

S1 Fig. The PBAP complex is required for regenerative growth whereas the BAP complex is not.

(A) Pupariation rates of animals during normal development at 18°C. n = 79 pupae ($osa^{308}/+$) and 173 pupae (w^{1118}) from 3 independent experiments.

(B) Pupariation rates of animals after tissue damage (30° C) and regeneration (18° C). n = 101 pupae (osa^{308} /+) and 155 pupae (w^{1118}) from 3 independent experiments. Because the temperature shift to 30° C in the ablation protocol increases the developmental rate, the pupariation timing of regenerating animals (B) cannot be compared to the undamaged control animals (A).

(C) Wild-type (w^{1118}) regenerating wing disc at R24 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.

(D) *brm*²/+ regenerating wing disc at R24 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.

(E) Comparison of regenerating wing pouch size at 24 hours after imaginal disc damage in *brm*²/+ and wild-type (w^{1118}) animals. n = 11 wing discs (*brm*²/+) and 10 wing discs (w^{1118}).

(F) Wild-type (*w*¹¹¹⁸) regenerating wing disc at R24 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.
(G) *osa*³⁰⁸/+ regenerating wing disc at R24 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.

(H) Wild-type (*w*¹¹¹⁸) regenerating wing disc at R48 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.
(I) *osa*³⁰⁸/+ regenerating wing disc at R48 with wing pouch marked by anti-Nubbin (green) immunostaining. DNA (blue) was detected with Topro3.

(J) Comparison of regenerating wing pouch size at 24 and 48 hours after imaginal disc damage and regeneration in osa^{308} /+ and wild-type (w^{1118}) animals. At R24, n = 8 wing discs (osa^{308} /+) and 10 wing discs (w^{1118}). At R48, n = 6 wing discs (osa^{308} /+) and 8 wing discs (w^{1118}).

(K) Average number of mitotic cells (marked with PH3 immunostaining) per μ m² in the regenerating wing primordium at R24 in *bap170*^{Δ 135}/+ and wild-type (*w*¹¹¹⁸) animals. n = 8 wing discs (*bap170*^{Δ 135}/+) and 10 wing discs (*w*¹¹¹⁸).

(L) Wild-type (w^{1118}) regenerating wing disc at R24 with Nubbin (green) (L') and cleaved Caspase 3 (red)(L") immunostaining. DNA (blue)(L") was detected with Topro3.

(M) brm^2 /+ regenerating wing disc at R24 with Nubbin (green)(M') and cleaved Caspase 3 (red)(M'') immunostaining. DNA (blue)(M''') was detected with Topro3. (N-O) Wild-type (w^{1118}) (N) and brm^2 /+ (O) regenerating wing discs at R24 with Myc immunostaining.

(P) Quantification of anti-Myc immunostaining fluorescence intensity in the wing pouch in *brm*²/+ and wild-type (w^{1118}) regenerating wing discs at R24. n = 11 wing

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discs (brm^2 /+) and 12 wing discs (w^{1118}).

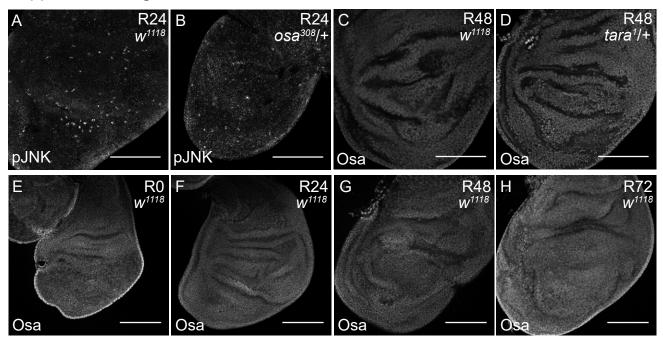
(Q-R) Wild-type (w^{1118}) (Q) and osa^{308} /+ (R) regenerating wing discs at R24 with Myc immunostaining.

(S) Quantification of anti-Myc immunostaining fluorescence intensity in the wing pouch in osa^{308} /+ and wild-type (w^{1118}) regenerating wing discs at R24. n = 6 wing discs (osa^{308} /+) and 8 wing discs (w^{1118}).

Error bars are SEM. Scale bars are 100µm for all wing discs images. *** p <

0.01, Student's *t*-test.

Supplemental Figure 2



S2 Fig. The function of BAP in preventing P-to-A transformation.

(A-B) Wild-type (w^{1118}) (A) and osa^{308} /+ (B) regenerating wing discs at R24 with phospho-JNK immunostaining.

(C-D) Wild-type (w^{1118}) (C) and $tara^{1/+}$ (D) regenerating wing discs at R48 with

Osa immunostaining.

(E-H) Wild-type (w^{1118}) regenerating wing discs at 0, 24, 48, and 72 hours after

imaginal disc damage and regeneration with Osa immunostaining.

Scale bars are $100\mu m$ for all wing discs images.