Cell density-dependent stimulation of PAI1 and hyaluronan synthesis by TGF-β in orbital fibroblasts

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Key Words
- Graves’ orbitopathy
- orbital fibroblasts
- PAI1
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- cell density
- TGF-β

Abstract
During the course of Graves’ orbitopathy (GO), orbital fibroblasts are exposed to factors that lead to proliferation and extracellular matrix (ECM) overproduction. Increased levels of tissue plasminogen activator inhibitor type 1 (PAI1 (SERPINE1)) might promote the accumulation of ECM components. PAI1 expression is regulated by cell density and various cytokines and growth factors including transforming growth factor β (TGF-β). We examined the effects of increasing cell densities and TGF-β on orbital fibroblasts obtained from GO patients and controls. Responses were evaluated by the measurement of proliferation, PAI1 expression, and ECM production. There was an inverse correlation between cell density and the per cell production of PAI1. GO orbital, normal orbital, and dermal fibroblasts behaved similarly in this respect. Proliferation rate also declined with increasing cell densities. Hyaluronan (HA) production was constant throughout the cell densities tested in all cell lines. In both GO and normal orbital fibroblasts, but not in dermal fibroblasts, TGF-β stimulated PAI1 production in a cell density-dependent manner, reaching up to a five-fold increase above baseline. This has been accompanied by increased HA secretion and pericellular HA levels at high cell densities. Increasing cell density is a negative regulator of proliferation and PAI1 secretion both in normal and GO orbital fibroblasts; these negative regulatory effects are partially reversed in the presence of TGF-β. Cell density-dependent regulation of PAI1 expression in the orbit, together with the local cytokine environment, may have a regulatory role in the turnover of the orbital ECM and may contribute to the expansion of orbital soft tissue in GO.

Introduction
Graves’ orbitopathy (GO) is the extrathyroidal manifestation of Grave’s disease characterized by an autoimmune inflammation resulting in an increased volume of the orbital connective tissue and enlargement of the extraocular muscles (Bahn 2010, Wang & Smith 2014). Orbital fibroblasts (OFs) are the main target cells in GO due to their expression of autoantigens specific to GO (Otto et al. 1996, Bahn 2010, Wang & Smith 2014). Macrophages, T and B lymphocytes, and mast cells infiltrate the orbit and activate the OFs (Bahn 2010).
Cytokine-dependent activation of OFs augments the inflammatory and autoimmune processes and accounts for orbital tissue remodeling (Smith 2005).

Altered synthesis and degradation of extracellular matrix (ECM) components by activated OFs play a key role in the pathogenesis of GO (Smith 2005). Hyaluronan (HA) is the major ECM glycosaminoglycan, a high-molecular-weight polysaccharide with essential role in cell–ECM and cell–cell interactions. HA is synthesized at the plasma membrane by three isoenzymes named hyaluronan synthase (HAS) 1, 2, and 3, which possess different biochemical properties (Vigetti et al. 2014). HA has a high turnover rate; hyaluronidase (HYAL) 1 and 2 are the major hyaluronidases expressed in human tissues (Girish & Kemparaju 2007). HA is retained as a pericellular coat after its synthesis, anchored to the cell surface via the synthase enzyme or through binding to a surface receptor, and certain amount cleaved by hyaluronidase is released from the pericellular matrix and incorporated as an integral component of the ECM. Alterations in this process could affect the physiological role of HA in the surrounding tissue (Monslow et al. 2015). In GO, OFs secrete large amounts of HA in response to, as of now, only partially characterized cytokines and antibodies (Bahn 2010). Due to its hydrophilic nature, HA accumulation accelerates expansion of the orbital connective tissues (Wang & Smith 2014). In addition, the interaction between HA and its receptor on the surface of most immune cells is important for the infiltration of leukocytes into the inflamed tissue (DeGrendele et al. 1997).

In addition to ECM molecules, fibroblasts synthesize and secrete proteases capable of degrading the ECM (McAnulty 2007). The plasminogen activator/plasmin system and its main negative regulator, the plasminogen activator inhibitor type 1 (PAI1 (SERPINE1)), have important roles in ECM degradation and remodeling (Smith & Marshall 2010). PAI1 is a single-chain glycoprotein and a member of the serine protease inhibitor family. PAI1 inhibits tissue and urokinase-type plasminogen activators, thus inhibiting the plasminogen-plasmin conversion as well as plasmin-dependent matrix metalloprotease activation. Therefore, PAI1 is a primary regulator of plasmin-driven proteolysis of the ECM. The increased expression of PAI1 results in ECM accumulation (Małgorzewicz et al. 2013). The expression of PAI1 is regulated by various cytokines and growth factors including transforming growth factor-β (TGF-β) (Ghosh & Vaughan 2012). Strong TGF-β expression has been found in the orbital tissue of patients with mild and severe GO, which correlated positively with the clinical activity score (CAS) (Pawlowski et al. 2014). It has been demonstrated that OFs synthesize PAI1, and that PAI1 expression is up-regulated by TGF-β, interferon-γ (IFN-γ), and leukoregulin (Smith et al. 1992, Cao et al. 1995, Hogg et al. 1995); however, the role of PAI1 in the pathogenesis of GO is still unclear.

PAI1 expression has been shown to be dependent on cell density in various cell types including human preadipocytes (Crandall et al. 1999), dermal fibroblasts (McFarland et al. 2011), and NIH/3T3 cells (Tanaka et al. 2013). Those studies revealed that PAI1 synthesis decreased during the process of growth toward confluency, which was consistent with the observation that PAI1 transcription occurred during the G0 to G1 transition (Qi et al. 2006). However, it is unknown how PAI1 expression is altered by the increased fibroblast proliferation during the course of GO.

The aim of this study is to investigate whether cell density alone or in combination with TGF-β affected the secretion of PAI1 and the accumulation of the ECM component HA in OF cultures originating from normal and GO orbits.

Materials and methods

Materials

Recombinant human TGF-β1, fetal bovine serum (FBS), Medium 199 with Earles’ salts and GlutaMAX supplement, penicillin/streptomycin, Dulbecco’s phosphate-buffered saline without calcium and magnesium (DPBS), trypsin–EDTA solution, freezing medium, and TrypLE Express were purchased from Gibco (Thermo Fisher Scientific). DuoSet ELISA Human Serpin E1/PAI-1 Kit and DuoSet Hyaluronan Kit were purchased from R&D Systems (R&D Systems). Cell proliferation ELISA and BrdU (colorimetric) were purchased from Roche (F. Hoffmann-La Roche Ltd). RNeasy Protect Cell Mini was purchased from QIAGEN (QIAGEN GmbH). High Capacity cDNA Reverse Transcription Kit and TaqMan Gene Expression Assays were purchased from Applied Biosystems (Thermo Fisher Scientific).

Tissue samples and cell cultures

Orbital connective tissue explants were obtained from five patients (three females and two males) undergoing orbital decompression surgery for GO. The mean age of patients at the time of surgery was 45.4 (range 37–55) years. The mean duration of Graves’ disease in GO was 5.6±4.2 (range 1–11) and 3.0±2.9 (range 1–8) years respectively.
All but one patient was in the inactive phase of GO. The patient with active GO and one patient with inactive GO were positive to thyroid-stimulating hormone (TSH) receptor antibody. Before surgery, two patients underwent thyroideectomy, two patients got radioactive iodine treatment, and all but one patient had been on antithyroid drugs for some time during the course of the disease. All patients had received corticosteroid treatment in the past, and two patients received orbital irradiation. During the last 2 months before orbital surgery, patients used only diuretics, β-blockers, thyroxine supplementation, and local measures. Patients at the time of surgery had suppressed TSH levels and high-normal thyroid hormone levels. Control normal orbital tissues (five patients: two females and three males, mean age: 63.4 years; range 47–72 years) were obtained during surgery (enucleation) for non-orbital eye diseases, and control dermal connective tissue (three patients: two females and one male, mean age: 64.7 years; range 50–80 years) were obtained during abdominal hernia operations from patients with no history of thyroid diseases. The study was approved by the Ethics Committee of the University of Debrecen. All patients gave their written informed consent.

Human OFs were cultured as described previously in detail by Bahn et al. (1987). Briefly, the tissues were cut into 2 × 2 mm pieces and inoculated into culture dishes containing Medium 199 with Earle's salts with 20% (v/v) FBS and penicillin–streptomycin, and cultured at 37°C, 5% CO₂, in a humidified incubator. Medium was changed every 3–4 days. After 2–3 weeks, depending on the proliferative rate of the culture, the tissue pieces were removed and the cell cultures were maintained in Medium 199 with 10% (v/v) FBS under the same conditions. After gentle treatment using TrypLE Express dissociating reagent, cells were stored in freezing medium in liquid nitrogen until needed. The cells were studied between passages 2 and 8.

For experiments, orbital and dermal fibroblasts were plated in 24-well plates, in various cell densities (6.24 × 10^4, 3.12 × 10^4, 1.56 × 10^4, 7.8 × 10^3, 3.9 × 10^3, and 1.95 × 10^3 cells/cm²) in Medium 199 supplemented with 10% (v/v) FBS. Densities of 1.56 × 10^4 cells/cm² and above represent confluent and post-confluent cultures respectively. The cultures were synchronized with serum starvation for 24 h (Khammanit et al. 2008) followed by treatment with medium containing 10% (v/v) FBS with or without TGF-β (1 ng/mL) for an additional 24 h. The conditioned media were collected and centrifuged at 3500 rpm for 5 min to remove any cellular debris and stored at −20°C until used. All experiments were performed at least three times and carried out in triplicate.

Cell proliferation assay

Cells were treated as described above. The assay was performed in 96-well plates according to the manufacturer's instructions. Briefly, the 5-bromo-2′-deoxyuridine (BrdU) solution was added to the cell cultures and incubated for 2 h. After removing the medium, cells were fixed using FixDenat solution for 30 min. Then, peroxidase-conjugated anti-BrdU antibody was added for 90 min. Finally, 3,3′,5,5′-tetramethylbenzidine substrate was introduced for 10 min, and after the addition of 2 N H₂SO₄, the absorbance at 450 nm (reference wavelength: 620 nm) was detected using a Beckman Coulter, DTX 880 Multimode Detector (Beckman Coulter Inc., Brea, CA, USA).

Quantitation of PAI1 protein and HA

Secreted PAI1 protein and HA levels in cell culture supernatants were measured by using DuoSet ELISA Human Serpin E1/PAI-1 Kit and DuoSet Hyaluronan Kit respectively. For pericellular HA measurement, cells were washed twice with DPBS and treated with 0.05% (w/v) trypsin–EDTA solution at 37°C for 20 min, and the reaction was stopped by addition of FBS (10% v/v final concentration). After centrifugation at 3500 rpm for 5 min, supernatants were used for HA determination. In each case, results were adjusted for the HA content of FBS. In all experiments, the PAI1 and HA productions per cell were expressed as ng/10^5 cells.

Real-time polymerase chain reactions (RT-PCR)

The supernatants were removed and cells were washed twice with DPBS. RNasey Protect Cell Mini Kit was used for the isolation of RNA from cells. We used the QIAcube robotic workstation (QIAGEN) for the isolation of RNA. The purified RNA samples were reverse transcribed by High Capacity cDNA Reverse Transcription Kit. The TaqMan Gene Expression Assays was used for the detection of the expression of PAI1, HAS1, HAS2, HAS3, HYAL1, HYAL2, and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (assay IDs: SERPINE1-Hs01126603_m1, HAS1-Hs00987418_m1, HAS2-Hs00193435_m1, HAS3-Hs00193436, HYAL1-Hs00201046_m1, HYAL2-Hs01117343_g1, and GAPDH-Hs02758991_g1 respectively). The reactions were performed by the StepOnePlus Real-time PCR System.
(Applied Biosystems). Results were normalized to GAPDH mRNA levels by the ΔC<sub>T</sub> method.

**Statistical analysis**

Statistical analysis was performed using the STATISTICA 12 software (Statsoft Inc. Tulsa, OK, USA). Data are expressed as mean±s.e.m. Repeated measures analysis of variance (ANOVA) with density as the within-subjects factor and origin of fibroblasts (dermal fibroblasts, normal OFs, and GO OFs) as between-subjects factor and LSD post hoc analysis were performed to evaluate the differences. Correlation analysis was performed to study the relationship between continuous variables. The level of statistical significance was set at P<0.05.

**Results**

Dermal fibroblasts, normal OFs, and GO OFs secreted comparable amounts of PAI1 protein at each density. Cell culture synchronization by serum starvation for 24 h before experiments resulted in slightly but significantly higher amounts of secreted PAI1 at the end of 24-h tests in medium re-supplemented with serum and led to less variance between repetitions (data not shown). PAI1 secreted by individual cells (expressed as ng/10<sup>5</sup> cells) decreased significantly (P<0.000001) with increasing cell densities (Fig. 1A); however, the origin of fibroblasts was not a significant factor; normal OFs and GO OFs behaved similarly in this respect. RT-PCR confirmed that PAI1 mRNA level decreased in parallel with PAI1 protein concentration with increasing cell density (P<0.0001) (Fig. 2A). PAI1 mRNA expression was different in cell lines with different origin (P<0.01): in descending order, dermal fibroblasts, normal OFs, and GO OFs (dermal fibroblasts vs normal OFs, P<0.01; dermal fibroblasts vs GO OFs, P<0.001; normal OFs vs GO OFs, P=0.03).

HA secretion into the medium and pericellular HA levels did not show cell density-dependent behavior (P=0.6 and P=0.2 respectively), and the origin of fibroblasts was the main predictor of HA synthesis (P<0.001) (Fig. 1B and C). Post hoc analysis showed that dermal fibroblasts secreted significantly higher amounts of HA than OFs (dermal fibroblasts vs normal OFs, P<0.01; dermal fibroblasts vs GO OFs, P<0.001). In OFs, higher HA producers (>500 ng/10<sup>5</sup> cells) only occur among normal OFs (two out of five), and lower HA producers (<300 ng/10<sup>5</sup> cells) only occur among GO OFs (three out of five). Similar results were found

![Figure 1](http://joe.endocrinology-journals.org)
in pericellular HA levels (dermal fibroblasts vs normal OFs, \(P=0.013\); dermal fibroblasts vs GO OFs, \(P<0.001\)), although there was a tendency toward lower pericellular HA in GO OFs than in normal OFs (\(P=0.06\)). We found significant positive correlation between HA released into the medium and HA retained in the pericellular matrix (dermal fibroblasts: \(r=0.72, P<0.01\); normal OFs: \(r=0.77, P<0.0001\); GO OFs: \(r=0.97, P<0.000001\)). Results of the RT-PCR showed that \(\text{HAS1}\) and \(\text{HAS3}\) expressions were in the same order of magnitude, whereas \(\text{HAS2}\) expression was the predominant HAS enzyme in all cell lines studied (Fig. 2B, C and D). The expression pattern of HAS enzymes was different in fibroblasts with distinct type of origin. Dermal fibroblasts had higher expression levels of \(\text{HAS1}\) and \(\text{HAS2}\) mRNA than OFs (dermal fibroblasts vs normal OFs, \(P=0.05\) and \(P<0.0001\), dermal fibroblasts vs GO OFs, \(P=0.02\) and \(P<0.0001\), respectively), whereas OFs had higher \(\text{HAS3}\) expression than dermal fibroblasts (dermal fibroblasts vs normal OFs, \(P=0.03\), dermal fibroblasts vs GO OFs, \(P=0.0001\)). GO OFs had higher \(\text{HAS3}\) expression than normal OFs (\(P<0.001\)). The expression of \(\text{HAS1}\) and \(\text{HAS2}\) decreased with increasing cell density in dermal fibroblasts. The expression of \(\text{HYAL1}\) was more diverse than the expression of \(\text{HYAL2}\) (Fig. 2E and F). Dermal fibroblasts had the lowest \(\text{HYAL1}\) expression, and GO OFs had lower \(\text{HYAL1}\) expression than normal OFs (dermal fibroblasts vs normal OFs, \(P<0.001\), dermal fibroblasts vs GO OFs, \(P<0.01\), normal OFs vs GO OFs, \(P<0.001\)). There was a tendency for lower \(\text{HYAL1}\) and \(\text{HYAL2}\) expression in confluent cultures than in pre- and post-confluent cultures.

The proliferation rate of fibroblasts declined significantly (\(P<0.000001\)) with increasing cell densities in all fibroblasts tested, irrespective of the site of origin (Fig. 1D). Accordingly, significant positive correlation was found between proliferation and PAI1 secretion in all fibroblasts studied (dermal fibroblasts: \(r=0.81, P<0.01\); normal OFs: \(r=0.71, P<0.0001\); GO OFs: \(r=0.70, P<0.0001\)).
The proportion of proliferating cells was significantly higher in OFs than in dermal fibroblasts at all but one density (normal OFs vs dermal fibroblasts, \( P=0.03 \); GO OFs vs dermal fibroblasts, \( P=0.02 \)). No differences were observed between OFs derived from normal and GO orbital connective tissues (\( P=0.8 \)) according to the post hoc tests.

Next, our model was completed with TGF-\( \beta \) treatment. Dose–response experiments were performed in a range of 0.01–10 ng/mL; 1 ng/mL TGF-\( \beta \) had the maximal effect on PAI1 secretion during the 24-h treatment, and this concentration was selected for further experiments.

At each density, TGF-\( \beta \) stimulated PAI1 secretion in all tested cell lines, irrespective of the site of origin (\( P<0.000001 \)). In both GO and normal OFs, but not in dermal fibroblasts, the extent of the stimulation by TGF-\( \beta \) strongly depended on cell density (\( P<0.000001 \)); more marked stimulation of PAI1 secretion was seen with increasing densities (Fig. 3A). The same stimulation pattern was detected when PAI-1 mRNA levels were examined under the effect of TGF-\( \beta \) (Fig. 4A). Thus, the PAI-1-lowering effect of high cell densities (Fig. 1A) has been partially reversed by TGF-\( \beta \).

Significant stimulatory effect of TGF-\( \beta \) on HA secretion into the medium was seen at the highest cell densities, irrespective of the origin of fibroblasts (\( P<0.0001 \)) (Fig. 3B). The same was true for pericellular HA of OFs (Fig. 3C), whereas dermal fibroblasts responded to TGF-\( \beta \) with a pericellular HA rise at all densities. The positive correlation between HA in the medium and pericellular HA seen in unstimulated cultures (Fig. 1B and C) remained significant in cell cultures after 24 h TGF-\( \beta \) treatment (dermal fibroblasts: \( r=0.58, P=0.02 \); normal OFs: \( r=0.95, P<0.000001 \); GO OFs: \( r=0.89, P<0.0000001 \)). The expression of HAS2, HAS3, HYAL1, and HYAL2 did not change or slightly decreased in TGF-\( \beta \)-treated cells (data not shown), whereas HAS1 mRNA expression increased markedly in a cell density-dependent manner (\( P=0.03 \)), especially in OFs (Fig. 4B).

In OF cultures, TGF-\( \beta \)-induced increase in PAI1 secretion was directly proportionate to the changes of both HA secretion (normal OFs: \( r=0.78, P<0.000001 \); GO OFs: \( r=0.56, P<0.01 \)) and pericellular HA level (normal OFs: \( r=0.67, P<0.0001 \); GO OFs: \( r=0.73, P<0.00001 \)).

There was only a minor effect of TGF-\( \beta \) on the proliferation rate (Fig. 3D), which was dependent on cell density (\( P<0.01 \)). A slight non-significant increase was only observed at the highest density of orbital cultures (\( P=0.051 \) in normal and \( P=0.22 \) in GO OFs). At other densities, mild inhibitory effect of TGF-\( \beta \) was detected (most pronounced in dermal cultures at the lowest density

\( P<0.0001 \)).
Discussion

Orbital fibroblasts are considered to be the primary targets of the autoimmune process in GO, activated by cytokines released by immune cells infiltrating the orbital connective tissue (Bahn 2010, van Steensel et al. 2012). Stimulated OFs deposit ECM components and proliferate in an unregulated manner (Wang & Smith 2014). HA is the major overproduced glycosaminoglycan in the GO orbit (Hufnagel et al. 1984, Smith et al. 1989, Bahn 2010). HA accumulation in the orbital connective tissue causes edematic swelling (Natt & Bahn 1997) and facilitates inflammatory cell infiltration (Guo et al. 2011, Evanko et al. 2012), thereby promoting disease progression. ECM remodeling requires proteolytic enzyme activity (Lu et al. 2011). The plasminogen activator system plays a key role in ECM remodeling in many physiological and pathophysiological processes (Mignatti & Rifkin 1993), as PAI1 is the main negative regulator of the conversion of plasminogen to plasmin; inhibits fibrinolysis and plasmin-dependent pericellular proteolysis (Ghosh & Vaughan 2012); modulates cellular adhesion, migration, and wound healing. As a consequence, the balance between ECM synthesis and degradation is regulated, to a great extent, by PAI1. However, cell–cell contacts regulate proliferation and PAI1 expression in non-transformed cells (Comi et al. 1995, Tanaka et al. 2013). During the course of GO, overproduction of HA creates a loose, highly hydrated environment, which has an impact on cell–cell contacts and cell–ECM interactions.

In this study, we have shown that cell density is a negative regulator of both proliferation and PAI1 production in OFs, regardless of whether they are derived from normal or GO orbital tissue. Contact inhibition of proliferation is typical of non-transformed cells (Küppers et al. 2010) and is considered to be continuously active, regulating cell proliferation and organ size in adult tissues (Zeng & Hong 2008). Therefore, we have hypothesized that confluent cultures correspond to the healthy orbit, whereas pre-confluent cultures represent the expanding, high ECM orbital tissue with higher fibroblast proliferation rate. We found that OFs, both from control and GO orbits, had higher proliferative capacity than dermal fibroblasts (Fig. 1D). This means that OF cultures are characterized by higher cell turnover compared with dermal fibroblasts. Others made similar observations under the same conditions in confluent cultures of OFs (Heufelder & Bahn 1994). In our experiments, a progressive transcriptionally regulated reduction in PAI1 synthesis with increasing cell densities was detected using fibroblasts of both dermal and orbital origin; PAI1 synthesis per cell was inversely proportional to cell density. This is in agreement with similar studies in non-orbital cultures (Crandall et al. 1999, Tanaka et al. 2013). However, we are the first to show that OFs behave in the same manner. We assume that the growth state of OFs at different cell densities is the main determinant of PAI1 expression. Elevated PAI1 synthesis during G0 to G1 transition maintains a supporting scaffold for proliferation by limiting pericellular proteolysis. It was found in human keratinocytes that a dynamic occupancy of the E box site in the PE2 region of PAI-1 gene promoter by upstream stimulatory factor (USF) subtypes (USF1 vs USF2) determine the transcriptional status of the PAI-1 gene in quiescent versus cycling cells (Qi et al. 2006). Further studies are needed to confirm whether these
transcription factors play a role in the proliferation rate-dependent PAI1 expression of OFs.

We assumed that TGF-β may be one of the cytokines that is responsible for the connective tissue changes in the orbit. TGF-β is known to transcriptionally regulate cell growth, motility, matrix remodeling, and pericellular proteolytic activity (Samarakoon et al. 2013); TGF-β is also known to stimulate PAI1 transcription and to up-regulate the expressions of certain genes that encode elements of the ECM, such as fibronectin, collagen I, and proteoglycans in non-orbital tissues (Matrisian & Hogan 1990, Wight & Potter-Perigo 2011). Furthermore, TGF-β has been shown to be present in the orbits of GO patients (Pawlowski et al. 2014). Indeed, we found that TGF-β was a potent stimulator of PAI1 synthesis at all densities in OFs, resulting in up to a fivefold PAI1 increase above baseline at high cell densities (Fig. 3A). The cell density-dependent stimulation of PAI1 expression by TGF-β may occur due to the interference or cooperativity of transcription factors (Samarakoon & Higgins 2008). Only mild and non-density-dependent effect was seen in dermal fibroblasts (Fig. 3A). We think that the high PAI1 concentration in the GO orbit acts toward increased matrix stability, which may contribute to matrix expansion.

Enhanced HA synthesis was described in OFs cultured in serum-free medium with TGF-β (Wang et al. 2005, Guo et al. 2011). We found that under basic circumstances, HA production per cell did not show a cell density-dependent pattern but was highly dependent on the origin of fibroblasts (Fig. 1B and C). This was supported by the different expression patterns of HAS enzymes in the cell lines with different origins. The three HAS isoforms have distinct enzymatic properties, such as rate of synthesis (HAS1 < HAS2 < HAS3), produced HA size (HAS1 and HAS3, 200–2000 kDa; HAS2 over 2000 kDa), and capability for retaining HA in the pericellular coat (HAS1 overexpression leads to smaller pericellular coat); however, changes in the transcriptional level of HAS do not always correlate with changes in the HA secretion (Itano & Kimata 2002). During the turnover of HA, HYAL2 initiates the degradation of HA at the cell surface to smaller chains, which are then further degraded by HYAL1 (Girish & Kemparaju 2007). Despite the higher expression of the HAS3, the isoform with highest activity, and lower expression of HYAL1, we observed lower HA production of GO OFs compared with normal OFs, an observation similar to that of others (Krieger & Gershengorn 2014). When exposed to TGF-β, the same cells responded with increased HA secretion at high cell densities, in parallel with PAI1 rise (Fig. 3B and C). We confirmed that TGF-β treatment up-regulates HAS1 mRNA expression. Elevated HA synthesis induced by HAS1 overexpression is known to increase HA cable formation and promote HA-dependent monocyte binding in vascular smooth muscle cells (Wilkinson et al. 2006).

From the point of disease course, the matrix overproducing responses of OFs to TGF-β at high cell densities may be detrimental in GO. HA overproduction can diminish contact inhibition and lead to elevated proliferation (Itano et al. 2002). HA increases PAI1 expression in a concentration-dependent manner in human vascular smooth muscle cells (Marutsuka et al. 1998) and a positive correlation exists between HA and PAI1 produced by human aortic endothelial cells incubated with C-reactive protein (Devaraj et al. 2009). In human umbilical vein endothelial cells, high-molecular-weight HA induces activation of the TGF-β receptor I and expression of PAI1 (Park et al. 2012). In combination with these previous findings, our data suggest that the TGF-β-induced HA secretion at high cell densities may facilitate the cell density-dependent PAI1 stimulation in OFs.

Based on our results, contact inhibition is an important negative regulator of PAI1 synthesis in OFs. High proliferation rate is associated with high PAI1 synthesis (Fig. 1A and D). As HA production of OFs did not depend on contact inhibition and/or proliferation, its excessive production in GO is most probably under the influence of locally produced cytokines. OFs may respond to certain cytokines by elevated HA production and increased proliferation; we found that TGF-β had such effect at high cell densities. HA accumulation provides a stimulating environment for further fibroblast proliferation and leukocyte infiltration (Wight & Potter-Perigo 2011). Our results show that the diminution of PAI1 synthesis caused by increasing cell density is partially reversed by TGF-β (compare Figs 1A and 3A). From the point of the pathogenesis of GO, it is remarkable that PAI1 can directly promote the infiltration of macrophages and T cells (Samarakoon et al. 2013). However, the plasminogen-plasmin conversion decreases; therefore, the degradation of ECM is also reduced. The ECM stabilization by PAI1 and, as of now, unidentified factors may alter local cytokine levels by trapping cytokines and further influencing the course of GO. The ability of PAI1 to inhibit the conversion of the latent form of TGF-β to its active form (Ghosh & Vaughan 2012) may act in the opposite direction.

The data presented here are preliminary and may not be related to the initiating factors of orbital autoimmunity. Connective tissue expansion is most probably several steps down from the early immune regulatory derailment.
of the autoimmune process. However, connective tissue expansion is a major contributor to disease outcome, and therapeutic interventions that could interfere with ECM overproduction may have a beneficial effect in GO.

Here, we have shown for the first time that increasing cell density is a negative regulator of both proliferation and PAI1 secretion in OFs, regardless of whether they are derived from normal or GO orbital tissues, and these negative regulatory effects are partially inhibited in the presence of TGF-β. We conclude that TGF-β is a potent inducer of PAI1 and HA expression in OFs in culture. This effect of TGF-β on PAI1 synthesis is specific to fibroblasts of orbital origin. As both GO and normal OFs responded the same way to TGF-β, we assume that it is rather the presence of immune cells and cytokines in the GO orbits, than any inherent difference between GO and non-GO orbital tissues, that contributes to the development of the disease in GO.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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