RESEARCH ARTICLE

Protein phosphatase 2A promotes the transition to G0 during terminal differentiation in Drosophila

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ABSTRACT

Protein phosphatase type 2A complex (PP2A) has been known as a tumor suppressor for over two decades, but it remains unclear exactly how it suppresses tumor growth. Here, we provide data indicating a novel role for PP2A in promoting the transition to quiescence upon terminal differentiation in vivo. Using Drosophila eyes and wings as a model, we find that compromising PP2A activity during the final cell cycle prior to a developmentally controlled cell cycle exit leads to extra cell divisions and delays entry into quiescence. By systematically testing the regulatory subunits of Drosophila PP2A, we find that the B56 family member widerborst (wdb) is required for the role of PP2A in promoting the transition to quiescence. Cells in differentiating tissues with compromised PP2A retain high Cdk2 activity when they should be quiescent, and genetic epistasis tests demonstrate that ectopic Cyclin E/Cdk2 activity is responsible for the extra cell cycles caused by PP2A inhibition. The loss of wdb/PP2A function cooperates with aberrantly high Cyclin E protein levels, allowing cells to bypass a robust G0 late in development. This provides an example of how loss of PP2A can cooperate with oncogenic mutations in cancer. We propose that the PP2A complex plays a novel role in differentiating tissues to promote developmentally controlled quiescence through the regulation of Cyclin E/Cdk2 activity.

KEY WORDS: Cell cycle, Quiescence, Terminal differentiation

INTRODUCTION

In adult metazoans, most terminally differentiated cells exit from the cell cycle and lie in a state of prolonged or permanent quiescence. The transition from active proliferation to quiescence in vivo is robust, often irreversible, and ensured by redundant cell cycle regulatory mechanisms (Buttitta et al., 2007; Firth and Baker, 2005; Nicolay et al., 2010; Pajcini et al., 2010; Wirt et al., 2010). By comparison, most studies of quiescence have been performed in cell culture, where contact inhibition, drug treatments or withdrawal of mitogens induce a quiescent state that is most often readily reversible (Coller, 2011). Although some of the key cell cycle regulators promoting quiescence in these contexts overlap [e.g. Retinoblastoma (RB) family members, Cyclin-dependent kinase inhibitors (CKIs)], there must be key differences between the reversible quiescence in cell culture and developmentally controlled robust cell cycle exit in vivo.

Recent work in mammalian cell culture has demonstrated that the level of Cdk2 activity after mitosis impacts the proliferation versus quiescence decision for the next cell cycle, such that cells with low Cdk2 activity enter a quiescent ‘G0-like’ state (Spencer et al., 2013). This suggests that mechanisms regulating Cyclin/Cdk2 activity during the final cell cycle in vivo could impact the timing and robustness of cell cycle exit in tissues. Consistent with this hypothesis, the loss of CKIs that inhibit Cyclin E/Cdk2 complexes or loss of the F-box protein Fbw7 (Fbxw7), which regulates Cyclin E stability, can partially delay proper cell cycle exit in certain tissues (Chen and Segil, 1999; de Nooij et al., 1996; Fero et al., 1996; Kiyokawa et al., 1996; Lane et al., 1996; Minella et al., 2008; Moberg et al., 2001; Tate et al., 2014). But even in the presence of aberrantly high Cyclin E/Cdk2, cell cycle exit is most often only delayed by one or two cell cycles in vivo, demonstrating the robustness of developmentally controlled quiescence (Baumgardt et al., 2014; Buttitta et al., 2010; Loeb et al., 2005; Nakayama et al., 1996).

Determining which cell cycle regulators are required for developmentally controlled cell cycle exit in vivo has posed some challenges. Redundancy in the function of multiple paralogs for each cell cycle regulator makes genetic analysis complicated, with studies often encompassing double or triple mutants (Gui et al., 2007; Wirt et al., 2010; Zindy et al., 1999). In addition, the late stage of development at which cell cycle exit occurs and the asynchronous nature of cell cycle exit in many tissues requires conditional genetic manipulations and timecourse analysis of samples. Drosophila eyes and wings have provided an advantageous system with which to study this process, as they undergo a relatively synchronized cell cycle exit during metamorphosis, have fewer paralogs and there are tools for precise conditional genetic manipulations. We used this system to examine cell cycle exit in terminally differentiating tissues and found that even in RB family member Rbf1 (Rbf–FlyBase)-deficient cells, Cyclin E/Cdk2 overexpression delays but cannot bypass cell cycle exit (Buttitta et al., 2007), suggesting that additional downstream mechanisms ensure the transition from proliferation to quiescence in vivo (Ehmer et al., 2014; Nicolay et al., 2010; Simon et al., 2009).

To identify additional mechanisms ensuring cell cycle exit, we examined Drosophila homologs of several tumor suppressor proteins expected to play a role in promoting quiescence. PP2A has been recognized as a tumor suppressor for over two decades (Janssens et al., 2005), but the molecular mechanism for PP2A in tumor suppression remains unknown. PP2A dephosphorylates RB family members to inhibit cycling (Kolupaeva and Janssens, 2013; Kurimchak and Grana, 2013), and removes an essential activating phosphorylation on the Cdk2 T-loop in vitro (Poon and Hunter, 1995). We therefore examined whether PP2A plays multiple, redundant roles to promote the developmentally controlled robust cell cycle exit in vivo.

Here, we show that cells with reduced PP2A function fail to transition to a quiescent state at the normal developmental time. Loss of PP2A function specifically during the final cell cycle leads ~10% of cells to perform an extra cycle before entry into permanent quiescence. Cells with compromised PP2A exhibit increased Cdk2 activity...
activity and aberrant E2F transcriptional activity. In the presence of high, oncogenic Cyclin E levels the loss of PP2A function allows cells to bypass a robust G0 mechanism during late stages in fly development. The PP2A enzyme is directed to distinct substrates via associations with different regulatory subunits, which can be highly dynamic during development. We show that the PP2A B56 regulatory subunit widerbost (wdb) is specifically required for the PP2A-mediated transition between proliferation and quiescence. Furthermore, this new function for PP2A/Wdb occurs even in the complete absence of RB/E2F/DP function, suggesting that it acts through downstream targets directly on the cell cycle machinery to promote quiescence in vivo.

RESULTS
Loss of PP2A delays the transition to quiescence in vivo
We performed a small-scale RNAi-based screen of ~500 randomly chosen Harvard Transgenic RNAi Project (TRiP) lines to identify new potential tumor suppressor genes involved in the proper timing of the developmentally controlled quiescence in the Drosophila eye. The primary screen was an adult eye color-based screen, an adaptation of the method described by Bandura et al. (2013). This was followed by a secondary, dissection-based screen to determine which hits from the initial screen effectively compromising cell cycle exit. Normally, the Drosophila eye becomes completely quiescent by 24 h after pupa formation (APF) (Buttitta et al., 2007). We therefore looked for RNAis that compromised quiescence, leading to ectopic cell cycles at 24 h APF. We used the Glass Multimer Reporter (GMR)-Gal4 (Ellis et al., 1993) driver to express UAS-controlled RNAi lines, and assayed for ectopic S phases by EdU incorporation (Buck et al., 2008) and ectopic expression of a cell cycle transcriptional reporter PCNA-GFP (Thacker et al., 2003) in eyes after normal quiescence from 24-30 h APF (supplementary material Fig. S1). Importantly, the GMR-Gal4 driver activates the UAS-RNAi specifically during the final cell cycle in the eye, thereby avoiding earlier deleterious effects. We found that two independent RNAi lines to the Drosophila PP2A catalytic subunit microtubule star (mts) and one to the sole PP2A scaffold A subunit Pp2A-29B caused ectopic S phases and cell cycle gene expression, at time points when the Drosophila eye should be fully quiescent (supplementary material Fig. S1A-C).

To confirm the RNAi results, we overexpressed a dominant-negative form of mts (mts<sup>DN</sup>) during the final cell cycle in eye muscles and found that it fully recapitulated the RNAi phenotypes. mts<sup>DN</sup> is a truncation that interacts non-productively with PP2A scaffolding (A) and regulatory (B) subunits, and serves as an effective competitive inhibitor when overexpressed (Evans et al., 1999). To test whether the role of PP2A in quiescence is eye specific, we overexpressed mts<sup>DN</sup> in the posterior wing during the final one to two cell cycles using the engrailed-Gal4, modified with a temperature-sensitive Gal80<sup>TS</sup> (en<sup>TS</sup>; see Materials and Methods for details). Similar to the eye, we observed ectopic S phases by EdU incorporation and ectopic mitoses by staining for phosphorylation of serine 10 on Histone H3 (PH3) at 24-28 h APF, time points when few cell cycles are normally observed in the wing (Fig. 1A,B). Overexpression of a functional wild-type mts (mts<sup>WT</sup>) had no observable effect on quiescence in these tissues (Fig. 1C,D), confirming that the observed phenotype is due to the loss of PP2A function.

We performed a timecourse and quantification of the mitoses in wings expressing mts<sup>DN</sup> (Fig. 1E; supplementary material Fig. S1D) or PP2A RNAi (Fig. 1F), which revealed continued mitoses in eyes and wings until 37 h APF, 13 h after the normal cell cycle exit in these tissues (Buttitta et al., 2007; Milan et al., 1996; Schubiger and Palka, 1987). To confirm the staining results, we measured the DNA content of pupal eyes expressing mts<sup>DN</sup> by flow cytometry. As expected from the ectopic cell cycle markers, an increased proportion of cells containing greater than 2C DNA content...
was observed in *mtsDN*-expressing eyes as compared with stage-matched controls without transgene expression (Fig. 1G). However, after 37 h APF, eyes expressing *mtsDN* exit the cell cycle with a normal G1 DNA content. Altogether, our data suggest that inhibition of PP2A during the final cell cycle causes a temporary delay of the transition to quiescence in a compartment-autonomous manner.

**Inhibition of PP2A leads to an extra cell cycle during the delay of quiescence**

We next investigated whether the delayed transition to quiescence caused by loss of PP2A leads to additional cell cycles or whether it is the result of a prolonged final cell cycle. We performed a clonal analysis to count the number of cells per clone, which reflects the number of cell divisions before quiescence, using the *heat shock (hs)-flip actin-stop>Gal4/UAS* ‘flip-out’ system. In brief, with this system a precisely timed heat shock leads to random cis-recombination between FRT sites (indicated by >) flanking a stop codon. Cells in which recombination occurs flip-out the stop codon to allow Gal4-mediated gene expression, which continues permanently in all daughter cells (Pignoni and Zipursky, 1997). In this manner, the number of daughter cells can be counted after the delayed quiescence at 37 h APF. Non-overlapping clones were induced at 0 h APF (just prior to the final cycle) by a short heat shock at 37°C. Transgenes to manipulate PP2A activity, as well as GFP to mark clones, and an apoptosis inhibitor (P35; to prevent loss of daughters, which confounds clonal cell counts) were expressed and cells per clone were counted blind for at least 100 clones in the wing blade at 42-44 h APF (Fig. 2A; supplementary material Fig. S1G). Most (95%) control clones expressing GFP and P35 contain two cells or fewer per clone, as the induction of the recombination occurs during or just prior to the final cycle. However, 15% of clones expressing *mtsDN* contain more than two cells per clone, which indicates that ~10% of *mtsDN*-expressing cells undergo an extra cell cycle before entering G0. For comparison, when the G1 cyclin complex Cyclin D/Cdk4 is directly overexpressed via *Gal4/UAS*, we observe 50% of cells performing an extra cell cycle, before becoming quiescent at 36 h APF (Buttitta et al., 2007).

To determine whether PP2A inhibition also causes extra cell cycles in the eye, we examined the morphology of the fly eye at late...
pupal stages (40–42 h APF). In wild-type or control retinas (Fig. 2B), the apical ommatidial structure consists of four cone cells in the center surrounded by interommatidial cells (IOCs) (Tomlinson and Munby, 2007; Hahn et al., 2010).

PP2A counteracts the phosphorylation of S6K, which we used as an assay to confirm the activity of our mts transgenes (supplementary material Fig. S2A). To test whether the increase in active phospho-S6K impacts the transition to quiescence, we overexpressed the GTPase Rheb, which increases cellular growth, TOR signaling and phospho-S6K (Saucedo et al., 2003). We did not observe any delay in cell cycle exit in the pupal wing, nor extra IOCs in the retina when Rheb was overexpressed, despite increased phospho-S6K (supplementary material Fig. S2). We therefore suggest that the function of PP2A in the transition to quiescence is independent of its role in regulating phospho-S6K.

**Inhibition of PP2A leads to ectopic Cdk2 activity**

Proper cell cycle exit in *Drosophila* eyes and wings is ensured by inhibition of E2F/DP-mediated transcription and suppression of Cyclin E (CycE)/Cdk2 activity (Bottitita et al., 2007; Firth and Baker, 2005). To examine whether cells with inhibited PP2A retain high Cdk2 activity, we used MPM2 antibody staining as an *in vivo* readout for ectopic Cdk2 activity at time points after normal cell cycle exit. MPM2 antibodies detect nuclear Cdk2 phospho-epitopes on the histone locus body (HLB) that normally occur only during S phase in proliferating cells, in addition to the well-described cytoplasmic epitopes present during mitosis (White et al., 2011, 2013). The size of the adult eye is also increased when PP2A is inhibited (Fig. 2H). Our cell count data suggest that PP2A inhibition enlarges the eye partly by ectopic cell proliferation, but we also consistently observed an increase in cell size. An increase in cell size is consistent with known functions of PP2A in the TOR/S6 kinase (S6K) pathway (Bielinski and Munby, 2007; Hahn et al., 2010).

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**Inhibition of PP2A function does not delay cell cycle exit by preventing APC/C activity**

The Anaphase-promoting complex/Cyclosome (APC/C) promotes timely cell cycle exit in *Drosophila* eyes and wings by degrading residual Cyclin A (CycA) and Cyclin B (CycB) during the final G1 (Bottitita et al., 2010; Ruggiero et al., 2012; Tanaka-Matukatsu et al., 2007). Furthermore, the APC/C complex can cooperate with RB proteins to reinforce cell cycle exit by promoting degradation of Skp2, which targets CKIs for destruction (Binné et al., 2007). PP2A can impact APC/C indirectly by counteracting CycB/Cdk1 phosphorylations (Hunt, 2013), as well as regulating the binding and stability of the APC/C inhibitor Emi (Wu et al., 2007), which functions similarly to *Drosophila* Regulator of cyclin A1 (Rca1) (Grosskortenhaus and Sprenger, 2002). We therefore examined whether APC/C function is inhibited when PP2A is compromised, leading to a delay in cell cycle exit. As a read-out for APC/C activity, we examined the levels of the known APC/C target CycB by immunohistochemistry. GFP-marked clones with transgene expression or PP2A gain-of-function with *mts* expression or *mts* expression created by PP2A loss, we used the MARCM system (Lee and Luo, 2001) to create *Dp* homozygous null mutant clones (supplementary material Fig. S3), and we verified that *Dp* mutant clones of this size in the pupal wing lack Dp protein (Fig. 3H,I). Dp null mutant clones expressing *mts* in pupal wings exhibit ectopic mitoses in wings at time points after normal cell cycle exit, whereas no mitoses were observed in any stage-matched *Dp* null mutant clones (Fig. 3G,H). This suggests that the delay of cell cycle exit upon inhibition of PP2A is epistatic to E2F/DP function, and reveals an additional role for PP2A in promoting quiescence that is independent of RB/E2F/DP complexes *in vivo*.

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cell cycle exit. However, we did not observe significant effects on Cdk1 inhibitory phosphorylations upon genetic manipulations of PP2A activity (Fig. 4F), in contrast to the ectopic expression of string, which strongly reduces Cdk1 inhibitory phosphorylation, as expected (Fig. 4F).

**PP2A interacts with negative regulators of CycE/Cdk2 activity in vivo**

Consistent with the evidence of ectopic Cdk2 activity when PP2A is compromised (Fig. 3A-C), we also observed functional genetic interactions between known negative regulators of Cdk2 activity and PP2A in the fly eye (Fig. 5). The sole p21/p27 (Cdkn1a/b) CKI in *Drosophila, dacapo* (*dap*), is a major inhibitor of the CycE/Cdk2 complex upon cell cycle exit (de Nooij et al., 1996; Lane et al., 1996; Sukhanova and Du, 2008). We examined whether loss of *dap* cooperates with inhibition of PP2A to delay quiescence by quantifying IOCs in late pupal stages as described previously for Fig. 2. The loss of one copy of *dap* (using the *dap* null allele) enhanced the effect of mtsDN expression (driven by GMR-Gal4) on the number of IOCs (18.7±0.3), compared with PP2A inhibition alone (17.3±0.2) (Fig. 5A-D). We also observed a 15% increase in adult eye size in *dap* heterozygotes expressing mtsDN compared with...
siblings with normal dap, while mtsDN expression alone caused a ~8% increase in adult eye size (Fig. 5E). In addition, we used the MARCM system to create dap homozygous null mutant clones, with and without PP2A inhibition via expression of mtsDN. In dap null mutant clones expressing mtsDN we also observed an increase in extra cells, including an increase in lens-producing cone cells, which is rarely seen in wild-type clones expressing mtsDN (Fig. 5G-I; supplementary material Fig. S4I). In a reciprocal experiment, we overexpressed dap together with mtsDN during the final cell cycle using the GMR-Gal4 driver, and observed a partial suppression of the enlarged eye phenotype caused by PP2A inhibition alone (Fig. 5F). This indicates that high Cdk2 activity is, at least in part, required for the enlarged eye phenotype resulting from PP2A inhibition.

To confirm that the enhancement of the dap eye phenotypes by mtsDN was due to impacts on CycE/Cdk2 function, we next examined a different negative regulator of CycE for genetic interactions with PP2A. The CycE protein level is controlled by the SCF complex with the ubiquitin ligase Fbw7, termed Archipelago (Ago) in Drosophila. Loss of ago in the fly leads to aberrant accumulation of CycE protein and temporally delays cell cycle exit of the bristle precursors in the eye (Moberg et al., 2001). Consistent with our results from loss of one copy of ago, we found that loss of one copy of ago (using the ago1 allele) also enhanced the mtsDN large eye phenotype (supplementary material Fig. S4).

We next examined whether modulation of PP2A activity itself could impact CycE protein levels or stability during the final cell cycle. We used GMR-Gal4 to drive expression of mtsDN or mtsWT during the final cell cycle in the eye and extracted protein from larval eyes for western blot analysis. We observed no significant increase in CycE protein levels when PP2A was compromised (supplementary material Fig. S4). Altogether, our genetic data indicate that PP2A acts through a pathway that is independent of RB/E2F/DP, and possibly in parallel to dap or ago to regulate CycE/Cdk2 activity.

PP2A inhibition cooperates with high CycE levels to bypass robust cell cycle exit

High CycE/Cdk2 activity during the final cell cycle in fly tissues delays cell cycle exit, but after only a few extra cell cycles a robust cell cycle exit mechanism ensures permanent quiescence (Baumgardt et al., 2014; Buttitta et al., 2010, 2007). We asked whether PP2A inhibition could promote cells with aberrantly high CycE expression to override the robust transition to quiescence and maintain proliferation during later stages of development, as suggested by its known role as a tumor suppressor. We used GMR-Gal4 to drive UAS-CycE expression together with the baculoviral apoptosis inhibitor P35 (to minimize corrective apoptosis) with or without PP2A inhibition via mtsDN. We examined proliferation in the eye at late pupal stages, several hours after the normal robust exit that occurs even in the presence of dysregulated CycE (Fig. 6A-D).

Pupal eyes expressing CycE without any PP2A modulation exhibit few S phases and mitoses at this late stage of development, whereas eyes expressing CycE together with mtsDN maintain high proliferation even after the stage normally associated with robust, permanent cell cycle exit. To confirm this, we isolated late pupal eyes and performed flow cytometry to examine their DNA content at 46 h APF. When eyes overexpress CycE, only ~9% of cells from the entire retina exhibit an abnormal S/G2 DNA content. By contrast, when PP2A is compromised in stage-matched eyes overexpressing CycE, 27% of cells exhibit abnormal S/G2 DNA contents (Fig. 6F). This suggests that PP2A normally functions as a barrier to limit the bypass of cell cycle exit when CycE is dysregulated in vivo.

The PP2A B56 subunit Widerborst regulates the transition to quiescence in vivo

To identify the PP2A regulatory subunit responsible for promoting quiescence in differentiating tissues, we systematically tested each
PP2A regulatory subunit in *Drosophila* (supplementary material Table S1) for phenotypes in the eye. We used RNAi to knockdown regulatory subunits during the final cell cycle, and compared the adult eye sizes of progeny (supplementary material Fig. S5A). Inhibition of the *Drosophila* B56 epsilon homolog (also called PPP2R5E) *wdb* contributed to the role of PP2A as a barrier to limit proliferation when CycE is dysregulated in terminally differentiating tissues. Consistent with our previous tests of genetic interactions between PP2A and negative regulators of CycE/Cdk2 activity, we observed that adult eye size is increased by 15% in *wdb* mutants (Fig. 5A). We observed that adult eye size is increased by 15% in *wdb* mutants (Fig. 5A). We observed that adult eye size is increased by 15% in *wdb* mutants (Fig. 5A). We observed that adult eye size is increased by 15% in *wdb* mutants (Fig. 5A).

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Inhibition of PP2A increases the T-loop phosphorylation of Cdk2

Our data suggest that PP2A may promote quiescence by limiting Cdk2 activity during the final cell cycle in vivo to restrict proliferation in terminally differentiating tissues. To test this hypothesis, we compromised PP2A function in vivo by expressing mtsDN and CycE in the posterior larval eye under the control of the GMR-Gal4 promoter, followed by immunoprecipitation of CycE to measure effects on CycE/Cdk2 kinase activity. When mtsDN is expressed in the posterior larval eye, we observe a 20-40% increase in CycE/Cdk2 kinase activity after normalization to the amount of CycE pulled down (Fig. 8A). One interpretation of this result is that PP2A knockdown leads to an increase in CycE/Cdk2 activity; however, it is also possible that the observed increase in CycE/Cdk2 kinase activity is a result of the increased proliferation that we observe when CycE is expressed under conditions in which PP2A is compromised (e.g. Fig. 6B,D) and not a direct effect of PP2A on CycE/Cdk2 activity.

To distinguish whether the increased Cdk2 activity is due to an immediate effect of PP2A inhibition on CycE/Cdk2 activity, we performed a kinase assay in Drosophila S2R+ cultured cells, where we can use short-term treatments with the pan-PP2A inhibitor okadaic acid (OA) to discern immediate versus indirect effects of PP2A inhibition on CycE/Cdk2 activity. We performed a timecourse and dosage test of OA treatment in S2R+ cells and confirmed that, with 30 min of OA treatment, PP2A activity is inhibited as assessed by increased phosphorylation of S6K. We performed a timecourse of OA treatment on S2R+ cells transiently transfected with a CycE expression vector and undertook CycE/Cdk2 kinase assays as described above. We found that with 30 min of OA treatment S2R+ cells exhibit a mild increase in CycE/Cdk2 activity (Fig. 8C), consistent with a direct effect of PP2A on CycE/Cdk2 activity. However, upon longer OA treatments (2 h shown), cells exhibit a reduction in CycE/Cdk2 kinase activity and a slower migrating form of CycE protein is immunoprecipitated (Fig. 8D). In mammalian cells, PP2A/B55 can dephosphorylate the N- and C-terminal phosphodegrons of CycE1 (Tan et al., 2014). Thus, the slower migrating form of CycE that we observe might be due to inhibition of PP2A/Twins by OA in Drosophila, which impacts the measured CycE/Cdk2 activity.

Altogether, our data suggest that short-term inhibition of PP2A can increase CycE/Cdk2 activity, whereas prolonged loss of PP2A function impacts CycE/Cdk2 in a complex manner due to differing functions of multiple PP2A complexes. We suggest there might be a smaller contribution of PP2A/Twins complexes to the overall PP2A activity during the final cell cycle in vivo, as compared with actively proliferating S2R+ cells in vitro.

PP2A can bind and remove an activating phosphate on the T-loop of human CDK2 in vitro (Poon and Hunter, 1995) and the T-loop and critical activating phosphorylation sites are conserved between mammals and Drosophila. To test whether PP2A complexes limit CycE/Cdk2 activity after mitosis by removing the Cdk2 T-loop phosphorylation, we turned to murine cell lines in which cells can be synchronized in M phase and Cdk2 phospho-T-loop-specific antibodies are available. We synchronized NIH 3T3 mouse embryonic fibroblasts (MEFs) in M phase by nocodazole treatment to depolymerize microtubules. We then released cells from the mitotic arrest and performed a timecourse analysis of Cdk2 T-loop phosphorylation in cells treated with the pan-PP2A inhibitor OA versus vehicle only. We observed that, 8 h after release from a mitotic arrest, treatment with OA for 30 min doubles T-loop phosphorylation compared with a vehicle-treated control (Fig. 8F).
We next tested whether a similar OA treatment in asynchronously proliferating MEFs could lead to an increase in Cdk2 T-loop phosphorylation. We observed a mild increase (20%) in Cdk2 T-loop phosphorylation in NIH 3T3 MEFs, whereas we detected no effect on Cdk2 T-loop phosphorylation in primary asynchronous MEFs. This suggests that redundant mechanisms may limit the effect of PP2A on the T-loop in primary cells. However, we observed a 50% increase in Cdk2 T-loop phosphorylation in p27 (Cdkn1b) knockout (p27KO) primary MEFs treated with OA (Fig. 8G), suggesting that PP2A might preferentially act on Cdk2 complexes that are not bound to Cdk inhibitors. We also observed increased levels of Cdk2 in p27KO MEFs, suggesting the role of PP2A might be fully revealed under conditions in which CycE/Cdk2 levels are high, but need to be rapidly inhibited. This is consistent with the genetic interactions we observed in Drosophila between PP2A and the p27 homolog dap.

An interaction between Drosophila Wdb and Cdk2 in a yeast two-hybrid assay has been reported (Stanyon et al., 2004). To confirm whether PP2A/Wdb complexes interact with CycE/Cdk2 complexes, we performed an immunoprecipitation of endogenous CycE with a V5-tagged Wdb in proliferating S2R+ cells (supplementary material Fig. S6A). We observe a mild enrichment of CycE in samples with Wdb-V5 pulled down, compared with controls and mock precipitations. The enrichment of CycE might be mild because PP2A/Wdb interacts with many substrates in a transient manner throughout the cell cycle. We propose that only a fraction of the precipitated Wdb-V5 complexes at a given time from asynchronously proliferating cells will therefore contain endogenous CycE.

To examine this in more detail, we next transfected CycE and Wdb-V5 expression vectors into S2R+ cells, and looked for colocalization of the proteins during the cell cycle. We found that CycE and Wdb-V5 colocalize in the cytoplasm during mitosis (supplementary material Fig. S6B). CycE is predominantly nuclear, but becomes dispersed in the cytoplasm during nuclear envelope breakdown in mitosis, whereas Wdb is predominantly cytoplasmic. This suggests that PP2A/Wdb complexes most likely interact with CycE/Cdk2 complexes transiently during or just after mitosis, before nuclear envelope reformation. This is consistent with our results in NIH 3T3 cells, which suggest that the maximal effect of PP2A on the Cdk2 T-loop occurs ∼8 h after a mitotic release.

Altogether, our studies suggest that B56/PP2A can act to restrict CycE/Cdk2 activity after mitosis to promote quiescence in vivo.

**DISCUSSION**

We identify a new role for PP2A in promoting quiescence during the transition to a permanently postmitotic state in Drosophila wings and eyes. We observe that ∼10% of cells undergo an extra cell cycle when PP2A functions are compromised. Although this effect may appear small, the cell cycle exit mechanism in vivo is so robust that cells completely lacking major cell cycle regulators such as the RB family member Rbf1 or the sole p21/p27-type CKI dap only exhibit a mitotic or S-phase index of ∼9% or less in eyes and wings (Butitta et al., 2007; Sukhanova and Du, 2008). As with other cell cycle regulators that act redundantly to promote cell cycle exit, we see synergism when PP2A is compromised under conditions that dysregulate the G1-S Cyclin, CycE (Firth and Baker, 2005).

Cells with reduced PP2A function exhibit ectopic cell cycle markers until 13 h after normal exit timing, which is roughly consistent with the one extra cell division that we measure by clonal lineage analysis. Importantly, the ectopic proliferation phenotypes we observe areтать the result of manipulating PP2A functions specifically during the final one to two cell cycles, without disturbance of prior PP2A mitotic functions during active proliferation.

**PP2A impacts the proliferation-quiescence decision in vivo**

Recent studies on PP2A in the proliferation-quiescence decision have revealed that PP2A activates the RB related family member p107 by dephosphorylation to promote growth arrest in chondrocytes (Kolupaeva et al., 2008; Kurimchak et al., 2013). Another group
recently found a second mechanism for PP2A to promote quiescence, whereby PP2A/B56 inhibits Ras signaling during G2 phase, which limits subsequent Myc expression and reduces CycE expression in the following G1 (Naetar et al., 2014). This promotes quiescence by limiting CycE, which would otherwise disrupt the association of RB family members with E2F/DP complexes by phosphorylation. Our data, however, suggest that there is yet another mechanism during the final cell cycle in vivo, independent of Ras/ERK signaling (supplementary material Fig. S7), Myc (supplementary material Fig. S7), CycE levels (supplementary material Fig. S4) and RB/E2F/DP function (Fig. 3), which promotes timely entry into quiescence. This additional mechanism acts directly on the cell cycle machinery, downstream or in parallel to RB/E2F/DP function, and appears to be crucial for the extremely robust type of developmentally controlled G0 observed in vivo.

**B56 regulatory subunits promote quiescence in vivo**

The emergence of multiple pathways for PP2A to promote quiescence might be a consequence of its broad functions, with impacts on various substrates in different cell cycle phases (Janssens et al., 2005; Mumby, 2007; Westermarck and Hahn, 2008; Yang and Phiel, 2010). In normal development, cells enter into the quiescent state in response to developmental signals, while in cell culture serum deprivation is most often used for synchronization in G0, via disrupted metabolic signals (Naetar et al., 2014). It might be that in these different biological contexts PP2A acts upon different targets to influence the proliferation-quiescence decision. PP2A is directed to distinct targets via the regulatory subunit, which has dynamic associations and localizations during the cell cycle. It is therefore important to note that, consistent with the recent work of Naetar et al. (2014), we also independently identified a B56 subunit (Wdb) as the main PP2A regulatory subunit promoting quiescence in vivo. However, our data demonstrate that wdb acts via a different mechanism to promote permanent cell cycle exit in vivo.

Most known cell cycle functions for PP2A in *Drosophila* involve the B55 regulatory subunit Twins and its roles in regulating mitotic entry and exit (Brownlee et al., 2011; Chabu and Doe, 2009; Chen et al., 2007; Wang et al., 2013, 2011). Consistent with this, when we...
manipulate PP2A activity in early tissues, such as the actively proliferating larval wing or eye, we also observe disruptions of mitosis. An inhibitory role for PP2A in the Hippo signaling pathway, which regulates tissue growth, survival and proliferation, has also been described (Ribeiro et al., 2010). However, when PP2A inhibits Hippo signaling it acts via B" regulatory subunits and has exactly the opposite effect to the growth and cell cycle phenotype that we observe here. The requirement for wdb during the final cell cycle to promote quiescence implies that the PP2A enzyme complexes may switch from predominantly B55 (Twins) to B56 (Wdb) during the final cell cycle, mitotic exit and the subsequent G0 arrest. Understanding how the switches in PP2A regulatory subunits are regulated during the cell cycle and in response to developmental signals will be an important area for future study.

**PP2A inhibits CycE/Cdk2 activity during the final cell cycle to promote quiescence**

A recent study monitoring single-cell cycle dynamics in cell culture demonstrated that thresholds of Cdk2 activity after the completion of mitosis regulate the subsequent proliferation-quiescence decision (Spencer et al., 2013). Our data suggest a role for PP2A in limiting Cdk2 activity during the final cell cycle in vivo to restrict proliferation in terminally differentiating tissues. Inhibition of PP2A during the final cell cycle leads to ectopic Cdk2 activity, as detected via the MPM2 epitopes at the HLB (Fig. 3), and genetically cooperates with the CycE inhibitors ago and dap. In mammalian cells, PP2A inhibition after mitosis leads to an increase in the activating Cdk2 T-loop phosphorylation. It is possible that PP2A and CycE/Cdk2 also share downstream targets in cell cycle regulation, similar to the role of PP2A/B55 complexes in reversing Cdk1 phosphorylation of mitotic targets. However, we could not confirm any effect of PP2A genetic manipulations on the phosphorylation of Drosophila Rbf, an important target of CycE/Cdk2 activity for cell cycle exit in flies (Meyer et al., 2000) (supplementary material Fig. S6C). We suggest that PP2A/Wdb acts to modulate CycE/Cdk2 activity during the final cell cycle to help promote rapid and timely induction of G0 during development.

**MATERIALS AND METHODS**

**Fly stocks**

For a list of fly stocks, see supplementary material Table S2.

**Histology and antibodies**

Pupae, staged from white pre-pupae (0 h APF) at 25°C, were dissected and fixed as described (Butitta et al., 2007). Pupal cuticle was removed from wings post fixation. Note that the wing hinge and notum were excluded from our quantifications. Hoechst 33258 (Molecular Probes, 1 µg/ml) was used to label nuclei. Antibodies used: rabbit anti-phospho-Histone H3 (Ser10) (Upstate, #06-570; 1:4000), mouse anti-MPM2 (Upstate, #05-368; 1:200), rabbit anti-GFP (Molecular Probes, A1122; 1:1000), mouse anti-CycB (DSHB, F2F4; 1:100), mouse anti-Discs large (DSHB, 4F3; 1:100) and anti-Drosophila Dp (gift from Dr M. Frolow; Frolow et al., 2005). Appropriate secondary antibodies were Alexa 488, 568 or 633 conjugated (Molecular Probes) or HRP conjugated (Jackson ImmunoResearch) and used at 1:4000. 5-ethyl-2′-deoxyuridine (EdU) incorporation was performed using the Click-iT EdU Alexa Fluor 555 Imaging Kit (Life Technologies).

**IOCs counting**

IOCs are shared by adjacent ommatidial cores, and the number of IOCs is quantified within an ommatidial group (OG) that covers a defined hexagonal area (bordered by the yellow dots in Fig. 2) with its apices being the adjacent ommatidial centers. Those secondary pigment cells crossed by the dashed lines were counted as half. At least 15 OGs were scored from independent samples for each genotype, following the method of Ou et al. (2007).

**Clonal analysis**

Clones were induced by heat shock for 7 min at 37°C between 48 and 70 h after egg deposition in a hsflp; tub-Gal4, UAS-GFP; tub::Gal80TS, UAS-Diap background. Animals were aged at 18°C (permissive for Gal80TS, Gal4 OFF), and shifted to 28°C (non-permissive for Gal80TS, Gal4 ON) at late L3 larval stage, collected at 0 h APF and aged to different stages in metamorphosis. Experiments using engrailed-Gal4 with Gal80TS were carried out in the same way, except that experiments restricting transgene expression to the final cell cycle were shifted to 28°C at 0 h APF. By phenotypic analyses and GFP visualization, we confirmed complete inhibition of Gal4 in the lines used here with Gal80TS at 18°C, and we detected activation of target genes within 6 h of shifting to 29°C. Development at 28°C proceeds 1.15 times faster than at 25°C, and at 45% of the 25°C rate at 18°C (Ashburner, 1989). All incubation times were adjusted accordingly. Hours APF are presented as the equivalent time at 25°C for simplicity. Loss-of-function clones were generated using MARCM (Lee and Luo, 2001). Larvae were heat shocked for 20 min at 37°C at early third larval instar, collected for staging at 0 h APF, aged at 25°C and dissected at the times indicated.

**Clonal cell counts to quantify cell divisions**

Non-overlapping clones labeled with GFP and expressing the indicated transgenes were induced at 0 h APF (white prepupa) with 2 min heat shock at 37°C. Wings were dissected and fixed 40-42 h later, nuclei were labeled with 1 µg/ml Hoechst 33258, and GFP-positive cells per clone were scored blind on a Leica DMi6000 microscope. Cells per clone were counted blind for at least 100 clones in the wing blade and the average cell number per clone reflects the number of cell divisions undergone before exit. We excluded clones in the wing margin, hinge, notum area and hemocytes in the veins. Transgenic expression of an apoptosis inhibitor (UAS-P35) was used in the clonal cell count experiments, including all controls.

**Flow cytometry**

Dissociation of cells from staged, dissected pupae and FACS were carried out as previously described (Flegel et al., 2013). All experiments were carried out at least three times; representative examples are shown.

**Western blotting and kinase assays**

Antibodies used: rabbit anti-CycE (Santa Cruz, sc-33748; 1:1000), goat anti-CycE (Santa Cruz, sc-15905; 1:200), mouse anti-CycA (DSHB, A12; 1:1000), mouse anti-CycB (DSHB; 1:1000), rabbit anti-phospho-Cdc2 (Cell Signaling, #9111; 1:1000), rabbit anti-Cdc2 (Upstate, #06-923; 1:1000), anti-phospho-S6K (Thr398) (Cell Signaling, #2920; 1:333), anti-Drosophila Myc (Santa Cruz, sc-28207; 1:1000), anti-dpERK (Sigma, M8159; 1:500), anti-pERK (Cell Signaling, #4370; 1:1000), anti-HA probe (Santa Cruz, sc-805; 1:1000), anti-mouse phospho-Cdk2 (Cell Signaling, #2561; 1:500), anti-mouse Cdk2 (Santa Cruz, M2, sc-163; 1:1000), anti-Drosophila Rbf (DX3; Lee et al., 2012), anti-α-tubulin (DSHB, 12G10; 1:1000), anti-β-tubulin (DSHB, E7; 1:1000), anti-γ-tubulin (Sigma, T6557; 1:1000) and anti-mouse GAPDH (Cell Signaling, 14C10; 1:2000) were used as loading controls with the appropriate HRP-conjugated secondary antibody. Enhanced chemiluminescence (ECL) detection (Amersham) followed by digital imaging (to prevent signal saturation, Bio-Rad) was performed and band signal intensity was quantified using ImageJ (NIH).

For kinase assays, cell lysates were collected from late L3 imaginal discs or S2R+ cells transfected with pMT-Cyclin E. For the S2R+ cells, 30 min or 2 h OA (50 nM) treatment was performed before cell harvest. For further details, see Guest et al. (2011).

**Microscopy**

Images were obtained using a Zeiss LSM 510 confocal or Leica DMi6000 epifluorescence system with deconvolution (ImageQuant). All images were...
cropped, rotated and processed using Adobe Photoshop. For brightness/contrast the Auto Contrast function was used. All brightness/contrast adjustments were applied equally on the entire image. Adult eye images were obtained using a Leica MZ10F microscope and a Nikon Ds-Vi1 digital camera. All adult eye images were measured using Nikon NIS Elements D software and processed with Adobe Photoshop.

Cell culture

_D. melanogaster_ S2R+ cells were cultured at 25°C in Schneider’s Insect Medium supplemented with 10% fetal bovine serum (FBS). NIH 3T3, p27<sup>Wdb-V5</sup> and p27<sup>2H10</sup> MEFs were cultured at 37°C, 5% CO2 in Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% FBS. For cell cycle synchronization, NIH 3T3 cells were treated with 200 ng/ml nocodazole for 18-20 h. The constructs pMT-Wdb-V5 and pMT-Cyclin E were transiently transfected using FuGENE (Roche) and expressed by copper induction (0.5 mM) in S2R+ cells.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

D.S. and L.B. developed the approach, performed the experiments and prepared the manuscript.

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Supplementary material

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