DaPKC-dependent phosphorylation of Crumbs is required for epithelial cell polarity in *Drosophila*

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Both in *Drosophila* and vertebrate epithelial cells, the establishment of apicobasal polarity requires the apically localized, membrane-associated Par-3–Par-6–aPKC protein complex. In *Drosophila*, this complex colocalizes with the Crumbs–Stardust (Sdt)–Pals1-associated TJ protein (Patj) complex. Genetic and molecular analyses suggest a functional relationship between them. We show, by overexpression of a kinase-dead *Drosophila* atypical PKC (DaPKC), the requirement for the kinase activity of DaPKC to maintain the position of apical determinants and to restrict the localization of basolateral ones. We demonstrate a novel physical interaction between the apical complexes, via direct binding of DaPKC to both Crb and Patj, and identify Crumbs as a phosphorylation target of DaPKC. This phosphorylation of Crumbs is functionally significant. Thus, a nonphosphorylatable Crumbs protein behaves in vivo as a dominant negative. Moreover, the phenotypic effect of overexpressing wild-type Crumbs is suppressed by reducing DaPKC activity. These results provide a mechanistic framework for the functional interaction between the Par-3–Par-6–aPKC and Crumbs–Sdt–Patj complexes based in the posttranslational modification of Crb by DaPKC.

Introduction

Epithelial cells are polarized along their apicobasal axis. Their plasma membranes are subdivided into apical and basolateral domains that are separated by specialized junctional structures, such as tight junctions (TJ) and the zonula adherens (ZA; for reviews see Drubin and Nelson, 1996; Müller, 2000; Tepass et al., 2001; Roh and Margolis, 2003). Formation and stabilization of these junctions are paramount for the maintenance of epithelial polarity, which, in turn, is required for proper epithelial cell physiology, cell motility, asymmetric division, and intercellular signaling (for review see Tepass et al., 2001). Hence, the functional analysis of the factors that establish and/or maintain epithelial polarity is a major issue in developmental biology.

The mechanisms that underlie the establishment and maintenance of epithelial polarity have been extensively studied in *Drosophila*. Thus, three membrane-associated multiprotein complexes, the Bazooka (Baz), Crumbs, and Discs large (Dlg) complexes, are required to organize apicobasal polarity in the epithelial cells of the *Drosophila* embryo (for reviews see Müller, 2000; Ohno, 2001; Tepass et al., 2001; Knust and Bossinger, 2002; Henrique and Schweisguth, 2003; Roh and Margolis, 2003). The Baz complex is formed by the PSD-95/Dlg/ZO-1 domain containing proteins Baz and DPar-6, and the *Drosophila* atypical PKC (DaPKC), a Ser/Thr kinase (Kuchinke et al., 1998; Wodarz et al., 2000; Petronczki and Knoblich, 2001). Embryos deficient in Baz or DPar-6 cannot properly assemble the ZA and lack several apical markers of the cell membrane (Müller and Wieschaus, 1996; Petronczki and Knoblich, 2001). Similarly, epithelial cells of *DaPKC* mutant imaginal discs display disrupted apicobasal polarity in the *Drosophila* embryo.

Abbreviations used in this paper: ASIP, atypical PKC isotype-specific interacting protein; Baz, Bazooka; Crbi, intracellular domain of Crumbs; DaPKC, *Drosophila* atypical PKC; Dlg, Discs large; DN, dominant negative; Lgl, Lethal giant larvae; Nrt, Neurotactin; Patj, Pals1-associated TJ protein; Scrib, Scribble; Sdt, Stardust; SP, signal peptide; TJ, tight junctions; TM, transmembrane; ZA, zona adherens.
(Rolls et al., 2003). Thus, each of the components of the Baz complex is essential to establish epithelial polarity.

The Crumbs complex, which is assembled at gastrulation, is also localized at the marginal zone. It consists of the transmembrane (TM) protein Crumbs (Crb) and the cytoplasmic PDZ-containing proteins Stardust (Sdt) and Pals1-associated TJ protein (Patj; formerly known as Discs lost; Tepass et al., 1990; Bhat et al., 1999; Bachmann et al., 2001; Hong et al., 2001; Pielage et al., 2003). Crb contains 30 epidermal growth factor–like and four laminin A G-domain–like repeats in its extracellular region and a short intracellular domain (Tepass et al., 1990). Expression of this intracellular domain partially normalizes mutant crb embryos (Wodarz et al., 1995; Klebes and Knust, 2000).

This suggests that Crb exerts its function, at least in part, by protein–protein interactions mediated by this domain. The intracellular domain has been subdivided into proximal juxtamembrane and COOH-terminal subdomains. The latter contains the binding site for Sdt (Bachmann et al., 2001; Hong et al., 2001; Roh et al., 2002), which on its turn binds to Patj (Roh et al., 2002). The juxtamembrane domain is required for Crb to form a complex with DMoesin and β-heavy spectrin and thus mediates the interaction of the Crb complex with the apical spectrin cytoskeleton (Medina et al., 2002). Genetic studies have shown that the Crb complex is, similarly to the Baz complex, indispensable for the establishment of epithelial apicobasal polarity and stabilization of the ZA (Knust et al., 1993; Grawe et al., 1996; Müller and Wieschaus, 1996; Tepass, 1996; Bachmann et al., 2001; Hong et al., 2001).

The third complex, formed by the PDZ proteins Dlg (Woods and Bryant, 1991) and Scribble (Scrib; Bilder and Perrimon, 2000), and the Myosin type II binding protein Lethal giant larvae (Lgl; Mechler et al., 1985), is found at the basolateral domain of the cell membrane, basal to the ZA. These proteins are required to restrict the Baz and Crb complexes to the apical cell membrane and for correct positioning of the ZA (Bilder et al., 2000, 2003; Bilder and Perrimon, 2000; Tanentzapf and Tepass, 2003).

The three polarity complexes are evolutionarily conserved (for reviews see Ohno, 2001; Roh and Margolis, 2003; Macara, 2004). Thus, in mammalian epithelial cells, the homologues of DaPKC, Baz, and DPar-6, namely aPKC, Par-3/atypical PKC isotype-specific interacting protein (ASIP), and Par-6, and the corresponding homologues of Crb, Sdt, and Patj, that is, Crb3, Pals1 (protein associated with Lin seven1), and Patj, respectively, form two complexes that are localized at the TJ and regulate the assembly of these junctions. The basolateral proteins Scrib, mDlg, and mLgl also play important roles in epithelial cell polarity (Izumi et al., 1998; Gao et al., 2002; Hirose et al., 2002; Lemmers et al., 2002; Roh et al., 2002, 2003; Suzuki et al., 2001, 2002; Hurd et al., 2003; Straight et al., 2004).

In Drosophila, the Baz and Crb complexes not only colocalize in the apicolateral region of the cytocortex (Hong et al., 2001; Bilder et al., 2003), but they interact physically by means of DPar-6 and Patj (Nam and Choi, 2003). Furthermore, absence of either complex similarly disrupts epithelial polarity (for review see Tepass et al., 2001), which suggests a functional relationship between them. Here, we report a novel physical connection between the Baz and Crb complexes mediated through direct binding of DaPKC to both Crb and Patj. We find that the kinase activity of DaPKC is necessary to maintain the apical localization of the components of the Crb complex and to prevent the apical localization of basolateral markers. Furthermore, Crb is a phosphorylation target of DaPKC. These results provide a molecular mechanism for the functional interaction between the Crb and Baz complexes based on DaPKC-dependent phosphorylation of Crb.

**Results**

**DaPKC kinase activity is required for epithelial polarity in the Drosophila embryo**

To gain insight into the role of DaPKC in epithelial cells, we first examined whether or not its kinase activity is required for epithelial polarity, as it is the case in cultured mammalian epithelial cells. In these cells, overexpression of a kinase-defective aPKC disrupts the assembly of the TJ (Suzuki et al., 2001, 2002). We interfered with DaPKC signaling in vivo by overexpressing a form of DaPKC that was targeted to the cell membrane (by addition of the CAAX sequence), and that harbored a mutation (K293W) in the ATP binding site (DaPKC<sup>CAAXDN</sup>; see Materials and methods). Similar modifications in Xenopus aPKCa converted it into a kinase-
The Baz and Crb apical complexes directly interact

The colocalization of the Baz and Crb complexes (Hong et al., 2001; Bilder et al., 2003) suggested that components of these two complexes might physically interact.

Figure 2. Physical interaction between the apical complexes.

(A) Drosophila embryonic extracts were incubated with GST or with the indicated GST-fusion proteins (GST-Crb wild-type intracellular domain, GST-Crb, and GST-Patj) bound to a glutathione matrix. Immunoblotting of bound proteins, resolved by SDS-PAGE, with anti-aPKC antibody, indicates the interaction of a protein complex containing DaPKC with Crbi and Patj. First lane (Lysate) shows 13% of the extract used in the assay. (B) Direct binding of DaPKC to Crbi and Patj. His-tagged DaPKC was bound to Ni-NTA-agarose and incubated with purified GST, GST-Crb, and GST-Patj. Bound proteins were revealed by immunoblotting with anti-GST antibody. Bottom panel is a loading control. Half the amount of His-tagged DaPKC used in the experiment was run in a parallel gel and revealed with anti-Xpress epitope antibody. (C) Pull-down experiments of embryonic extracts were performed as in A with GST or with the indicated GST-fusion proteins: GST-Crb, GST-CrbT6A, GST-CrbT9A, GST-CrbT6A,T9A, and GST-CrbT6A,T9A,S11A,S13A. Bound proteins were resolved by SDS-PAGE and immunoblotted with anti-aPKC and anti-Patj antibodies. First lane (Lysate) shows 10% of the extract used in the assay. Molecular weight markers are shown on the left.

Indeed, in GST pull-down assays, the intracellular domain of Crumbs (Crb) was able to precipitate DaPKC and Baz from embryonic extracts (Fig. 2 A and not depicted). Similarly, Patj was able to pull down DaPKC (Fig. 2 A). To identify directly interacting proteins, we performed binding assays with bacterially purified His-tagged DaPKC and GST-Crb and GST-Patj fusion proteins. As shown in Fig. 2 B, DaPKC was able to bind directly both GST-Crb and GST-Patj.

To investigate further if Patj or Sdt bound to Crb are required in vivo for the interaction between the two apical complexes, we performed pull-down assays with CrbiT6A,T9A,S11A,S13A, a truncated form of Crb devoid of the four COOH-terminal amino acids (ERLI) which constitute the Sdt binding domain (Bachmann et al., 2001; Hong et al., 2001). As expected, CrbiT6A,T9A,S11A,S13A was unable to pull down Patj (Fig. 2 C). Interestingly, it was also unable to pull down DaPKC (Fig. 2 C). These results suggest that although Crb on its own is able to bind DaPKC, in vivo Sdt and/or Patj may also intervene in the interaction between the two apical complexes.

Crb is a phosphorylation target of DaPKC

Because the kinase activity of DaPKC was required for the apical localization of the Crb complex and DaPKC directly binds Crb and Patj, we analyzed if these proteins were phosphorylation substrates of DaPKC. We found that in an in vitro assay recombinant aPKCζ, the human homologue of DaPKC, phosphorylated a GST-Crb fusion protein, either with or without the ERLI domain (Fig. 3 A, arrow). Interestingly, Patj, which was not phosphorylated by aPKCζ, strongly decreased Crb phosphorylation when present in the reaction mixture (Fig. 3 A). Note however that Patj was not a general inhibitor of aPKCζ, as it did not block aPKCζ autophosphorylation (Fig. 3 A, asterisk) or phosphorylation of another substrate, the myelin basic protein (not depicted).

The intracellular domain of Crb contains four Ser/Thr residues (T6, T9, S11, and S13) in an evolutionarily conserved region (Fig. 3 B), which are putative targets for DaPKC phosphorylation in vivo. To analyze the possibility of a functional regulation of Crb by phosphorylation, first we replaced with Ala all the putatively phosphorylatable sites of Crb, either individually or all together (CrbT6A,T9A,S11A,S13A) and performed in vitro kinase assays using the mutated GST-Crb fusion proteins as substrates. Mutation of the four putative phosphorylatable residues in Crb abolished its phosphorylation by DaPKC (Fig. 3 C, left), although interestingly, CrbT6A,T9A,S11A,S13A was still able to interact with DaPKC, Patj, and DMoesin in pull-down assays (Fig. 2 C and not depicted). Individual point mutations did not allow unambiguous identification of the phosphorylation site(s) (not depicted). To clarify this point, wild-type amino acids were added back to the Crb protein mutated in the four Ser/Thr residues and the corresponding Crb proteins were assayed for aPKCζ-dependent phosphorylation. This experiment identified T6 and T9 as substrates of aPKCζ (Fig. 3 C, middle). Accordingly, the double mutant CrbT6A,T9A was no longer phosphorylated by aPKCζ (Fig. 3 C, right).
In vivo activity of Crb is modulated by its state of phosphorylation

To examine the functional relevance of Crb phosphorylation, we compared the in vivo activity of wild-type and nonphosphorylatable Crb proteins. Wild-type and mutated Crb intracellular domains, either Crbi<sup>T6DT9D</sup> wt or Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup>, were fused to Crb signal peptide (SP) and TM domains (see Materials and methods). These proteins were overexpressed in embryos under the control of the Ubx-Gal4 driver line. Use of this driver enables direct comparison, in a single embryo, of the wild-type nonexpressing and the Crb-overexpressing domains (Fig. 4 A). As described previously (Wodarz et al., 1995; Klebes and Knust, 2000), overexpression of membrane-tethered Crbi conferred partial apical character to the whole cell membrane, as shown by the distribution of DaPKC and Patj around the whole cell contour, coincident with the ectopic distribution of Crb (Fig. 4 B and not depicted). However, localization of the basolateral markers Nrt and Scrib was not affected (Fig. 4 C and not depicted). In contrast, in embryos overexpressing nonphosphorylatable Crbi constructs (either UAS-Crbi<sup>T6DT9D wt</sup> or UAS-Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup>), both the endogenous and overexpressed Crb as well as Patj and DaPKC failed to associate with the membrane and were found diffusely in the cytoplasm (Fig. 4, D and E; not depicted). In addition, the basolateral marker Scrib appeared mislocalized in the apical membrane (Fig. 4 E). Thus, overexpression of a nonphosphorylatable Crbi had the opposite effect than overexpression of wild-type Crbi and resembles the polarity phenotype of strong crb mutants, where Patj and DaPKC are lost from the plasma membrane (Bhat et al., 1999; Klebes and Knust, 2000; Bilder et al., 2003). This suggested that the nonphosphorylatable Crbi acted as a DN. Indeed, nonphosphorylatable Crbi was still able to pull down DaPKC, Patj, and DMoesin from embryonic extracts (Fig. 2 C and not depicted) and thus it may compete with the endogenous Crb for its interacting partners.

To further analyze the role of Crb phosphorylation, T6 and T9 were mutated to Aspartate to mimic a constitutively phosphorylated state. This protein, Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup> wt, was still able to pull down DaPKC, Patj, and DMoesin (Fig. 2 C and not depicted) and its ectopic overexpression caused redistribution of the apical proteins DaPKC and Patj to most of the cytocortex (Fig. 4 F and not depicted). Furthermore, and similarly to Crbi, it did not affect the localization of the basolateral marker Nrt (Fig. 4 G). These results support that Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup> mimics wild-type Crbi.

As a further functional test, we analyzed the consequences of overexpressing the different forms of Crb in the imaginal disc epithelia using the en-Gal4 line. This line drives expression of UAS transgenes in the posterior compartment of the wing disc, which gives rise to the posterior wing (Fig. 4 H). Overexpression of Crbi<sup>wt</sup> caused severe tissue disorganization in the wing, an effect that has been attributed to apicalization of the wing disc cells (Fig. 4 J; Bilder et al., 2003; Tanentzapf and Tepass, 2003). Although the overexpression of nonphosphorylatable Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup> generally lead to embryonic lethality, in a few cases, surviving flies were obtained and displayed only mild defects in the posterior crossvein (not depicted). In contrast, overexpression of Crbi<sup>T6DT9D</sup> in the imaginal discs disrupted the wing epithelium in a way similar to that caused by Crbi<sup>wt</sup> (Fig. 4 K). This supports the results obtained in the embryo.

The above experiments indicate that the phosphorylation of Crb by DaPKC is important for its function. Hence, reducing the DaPKC activity should decrease the effect of the overexpression of Crbi<sup>wt</sup>. This was the case. Expression of UAS-DaPKC<sup>CAAXDN</sup> largely normalized the effect caused by overexpression of wild-type Crbi (Fig. 4, L and M). As expected, the effect of overexpressing Crbi<sup>T6DT9D</sup> <sup>TGAT9A</sup> was less sensitive to the decrease of DaPKC activity. Thus, whereas 100% of flies overexpressing simultaneously UAS-Crbi<sup>wt</sup> and UAS-DaPKC<sup>CAAXDN</sup> showed essentially normal wings (Fig. 4 M),...
Discussion

In the *Drosophila* embryo, several regulators of epithelial polarity are sequentially associated, in the form of protein complexes, with the apicolateral (Crb and Baz complexes) and basolateral (Dlg complex) domains of the plasma membrane. The concerted activities of these protein complexes establish and maintain apicobasal polarity. Thus, the Baz and Crb complexes are indispensable for the formation of the ZA and the establishment of the apical domain. The concerted activities of these protein complexes result in DaPKC-dependent phosphorylation of Crumbs and Patj concomitant with an apical expansion of basolateral complexes. Thus, DaPKC-dependent phosphorylation of Crumbs | Sotillos et al. 553

Figure 4. Regulation of Crbi activity by DaPKC-dependent phosphorylation. Different forms of Crbi were overexpressed in embryos (B–G, Ubx-Gal4 line) and in the wing disc (J–N, en-Gal4 line). (A) Embryos expressing UAS-LacZ under the control of Ubx-Gal4 line were stained with anti-β-galactosidase antibody to reveal the domain of expression of the Ubx-Gal4 line. Similar regions to the one marked in A are shown in B–G. In these panels, the extent of the Ubx-Gal4 expressing domain is indicated by the white bar. (B–G) Embryos of the indicated genotype were stained with an antibody raised against the intracellular domain of Crb to reveal the distribution of both the endogenous and overexpressed Crb (purple) and costained (green) for DaPKC (B and F), Nrt (C and G), Patj (D), and Scrib (E). (B and C) Embryos expressing wild-type Crbi showed extensive relocation of DaPKC to the whole cell contour in the cells overexpressing Crb (B). The localization of the basolateral marker Nrt was not affected (C). (D and E) On the contrary, overexpression of *DaPKC* or *Crbi*, causes the loss of membrane associated Crb activity of DaPKC by overexpressing a DN, kinase-dead aPKC kinase activity is also needed to counteract the activity of these proteins (Bilder and Perrimon, 2000; Bilder et al., 2000, 2003; Tanentzapf and Tepass, 2003). Recently, another basolateral protein, the Par-1 kinase has been involved in epithelial polarity. Par-1 inhibits formation of the Baz complex at the basolateral domain by blocking the oligomerization of Baz and its binding to DaPKC (Benton and Johnston, 2003). In addition to these regulatory interactions, ours and previous data indicate a functional reciprocal interaction between the two apical complexes. Thus, DaPKC is required and instructive for membrane localization of Crb and Patj (this work). Conversely, the Crb complex is required to maintain the Baz complex at the apical membrane, as shown by the mislocalization of Baz and DaPKC in *sdtscrb* embryos (Bachmann et al., 2001; Hong et al., 2001; Bilder et al., 2003). This functional interaction is supported by the synergism between *sdtscrb* and *bazz* mutations (Müller and Wieschaus, 1996).

Thus, the similar mutant phenotypes associated with the absence of either of the apical complexes (Tepass et al., 2001) may then be accounted for by their functional relationship. Here, we show that physical interaction between the apical complexes results in DaPKC-dependent phosphorylation of Crb, which is a requisite for the proper activity of Crb in the control of apicobasal polarity.

Requirement of aPKC kinase activity in epithelial polarity

We have addressed the question of whether the kinase activity of DaPKC is required for epithelial polarity and have found that this is indeed the case. Thus, diminishing the activity of DaPKC by overexpressing a DN, kinase-dead DaPKC, causes the loss of membrane associated Crb and Patj concomitant with an apical expansion of basolateral markers. Conversely, membrane tethered wild-type DaPKC, located all around the cell cortex, targets apical proteins (Crb, Patj, and Baz) to the basolateral domain. These polarity defects are not due to nonspecific effects of DaPKC (unpublished data). aPKC kinase activity is also needed to establish epithelial polarity in zebra fish. Thus, the zebra fish mutant Heart and soul encodes an aPKCΔ mutated in a do-
main critical for the activation of its kinase function and this results in a defective ZA (Horne-Badovinac et al., 2001). All these results show that the kinase activity of aPKC is necessary to establish epithelial polarity in different animal phyla and poses the question as to the identification of the phosphorylation targets of DaPKC in epithelial cells.

**Crb is a phosphorylation target of DaPKC in epithelial cells**

The identified targets of aPKC belong to two groups: basolateral proteins, negatively regulated by aPKC (Lgl, both in vertebrate and invertebrate cells [Betschinger et al., 2003; Plant et al., 2003; Yamanaka et al., 2003] and mammalian Par-1 [Hurov et al., 2004]), and apical proteins, namely Par-3 (Hirose et al., 2002) and Crb (this work), which require such phosphorylation for their proper activity.

Our results strongly suggest that interaction of DaPKC with Crb and the subsequent phosphorylation of the latter are most relevant for epithelial polarity. This conclusion is based in several experimental observations. First, aPKC phosphorylates Crb in vitro at its intracellular domain. Second, this phosphorylation appears to be essential in vivo because mutation of the putatively phosphorylatable Thr/Ser of the Crb intracellular domain into Ala inactivates Crb. Nonphosphorylatable Crbi still binds DaPKC, Patj, and DMOesin. Thus, it should act as a DN form competing with wild-type Crb and we find that this is indeed the case. Third, mutation of the putatively phosphorylatable residues of Crbi into negatively charged Asp maintains its biological function. This suggests that the DN character of the CrbiT6A,T9A is not due to a mere change of T6 and T9 to Ala but to the conversion of Crb into a nonphosphorylatable protein. Fourth, the phenotypic effects of overexpressing CrbiT6A,T9A were suppressed by a reduction of DaPKC kinase activity, supporting that DaPKC is required for Crb activity. CrbiT6A,T9A was not sensitive to a reduction in DaPKC activity than CrbiWT. However, there was still a partial recovery of the cuticular malformations caused by overexpression of CrbiT6A,T9A when DaPKC activity was compromised. This suggested that conversion of both Thr into Asp did not render Crb into a completely constitutively active protein. Note also that CrbiT6A,T9A still retains Ser11 and Ser13. Although our in vitro experiments do not point at these residues as DaPKC-dependent phosphorylation sites, we cannot totally exclude their functionality in vivo, which could be modified by reduced DaPKC activity. Other alternative, but not mutually excluding explanations, may be that the partial rescue of CrbiT6A,T9A by DaPKCΔCAAXΔN results from the interference of DaPKCΔCAAXΔN with the endogenous Crb and/or may be caused by the functional antagonism between Crb and the basolateral proteins that colocalize on the whole cytocortex in these experimental conditions.

Together, these results indicate that DaPKC-dependent phosphorylation of Crb is essential for its wild-type activity. Because the putatively aPKC-dependent phosphorylatable residues of Crb are conserved in mammalian homologues, one of the main functions of aPKC in epithelial cells may be to maintain apicobasal polarity through posttranslational modification of Crb. Note, however, that the genetic control of epithelial polarity may be different in *C. elegans*, because the putatively phosphorylatable residues are absent from *C. elegans* Crb-1 and inactivation of *crb-1* by dsRNAi did not disturb formation of the *C. elegans* apical junction (Bosssinger et al., 2001).

Additionally, and although it has not been demonstrated, it is conceivable that DaPKC-dependent phosphorylation of Baz may be required for apicobasal polarity in *Drosophila*, as it is the case in cultured mammalian epithelial cells where phosphorylation of the mammalian homologue of Baz, ASIP/Par-3, by PKC-3 is indispensable for the formation of the TJ (Hirose et al., 2002).

aPKC plays a dual role in the control of epithelial polarity. In addition to its requirement for the proper activity of the apical proteins, aPKC negatively regulates the activity of the basolateral proteins. Thus, in mammalian epithelial cells aPKC-dependent phosphorylation of Lgl and Par-1b prevents the association of these proteins with the plasma membrane (Yamanaka et al., 2003; Hurov et al., 2004). In the apical side of the *Drosophila* neuroblasts, DaPKC counteracts the activity of Lgl releasing it from its association with the cell membrane (Betschinger et al., 2003). A similar antagonistic effect between DaPKC and Lgl and Dlg appears to take place in the imaginal discs (Rolls et al., 2003). In agreement with these observations, we have found that the kinase activity of DaPKC is required to exclude Scrib from the apical region of the embryonic epithelial cells. This function of DaPKC would be independent of its role in Crb activation because ectopic basolateral localization of Crb on its own did not affect the localization of basolateral markers (Fig. 4 C). Although ectopic accumulation of Crb is associated with relocation of endogenous DaPKC to the whole cell contour, conceivably that amount of DaPKC is insufficient to affect the localization of basolateral markers.

**A molecular mechanism for the functional interaction between the apical complexes**

It has been proposed that the Baz, Crb, and Dlg complexes act sequentially to direct the maturation of epithelial cell polarity, the Baz complex being the first to be assembled and the most critical apical regulator (Bilder et al., 2003). Our findings provide a mechanistic framework for the regulation of the Crb complex by the Baz complex. The functional interaction between the *Drosophila* apical complexes relies in physical contact between some of their components. Thus, DPar-6 and Patj directly interact in vitro (Nam and Choi, 2003). Now we find that DaPKC directly binds to Crb and Patj. Although in vitro DaPKC interacts with and is able to phosphorylate a Crb protein devoid of the COOH-terminal ERLI domain (Fig. 3 A), pull-down experiments of embryonic extracts show that this domain is necessary for the interaction between the Baz and Crb complexes. This suggests that in vivo the interaction may be reinforced by proteins that, directly or indirectly, are bound to the ERLI domain. Sdt is a likely candidate because it binds to this domain of Crb (Bachmann et al., 2001; Hong et al., 2001; Roh et al., 2002) and its mammalian homologue Pals-1 binds Par-6 (Hurd et al., 2003). Additionally, human Crb-3 and Par-6 interact (Lemmers et al., 2004), but it is not yet known whether this also occurs in *Drosophila*. 
The interaction between the apical complexes results in DaPKC-dependent phosphorylation and activation of Crb. Although our results show that the phosphorylation state of Crb does not affect its interaction with several known binding partners, we speculate that such phosphorylation may be necessary for Crb to adopt a correct conformation and/or to interact with proteins required for Crb stabilization at the apical domain. Moreover, the finding that Patj inhibits the DaPKC-dependent phosphorylation of Crb, suggests the existence of alternative and excluding complexes. During early embryonic stages, Patj and DaPKC may directly interact, although such interaction would not affect the kinase activity of DaPKC (in fact, DaPKC kinase activity is required for the apical localization of Patj at blastoderm stage; unpublished data). At gastrulation, when the Crb complex accumulates at the plasma membrane, we hypothesize that a greater affinity of Patj for Sdt (bound to Crb) than for DaPKC would release Patj from its interaction with DaPKC and allow the phosphorylation of Crb. Note that the Baz complex may perform additional roles in relation to the Crb complex. Thus, previous data suggested the existence of Crb-dependent and -independent mechanisms for the localization of Patj and Sdt (Tanentzapf et al., 2000; Nam and Choi, 2003). The interaction between components of the apical complexes, as stated by Nam and Choi (2003), would account for these results.

In summary, the present results stress the intricacy of the relationships between the evolutionarily conserved proteins that establish and maintain epithelial polarity. There is a feedback mechanism between the Baz and Crb complexes. Thus, phosphorylation of Crb by DaPKC is a requisite for the ulterior Crb complex-dependent maintenance of the apical localization of the Baz complex. The wealth of data demonstrating the physical and functional interactions between both complexes allows them to be considered as part of a unique apical determinant complex, whose components are added stepwise. Additionally, there is a functional antagonism among the apical and the basolateral domain determining proteins (Bilder et al., 2003, Tanentzapf and Tepass, 2003; this work). Clearly, identification of additional targets of DaPKC and comprehensive molecular and genetic analyses of the functional interactions among the different complexes will allow further insight into the establishment and maintenance of epithelial polarity.

Materials and methods

Fly strains

Stocks used were: wild type (Oregon R). Transgenic lines expressing wild-type DaPKC (UAS-DaPKC), membrane targeted wild-type DaPKC (UAS-DaPKC), membrane targeted kinase-dead DaPKC (UAS-DaPKCCASINO), membrane targeted kinase-dead Xenopus aPKC (UAS-XaPKCCASINO) and wild-type and mutated membrane bound Crb-intra (UAS-Crb) were generated by subcloning the corresponding cDNAs (Berra et al., 1993; see Site directed mutagenesis) in the pUAST vector (Brand and Perrimon, 1993). The resulting pUAST-plasmids were used to transform w1118 embryos (Ashburner, 1989) using 0.3 mg/ml of pCISd2.3 as the source of transposase. Several lines were established and analyzed for each transgene.

Mis-expression studies

Mis-expression experiments were performed using the Gal4/UAS system (Brand and Perrimon, 1993) and en-Gal4, Ubx-Gal4 (Calleja et al., 1996; a gift from M. Calleja, CBMso, Madrid, Spain) and the maternal Gal4 line MatVP16V67-Gal4 (generated by D. St. Johnston, University of Cambridge, Cambridge, UK). Embryos expressing UAS-DaPKC, UAS-DaPKCCASINO, UAS-DaPKCCASINO, or UAS-XaPKCCASINO driven by the MatVP16V67-Gal4 line were collected after 6–16 h at 18 or 25°C, fixed, and stained. Embryos expressing UAS-Crb, UAS-Crb/SA13, UAS-Crb/SA1A, and UAS-Crb/SA1A driven by Ubx-Gal4 were collected for 16 h at 18 or 25°C. To assay the effect of a reduction of DaPKC signaling on the activity of Crb, either wild-type or mutated females were crossed with UAS-

DaPKC/SA13; UAS-Crb/SA13; UAS-Crb/SA1A males. This cross scheme allows unambiguous identification of en-Gal4/UAS-DaPKCCASINO, en-Gal4, UAS-Crb and en-Gal4/UAS-DaPKCCASINO, UAS-Crb flies in the progeny. The progeny of the above mentioned cross was maintained at 18°C during embryonic development and then transferred to 25°C for subsequent development. In order to minimize deleterious effects associated with the expression of Crb constructs at the embryonic stages. The effect of concomitant expression of UAS-DaPKCCASINO and UAS-DaPKCCASINO was examined in the progeny of the cross of en-Gal4 females with UAS-DaPKCCASINO; UAS-DaPKCCASINO males.

Immunohistochemistry

Embryos were fixed as in Tepass et al. (1990) and stained with rabbit anti-
Patz and rat anti-Crb-intra (Bhat et al., 1999; a gift from H. Bellen, Baylor College of Medicine, Houston, TX), mouse Mab cc4 anti-Crb (DSHB), rabbit anti-Baz (a gift from A. Wodarz, Heinrich-Heine-Universität Düsseldorf, Dusseldorf, Germany), rabbit anti-aPKC C20 (Santa Cruz Bio
technology, Inc.), mouse Mab BP106 anti-Nrt (Hortsch et al., 1990), rabbit anti-Scr (a gift from C. Doe, University of Oregon, Eugene, OR); rabbit anti-β-galactosidase (Cappel) and mouse anti-β-galactosidase (Promega). FITC and rhodamine red-X secondary antibodies were purchased from Jackson ImmunoResearch Laboratories. Embryos were mounted in Vectashield (Vector Laboratories). Staining of en-Gal4/UAS-lacZ wings with X-gal (5-bromo-4-chloro-3-indolyl-β-D-galactopyranoside) was performed as described in Calleja et al. (1996).

Confocal images were acquired in a MicroRadiance 2000 microscope (Bio-Rad Laboratories). Images of Fig. 1 were collected with a 63 X/1.4 oil Plan-Apochromat objective and magnified 2X. Images of Fig. 4 were collected with a 40 X/1.3 oil Plan-Neofluar objective and magnified 2.4X. The acquisition software was Lasersharp 2000 BioRad 5.0. Wing images shown in Fig. 4 and Fig. S1 were obtained in an Axioptophot microscope (Carl Zeiss Microimaging, Inc.) with 2.5X/0.075 Plan-Neofluar objective, 10X/0.25 oil, and acquired with a camera (model DFC300, Leica) using Fire Cam software (Leica). Images were imported into Adobe Photoshop 7.0 for cropping and contrast adjustment before being assembled into figures using the same software.

Protein binding and in vitro phosphorylation assays

GST-fusion proteins were obtained from subclones in pGEX-3X (Amer
sham Biociences) of full-length Patj (Bhat et al., 1999), a gift from H. Bellen, and Crb wild-type or mutated intracellular domain (see Site directed mutagenesis) and Crb/WP113 (Hong et al., 2001), a gift from Y.N Jan (University of San Francisco, San Francisco, CA). To perform pull-down experiments, protein extracts were prepared from 0-16 h Drosophila embryos by homogenizing them in lysis buffer (50 mM Tris-HCl, pH 7.5, 150 mM NaCl; 1 mM EDTA; 1% Triton X-100 containing protease inhibitors). Debris were removed by centrifugation at 12,000 g for 15 min at 4°C. 300 μg of Drosophila protein extract were incubated for 16 h at 4°C with 30 μg of GST or of the different GST-fusion proteins bound to glutathione beads as described in S anz et al. (1999). Bound proteins were separated by SDS-PAGE and immunoblotted with anti-aPKC C20, anti-

Baz (a gift from A. Wodarz), anti-Dmoein (from D. Kiehart, Duke University, Durham, NC) or anti-Patj antibodies. For in vitro binding assays, His-

tagged DaPKC was obtained by immunoblotting with anti-Xpress epitope antibody (Invitrogen). In vitro kinase assays were performed as in (Leitges et al., 2001) using 30 ng of baculovirus-expressed recombinant human aPKC (Calbiochem) and 3 μg of the indicated GST-fusion proteins or of myelin basic protein. This phosphorylation depended on aPKC activ-

ity because it was inhibited by previous denaturation of the kinase. Note that although a human aPKC was used, the high similarity between human aPKC and DaPKC (79% of identity and 90% of similarity in the kinase do-

main) makes it unlikely the possibility of obtaining different results with DaPKC instead of human aPKC.
Site directed mutagenesis

Mutagenesis were performed by the site directed Quick-Change system (Stratagene). Full length DaPKC cDNA, isolated from a cDNA Drosophila azap library (a gift from P. Burhan, Paradigm Genetics, Research Triangle Park, NC), was used as a template to incorporate to DaPKC the CAAX mitochondrial material is available at http://www.jcb.org/cgi/content/full/jcb.200311031/DC1.

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Online supplemental material

An overexpression assay was used to exclude the possibility of nonspecific effects of DN-DaPKC expression. Fig. S1 shows that the polarity defects, found in wings and associated with the overexpression of DaPKC alone are normalized by overexpression of UAS-DaPKC. The online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200311031/DC1.
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