Pak1 control of E-cadherin endocytosis regulates salivary gland lumen size and shape

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SUMMARY
Generating and maintaining proper lumen size and shape in tubular organs is essential for organ function. Here, we demonstrate a novel role for p21-activated kinase 1 (Pak1) in defining the size and shape of the Drosophila embryonic salivary gland lumen by regulating the size and elongation of the apical domain of individual cells. Pak1 mediates these effects by decreasing and increasing E-cadherin levels at the adherens junctions and basolateral membrane, respectively, through Rab5- and Dynamin-dependent endocytosis. We also demonstrate that Cdc42 and Merlin act together with Pak1 to control lumen size. A role for Pak1 in E-cadherin endocytosis is supported by our studies of constitutively active Pak1, which induces the formation of multiple intercellular lumens in the salivary gland in a manner dependent on Rab5, Dynamin and Merlin. These studies demonstrate a novel and crucial role for Pak1 and E-cadherin endocytosis in determining lumen size and shape, and also identify a mechanism for multiple lumen formation, a poorly understood process that occurs in normal embryonic development and pathological conditions.

KEYWORDS: Cadherin, Endocytosis, Pak, Rab5, Salivary gland, Lumen and tube morphogenesis, Cdc42, Merlin, Drosophila

INTRODUCTION
Tubular networks provide structure and function for many essential organs, such as the vasculature, gut, lung and kidney. Tube formation occurs through a complex series of events including cell shape changes, cell migration and changes in cell adhesion (Lubarsky and Krasnow, 2003; Martin-Belmonte and Mostov, 2008). Central to the morphogenesis of tubular organs is the formation of a lumen of the correct size and shape that mediates the delivery of gases and nutrients or the removal of waste. Failure to achieve and/or maintain correct lumen size and shape can lead to pathological conditions such as polycystic kidney disease and stenoses. Lumens can form by a number of mechanisms, such as through folding, as occurs in the mammalian and avian neural tube, or de novo through apoptosis and autophagy, as occurs in zebrafish and Jaskoll, 2000). Lumens can also form de novo by the formation of intracellular vesicles, as occurs in zebrafish and cultured endothelial cells (Kamei et al., 2006; Koh et al., 2008). Alternatively, lumens can form extracellularly between adjacent endothelial cells, as was recently demonstrated in the mouse aorta (Strilic et al., 2009).

Recent in vitro studies of lumen formation using cultured cells grown as cysts in three-dimensional matrices revealed an essential role for the Rho family GTPases Cdc42 in de novo lumen formation. In MDCK cysts, reduction of Cdc42 activity delayed single lumen formation and led to formation of cysts with multiple small lumens (Martin-Belmonte et al., 2007). Knockdown of Cdc42 in Caco-2 cells led to cysts with two or more lumens due to a defect in the orientation of the mitotic spindle (Jaffe et al., 2008). Although these studies provided mechanistic insight into Cdc42-dependent lumen formation in vitro, it is not known whether the same mechanism holds true for lumen formation in vivo in a developing tubular organ.

Pak proteins are a family of serine-threonine kinases known to bind and be activated by the Rho family GTPases Cdc42 and Rac to regulate a number of diverse biological processes, including cell migration and cytoskeletal rearrangements (Arias-Romero and Chernoff, 2008; Bokoch, 2003). Although much is known about the structure, biochemistry and cell biology of mammalian Paks, less is known about their biological role in vivo. Studies of cultured mammalian cells showed that Paks are required for lamellipodium extension and for disruption of E-cadherin-based cell-cell junctions (Kiosses et al., 1999; Lozano et al., 2008; Royal et al., 2000; Zegers et al., 2003). Mutations in zebrafish Pak2a cause loss of vascular integrity characterized by abnormally formed blood vessels (Buchner et al., 2007; Liu et al., 2007). In cysts formed from cultured endothelial cells, Pak proteins act downstream of Cdc42 to regulate de novo lumen formation (Koh et al., 2008; Koh et al., 2009). Thus, these studies suggest a crucial role for Pak kinases in tube and lumen morphogenesis; however, the mechanism by which Paks act is not known.

The Drosophila embryonic salivary gland is a genetically tractable model system for investigating tube and lumen morphogenesis. It consists of a pair of elongated epithelial tubes that are required for secretion during the larval stage. Each gland comprises a single layer of epithelial cells surrounding a central lumen. The gland lumen is formed during the process of invagination when primordial cells invaginate from the ventral surface of the embryo (Myat, 2005). Once internalized, gland cells undergo a phase of robust apical membrane growth that is regulated by the transcription factors Hairy, Huckebein (Hkb) and Ribbon, the apical membrane protein Crumbs, and a microtubule motor encoded by the klarsicht gene (Kerman et al., 2008; Myat and Andrew, 2002). As internalized gland cells begin to migrate...
posteriorly, the expanded apical membrane is re-absorbed and the apical domain of each cell becomes elongated with its longest axis along the proximal-distal (Pr-Di) axis of the gland through an unidentified mechanism (Myat and Andrew, 2002).

A role for the Drosophila Pak4 homolog Mbt in the regulation of E-cadherin (Shotgun – FlyBase)-based adherens junctions and a role for Drosophila Pak1 (Pak – FlyBase) in apicobasal polarity and F-actin assembly have been demonstrated (Menzel et al., 2008; Menzel et al., 2007). Here, we demonstrate a novel role for Drosophila Pak1 in the control of salivary gland lumen size and shape through Merlin and Rab5- and Dynamin (Shibire – FlyBase)-mediated E-cadherin endocytosis.

MATERIALS AND METHODS

Drosophila strains and genetics

Fly lines used were: UAS-Cdc42-GFP, UAS-Rab5WT, UAS-Rab7WT, UAS-Rab11WT, UAS-Rab11TMYFP; UAS-Rab53XN-YFP; UAS-Rab722ZN-YFP; UAS-Rab1122ZN-YFP; UAS-shild4A4, UAS-Rab5WT-YFP; UAS-Rac1141, Df(3R)Win11, shg (Bloomingtom Stockton Center); UAS-Rab5WT-GFP, UAS-Rab7WT-GFP, UAS-Rab11WT-GFP, UAS-LAMP1-GFP (C. Samakovlis, Stockholm University, Stockholm, Sweden); UAS-E-cadherin-GFP (H. Oda, Osaka University, Osaka, Japan); UAS-p35 (H. Stellar, The Rockefeller University, New York, NY, USA); fkh-GAL414, fkh-GAL414 (D. Andrew, Johns Hopkins University, School of Medicine, Baltimore, MD, USA); Pak114, Pak116, Pak16, UAS-Pak114, UAS-Pak116, Pak16, Pak116, Pak114 (N. Harden, Simon Fraser University, Burnaby, B.C. Canada); UAS-MerPxxP-myc, UAS-MerPxxP-myc (R. Fehon, The University of Chicago, Chicago, IL, USA). Canton-S flies were used as wild-type controls. The UAS-Gal4 system (Brand and Perrimon, 1993) was used with fkh-GAL4 to drive salivary gland-specific expression. Homozygous mutant embryos were distinguished by the absence of β-galactosidase staining, which detects the expression of lacZ from the Ubx-lacZ insert on the balancer chromosomes.

Antibody staining of embryos

Embryo fixation and staining were performed as described previously (Pirraglia et al., 2006). Primary antibodies used were: anti-Phalloidin (Sigma) and visualized on a Zeiss Axioplan 2 microscope with Axiovision Diaminobenzidine-stained embryos were mounted in methyl salicylate and analyzed on a Joel 100CX-11 Transmission Electron Microscope described previously (Myat and Andrew, 2002). Thin serial sections were generated and analyzed on a Joel 100CX-11 Transmission Electron Microscope at the Weil Cornell University Microscopy Core Facility (New York, NY, USA). At least three embryos were analyzed for each genotype.

RESULTS

Morphometric analyses

The apical domain area of individual gland cells was measured by tracing E-cadherin (E-cad) immunofluorescent staining at the level of the adherens junctions (AJs) using LSM 510 Image Browser software (Carl Zeiss). The elongation ratio was determined by calculating the ratio of apical domain length oriented along the Pr-Di axis to the apical domain length along the dorsal-ventral (D-V) axis using LSM 510 software. The basolateral perimeter and domain areas of individual gland cells were measured by tracing F-actin staining of laterally viewed stage 13 embryos using LSM 510 software. Lumen length was determined by tracing E-cad-labeled lumens, and lumen width was determined by measuring lumen diameter of the proximal, medial and distal regions of the gland with LSM 510 software. At least five embryos were analyzed for each genotype and a maximum of three cells were measured per gland. Statistical analysis was performed using Microsoft Excel.

Fluorescence intensity analyses

Total fluorescence intensity of E-cad was quantified in embryos double-stained for E-cad and Crb or triple-stained for E-cad, aPKC and Nrt. Total fluorescence intensity of E-cad per apical domain was measured using NIH Image J software (http://rsb.info.nih.gov/ij/) and normalized against the total fluorescence intensity of Crb or aPKC. Total fluorescence intensity of E-cad per basolateral membrane was normalized against the total fluorescence intensity of Nrt. At least five embryos were analyzed for each specified genotype and a maximum of three cells were measured per gland. Statistical analysis was performed using Microsoft Excel.

Analysis of recombinant lines

Recombinant lines carrying different UAS-transgene insertions were confirmed by PCR analysis as described previously (Pirraglia et al., 2006).

Transmission electron microscopy of embryos

Embryos were processed for transmission electron microscopy (TEM) as described previously (Myat and Andrew, 2002). Thin serial sections were generated and analyzed on a Joel 100CX-11 Transmission Electron Microscope at the Weil Cornell University Microscopy Core Facility (New York, NY, USA). At least three embryos were analyzed for each genotype.
migration. The lumens of Pak114 mutant dorsal trunk and visceral branches were stalled and not extended like those of heterozygous siblings, suggesting a defect in tracheal migration and/or lumen extension; however, tracheal lumens were not expanded as in the salivary gland (see Fig. S2 in the supplementary material).

To understand how gland lumen size changes over the course of gland development, we measured lumen length and width in glands of wild-type embryos between stage 11 (after invagination is complete) to stage 13 (when glands have turned posteriorly). Lumen length more than tripled during this timecourse, whereas changes in lumen width varied; lumen width in the proximal region of the gland decreased the most over time, whereas no changes in lumen width were observed in the medial and distal portions of the gland (see Fig. S3B,C in the supplementary material).

During early development of wild-type glands, apical membrane growth accompanied elongation of the apical domain of individual gland cells with their longest axis being the Pr-Di axis (Myat and Andrew, 2002). To assess how changes in the apical domain contribute to the overall shape and size of the gland lumen, we measured changes in apical domain area and elongation.

![Diagram of a lateral (a) and apical (b) view of a gland cell showing the apical (red) and basolateral (blue) membranes, adherens junctions (green) and the plane in which optical sections were acquired (gray) to measure the apical domain area and elongation ratio of individual cells.](image)

**Fig. 1. Pak1 is required for correct lumen shape and size during salivary gland migration in Drosophila.** (A-F') Embryos were stained for Crb to label the apical membrane. In Pak114 heterozygous embryos (A,C), glands migrate posteriorly to form elongated tubes (arrows), whereas in homozygous embryos (B,D), the gland lumen is irregular in shape and expanded (arrowheads). In Pak114 heterozygous embryos (E,E'), Crb outlines the gland lumen and the apical domains of individual cells (solid white outline) which are smaller and more elongated in the Pr-Di axis, compared with homozygous siblings (F,F'; solid white outline). Dotted white lines in E and F outline the gland. E' and F' are magnified views of the dotted boxed regions in E and F, respectively. (G) Diagram of a lateral (a) and apical (b) view of a gland cell showing the apical (red) and basolateral (blue) membranes, adherens junctions (green) and the plane in which optical sections were acquired (gray) to measure the apical domain area and elongation ratio of individual cells. (H,I) Measurements of apical domain area (H) and elongation ratio (I) in wild-type and Pak114 salivary gland cells. n represents the total number of cells scored per genotype, with a maximum of three cells measured per gland. Data are mean ± s.d. **P<0.005. (J-L) In Pak114 heterozygous glands (J), the apical membrane facing the lumen consists of numerous membrane extensions containing vesicular structures (small arrows), some of which are in contact with the luminal space (arrowhead). In Pak114 mutant glands (K), and glands expressing shiK44A (L), the apical surface is devoid of membrane extensions and lacks vesicular structures, and instead is studded with electron-dense patches (arrows within insets in K and L) compared with wild type (arrow within inset in J), which has none. A, apical; B, basal; D, dorsal; Di, distal; L, lumen; Pr, proximal; V, ventral. Scale bars: 20 µm in A; 10 µm in E and E'; 1 µm in J.
stages 12 and 13), cells progressively reduced their apical domain size (see Fig. S3D in the supplementary material). We also observed changes in apical domain shape during gland turning. In early stage 11 gland cells, the apical domain was isotropic, whereas during gland turning, the apical domain became anisotropic and was three times as long in the Pr-Di direction as in the D-V axis (see Fig. S3E in the supplementary material). After turning was complete, the apical domains became less anisotropic and were twice as long in the Pr-Di direction as in the D-V direction (see Fig. S3E in the supplementary material).

Loss of Pak1 function resulted in changes in gland lumen size. The gland lumen was wider in the medial and distal regions of Pak1\textsuperscript{14} mutant embryos compared with wild-type glands although lumen length was unchanged (see Fig. S4E,F in the supplementary material). Expanded gland lumens of Pak1\textsuperscript{14} mutant embryos did not arise from abnormal cell proliferation as determined by cell counting and staining for phosphorylated histone 1B (data not shown). Instead, increased gland lumen width was accompanied by a two- to three-fold increase in the apical domain area of individual gland cells with a concomitant failure to elongate their apical domains in the Pr-Di axis (Fig. 1E-H). Although apical domain size and elongation were dramatically altered in Pak1\textsuperscript{14} mutant gland cells, the perimeter of the basolateral membrane and the area of the basolateral domain were not affected (see Fig. S3F in the supplementary material).

To determine whether loss of Pak1 affected apical membrane topography, we performed transmission electron microscopy (TEM) of Pak1\textsuperscript{14} homozygous embryos. In the glands of Pak1\textsuperscript{14} heterozygous embryos that have completed invagination, highly irregular membrane extensions protruded into the lumen and contained vesicular structures, some of which were in contact with the lumenal space (Fig. 1J). By contrast, in the glands of Pak1\textsuperscript{4} homozygous embryos, the apical surface was devoid of membrane extensions and vesicular structures, and instead was characterized by electron-dense patches that resembled Dynamin-positive flat-coated surfaces at the plasma membrane of cultured mammalian cells (Fig. 1K) (Damke et al., 1994). Thus, in Pak1 mutant gland cells, not only were the apical domains expanded and failed to elongate, but the topography of the apical surface membrane was also dramatically altered.

### Pak1 functions in salivary gland cells to control E-cadherin localization

To determine whether remodeling of E-cadherin (E-cad)-based adherens junctions (AJs) plays a role in reorganization of the apical domain function, we analyzed E-cad localization in wild-type and Pak1\textsuperscript{14} mutant gland cells. In contrast to wild-type gland cells where E-cad localized to the AJs and to the basolateral membrane (Fig. 2A), in Pak1\textsuperscript{14} mutant glands (Fig. 2B), E-cad localization at the basolateral membrane was lost, although it was still present at the AJs. Measurement of fluorescence intensity demonstrated that Pak1\textsuperscript{14} mutant gland cells contained more E-cad at the AJs and less at the basolateral membrane compared with heterozygous siblings (Fig. 2C,D); however, Crb levels were similar in Pak1\textsuperscript{14} homozygous and heterozygous embryos (data not shown), indicating that Pak1 specifically regulates E-cad at the AJs. The increased expression of E-cad in the AJs in Pak1\textsuperscript{14} mutant gland cells suggested that increased E-cad localization at the AJs might be responsible for the observed defects in apical domain size and elongation. Thus, we analyzed embryos homozygous for a zygotic hypomorph allele of shotgun, the gene which encodes E-cad, shg\textsuperscript{2}. In shg\textsuperscript{2} homozygous embryos, gland lumens were mis-shapen and expanded (data not shown); however, apical domain size was only slightly more expanded than that of wild-type cells and not expanded to the same extent as Pak1\textsuperscript{14} mutant cells (Fig. 2E). Moreover, E-cad was not localized at the basolateral membrane in shg\textsuperscript{2} mutant gland cells (data not shown) and apical domains were not elongated in the Pr-Di axis (Fig. 2F). In embryos trans-heterozygous for Pak1\textsuperscript{14} and shg\textsuperscript{2}, E-cad localized to the AJs and apical domain size was comparable with that of wild-type gland cells; however, E-cad was not found at the basolateral membrane and apical domains failed to elongate (Fig. 2E,F; data not shown). To test whether expansion of the apical domain in Pak1\textsuperscript{14} mutant gland cells was dependent on having normal levels of shg, we analyzed Pak1\textsuperscript{14} mutant glands that were also heterozygous for shg\textsuperscript{2} (Pak1\textsuperscript{14};shg\textsuperscript{2/–}). The apical domain area of Pak1\textsuperscript{14};shg\textsuperscript{2/–} gland cells was significantly reduced compared with Pak1\textsuperscript{14} homozygous mutants; however, the apical domains of Pak1\textsuperscript{14};shg\textsuperscript{2/–} gland cells were not elongated (Fig. 2E,F). From these data, we conclude that Pak1 limits the pool of E-cad at the AJs in order to restrict apical domain size, and promotes E-cad at the basolateral membrane, which correlates with elongation of the apical domain along the Pr-Di axis. These Pak1-mediated events are necessary for establishing the correct lumen size and shape in the salivary gland.

Endogenous Pak1 was expressed ubiquitously in the embryo and in gland cells, and was strongly expressed in the apical membrane and in sub-apical vesicles where it colocalized with the early endosome marker protein Avalanche (Aval; see Fig. S5 in the supplementary material). To demonstrate that Pak1 functions cell-autonomously in gland cells to regulate E-cad localization, we expressed wild-type Pak1 (Pak1\textsuperscript{WT}) specifically in the glands of Pak1\textsuperscript{14}Pak1\textsuperscript{6} trans-heterozygous (Pak1\textsuperscript{14/+};Pak1\textsuperscript{6/+}) embryos. In Pak1\textsuperscript{14/+};Pak1\textsuperscript{6/+} embryos, apical domain size was expanded and apical domains failed to elongate, as was observed also in Pak1\textsuperscript{14} homozygous embryos (see Fig. S3G,H in the supplementary material). We observed a significant rescue of apical domain size and elongation, and basolateral localization of E-cad in Pak1\textsuperscript{14/+};Pak1\textsuperscript{6/+} mutant glands expressing Pak1\textsuperscript{WT} compared with Pak1\textsuperscript{14/+};Pak1\textsuperscript{6/+} mutant glands (see Fig. S3G,H in the supplementary material; data not shown). Thus, Pak1 is required in gland cells for generating proper lumen size and shape through differential localization of E-cad.

### Cdc42 controls salivary gland lumen size

Since Pak proteins are activated by both Cdc42 and Rac small GTPases, we tested whether Cdc42 and/or Rac were required for correct gland lumen size and shape. We inhibited Cdc42 and Rac1 function specifically in the gland by expressing either dominant-negative Cdc42 (Cdc42\textsuperscript{DN}) or Rac1 (Rac1\textsuperscript{DN}) with the fork head (jkh)-GAL4 driver. Cdc42\textsuperscript{DN} expression led to expanded gland lumen width in the medial and distal regions similar to that observed as a result of loss of Pak1 function, whereas Rac1\textsuperscript{DN} expression had no effect (see Fig. S4B,D,E in the supplementary material). Cdc42\textsuperscript{DN} expression had no effect on lumen length, as with loss of Pak1 function; however, Rac1\textsuperscript{DN} expression led to formation of a shorter lumen that is likely to be a result of the role of Rac1 in gland migration that we reported previously (see Fig. S4F in the supplementary material) (Pirrilla et al., 2006). E-cad failed to localize to the basolateral membrane of gland cells expressing Cdc42\textsuperscript{DN}, as observed also in Pak1\textsuperscript{14} mutant gland cells (see Fig. S4B in the supplementary material). By contrast, E-cad localized prominently to the basolateral membrane of gland cells
expressing Rac1DN (see Fig. S4D in the supplementary material). Measurements of apical domain size and elongation showed that Cdc42 inhibition in the gland resulted in expansion of the apical domain and loss of elongation, similar to that induced by loss of Pak1 function; however, Rac1DN had no effect (see Fig. S4G,H in the supplementary material).

Given the similar gland lumen phenotypes of Pak1 and Cdc42 mutant glands, we tested whether Pak1 and Cdc42 could act in a single linear pathway to control lumen size. Inhibition of Cdc42 specifically in glands of Pak114 mutant embryos phenocopied inhibition of Cdc42 or loss of Pak1 alone in terms of gland lumen width, length, apical domain size and elongation (see Fig. S4E-H in the supplementary material), suggesting that Cdc42 and Pak1 act in a single linear pathway. In contrast to Cdc42, expression of Rac1N17 specifically in Pak114 mutant glands decreased lumen length, like inhibition of Rac1 alone, and widened the lumen, like loss of Pak1 alone. Thus, simultaneous loss of Rac1 and Pak1 function altered both lumen width and length suggesting that Rac1 and Pak1 act in parallel pathways. Although Cdc42 was shown to affect apical domain size through regulation of the F-actin cytoskeleton in the Drosophila notum (Georgiou et al., 2008), F-actin structure in gland cells of Pak114 mutant embryos was normal, and embryos mutant for regulators of the actin cytoskeleton, such as WASp, SCAR and Arp2/3 (Arp66B – FlyBase), showed no defects in gland lumen size (data not shown).

Pak1 regulates lumen size and shape through Rab5- and Dynamin-mediated endocytosis

A number of membrane trafficking pathways have been shown to regulate the subcellular localization of E-cad (Delva and Kowalczyk, 2009). For example, E-cad is known to be transported through the recycling endosomes to the plasma membrane in both Drosophila and mammalian cells (Desclozeaux et al., 2008; Langevin et al., 2005; Lock and Stow, 2005). In internalized salivary gland cells that are beginning to undergo apical membrane growth and elongation, E-cad localized to the apicolateral membrane (arrowheads). In Pak114 mutant glands (B,B'), E-cad was absent from the basolateral membrane (arrows) but still localized to the AJs (arrowheads). (C,D) Total fluorescence intensity measurements of E-cad at the AJs (C) and at the basolateral membrane (D) in Pak114 heterozygous and homozygous embryos. (E,F) Measurements of apical domain area (E) and elongation ratio (F). n represents total number of cells scored per genotype, with three cells measured per gland. Data are mean ± s.d. **P<0.0001. Scale bar: 10 μm.
stage 12 wild-type glands when apical domain size was decreasing and elongating. E-cad colocalized with Aval at the AJs, in subapical puncta and at the basolateral membrane (see Fig. S6B in the supplementary material). In stage 13 wild-type glands, E-cad continued to localize to puncta at the lateral membrane and at the AJs (see Fig. S6C in the supplementary material). To determine how levels of E-cad at the AJs compared with that at the basolateral membrane, we measured the fluorescence intensity of E-cad at these two locations. Between stages 12 and 13 when the gland turns posteriorly, E-cad levels decreased at the AJs and increased along the basolateral membrane (see Fig. S6D,E in the supplementary material). These data demonstrate that E-cad localization is dynamic during gland turning and suggest that E-cad traffics through the early endosome during this process.

Since E-cad and Pak1 localized to early endosomes in gland cells (see Figs S5 and S6 in the supplementary material), we tested whether Pak1 could regulate E-cad localization in gland cells through endocytosis. Inhibition of Rab5- or Dynamin-dependent endocytosis, specifically in gland cells, through expression of Rab5S43N (dominant-negative Rab5) or shiK444 (dominant-negative shibire, the gene encoding Dynamin) resulted in expansion and irregular outpouching of the gland lumen (Fig. 3B,C). Similar luminal defects were observed in embryos homozygous for a hypomorph allele of Rab5, Rab5EV10619 and embryos transheterozygous for Pak114 and Rab5EV10619 (Fig. 3D,F). At the cellular level, apical domains were expanded and not elongated, and E-cad was enhanced at the AJs but reduced or absent at the basolateral membrane of shiK444- and Rab5S43N-expressing gland cells (Fig. 3H,I,M,N; see Fig. S7 in the supplementary material), with the effect of shiK444 being more severe than that of Rab5S43N. At the ultrastructural level, the apical surface of shiK444-expressing gland cells looked identical to that of Pak114 mutant gland cells, in which membrane protrusions into the lumen were absent and instead electron-dense patches at the apical surface membrane were evident (Fig. 1L). To determine to what extent inhibition of endocytosis phenocopied the Pak1 mutant gland phenotype, we analyzed the effect of reducing shg (which encodes E-cad) by half in gland cells expressing shiK444. Reduction of shg levels in glands expressing shiK444 restored the correct apical domain size but had no effect on apical domain elongation (see Fig. S7C,D in the supplementary material). This outcome was identical to the effect of reducing shg in Pak114 mutant glands (Fig. 2E,F).

In gland cells of Pak114 homozygous embryos expressing either Rab5S43N (Pak114Rab5S43N) or shiK444 (Pak114shiK444), specifically in the gland cells, apical domains were expanded and failed to elongate (Fig. 3M,N), and E-cad was absent from the basolateral membrane (data not shown), suggesting that Pak1 acts in a single linear pathway with Rab5 and Dynamin. In contrast to inhibition of Rab5, expression of an activated form of Rab5, Rab5Q88L, caused reduction of apical domain size and accumulation of E-cad in sub-apical Aval-positive compartments, compared with wild type (Fig. 3J,M,N; data not shown). Expression of Rab5Q88L in Pak114 mutant gland cells restored apical domain size and elongation, and E-cad localization at the basolateral membrane was similar to that of wild-type gland cells (Fig. 3L,M,N). From these data, we conclude that Pak1 control of Dynamin- and Rab5-dependent endocytosis allows differential localization of E-cad to the AJs and the basolateral membrane, which is important for establishing proper apical domain size and orientation, respectively.

In cultured mammalian cells and in Drosophila epithelia, Rab11-dependent recycling of E-cad regulates E-cad localization at the AJs (Delva and Kowalczyk, 2009). To test whether Rab11- or Rab4-dependent recycling could affect E-cad localization at the AJs and/or the basolateral membrane, we analyzed glands expressing dominant-negative Rab11 (Rab11T21N) or dominant-negative Rab4 (Rab4Q22N). In contrast to loss of Pak1 or Shi inhibition, E-cad still localized to the basolateral membrane of gland cells expressing Rab11T21N or Rab4Q22N (see Fig. S8B,C in the supplementary material), and in Rab11T21N-expressing gland cells we observed an increase in the number of E-cad puncta in the sub-apical region (see Fig. S8C in the supplementary material). Inhibition of either Rab4 or Rab11 doubled apical domain size and only mildly affected apical domain elongation (see Fig. S8D,E in the supplementary material). Thus, in addition to Rab5- and Shi-dependent endocytosis, Rab4- and Rab11-dependent recycling also play a role in determining apical domain size, but do not appear to be required for basolateral localization of E-cad and apical domain elongation.

**Pak1 activation promotes E-cadherin endocytosis and formation of multiple lumens**

Since our studies of Pak1 mutant glands indicated a role for Pak1 in E-cad endocytosis, we next tested whether Pak1 activation could enhance E-cad endocytosis. We expressed a dominant active form of Pak1, in which the myristylation sequence of Src1 (Src64B – FlyBase) was fused to the N terminus of Pak1 (Pak1myr) (Hing et al., 1999). For gland-specific expression, we used either the second chromosome driver, fkh-GAL411, or the third chromosome driver, fkh-GAL411, the latter driving higher levels of Pak1 expression compared with the former (see Fig. S9 in the supplementary material). In glands in which Pak1myr was expressed at moderate levels with fkh-GAL411, a few intercellular lumens containing the lumen protein Pippio formed at sites of cell-cell contact (Fig. 4B), in contrast to the single continuous lumen of wild-type glands (Fig. 4A). Intercellular lumen formation in Pak1myr glands was preceded by loss of the integrity of the single central lumen. In invaginating control glands, E-cad colocalized with the early endosome markers Rab5-GFP (Fig. 4E) and Aval (data not shown) at the apical membrane. In invaginating Pak1myr glands, a single central lumen was no longer apparent; however, strong colocalization of E-cad and Rab5-GFP (Fig. 4F), and E-cad and Aval (data not shown) were observed in the disorganized apical domain and in discrete puncta. All apical marker proteins tested, such as Bazooka, Crb and aPKC, and the AJ proteins E-cad and β-catenin, localized to the intracellular vesicles of Pak1myr glands, whereas the basolateral proteins Neurotactin (Nrt) and Discs Large were excluded (data not shown). E-cad was the earliest protein to be internalized in Pak1myr-expressing gland cells whereas the localization of Crb remained intact (Fig. 5C). In contrast to the apical membrane, the basolateral membrane did not lose integrity and intracellular vesicles were not observed in the vicinity of the basolateral membrane upon expression of Pak1myr (data not shown).

In glands expressing Pak1myr at high levels with fkh-GAL411, formation of multiple lumens was followed by apoptosis and engulfment of gland cells by macrophages (see Fig. S9H,K in the supplementary material). Expression of the baculovirus caspase inhibitor p35 prevented apoptosis of glands induced by Pak1myr (see Fig. S9F,I in the supplementary material). In Pak1myr,p35 glands, the single continuous lumen became disorganized and was replaced with multiple intercellular lumens in the same temporal manner as glands expressing Pak1myr alone. Pak1myr expression with fkh-GAL411 resulted in a more pronounced phenotype, in which the entire gland consisted of multiple intercellular lumens outlined by apical membrane proteins, early endosome marker
proteins and AJ proteins but not basolateral proteins (Fig. 4D and data not shown). We confirmed that the multiple lumens formed by Pak1myr were indeed intercellular and not intracellular by TEM, which showed between six and eight cells surrounding a small central lumen in Pak1myr p35-expressing glands in contrast to wild-type glands, in which the single continuous lumen was surrounded by two rows of cells (Fig. 4G,H). Electron-dense secreted products were found in wild-type and mutant gland lumens demonstrating that loss of normal lumen morphology by Pak1myr did not abrogate the gland’s secretory function.
In Pak1\textsuperscript{myr} and control glands, a subpopulation of E-cad colocalized with the late endosome marker, Rab7-GFP, in the disorganized apical domain but not in the late endosomes in more basal regions of the cells (see Fig. S10B in the supplementary material). We did not observe colocalization of internalized E-cad with lysosomal protein, LAMP1 (CG3305 – FlyBase)-GFP (data not shown).

By TEM, we observed membrane-enclosed structures and deep invaginations of the apical membrane in the apical domain of Pak1\textsuperscript{myr} p35 gland cells, in contrast to wild-type cells that had few or none during invagination (see Fig. S11A,B in the supplementary material). Some structures contained cytoplasm enclosed within double membranes, whereas others contained electron-dense materials and vesicles of varying sizes and shapes (see Fig. S11B in the supplementary material). By stage 12, a central lumen was no longer apparent in Pak1\textsuperscript{myr} p35 glands which also lacked membrane protrusions characteristic of those in wild-type glands (see Fig. S11C-E in the supplementary material). Instead, intercellular vesicles surrounded by electron-dense material were apparent in Pak1\textsuperscript{myr} p35 glands (Fig. S11D,E in the supplementary material).

Fig. 4. Pak1 activation causes the formation of multiple intercellular lumens in Drosophila salivary glands. (A-B*) Triple fluorescence immunostaining of E-cad (white), Neurotactin (Nrt; green) and Pio (red) in salivary glands of stage 13 embryos. For clarity, E-cad staining alone is shown in A’ and B’. Insets in A’ and B’ are magnified views of boxed areas in A and B, respectively. In wild-type embryos (A-A*) salivary glands contain a single continuous lumen filled with Pio whereas E-cad localizes to the apical membrane (arrowheads) and to the basolateral membrane (arrows) labeled with Nrt. In stage 13 embryos expressing Pak1\textsuperscript{myr} p35 specifically in the gland with fkh-GAL4\textsuperscript{II} (B-B*), E-cad localized to intercellular lumens (arrowheads) adjacent to the lateral membranes labeled with Nrt (arrows) that contained Pio. (C-D*) Double fluorescence immunostaining of E-cad (red) and Avalanche (Aval; green) in salivary glands of stage 13 embryos. For clarity, E-cad staining alone is shown in C’ and D’. In wild-type embryos (C,C*), E-cad colocalizes with Aval at the apical domain (arrowheads) and basolateral membrane (arrows). In stage 13 embryos expressing Pak1\textsuperscript{myr} p35 specifically in the gland with fkh-GAL4\textsuperscript{II} (D,D*), E-cad localized to intercellular lumens (arrows) outlined with Aval.

(E-F*) Double fluorescence immunostaining of E-cad (red) and GFP (green) in salivary glands of stage 11 embryos. For clarity, E-cad staining alone is shown in E’ and F’ and GFP staining alone in E” and F”. In wild-type embryos expressing wild-type Rab5-GFP specifically in the gland (E-E*) E-cad colocalized with Rab5-GFP (arrowheads). In Pak1\textsuperscript{myr} glands expressing Rab5-GFP (F) E-cad colocalized with Rab5-GFP in intracellular vesicles (arrowheads). Insets in F-F” are magnified views of the boxed region in F-F”. Dashed white lines in A-F outline the gland. (G,H) In stage 14 wild-type embryos (G), microvilli (arrow in G) protrude into the lumen containing electron-dense secretory products (arrowhead in G), whereas Pak1\textsuperscript{myr} p35 glands have fewer and shorter microvilli (arrow in H), although secretory products were found (arrowhead in H). Adherens junctions (AJs) in G and H are marked in red. L, lumen. Scale bars: 10 μm in A,C,E; 5 μm in A*; 1 μm in G.
Pak1 control of lumen size and shape is dependent on E-cad levels and endocytosis

The internalization of apical membrane and AJ proteins into early endosomes in Pak1\textsuperscript{myr}/p35 glands suggests that endocytosis might play a key role in the formation of intercellular lumens. To test this hypothesis, we inhibited endocytosis by expressing dominant-negative Rab5 (Rab5\textsuperscript{S43N}) or dominant-negative Shi (shi\textsuperscript{K44A}) in Pak1\textsuperscript{myr} glands. Expressing Rab5\textsuperscript{S43N} or shi\textsuperscript{K44A} in Pak1\textsuperscript{myr} glands completely inhibited the formation of multiple intercellular lumens, whereas wild-type Rab5 (Rab5\textsuperscript{WT}) had no effect (Fig. 5A). By contrast, inhibition of early to late endosome maturation with dominant-negative Rab7 (Rab7\textsuperscript{T22N}) or inhibition of early endosome recycling to the plasma membrane with dominant-negative Rab11 (Rab11\textsuperscript{S25N}) in Pak1\textsuperscript{myr} glands still resulted in multiple intercellular lumens (data not shown).
As E-cad endocytosis in Pak1myr glands occurred prior to that of apical membrane proteins, such as Crb (Fig. 5C), we investigated whether expression of wild-type E-cad fused to GFP (E-cad-GFP) using a heterologous promoter would be sufficient to prevent multiple lumen formation in Pak1myr glands. Indeed, 100% of Pak1myrE-cad-GFP glands formed a single central lumen like wild-type glands (Fig. 5A). In the rescued Pak1myrE-cad-GFP glands, vesicles containing E-cad and Aval but not Crb formed between lateral membranes of neighboring gland cells (Fig. 5E,F); however, such vesicles were not replaced by intercellular lumens (data not shown). Thus, providing excess E-cad in Pak1myr glands was sufficient to prevent activated Pak1 from internalizing AJ and apical membrane proteins and hence prevent multiple lumen formation in the gland. These data indicate that endocytic internalization of E-cad is a key regulatory step in the formation of multiple intercellular lumens.
Merlin is an effector of Pak1 in salivary gland lumen size control

The ERM protein Merlin, also known as Neurofibromatosis 2 (NF2), is a substrate of Pak1 (Curto and McClatchey, 2008). Merlin localizes to endosomes in cultured Schwann cells (Scoles et al., 2000) and is required for endocytosis of cell surface proteins, including E-cad, in Drosophila wing tissue (Maitra et al., 2006). Thus, we tested whether Merlin could be a downstream effector of Pak1 in control of gland lumen size. Gland-specific expression of wild-type Merlin (MerWT) did not affect gland width; however, expression of dominant-negative Merlin (MerBBA) (LaJeunesse et al., 1998), expanded the lumen to a degree similar to that observed in Pak1 and Rab5 mutant glands (Fig. 6C). Expression of MerBBA also expanded the apical domain area and reduced the elongation ratio concomitant with loss of E-cad from the basolateral membrane, whereas MerWT had no effect (Fig. 6D-G). Pak1L44 mutant glands expressing either MerWT or MerBBA still had expanded apical domains that failed to elongate, suggesting that Pak1 and Mer act in a single pathway to regulate apical domain size and shape (Fig. 6F,G). Additionally, co-expression of MerBBA with Pak1myr prevented the formation of multiple lumens normally observed with Pak1myr alone, whereas co-expression of MerWT with Pak1myr had no effect (Fig. 6LJ). Thus, normal Merlin activity is required for activated Pak1 to induce E-cad and apical membrane endocytosis and to form intercellular lumens.

To test whether the control of apical domain size by Merlin is dependent on Rab5-mediated endocytosis, we expressed activated Rab5 (Rab5Q88L) or dominant-negative Rab5 (Rab5S43N) simultaneously with MerBBA specifically in the gland. Co-expression of MerBBA with Rab5Q88L restored the correct apical domain area and elongation, whereas co-expression of MerBBA with Rab5S43N had no effect (Fig. 6F,G). Thus, Merlin controls apical domain size and the shape of gland cells in a Rab5-dependent manner.

DISCUSSION

We demonstrate a novel function for Drosophila Pak1 in the control of lumen size and shape in the embryonic salivary gland by regulating E-cad localization at the AJs and basolateral membrane through Rab5- and Dynamin-dependent endocytosis. After gland cells have invaginated, they undergo a phase of robust apical membrane growth and apical domain elongation along the Pr-Di axis of the lumen. This activity of Pak1 is dependent on Merlin, and Rab5- and Dynamin-dependent endocytosis. Constitutively active Pak1 promotes internalization of E-cad and apical membrane proteins into early endosomes, which persist at sites of cell-cell contact and mature into intercellular lumens. This activity of Pak1 is dependent on Merlin, and Rab5- and Dynamin-dependent endocytosis.

Fig. 7. Model for Pak1 function in salivary gland lumen size control. (A) Diagram depicting the lateral view of a gland cell. In wild-type gland cells, Pak1 mediates endocytosis of E-cad from the adherens junctions (AJs) and localization of E-cad to the basolateral membrane, which reduces apical domain size and elongates the apical domain along the Pr-Di axis of the lumen. Pak1 mediates these events through Merlin and Rab5- and Dynamin-dependent endocytosis. A, apical; B, basal; D, dorsal; Di, distal; Pr, proximal; V, ventral. (B) Diagram depicting a lateral view of the gland. Constitutively active Pak1 promotes internalization of E-cad and apical membrane proteins into early endosomes, which persist at sites of cell-cell contact and mature into intercellular lumens. This activity of Pak1 is dependent on Merlin, and Rab5- and Dynamin-dependent endocytosis. A, anterior; D, dorsal; P, posterior; L, lumen; V, ventral. Diagrams are not drawn to scale.
axis of the gland. As gland cells turn and migrate, they reduce the size of the apical domain while maintaining its elongated shape. Our studies show that Pak1 regulates apical domain size and elongation during this step through Rab5- and Dynamin-dependent endocytosis of E-cad from the AJs and its subsequent localization at the basolateral membrane (Fig. 7A). A role for Pak1 in the endocytic control of E-cad localization is further supported by our studies with activated Pak1, which demonstrate that Pak1 activation leads to formation of multiple intercellular lumens in addition to, or instead of, a single central lumen (Fig. 7B). Pak1 activation led to Rab5- and Dynamin-mediated endocytosis of E-cad and apical membrane constituents into a single enlarged early endosome located near the cortex of the cell. This cortical early endosome persisted, localized laterally and subsequently expanded to become the new apical domain facing an intercellular lumen between sites of cell-cell contact (Fig. 7B).

Based on our studies, we propose a model for how Pak1 regulation of E-cad controls salivary gland lumen size through modulation of apical domain size and shape. We propose that E-cad destabilization at specific apical interfaces by endocytosis leads to changes in apical tension such that constriction only occurs along the axis of reduced apical tension, i.e. the D-V axis of the apical domains of individual gland cells. This selective constriction of the apical domain then allows the apical domain to elongate and become anisotropic. Pak1 could promote polarized destabilization of E-cad by directly regulating the endocytic machinery or indirectly through the activity of Myosin II and the actomyosin cytoskeleton, which mediates polarized remodeling of E-cad-based cellular junctions during Drosophila germband extension (Bertet et al., 2004). Although we did not observe changes in F-actin structure in Pak1 mutant gland cells and mutations in actin regulators did not perturb gland lumen size, it is still possible that increased localization of E-cad at the AJs in Pak1 mutant gland cells could lead to an increased apical stiffness, potentially through the association of the AJs with the actin cytoskeleton. Increased apical stiffness could then prevent anisotropic changes of the apical domain. Indeed, previous studies in the Drosophila salivary gland have shown that modulation of apical stiffness can result in luminal changes (Cheshire et al., 2008; Kerman et al., 2008).

Recent studies showed a role for the Cdc42-Par-aPKC pathway in E-cad endocytosis from the AJs in Drosophila tissues through regulation of the actin cytoskeleton (Georgiou et al., 2008; Leibfried et al., 2008). The E-cad internalization defects reported in these studies differ significantly from data presented here. For example, inhibition of Cdc42 or Dynamin formed disorganized AJs (Georgiou et al., 2008) or large endocytic structures tethered to the AJs (Leibfried et al., 2008), neither of which was observed in Pak1 mutant glands. Moreover, Cdc42 inhibition disrupted F-actin structure (Georgiou et al., 2008), which was also not observed in Pak1 mutant cells. These differences might, in part, be due to tissue-specific requirements for E-cad regulation at different developmental times. In the embryonic neuroectoderm, Cdc42 inhibition leads to loss of apical membrane and AJs proteins, which also contrasts with our findings in Pak1 mutant glands (Harris and Teppas, 2008). Thus, Pak1 probably controls E-cad endocytosis from the AJs by a mechanism distinct from that of Cdc42 reported in these studies. Based on the studies presented here, we propose that endocytosis in the salivary gland is important for the dynamic localization of E-cad to allow remodeling of the apical domain, which, in turn, is required for achieving proper lumen size and shape.

Our studies provide the first evidence that constitutive activation of Pak1 leads to formation of an epithelial tube with multiple intercellular lumens in a process dependent on endocytosis and E-cad levels. Multiple lumens have been observed during normal organogenesis and in pathological conditions. For example, in the zebrafish intestine, the formation of a tube with a single central lumen is preceded by the formation of multiple intercellular lumens that then coalesce (Bagnat et al., 2007; Home-Badovinac et al., 2001). In the zebrafish neural tube and Drosophila dorsal vessel, Par6βγ and Slit regulate single lumen character, respectively (Munson et al., 2008; Santiago-Martinez et al., 2008). Moreover, formation of multiple lumens is a histological feature of breast ductal carcinoma in situ, a pre-invasive form of breast cancer (Debnath and Brugge, 2005). Although a direct role for Pak1-induced endocytosis in these cases of multiple lumen formation has not been demonstrated, it is possible that modulation of cadherin-mediated cell-cell adhesion through membrane transport is a common mechanism by which lumen type and number are determined both in normal developmental processes and in pathological conditions.

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The authors declare no competing financial interests.

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Supplementary material
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