with IL-4 treatment, and was reduced or ablated upon pre-treatment with tofacitinib (Fig 8). Similarly, phosphorylation of STAT6 is concordant with IL-4-mediated collagen 1 or 3 induction and is robustly maintained by IL-4-treated cells for at least 48 hr in the absence of tofacitinib (Fig 8A–8D) (S8 Fig). These studies show that IL-4 induced expression of collagen 1 and 3 is coupled to JAK/STAT signaling in human prostate fibroblasts.
Aging- and inflammation-associated fibrotic changes in tissue architecture contribute to dysfunction and disease in multiple organ systems [33, 34]. Our studies recently showed that periurethral prostatic fibrosis is associated with lower urinary symptoms and urinary voiding.

Fig 8. IL-4-mediated induction of collagen 1 and collagen 3 expression is coupled to JAK/STAT signaling. N1 immortalized human prostate fibroblasts were serum-starved for 48 hr then treated with vehicle or IL-4 (20ng/ml) with or without 2 hr pre-treatment with 5um tofacitinib, a JAK2/JAK3 inhibitor, and maintained in serum-free media for an additional 24 or 48 hr. IL-4 significantly induced expression of collagen 1 (COL1) and collagen 3 (COL3) concurrent with induction of STAT6 phosphorylation, all of which was repressed upon pre-treatment with tofacitinib (A). Western blots shown in A are representative of 3 replicate experiments. Densitometric data from the replicate experiments is graphed in B (COL1), C (COL3), and D (pSTAT6/STAT6) (S8 Fig). Significant differences are indicated as * p < .05; ** p < .01; *** p < .001; **** p < .0001.

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Discussion

Aging- and inflammation-associated fibrotic changes in tissue architecture contribute to dysfunction and disease in multiple organ systems [33, 34]. Our studies recently showed that periurethral prostatic fibrosis is associated with lower urinary symptoms and urinary voiding.
dysfunction in men \[10, 12\]. Fibrosis in the lower urinary tract is associated with aging and likely results from complex interactions between prostate epithelial cells, resident fibroblasts, circulating fibrocytes, and infiltrating inflammation.

Few studies have sought to determine which pro-fibrotics may play predominant roles in lower urinary tract fibrosis. The most studied to-date is TGFβ, which has been shown to induce prostatic fibrosis and alter extracellular matrix deposition in mouse models \[35\]. Some of this activity may be stimulated by an imbalance between estrogen and androgen levels in aging men \[36, 37\]. A recent study utilized medical record data, patient samples, and in vivo models to evaluate the impact of inflammation, and TNFα in particular, within the context of autoimmune disease, on lower urinary tract disorders. Previous studies from our group showed that pro-inflammatory proteins, including chemokines and interleukins, are secreted by aging prostate fibroblasts, and that one of these, CXCL12, acts as a pro-fibrotic in vitro and in mouse models in vivo \[38–40\]. The Campisi group showed that many inflammatory mediators, including those mentioned above, are secreted as part of the senescence associate secretory phenotype (SASP) induced by the aging process or, experimentally, by exposure to H\(_2\)O\(_2\), or irradiation \[41\]. Several studies have shown that IL-4 and IL-13, which share a common JAK/STAT6 signaling axis, promote fibrillar collagen secretion in multiple organ systems (reviewed in \[16\], most notably in the lung \[17, 18\], the kidney \[19, 20\], and the liver \[17, 21\].

Immunohistochemical data reported here show that the IL4R\(\alpha\) and IL13R\(\alpha\)1 receptors are up-regulated in peri-urethral prostate tissues in conjunction with high tissue collagen expression and lower urinary tract symptoms (LUTS). Data from the Human Protein Atlas (proteinatlas.org) shows that both IL4R\(\alpha\) and IL13R\(\alpha\)1 are robustly expressed at the RNA and protein levels by immune cells but also by diverse cell types, including fibroblasts and mesenchymal cells, and by diverse tissues, including the human prostate. Combined with the studies cited above showing that IL-4 and IL-13 promote fibrosis in multiple organ systems, the elevated expression of IL4R\(\alpha\) and IL13R\(\alpha\)1 concurrent with prostate tissue fibrosis and lower urinary tract voiding dysfunction led to the hypothesis that the IL-4/IL-13 signaling axis may promote prostatic fibrosis as well.

ELISA and immunofluorescence data shown in the current study provide evidence that IL-4 and IL-13 can up-regulate their own expression levels as well as those of their cognate receptors in prostate fibroblasts. This is consistent with data acquired from the study of Th2 cells which shows that Th2 differentiation occurs through a sequential series of events initiated by IL-4 or IL-13 binding to heterodimeric IL-4 receptors. These ligand/receptor interactions activate receptor bound kinases JAKs or Tyk1, which phosphorylate STAT6. pSTAT6 dimerizes and translocates to the nucleus where it promotes its own expression and that of the GATA-3 transcription factor. After phosphorylation by diverse kinases, including p38, Akt, and Erk, pGATA-3 translocate to the nucleus where both pSTAT6 and pGATA-3 transcriptionally \[42–45\] activate IL-4 and IL-13 gene expression \[46\]. pSTAT6 may also directly activate transcription of the IL-4R\(\alpha\), COL1A1, and COL1A2 genes \[43\]. Our data shows that IL-4 and IL-13 self-upregulate their expression as well as that of their cognate receptors, IL-4R\(\alpha\) and IL-13R\(\alpha\)1, in prostate stromal fibroblasts. Both IL-4 and IL-13 promote STAT6 phosphorylation in prostate stromal fibroblasts, and pSTAT6 persists for at least 48 hr post-induction. These combined data suggest a mechanism whereby signaling through the IL-4/IL-13 axis up-regulates and activates the STAT6 transcription factor, which then promotes the expression of IL-4 and IL-13 and their cognate receptors, IL-4R\(\alpha\) and IL-13R\(\alpha\)1, and promotes the expression of collagens 1 and 3, contributing to activated fibroblast accumulation and persistence in the prostate.

Studies intended to examine the mechanism of IL-4/IL-13 mediated activation of STAT6 showed that IL-4 stimulation of STAT6 phosphorylation was repressed upon pre-treatment...
with antibodies against IL4Rα but not IL-13Rα1, whereas IL-13 stimulation of STAT6 phosphorylation was repressed upon pre-treatment with either IL-13Rα1 or IL4Rα antibodies. These data suggest that IL-4, rather than IL-13, is the ‘driver’ of JAK/STAT signaling promoting STAT6 phosphorylation in human prostate fibroblasts, likely due to its ability to signal through both the Type 1 γc/ IL4Rα and Type 2 IL4Rα/ IL-13Rα1 receptor complexes. Tofacitinib, a robust but non-cytotoxic JAK2/JAK3 inhibitor commercially known as Xeljanz, completely repressed IL-4-mediated collagen 1 and collagen 3 expression by these same cells.

A recent study showed that mouse models expressing STAT6 developed dorsolateral prostate lobe fibrosis, whereas those lacking STAT6 did not, upon infection with the uropathogenic E. coli strain CP1 [47]. Together these studies confirm the importance of the JAK/STAT axis, and of STAT6-dependent mechanisms, for IL-4-mediated prostate fibroblast activation and ECM accumulation [47].

Although IL-4 and IL-13 clearly up-regulated collagen expression, neither significantly stimulated αSMA expression or adoption of a myofibroblast morphology by prostate stromal fibroblasts. This suggests that activation of the IL-14/IL-13 axis induces prostate fibroblasts to express ECM but not to differentiate completely to myofibroblasts. Fibroblast cell types that secrete ECM but lack αSMA expression or contractile activity have been described as ‘activated fibroblasts’ or ‘proto-myofibroblasts’ [48]. More recently, single cell RNA sequencing studies have identified fibroblast cell populations that lack αSMA expression but highly express collagen-encoding genes in mouse models of pulmonary fibrosis [49], and lung and kidney fibrosis [50], and in human skin [51] and human lung fibrosis [52]. Taken together, this data suggests that ECM secreting cells associated with fibrosis may comprise a continuum of fibroblastic cell phenotypes and multiple sub-populations.

In conclusion, these studies show that IL-4 and IL-13 signal through the IL-4Rα receptor to activate JAK/STAT signaling, thereby promoting their own expression, that of their cognate receptors, and collagens. The concurrent high expression levels of the IL-4Rα and IL-13Rα1 receptors and collagen in peri-urethral tissues from men suffering from LUTS is certainly intriguing. This should be further investigated, along with elucidation of the relative contributions of immune cells and prostate fibroblasts in promoting JAK/STAT signaling and fibrosis, perhaps through using organoid or in vivo mouse models. Taken together, the studies presented here suggest that the IL-4/IL-13 signaling axis is a powerful, but therapeutically targetable, pro-fibrotic mechanism in the lower urinary tract.

**Material and methods**

**Institutional review board approval**

In all cases, written informed consent was obtained from patients all subjects under protocols approved by the University Institutional Review Board.

**Immunohistochemistry**

FFPE tissues were sectioned at five-micron thickness and labeled with IL-4Rα (MAB230, R&D Systems), IL-13Rα1 (AF152, R&D Systems), or IL-13Rα2 (AF146, R&D Systems) antibodies (1:100) using AEC solution and N-Histofine (universal immune-peroxidase polymer, anti-goat/anti-mouse, Nichirei Biosciences INC). IL-4Rα, IL-13Rα1, and IL-13Rα2 expression was detected using the universal immune-peroxidase polymer kit according to manufacturer’s protocol. Appropriate negative (no primary antibody) tissues were stained in parallel. CD8 expression levels were assessed on whole tissue sections using ImmunoMembrane [53], an ImageJ plugin that uses color deconvolution for stain separation and a customized algorithm for cell membrane segmentation. A quantitative score (IM-score, 0–20 points) is generated according
to the membrane staining intensity and completeness. Specimens are classified into 0/1+, 2 + or 3+ based on IM-score cut-offs. The classification and membrane segmentation are presented as a pseudo-colored overlay image (S1 Fig). ImmunoMembrane is freely accessible at: http://jvsmicroscope.uta.fi/immunomembrane/.

**Cell culture**

N1 cells were derived from prostate transition zone tissue explanted, grown as monolayer cells, and transduced with a recombinant LXSN E6E7 retrovirus [54]. SFT1 cells were derived from prostate transition zone tissue explanted and grown as monolayer cells from a solitary fibrous tumor of the prostate characterized by a NAB2/STAT6 gene fusion [55]. The cells were grown in 5% HIE culture media (Ham’s F-12, 5% FBS, Insulin [5 μg/mL], EGF [10 ng/mL], Hydrocortisone [1 μg/mL], Fungizone [0.5 μg/mL], Gentamicin [0.05 mg/mL]), Plasmocin [0.5 μg/mL]). Prior to treatment, cells were serum starved for 24–48 hr using SF HIE (Ham’s F12, EGF [50 ng/mL], 0.1% BSA, Insulin [5 μg/mL], Transferrin [5 μg/mL], 50 μM sodium selenite, 10 mM 3,3′,5-triiodo-L-thyronine, HEPES [10mM], Hydrocortisone [1 μg/mL], Fungizone [0.5 μg/mL], Gentamicin [0.05 mg/mL]) Ethanolamine [5mM]. Lung fibroblasts were explanted and grown from lung tissues obtained from patients undergoing thoracic surgery for non-fibrotic lung diseases (normal lung fibroblasts) and were maintained in Dulbecco’s modified Eagle medium (DMEM) with 20% fetal bovine serum (FBS), penicillin, streptomycin and fungizone as described previously [56]. Average cell numbers and standard deviations were calculated.

**Proliferation assays**

Proliferation (WST) assays were conducted as previously described [39]. Briefly, cells were seeded at 10000 cells/well in 96 well plates in complete media, washed in 1X PBS and switched to serum-free media (except for a positive control well), treated with increasing doses of IL-4 (0–200 ng/ml) or IL-13 (1–100 ng/ml) for 24 or 48 hours. These studies identified concentrations of IL-4 (20ng/ml) and IL-13 (20 ng/ml) that exhibited robust cellular proliferation. Inhibition studies were conducted by pre-treating cells plated as described above for 2hr with increasing concentrations IL-4Rα (MAB230, R&D Systems) or IL-13Rα1 (AF152, R&D Systems) antibodies followed by stimulation with 20ng/ml IL-4 or IL-13 for 24 hr. Each study was conducted in triplicate and included 9 technical replicates for each condition. Proliferation was assessed by WST assay (Roche, USA, Cat. No.11644807001) (S2 Fig).

**ELISA assays**

Cells were cultured as described above and supplemented with vehicle or 20ng/ml IL-4, IL-13, IFN-γ, TNFα, or 4ng/ml TGF-β, for 24 hours with or without 2 hr pre-treatment with 400 ng/ml IL-4Rα (MAB230, R&D Systems) or 40 ng/ml IL-13Rα1 (AF152, R&D Systems) antibodies. The cells were washed with 1XPBS and the cells were cultured for another 24 hrs in serum-free media. 4mls of conditioned media collected from treated cells were spun for 40 minutes at 4,000rpm at 4˚C using Amicon Ultracel 3K (Millipore #UFC800324) to concentrate the samples. The samples were then subjected to ELISA human IL-4, IL-13, IFN-γ, TNFα and TGF-β1 immunoassays ELISA Kits (cat. no. D4050, D1300B, DFNAS0, DTA00D, R&D Systems, # MBS761127, Biosource, respectively) (S3 Fig).

**RNA extraction and quantitative real time PCR (qRT-PCR)**

Cells were cultured as described above and supplemented with vehicle or 20ng/ml IL-4, IL-13, with or without 2 hr pre-treatment with 400 ng/ml anti-IL-4Rα (MAB230, R&D Systems) or
40 ng/ml anti-IL-13Rα1 (AF152, R&D Systems) antibodies, for 24 hrs. RNA was extracted using Trizol reagent (Invitrogen, Carlsbad, CA), assessed for purity by A260/A280 ratio and quantified using a Nanodrop spectrophotometer. 1 ug of RNA was reverse transcribed using a High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Carlsbad, CA). qRT-PCR was performed using a QuantStudio 12K Flex Real-Time PCR System, reagents and software (Applied Biosystems, Carlsbad, CA). Reactions were performed in triplicate, including no template controls and amplification of an endogenous control transcript, Larger Ribosomal Protein (RPLPO), to assess template concentration and loading precision. Cycle number to threshold was calculated for each gene by subtracting the average control value from each average experimental value and normalized to RPLPO (loading control) using the Pfaffl method [57]. Molecular probes used were Hs0016400_m1 for COL1α1 and Hs99999902_m1 for RPLPO (Applied Biosystems, Carlsbad, CA) (S4 Fig).

Sircol assays

Cells were cultured as described above and supplemented with vehicle or 20ng/ml IL-4, IL-13, TNFα, or 4ng/ml TGF-β, for 48 hours with or without 2 hr pre-treatment with 400 ng/ml IL-4Rα (MAB230, R&D Systems) or 40 ng/ml IL-13Rα1 (AF152, R&D Systems) antibodies, then assessed for secreted soluble collagen types I-V using Sircol assay reagents, collagen standards, and procedures as described by the manufacturer (S1000, Life Science) (S4 Fig).

Immunofluorescence assays

For the detection of COLA1A and αSMA proteins, cells were plated on chamber slides coated with 10 μg/ml fibronectin (Sigma-Aldrich, St. Louis, MO) in complete media washed, switched to serum-free media, and supplemented with vehicle or 20ng/ml IL-4 or IL-13, or 4ng/ml TGF-β, for 48 hours with or without 2 hr pre-treatment with 400 ng/ml IL-4Rα (MAB230, R&D Systems) or 40 ng/ml IL-13Rα1 (AF152, R&D Systems) antibodies. After 48 hr the cells were fixed, permeabilized, and subjected to immunofluorescence as previously described [14], using FITC-conjugated mouse monoclonal anti-αSMA (F3777, Sigma-Aldrich, St. Louis, MO), and biotin conjugated rabbit polyclonal anti-COL1A1 (600-406-103, Rockland Immunoc hemicals, Gilbertsville, PA), PE-Cy 5 streptavidin (BD Pharmingen San Diego, CA) conjugated secondary antibodies, and control mouse IgG2a (Sigma-Aldrich, St. Louis, MO). Cells were counterstained for 5 min with 1 mg/ml DAPI (Molecular Probes, Eugene, OR) in Tris-Buffered Saline/Tween 20, washed three times for 5 min each with TBST, and mounted in an Aqua-mount (Lerner Laboratories, PA) (S5 Fig)

For the detection of IL-4Rα or IL-13Rα1, cells were grown in 24-well plates in complete media washed, switched to serum-free media, and supplemented with vehicle or 20ng/ml IL-4 or IL-13 for 48 hours. After 48 hr the cells were fixed, permeabilized, blocked using 20% and subjected to immunofluorescence as previously described [14] but using mouse monoclonal IL-4Rα primary antibody (MAB230, R&D Systems) detected with goat anti-mouse IgG H&L (Alexa Fluor 594 (Abacan, ab150116) or goat polyclonal IL-13Rα1 primary antibody (AF152, R&D Systems) detected with donkey anti-goat IgG (H+L) Cross-Adsorbed Secondary Antibody, Alexa Fluor 488 (ThermoFisher A-11055) (S6 Fig).

Photomicrographs of were taken on an Olympus BX53 fluorescence microscope or EVOS Cell Imaging System. Signal density was quantified using Image Pro Plus software.

Western blot

Cells were treated as described above and lysed in radioimmunoprecipitation assay (RIPA) Buffer. Protein quantification was carried out using Bio-Rad OneStep Bradford reagent and an
Elx800 Microplate Reader (Bio-Tek) with Gen5 software. Protein lysates were prepared for electrophoresis as previously described [39]. Membranes were blocked using a 5% BSA in TBS-T solution for detection of phospho antibodies or 5% milk for detection of total antibodies in TBS-T solution for one hour. Primary antibodies incubation was performed using a 5% BSA TBS-T solution with IL-4Rα mouse monoclonal antibody (#25463), IL-13Rα1 goat IgG (#AF146), and anti-IL-13Rα2 goat IgG (#AF152) from R&D Systems, anti-Stat6 (#9362S), Y641 anti-pStat6 (#9361S), Smad3 (#9523), S423/425 pSmad3 (#9520S), anti-Collagen 3 (#30565) and GAPDH (#2118) from Cell Signaling Technologies, and anti-Collagen 1 (#EPR7785) from Abcam. All primary antibodies were incubated overnight at a 1:1000 concentration, except for GAPDH which was used at 1:5000, followed by 3 times washing with TBS-T. Secondary antibody incubations using Horse Radish Peroxidase, HRP, conjugated goat anti-rabbit (Cell Signaling, #7074) or chicken anti-goat IgG HRP conjugated (#SC2953), goat anti-mouse IgG HRP conjugated (#SC2005) from Santa Cruz Biotechnology. Secondary antibodies were used at a 1:5000 concentration for 1 hour at room temperature. Membranes were washed twice with TBS-T and for the detection of HRP-conjugated antibody, immunoblots were quantified and analyzed using the ImageStudio software suite. Immunoblots shown are representative of triplicate experiments (S7 and S8 Figs).

Statistical analysis
Statistical analysis was accomplished with GraphPad Prism v.9 using 1- or 2-way ANOVA using replicate study data as detailed in the supplemental information). In all tests, p≤.05 was considered statistically significant.

Supporting information
S1 Fig. Fig 1 supporting information. (XLSX)
S2 Fig. Fig 2 supporting information. (XLSX)
S3 Fig. Fig 3 supporting information. (XLSX)
S4 Fig. Fig 4 supporting information. (XLSX)
S5 Fig. Fig 5 supporting information. (XLSX)
S6 Fig. Fig 6 supporting information. (XLSX)
S7 Fig. Fig 7 supporting information. (XLSX)
S8 Fig. Fig 8 supporting information. (XLSX)

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