Anti-Fibrotic Activity of an Antimicrobial Peptide in a Drosophila Model

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Abstract
Fibrotic lesions accompany several pathological conditions, including tumors. We show that expression of a dominant-active form of the Ras oncogene in Drosophila salivary glands (SGs) leads to redistribution of components of the basement membrane (BM) and fibrotic lesions. Similar to several types of mammalian fibrosis, the disturbed BM attracts clot components, including insect transglutaminase and phenoloxidase. SG epithelial cells show reduced apicobasal polarity accompanied by a loss of secretory activity. Both the fibrotic lesions and the reduced cell polarity are alleviated by ectopic expression of the antimicrobial peptide drosomycin (Drs), which also restores the secretory activity of the SGs. In addition to extracellular matrix components, both Drs and F-actin localize to fibrotic lesions.

Introduction
Extracellular matrices (ECMs) are highly specialized three-dimensional, tissue- and stage-specific structures, which surround cells. Much of our understanding of the ECM structure and dynamics has been obtained by studying model organisms [1, 2]. The ECM is composed of fibrous as well as nonfibrous proteins and of proteoglycans. Fibrous collagen and laminin (lam) form 2 extracellular networks in part through self-assembly [3, 4]. Nonfibrous proteins of the ECM including nidogen (Ndg) and the proteoglycan perlecan (Pcan) mediate the interaction between the collagen- and lam networks [4]. Formation of the ECM is influenced by the underlying cells via receptors such as dystroglycan, syndecans, and integrins, which mediate contact between the cells and the ECM [3–5]. Mutual cell-ECM interactions influence cell adhesion, migration, proliferation, apoptosis, and cell differentiation during the development. In recent years, it has been increasingly recognized that instead of being a stable structure, the ECM is a highly dynamic matrix, which undergoes constant turnover both during the development and as part of physiologic adaptations [3–5]. During the development, the ECM directs morphogenetic processes, for example, in the intestine, the lungs, and the mammary and submandibular (salivary) glands [3, 5]. Physiological responses include rapid repair upon tissue damage, which restores cellular and ECM integrity. In contrast, if unchecked, ECM remodeling in humans may promote pathological states, such as osteoarthritis, fibrosis, and progression toward cancer [3, 5–7] and may impede the delivery of drugs [8]. Fibrosis affects different organs, including the skin, liver, kidney, heart, and lungs [9]. Fibrotic triggers include genetic disorders, mechanical stimuli, such as asbestos, poorly controlled diabetes.
and hypertension, tissue dysplasia, such as in tumors as well as persistent infections [10]. Pulmonary fibrosis may be of unknown origin (idiopathic, [11]) or a consequence of infections such as during severe cases of COVID-19 [12]. Fibrotic lesions are often characterized by an unabated activation of wound healing [10]. Chronic activation may occur during the (1) initial hemostatic phase, the subsequent (2) recruitment of the cells of innate and adaptive immunity, or (3) during the remodeling and restoration of tissue integrity. The last stage includes the migration and differentiation of fibroblasts into myofibroblasts [9]. Fibroblastic pathologies include increased production of ECM components, increased stiffness of the ECM, as well as chronic immune activation and inflammation [3, 4, 8, 9]. ECM dysregulation often affects tissue function and may lead to complete organ failure and ultimately to death [4, 10, 13, 14].

Effect mechanisms of insect innate immune systems comprise both soluble and cellular reactions [15, 16]. Soluble mediators are released from immune-competent organs, primarily the fat body but also the gut, the tracheae, and the salivary glands [15, 17–19]. Among the most strongly induced soluble mediators are several classes of antimicrobial peptides (AMPs), with both antibacterial and antifungal activities [15]. One example for the latter is the antifungal peptide Drosomycin (Drs, [20]). Cellular immunity relies on hemocytes and immune cells in the hemolymph, which in noninfected Drosophila melanogaster comprise plasmatocytes and crystal cells [15, 21, 22]. Plasmatocytes have been likened to macrophages, although recent single-cell RNA sequencing revealed a high level of diversity among plasmatocytes [16, 22–24]. They are also key players during insect hemostasis and release coagulation factors, which help to seal wounds and prevent dissemination of microbial intruders [25]. Coagulation factors in insects include transglutaminase, a homolog of the mammalian clotting factor XIIIa [22, 26, 27]. Crystal cells contain inactive precursors of the enzyme phenoloxidase (prophenoloxidase, PPO), which is activated upon infection and wounding and leads to the production of intermediates with antibacterial and cross-linking activity at wound sites, and ultimately to the production of melanin [22]. Melanization is also the final part of responses against large numbers of bacteria (nodule formation) and against larger intruders such as wasp eggs (encapsulation, [15, 16]). Nodule formation and encapsulation have been likened to the formation of granulomas in vertebrates [25, 28, 29]. Melanization is also observed as a consequence of deregulation of immunity and as part of the response against aberrant self-tissues similar to autoimmune reactions in mammals. Mutants that display endogenously driven melanization have been named “melanotic tumor mutants,” although the underlying mechanism is not necessarily linked to mutations in tumor-promoting genes [30].

Dysregulation of the ECM with similarities to fibrosis has been induced experimentally in Drosophila and the effects on immunity studied [31]. In flies, one particular kind of ECM, the basement membrane (BM) covers all internal organs and separates them from the hemolymph in the open circulatory system. The BM is created through the release of its components from the fat body into the hemolymph and subsequent deposition onto the basal site of tissues, including the fat body itself. Hemocytes contribute to BM formation or its repair [32–37], although the relative contribution of hemocytes versus fat body may be variable [38]. In one study, the formation of BM deposits with similarity to fibrotic lesions was induced in the fat body by the plasma membrane overgrowth or alternatively through increased secretion of immune effectors. Both scenarios led to a damage response, including melanization [31]. Melanization was also observed by simultaneous disruption of the BM and loss of cell polarity [39]. Similarly, the Drosophila ECM protein SPARC (Secreted Protein, Acidic and Rich in Cysteine, also called BM40) was found to contribute to age-related cardiac fibrotic deposits, associated with reduced life span in flies [40]. Here, we show that ECM components are dysregulated in flies that express a dominant-active oncogene (RasV12) in the salivary glands (SGs; [29]). SGs have so far been mostly

![Fig. 1. Dysplasia promotes formation of fibrosis and the loss of epithelial organization. a PBS and PBST-treated wild type and RasV12-expressing glands were stained with antibodies against SPARC, Lam, Ndg, Pcan, and actin. b–c Quantification of collagen IV (Vkg-GFP) thickness in wild type and RasV12 glands shows an increase in BM thickness (p < 0.0358). d Extracellular F-actin accumulates on the surface of the RasV12 glands (detected by Ph). e–g The intensity of SPARC and actin covary (e, f; R^2 0.4524) and (g) colocalize (R^2 0.3961). ROI is indicated in yellow and displayed as an unprocessed z-stack showing moderate colocalization of SPARC (red) and phalloidin (green). The scale bars (a, d, e) represent 100 μm. PBS, phosphate buffered saline; SPARC, Secreted Protein, Acidic and Rich in Cysteine; Lam, laminin; Ndg, nidogen; Pcan, perlecain; Ph, phalloidin; BM, basement membrane; ROI, region of interest.](For figure see next page.)
Supplementary Figure 2. (a) Representative images of the extracellular (left) and extracellular/intracellular (right) collagen IV layers across different genotypes. (b) Graph showing the comparison of collagen IV thickness between W1118 and RasV12 genotypes. (c) Representative images of Vkg-GFP expression in W1118 and RasV12 genotypes. (d) Representative images of SPARC localization in W1118 and RasV12 genotypes. (e) Representative images of SPARC-Ph co-localization in W1118 and RasV12 genotypes. (f) Graph showing the correlation between Phalloidin intensity and SPARC intensity in RasV12 genotypes. (g) Representative images of SPARC-Ph co-localization in a specific region of interest (ROI) for SPARC-Ph.
studied as models for organ development [41–44] and for innate immunity [45]. Proper secretion of salivary glue proteins and other mucin-type secretions in preparation for pupation is indicative of normal SG function [44] and is expected to be negatively affected by the formation of fibrosis [4], and this is what we observe. The fibrotic phenotype is strongest in the distal, secretory part of the SGs where ECM dysregulation coincides with the recruitment of transglutaminase and phenoloxidase and of hemocytes. In contrast, the proximal part of SGs displays an almost normal histology despite strong activation of AMPs, most notably the AMP Drs. Due to the similarities to fibrotic lesions in mammals, we refer to the regions with dysregulated ECM deposition concomitant with inflammatory reactions [46] as fibrotic deposits, although we acknowledge some differences to mammalian fibrosis [47]. Surprisingly, forced SG-wide expression of Drs, including distal SGs, strongly reverted fibrosis and restored tissue integrity in distal SGs and secretory activity to almost normal levels. This indicates that depending on the type of response, innate immune reactions may have pro- and anti-fibrotic consequences and establishes AMPs as regulators of tissue homeostasis.

Fig. 2. Activation of TG activity in RasV12-expressing glands. a, b Higher TG activity (TQ in red) was detected on RasV12-expressing SGs in comparison to w1118 SGs (quantified in b). Upper ROI insets (a: yellow) are magnified in the lower panel. c TG (red) activity covaries with Ph (green, $p < 0.0001$). d Colocalization analysis showed a moderate overlap between TQ and Ph (turquoise, $R^2 0.4725$). ROI inset (d: yellow) displays an unprocessed z-stack of TQ (red) and Ph (green). The scale bars in (a) correspond to 100 μm (2 upper panels) and 50 μm (ROI). TG activity, transglutaminase activity; ROI, region of interest; SGs, salivary glands; Ph, phalloidin.
Material and Methods

**Fly Strains and Sample Preparation**

*w^{118} Beadex^{Mbi1096-Gal4} (Referred here as Bx: 8860/Bl), w^{118}, w^{118}, UAS-Ras^{V12} (4847/Bl), w^{118}, UAS-Drs (Drs-OE; overexpression [46], w^{118}, Vkg^{50045/CyO.GFP} [48], UAS-Drs-HA [49], Bx; mCD8:RFP [46] and Sgs3-GFP (5884/Bl) (see also online suppl. Table 1; for all online suppl. material, see www.karger.com/doi/10.1159/000516104).

Flies were cultured in 25°C, 12-h dark/light cycle room. Female virgins were collected for 5 days and crossed to the respective male on day 7. Progeny larvae were grown as described in Ref [46]. Ten to twenty salivary gland pairs were fixed in 4% PFA for 20 min. Samples subjected for extracellular staining were washed 3 × 10 min with 1x phosphate buffered saline (PBS). Samples used to stain for intracellular expression were washed 3 × 10 min with PBS-T (PBS with Triton X-100: 1%) before staining.

**Production of Antibodies**

GM02366 (Flybase, [50]) coding for Drosophila SPARC was used as a template for PCR amplification using Takara Ex Taq polymerase (Takara Biomedicals) according to the manufacturer’s instructions. 5 and 3’ primers equipped with suitable restriction enzymes created a PCR fragment spanning the complete open reading frame lacking the SPARC signal peptide that was subsequently ligated in-frame downstream of the mouse BM-40 signal peptide of the episomal expression vector pCEP-Pu [51]. After verification of the sequence, the expression vector was used to transfected human 293-EBNA cells (Invitrogen), and serum-free
medium was collected for protein purification according to established methods [51]. Immunization of rabbits, affinity purification of antibodies, and ELISA titration followed the standard protocols [52].

**Immunohistochemistry**

Antibodies against SPARC (1:3,000: this report), Nidogen (1:2,000) [53], Lam (1:2,000) [54], and Pcan (1:2,000) [55] were incubated overnight (ON) at 4°C either in PBS (non-permeabilized) or PBS-T (permeabilized). PPO1 (1:500), PPO2 (1:250), and tiggrin (1:500 [56], a kind gift by A. Simmonds, Alberta) were visualized) or PBS-T (permeabilized). PPO1 (1:500), PPO2 (1:250), and Lam (1:2,000) [54], and Pcan (1:2,000) [55] were incubated overnight (ON) at 4°C, ON. Anti-HA (1:3,000, Thermofisher #5B1D10) was pre-stained in PBS at 4°C ON. Anti-β-integrin (1:200, DSHB #CF.6G11) and Anti-DIg (1:50, DSHB #4F3, [44]) was incubated at 4°C, ON. Anti-HA (1:3,000, Thermofisher #5B1D10) was pre-incubated with fixed RasV12 salivary glands for 1 h at room temperature (RT) in PBS-T and subsequently incubated with the samples for 1 h at RT. Samples were washed 3 × 10 min with PBS/PBS-T and incubated with secondary antibody Anti-Rabbit-568 (1:500, Thermofisher #A21069), Anti-Rabbit-488 (1:100, Thermofisher #A11008) or Anti-Mouse-546 (1:500, Thermofisher #A11030), DAPI (1:500, Sigma-Aldrich D95942), and Phalloidin-488/546 (1:500, Sigma-Aldrich #A12379 and #A22283, respectively) for 1 h at RT, washed 3 × 10 min with PBS or PBS-T before mounting in Fluoromount-G.

**Transglutaminase Activity**

Salivary glands were fixed in 4% PFA for 20 min and washed 3 × 10 min with PBS at RT. The glands were incubated with TQ, an antibody that recognizes isopeptide (ε-[ϒ-L-glutamyl]-L-lysine) generated by transglutaminase (1:100, Covalab #mab0012), in PBS at 4°C ON. Anti-β-integrin was incubated for 1 h at RT. Samples were washed 3 × 10 min in PBS. Primary antibody was detected with Anti-Mouse-546 (1:500, Thermofisher #A11030) for 1 h at RT. Thereafter, the samples were washed 3 × 10 min in PBS and subsequently mounted in Fluoromount-G. Image acquisition and analysis is described below.

**Image Acquisition and Analysis**

Whole SGs were photographed with a Zeiss AxioscopeII microscope and images were exported as TIFF files. The intensity was measured of the whole gland with ImageJ (ver. 1.52n). For co-localization of the intensity, the signal (SPARC, TQ, PPO1, PPO2, or HA) of the individual salivary glands was plotted against the phalloidin intensity signal of the same gland. Representative images were captured using a confocal Zeiss LSM780 microscope. Final figures were made on Affinity Designer (version 1.7.3). Z-stack images were taken with LSM780 (objective 40x/1.40 Oil DIC M27) and colocalization analysis was analyzed on Imaris (version 1.52n, coloc module). Statistics was performed in GraphPad (8.3.0), including D’Agostino for normal distribution, unpaired t test, Pearson’s correlation, and linear regression.

**Basal Membrane Thickness Analysis**

Z-stacked membranes were analyzed on a Y-Z plane in ImageJ (ver. 1.52n). The “straight line” tool was used to measure the thickness of Collagen IV (Vkg-GFP), presented in pixels. Ten measurements across a Y-Z plane were averaged per salivary gland. A minimum of three salivary glands per genotype were measured. An unpaired t test was performed in GraphPad (8.3.0).

**Reverse-Transcription Quantitative PCR**

RNA extraction, cDNA synthesis, and qPCR analysis were performed as described in Ref [46]. qPCR was performed on 3 independent replicates targeting Rpl32, Sgs3, and Eif73Ee. Primer sequence for Rpl32: 5′-GGATCGATATGCTA-3′ and 5′-CCGACCTCTGTGTTG-3′, Sgs3: 5′-GTGCTAAGAGGAGTGCACTTG-3′ and 5′-AGACGACATTGACGGATCTTGCG-3′, and Eif73Ee: 5′-CTAATGTGTCGTCGTTAGTGG-3′ and 5′-CAACGCCTTTCTCAATTACCTCCA-3′.

**Western Blot**

Salivary glands from 10-staged larvae were dissected and homogenized with a micropestle in Laemmli buffer supplemented with Complete (Roche) and DTT. Thereafter, the samples were incubated for 10 min at 80°C. Ten microliters of lysate was loaded onto a 6% SDS gel. Proteins were transferred to a 0.45-μm nitrocellulose membrane (Biorad: #1620115) and subsequently incubated with PNG Peroxidase (Sigma-Aldrich #L7759, 1:1,000) and antibody to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands, compared to collagen IV – its signal intensity increased in RasV12 glands.

**Results**

**A Fibrosis Model in Drosophila Salivary Glands**

We had previously shown that expression of a dominant-active version of Ras (RasV12) driven by Beadex-Gal4 (referred here as RasV12) in SGs [29, 57] leads to an increase in the signal intensity, redistribution, and disruption of collagen IV in the BM [46]. Here, we initially analyzed the expression of SPARC, which is known to directly interact with collagen IV and found that – similar to collagen IV – its signal intensity increased in RasV12 SGs (Fig. 1a; quantified in online suppl. Fig. 1a). The distribution of SPARC and 3 additional ECM components

![Fig. 4. Drs expression prevents fibrosis in RasV12 glands. a Drs-OE, RasV12 leads to reduced expression of SPARC and F-actin (Ph, quantified in b; p = 0.0004) compared to RasV12 SGs. c, d Collagen IV thickness was restored to control levels in Drs-OE, RasV12 glands, in comparison to RasV12 SGs (p < 0.001). e, f TG activity (TQ, quantified in f; p = 0.0093) and PPO1 (e, quantified in g; p > 0.0001), in comparison to RasV12 SGs was also reduced upon co-expression of Drs. Scale bars in (a, d) represent 100 μm. Drs-OE, RasV12, drosomycin overexpression in RasV12 glands; SPARC, Secreted Protein, Acidic and Rich in Cysteine; SGs, salivary glands; TG activity, transglutaminase activity, Ph, phalloidin.](image-url)

(For figure see next page.)
Figure 1: Effects of RasV12 expression and Drs-OE on collagen IV and TQ intensity.

(a) Representative images of collagen IV and TQ expression in different genotypes. (b) Graph showing phalloidin intensity in different genotypes (p<0.001 for Drs-OE vs. wild type, p=0.0032 for RasV12 vs. wild type, p=0.0004 for Drs-OE,RasV12 vs. wild type). (c) Graph showing collagen IV thickness in different genotypes (p<0.0001 for Drs-OE vs. wild type, p=0.0093 for RasV12 vs. wild type, p=0.0032 for Drs-OE,RasV12 vs. wild type). (d) Representative images of Vkg-GFP expression in different genotypes. (e) Representative images of PPO1 expression in different genotypes. (f, g) Graphs showing TQ and PPO1 intensity in different genotypes (p<0.0001 for Drs-OE vs. wild type, p<0.0001 for RasV12 vs. wild type, p=0.0004 for Drs-OE,RasV12 vs. wild type).
An Antimicrobial Peptide Reverts Fibrosis

RasV12, Drs-OE expressing glands (Fig. 1a, extra/intracellular). Under both conditions, the regular hexagonal pattern of all BM components in wild-type SGs was replaced by a more irregular distribution in RasV12-expressing glands. All 4 proteins displayed an increased signal intensity compared to wild-type controls (Fig. 1a). To exclude differences in accessibility to antibodies as an explanation for the difference in the signal intensity, we used GFP-tagged collagen IV both for quantification and to measure BM width. This confirmed that compared to control glands, the BM from RasV12 SGs contained more collagen IV and had also grown thicker (Fig. 1b, c). We conclude that the BM in RasV12 glands had become disorganized and more extensive, reminiscent of the fibrotic phenotype, which had been observed in Drosophila fat bodies [31] and of mammalian fibrotic lesions. Of note, the fibrotic phenotype was most pronounced in the distal parts of SGs similar to other dysplastic features in the same region.

Although initially used as a control to distinguish between normal and dysplastic SGs, F-actin staining with phalloidin largely mimicked the disorganized distribution of the BM components and was also detected in RasV12 SGs in the absence of detergent on surface sections, indicating that some F-actin may have obtained access to the extracellular compartment (Fig. 1d; quantified in online suppl. Fig. 1b). This was further confirmed by showing significant covariation between F-actin and SPARC on the surface of whole glands (Fig. 1e, f). Additionally, by analyzing Z-scanned SGs, we observed a moderate SPARC-phalloidin colocalization (Fig. 1g).

Thus, we find that in the distal part of RasV12 SGs, BM components are redistributed in concert leading to a fibrotic phenotype including F-actin, part of which may leak out from SG cells and associate with the BM.

Fig. 5. The epithelial character and production of secreted proteins is restored upon co-expressing Drs in RasV12 glands. a Staining of control (w1118; and Drs-OE), RasV12-expressing as well as Drs co-expressing glands (Drs-OE, RasV12) with integrin-specific antibodies, Ph, and Dlg shows a partial reversion to an epithelial phenotype. b The production of Sgs3 (Sgs3-GFP) is restored in Drs-OE, RasV12 glands compared to RasV12 glands (quantified in c: p < 0.0001) in line with the restoration of the SG lumen (detected with Ph). d Western blot analysis shows increased production of Ei-g71Ee (gp150, N = 3), in Drs-OE, RasV12 glands (d: p < 0.0114) (gp150, N = 3). e qRT-PCR analysis of Sgs3 and Eig71Ee shows increased expression in Drs-OE, RasV12 glands in comparison to RasV12. Scale bars represent 20 μm (a) and 100 μm (b). Drs-OE, drosomycin-overexpressing; Dlg, disc-large; SGs, salivary glands; Ph, phalloidin; RT-qPCR, reverse-transcription quantitative PCR; Sgs3, salivary glands secreted protein 3.

The Immune System Targets Dysplastic Salivary Glands

To assess the influence of hemostasis on fibrosis, we first measured the activity of Drosophila transglutaminase, which is a key clotting factor and has been shown to target intruders such as bacteria and insect pathogenic nematodes [26]. Mammalian TGs are known to be activated in fibrotic lesions in several organs [13, 14, 58, 59]. Activity of the single Drosophila TG was detected with an antibody against the covalent ε-(Y-1-glutamyl)-1-lysine links that are created by TG and, to focus on extracellular TG, we used non-permeabilized SGs and surface sections (Fig. 2). Compared to wild-type glands, significantly higher TG activity was detected on RasV12 SGs, indicating that – similar to microbial intruders – dysplastic glands are targeted by TG (Fig. 2a, b). In contrast to wild type, where it was faint, TG activity in RasV12 SGs was found to covary (Fig. 2c) and moderately colocalize with the phalloidin intensity (Fig. 2d). In addition to TG, tiggrin, which is also part of Drosophila hemolymph clots [60], was recruited to RasV12 SGs, further confirming that hemostasis is activated against dysplastic SGs (online suppl. Fig. 1c, d).

Ultimately, coagulation of hemolymph, the formation of nodules, and the encapsulation of larger objects result in the recruitment and activation of PPO. During coagulation, this follows TG activation [22]. To test whether PPOs are also recruited to dysplastic RasV12 glands, we used antibodies specific for PPO1 and PPO2, the 2 PPOs expressed by crystal cells and found that both of them were detected on RasV12-expressing glands (online suppl. Fig. 1e–g). Further analysis showed that PPO1 intensity and phalloidin staining covaried, whereas PPO2 did not (online suppl. Fig. 1h, i). In summary, these data show that beyond induction of antimicrobial peptides and hepcocyte recruitment [46], RasV12 SGs display local activation of the hemostatic system. This includes structural clot components, such as tiggrin, as well as enzymatic activities, such as transglutaminase, followed by recruitment of phenoloxidases.

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**Fig. 6.** Drs colocalizes with F-actin in fibrotic lesions. 

**a** Drs-HA expressed in wild-type and *Ras*V12*-expressing SGs was detected using HA-specific antibodies as well as Phalloidin. Then upper ROI insets (a: yellow) are magnified in the lower panels. 

**b** Partial overlap between the signals was confirmed using confocal microscopy including Z-stacks. ROI inset (b: yellow) displays an unprocessed Z-stack of Drs-HA (red) and Ph (green). Drs, drosomycin; SGs, salivary glands; ROI, region of interest; Drs-HA, HA-tagged Drs; Ph, phalloidin.
Defective Secretion in the Dysplastic Gland

Having observed fibrosis in dysplastic glands prompted us to investigate the integrity and secretory potential of SGs. As indicated by [39], in addition to a disrupted BM, a second precondition for activation of an inflammatory reaction against the aberrant tissue is a loss of apico-basal polarity. To test whether SG epithelial cells had undergone similar changes, we assessed the distribution of β-integrin, F-actin, and disc-large (Dlg) in permeabilized SGs. β-integrin, one of the receptors for BM components, shows an even surface expression and enrichment at the basal side in wild type SG cells. This pattern had been lost upon RasV12 expression, together with a complete loss of the SG lumen detected by phalloidin staining (Fig. 3a). Similarly, the regular hexagonal pattern detected with Dlg-specific antibodies was replaced by a weaker irregular pattern in RasV12 SGs. To test the secretory potential of RasV12 SGs, we assessed the secretion of 2 known SG secretory proteins, salivary glands secreted protein 3 (Sgs3) and Eig71Ee [61, 62]. We first investigated the intensity levels of Sgs3 using a GFP-tagged version under its endogenous promoter. In comparison to control SGs, the dysplastic glands displayed lower Sgs3 levels, in line with the absence of a lumen (Fig. 3b, c). Western blotting and Coomassie staining both revealed lower levels of Eig71Ee in RasV12 glands than in control glands (Fig. 3d; online suppl. Fig. 2a). Altogether, dysplastic glands displayed a loss of epithelial character and produced lower amounts of 2 main secretory proteins. Nevertheless, most proteins were still shared with wild type glands and no general degradation was observed (online suppl. Fig. 2a).

An Antimicrobial Peptide Reverts BM Dysplasia, Loss of Epithelial Character, and Secretion

We had previously found that Drs is expressed most strongly in the proximal region of RasV12 glands, a region which displayed fewer signs of fibrosis and looked more similar to wild type SGs. We also managed to revert collagen IV deregulation by expressing Drs across the whole glands including the strongly affected distal part [46], while expression of control GFP [29] and membrane-bound RFP [46] under the same UAS control had no such effects. Therefore, we wondered whether the fibrotic distribution of other BM components was rescued upon forced Drs expression across the whole glands. Indeed, in Drs-overexpressing (OE) RasV12 SGs, SPARC, Lam, Ndg, Pcan, and actin displayed a more regular localization, closer to the wild type situation in surface sections (Fig. 4a, b; online suppl. Fig. 2b). This was not observed upon RFP overexpression excluding a dilution effect due to the presence of dual UAS targets (online suppl. Fig. 2d). In line with the more regular distribution of BM components,
BM thickness in Drs-OE Ras\textsuperscript{V12} SGs were also reduced to wild type levels (Fig. 4c, d). Moreover, transglutaminase activity and PPO1 binding (Fig. 4e–g) decreased, although not completely, when compared to Ras\textsuperscript{V12}-expressing SGs, whereas PPO2 (online suppl. Fig. 2c) did not revert to wild type levels. At the same time, staining with phalloidin confirmed that F-actin had been reduced to almost wild type levels (Fig. 4b).

To examine the secretory activity in normal, Ras\textsuperscript{V12}-expressing, and Drs-OE SGs, we compared the SG histology and the production of Sgs3 and Eig71Ee. In line with the renewal of a more normal BM morphology and the restoration of the epithelial character, as assayed by integrin, Dlg, and phalloidin staining (Fig. 5a), the lumen of Drs-OE, Ras\textsuperscript{V12} SGs was also partially restored allowing production of both Sgs3 (Fig. 5b, c, e) and Eig71Ee (Fig. 5d, e), in comparison to the Ras\textsuperscript{V12} glands (Fig. 5b). Sgs3 levels in Ras\textsuperscript{V12} SGs were not restored when RFP was co-expressed as a negative control (online suppl. Fig. 2f). Both Western blots (Fig. 5d) confirmed higher levels of Eig71Ee in Drs-OE, Ras\textsuperscript{V12} glands.

Drs Localization in Fibrotic Lesions

To detect whether Drs is properly located to interfere with the formation of fibrotic lesions in Ras\textsuperscript{V12} SGs, we used an HA-tagged version (Drs-HA) to analyze surface sections (Fig. 6). Since it has previously been shown that AMPs have the potential to bind actin [63] and based on our own results (see above), we used phalloidin as a proxy to identify fibrotic lesions. Signals for both phalloidin and Drs-HA were detected in Ras\textsuperscript{V12} SGs and a weak signal upon expression of Drs-HA alone in a wild type background (Fig. 6a). In both cases, Drs-HA overlapped with the phalloidin signal, indicating partial colocalization with F-actin, which was confirmed in Z-sections (Fig. 6b). Taken together, the distribution of Drs in Ras\textsuperscript{V12} SGs is compatible with its inhibitory effect on fibrotic lesions.

Discussion

We show that expression of a dominant-active form of an oncogene induces a fibrotic phenotype at the BM of Drosophila SGs, a highly active secretory organ (summarized in Fig. 7). In contrast to other Drosophila fibrosis models, the redistribution of BM components, which are primarily produced in the fat body ([38], see also online suppl. Table 2) and deposited onto the internal organs, is most likely a secondary consequence of SG dysplasia. Although we observe an increase in signal intensity for all BM components in Ras\textsuperscript{V12} SGs, this appears not to be a consequence of transcriptional activation, since the BM components are only poor if at all induced in the fat body of Ras\textsuperscript{V12} larvae (see online suppl. Table 2). As a proxy for BM components, SPARC is found associated with extracellular F-actin, which appears to be released from the dysplastic cells either SG cells or rupturing crystal cells [22]. When analyzing, transglutaminase activity as a proxy for hemostasis, we found increased levels in Ras\textsuperscript{V12} SGs. Similar to what happens during wound closure in flies, TG activation was followed by recruitment of Phennoloxidase, although in this case only the PPO1 signal covaried with phalloidin staining of F-actin. Taken together we detect fibrotic lesions, which contain BM components, TG activity, PPO1, and extracellular F-actin. These findings are in line with the proposed model for fibrotic lesions as unabated wound healing. Although we were initially motivated by a possible hemostatic/immune function of TG, the single Drosophila TG may play a slightly different role during fibrosis similar to mammalian tissue transglutaminase (TG2), which has been shown to aggravate fibrosis in several organs due to its cross-linking activity [13, 14, 58, 59]. SG fibrosis affects the normal secretory function of SGs as shown here by: (1) a loss of the epithelial character of SG cells, (2) the collapse of the SG lumen, and (3) reduced production of 2 secretory proteins (Sgs3 and Eig71Ee).

Supporting our previous observations that single expression of the AMP Drs rescues several features of the dysplastic phenotype [46], we found that upon Drs overexpression, the formation of fibrotic lesions at the BM is substantially reduced and tissue integrity is largely restored including a partial reversion to an epithelial character of SG cells and restoration of secretory activity (summarized in Fig. 7). This is paralleled by a reduction in hemostatic/immune activation including (1) fewer TG cross-links and (2) a reduction in recruitment of PPO1, but not PPO2. These findings support the notion that early hemostatic components are recruited toward fibrotic foci, similar to wounds were PPO1 and TG activity have been shown to precede PPO2, which is subsequently released by rupture of crystal cells [22, 64].

Finally, we observe that Drs colocalizes with F-actin in fibrotic lesions similar to the mammalian AMP LL-37 which has previously been shown to interact with F-actin [63, 65] with different effects, including protection from microbial degradation [66], but also inhibition of the antimicrobial activity [67]. Due to high background staining, we were unable to determine whether Drs is also secreted apically into the SG lumen. Therefore, we do not
know whether Drs equally influences the apical and basal secretion of SG cells. For the basal localization, we propose a scenario where secreted Drs reduces the formation of fibrotic plaques by interfering with the aggregation of BM components. These may include particulate actin (F-actin), which, once released from cells, is known to have deleterious effects, for example, during sepsis [68], by acting as a procoagulant [69], and through inhibition of macrophage defenses [70]. These effects are counteracted by actin-scavenging systems [69], which include soluble gelsolin [71], vitamin D-binding protein [72], and – as proposed here – Drs. In line with potential role as damage-associated molecular patterns [73–75], cytosolic proteins have been shown to activate immunity in flies [76, 77]. Irrespective of its target in fibrotic foci, the anti-fibrotic effects of Drs are of clinical interest and may include other AMPs, such as the bee AMP melittin [78–80], which restores the epithelial character [81] similar to what we observe. Taken together, the data presented here support the concept that dysplasia of tumor cells supports the formation of fibrotic aggregates [6, 10]. The recruitment of both TG and PO, which are known players in hemolymph coagulation, lends further support to the concept that tumors induce a chronic procoagulatory state [82, 83] and adds the notion that AMPs have the potential to revert this phenotype.

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Statement of Ethics

No approval of studies involving animals was required.

Conflict of Interest Statement

The authors declare no conflict of interest.

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Author Contributions

Experiment design: D.K., S.B., and U.T.; experimental work: D.K., C.K., and S.B.; raising funding: S.B. and U.T.; manuscript drafting: D.K., S.B., and U.T.

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