TITLE

Octopamine signals coordinate the spatial pattern of presynaptic machineries in the *Drosophila* mushroom bodies

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SUMMARY

Neurons possess numerous synaptic terminals. Presynaptic structures in a single motor neuron exhibit heterogeneity that associated with distinct characters of synaptic vesicle release. However, such heterogeneity is scarcely reported in the central nervous system, and its regulatory mechanism remain unknown. Here, we explored the intracellular diversity of presynaptic structures within the Kenyon cells of the *Drosophila* mushroom bodies. Applying the CRISPR/Cas9-mediated split-GFP tagging, we devised cell-type specific fluorescent labeling of the endogenous active zone scaffold protein, Bruchpilot (Brp). Morphometry of individual Brp clusters revealed heterogenous accumulations among the axon terminal compartments. Mechanistically, the localized octopaminergic signaling along Kenyon cell terminals regulate the Brp heterogeneity via Octβ2R and cAMP signaling. We further found that acute food deprivation reduced the compartmental heterogeneity of Brp accumulation in an octopaminergic signaling-dependent manner. These findings are consistent with the mushroom body functions in integrating the signals of changing physiological states.

KEY WORDS

Split-GFP; mushroom bodies; Kenyon cell; presynaptic active zone; synaptic heterogeneity; compartments; octopamine; stress; cAMP signaling

INTRODUCTION

The molecular composition of chemical synapses determines the function of neurons, and therefore exhibits a remarkable diversity among cell types [1,2]. Studies in the motor neuron demonstrated that presynaptic molecular assemblies are highly heterogeneous even within a single cell [3,4]. Critically, this structural heterogeneity is correlated with synaptic functions such as release probability [5–7]. Given that a single neuron in the central nervous system (CNS) synapses onto multiple target cells, such intracellular

synaptic heterogeneity could generate complex computation of the circuit. However, the regulatory mechanisms of such synaptic heterogeneity in central neurons are largely unknown.

Neurons projecting to the *Drosophila* mushroom bodies (MBs) can serve as an excellent model system to study intracellular synaptic heterogeneity. Kenyon cells (KCs), the major MB intrinsic neurons, synapse onto five sets of post-synaptic partners in spatially segregated compartments [8,9]. Postsynaptic MB output neurons (MBONs) have distinct and compartmentalized dendrites within the MB lobes [9], and their activities collectively determine the MB output [10–14]. The compartmental distinctions in presynaptic assemblies within KCs are likely to be critical in controlling MB-guided behaviors.

We here aim to characterize the intracellular diversity of presynaptic structures in a KC-specific manner. Brp, the homolog of ELKS/CAST/ERC, is a major scaffold protein accumulated at active zones (AZs) and the main component of electron-dense projections decorating AZs [15–18]. Brp clusters determine functional characteristics of synapses such as the release probability of synaptic vesicles [5–7], therefore their structures serve as a proxy for estimating the synapse function. As with other neuropils in the central nervous system, MBs are composed of densely packed synapses originating from different types of neurons [19,20]. Therefore, it is challenging to characterize specific endogenous synapse proteins only in the target neuron without super-resolution microscopy [21,22]. Inspired by recent advances in *Drosophila* genetics to label endogenous Brp in designated cell types [23,24], we took advantage of the CRISPR/Cas9-mediated split-GFP tagging strategy for this purpose [25,26]. Profiling the character of individual Brp clusters only within KCs revealed heterogenous Brp accumulation along the MB compartments. Here we report the dynamics of this intracellularly heterogeneous AZ structure and approach its supervising regulatory mechanisms.

RESULTS

Cell-type specific visualization of endogenous active zone scaffold proteins Brp

Chemical tagging using Brp::SNAP [27] suggests endogenous Brp localizes heterogeneously within the MB (Figure S1), which is constituted by pre-synapses of various cell types (Figure S2). To visualize endogenous Brp in designated cell types, we inserted the GFP_{11} fragment (the eleventh β-strand of the super-folder GFP) just prior to the stop codon of *brp* using CRISPR/Cas9. The self-assembly and reconstitution of GFP is induced by expressing the $GFP_{1\text{-}10}$ fragments using the GAL4/UAS system (Figure 1A).

We validated this system with GAL4 driver lines that label 7 different MB-projecting neuronal populations: 3 KC subtypes, 2 dopamine neuron clusters (PPL1 and PAM) and 2 single interneurons (DPM and APL neurons) [9,28] (see STAR⍰METHOD for driver information). Confocal microscopy detected reconstituted GFP (rGFP) signals only at restricted areas, in contrast to the dense signals in Brp::SNAP or Anti-Brp immunostaining (Figure 1B, Figure S1). The Brp::rGFP signal appeared as individual puncta and matched the areas of axon terminals of the cell types (Figure 1B). Especially, Brp was labeled at single-cell resolution for the APL and DPM neuron [8,29]. Strikingly, the Brp::rGFP signal intensity appeared to be heterogenous along the axon terminals of KCs (Figure 1B). Since individual KCs arborize onto the entire lobe structure, these

results suggest distinct accumulations of Brp intracellularly.

To test whether Brp::rGFP labels dynamic synaptic structures, we benchmarked its performance with different live-imaging contexts. The axon arborizations of pleural muscle motor neurons (PM-Mns) can be imaged on the abdominal body wall during pupal development [30]. Live-imaging Brp::rGFP in the PM-Mn revealed the formation of individual clusters at 48 hours after puparium formation and the increasing density in the same branch following two days (Figure 1C). We further visualized Brp::rGFP dynamics in the ex-vivo preparation of the larval brain, and found multiple fusion and fission events of Brp clusters in the MB calyx (Figure 1D). These results are consistent with the AZ assembly through liquid-liquid phase separation [31] and demonstrated the immense potential to visualize the dynamics of endogenous AZ structures.

Intracellular active zone heterogeneity among MB compartments

To quantify the presynaptic Brp heterogeneity we observed in KCs (Figure 1B), an image-processing pipeline was developed to better characterize the signal intensity (total grey value) of individual Brp::rGFP puncta. Optimizing the parameter combination of 3D image deconvolution [32,33] and spot segmentation enabled to resolve Brp::rGFP clusters that represent single AZs using conventional confocal microscopy. We found that Brp::rGFP accumulation was heterogeneous among KC compartments, and it was especially pronounced in γ KCs (Figure 2A, Figure S3). The median Brp::rGFP signal intensity in the γ5 compartment was nearly three times as intense as that in γ1 (Figure 2C). These results suggest that individual KCs tune their presynaptic structures according to compartments (Figure 2B).

As different MB compartments are functionally coordinated and integrate internal states such as nutritional states, sleep need and aging [10,34–38], we examined whether KCs adapt the synaptic structures upon physiological changes. We quantified the median intensity of Brp::rGFP puncta in each compartment and calculated the variance of the 5 compartments as a measure of compartmental heterogeneity (Figure 2D). Strikingly, the compartmental heterogeneity in γ KCs significantly decreased upon food deprivation for 48 hours (Figure 2E, Figure S6F). In contrast, we did not find a significant effect by aging for 30 d (Figure S5). These results suggest that acute stress can drive local adjustment of AZ structure within KCs, reflecting reorganized compartmental activities.

Associations of Ca^{2+} channel and Brp clusters

In motor neurons, Brp is well characterized to serve as a scaffold protein accumulating synapse proteins such as voltage-gated Ca^{2+} channels [15,17,39]. Therefore, we hypothesized if Brp heterogeneity in γ KCs reflects the Ca^{2+} channel enrichment at AZs. To test the hypothesis, we double-labelled the endogenous α subunit of the voltage-gated Ca^{2+} channel, Cacophony (Cac) and Brp by Cac::sfGFP [6] and Brp::SNAP, respectively. We found that Cac::sfGFP and Brp::SNAP signals were co-localized in the γ lobe (Figure 3A-3B) and their intensities were correlated in Brp-positive areas (Figure 3C-E, Figure S4).

This correlation between Brp and Cac clusters suggest that the Ca^{2+} concentration is differentially set along the γ KCs. To measure the basal Ca^{2+} concentration nearby AZs, we expressed Brp^{short}::mCherry::GCaMP6s, a ratiometric Ca^{2+} sensor fused to mCherry and truncated Brp (Figure 3F) [40], in KCs. We found that

basal GCaMP signals were compartmentally distinct and gradually increased towards the γ5 compartment (Figure 3F). This compartmental pattern of GCaMP signals nicely mirrored the Brp heterogeneity. Taken together the colocalization of Brp and Cac, these results suggest that the Brp accumulation per AZ is set compartmentally, differentiating the basal Ca^{2+} influx via Ca^{2+} channels at AZs [41].

Octopamine input underlies the Brp compartmental heterogeneity

Each compartment receives dopamine projections from structurally and functionally distinct subsets of dopaminergic neurons (Figure 3A) [9]. We therefore examined whether dopamine inputs determine the compartmental Brp accumulation by knocking-down dopamine receptors (DopR1, DopR2 and D2R) in KCs using RNA interference (RNAi) [42,43]. None of these disruptions showed a significant effect on the compartmental heterogeneity (Figure 4B, Figure S6A).

The octopaminergic neurons (OANs), particularly OA-VPM3 and OA-VPM4 neurons, are known to project to the MB and concentrate their innervations in the γ1 compartment [9,44]. We characterized presynaptic components of OA-VPM3/4 visualized by Brp::rGFP and nSyb::CLIP (neuronal synaptobrevin, nSyb; synaptic vesicle marker) and found they were densely localized to γ1 compared to the other compartments, complementary to the Brp compartmental heterogeneity of KCs (Figure 5A-5D). These results suggest localized synaptic output of OANs along the γ lobe, potentially underlying the Brp compartmental heterogeneity.

To examine the contribution of octopamine to the Brp accumulation within KCs, we measured the Brp compartmental heterogeneity in the mutants of tyramine β-hydroxylase (*Tβh*), the enzyme that catalyzes the synthesis of octopamine from tyramine [45]. We generated two new *Tβh* null alleles: insertion mutant $T\beta h^{SK1\text{-}8}$ and deletion mutant $T\beta h^{SK2\text{-}4}$ using CRISPR/Cas9-mediated mutagenesis. We visualized Brp::rGFP in KCs of these mutants and found the significantly decreased compartmental heterogeneity of the γ KCs (Figure 5E, Figure S6B). This suggests that octopamine directly controls the AZ structure in KCs.

To identify the signaling pathway that regulates the Brp compartmental heterogeneity, we knocked-down octopamine/tyramine receptors (including OAMB, Octα2R, Octβ1R, Octβ2R, Octβ3R, OctTyrR, TyrR, TyrRII) in KCs using RNAi [42,46]. Among all the receptors tested, only Octβ2R knockdown significantly decreased the compartmental heterogeneity (Figure 6A, Figure S6C-6D). We could not test the effect of Octβ1R and Octβ3R because their deficiency in KCs caused abnormal development of the MB. Octβ2R is coupled to the Gs alpha subunit to stimulate cyclic adenosine monophosphate (cAMP) production [47]. To test whether cAMP plays a role in Brp accumulation, we knocked-down the adenylate cyclase Rutabaga (Rut) in KCs using RNAi [42,48–52]. The downregulation of Rut resulted in a significant decrease of the compartmental heterogeneity (Figure 6B, Figure S6E), suggesting that octopamine functions through Octβ2R-cAMP pathway.

Nutritional states adjust the Brp compartmental heterogeneity through octopamine signaling

The spontaneous firing of octopamine neurons was shown to be decreased upon starvation [53]. Therefore, we hypothesized state-dependent AZ remodeling through octopamine (Figure 7A). To test this hypothesis, we starved the *Tβh* mutants for 48

hours and measured the Brp::rGFP heterogeneity in γ KCs. Indeed, there was no significant state-dependent effect on the Brp compartmental heterogeneity in the *Tβh* mutants (Figure 7B, Figure S6G). Taken together, we propose that internal states adjust the synaptic output of MB compartments through octopamine signaling that supervise the synaptic structures within KCs.

DISCUSSION

By labeling endogenous AZ scaffold protein Brp, we have been able to resolve individual AZs at single-cell level. The substantial intracellular heterogeneity of Brp accumulation within γ KCs is found to be regulated by internal states of the animal. Other than the compartmental heterogeneity of Brp clusters within KCs, our experimental pipeline enables further profiling of other AZ characteristics such as the local AZ density.

How does octopaminergic signaling modify Brp accumulation? This study identified the requirement of Octβ2R and Rut in regulating the AZ structure in KCs (Figure 6). Octβ2R was shown to stimulate cAMP synthesis [47], suggesting that cAMP could play a determining role. Consistent with this idea, *rut* mutant displays abnormal AZ structures [54]. We propose that the localized innervations of octopamine neurons onto KCs (Figure 5) results in different cAMP concentrations along the γ lobe. In both KCs and motor neurons, cAMP concentration was shown to be locally regulated [55–57]. The affinity or stability of Brp to AZs among different compartments is perhaps distinctly set by the cAMP signaling compartmentation [58]. Interestingly, both DopR1 and Octβ2R stimulate cAMP biosynthesis, while lack of DopR1 had no significant effect on the Brp heterogeneity (Figure 4). This difference can be attributed to the differential targets the receptors activate downstream cAMP, because protein kinase A (PKA) is activated by octopamine but not dopamine in the γ lobe [59]. It is conceivable that PKA modifies the state of Brp to impact the AZ structure and synaptic function [60].

AZ structures, especially Brp clusters, frequently serve as a proxy to explain synaptic functions [5–7,15,17,39,61]. Consistent with previous studies, our data suggest that Brp regulates the localization of Ca^{2+} channels in KCs [15,17,62]. Brp heterogeneity may result in compartmental basal Ca^{2+} concentrations at AZs (Figure 3) and thus modulate spontaneous synaptic vesicle release via the stochastic activity of Ca^{2+} channels [41]. Indeed, the enrichment of the Ca^{2+} channel is associated to higher spontaneous release frequency in *Drosophila* NMJ [7,63] . Although the relationship between Brp accumulation and evoked release in KCs seems more complicated since it involves acute and compartmental dopaminergic modulations [13,64–66]. Unlike evoked activities, spontaneous release of synaptic vesicle persists in the absence of stimulation thereby affecting the states of the animal [10].

The Brp compartmental heterogeneity within KCs might directly modulate the MB output through controlling the activity of different MBONs. Each type of MBONs serves as an independent output unit, together regulating a variety of behaviors [11–14,37,67–69]. Starvation changes the activity balance between MBON-γ1 and MBON-γ5 thus enhances appetitive memory expression [67,69,70], and MBON- γ5 and MBON- γ2 oppositely control sleep amount [71]. Consistently, we found that the compartmental AZ heterogeneity within KCs, measured by Brp, decreased upon starvation (Figure 2E). Considering the associations between octopamine and feeding behaviors [53,72–78], KCs generates behavior adaptations in response to changing internal states and environments

by local AZ tuning [37,79,80].

SUPPLEMENTAL INFORMATION

Document S1. Figures S1-S6

ACKNOWLEDGEMENT

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AUTHOR CONTRIBUTIONS

Conceptualization, H.W., N.Y. and H.T.; Investigation, H.W., S.E., K.S., A.A., K.J.; Methodology, H.W., N.Y., S.E. and Y.M.; Writing – Original Draft, H.W, S.E.; Writing – Review & Editing, H.W., S.E. and H.T.; Funding Acquisition, H.W. and H.T.; Resources, K.J. and S.K.; Supervision, D.W., N.Y., S.K. and H.T.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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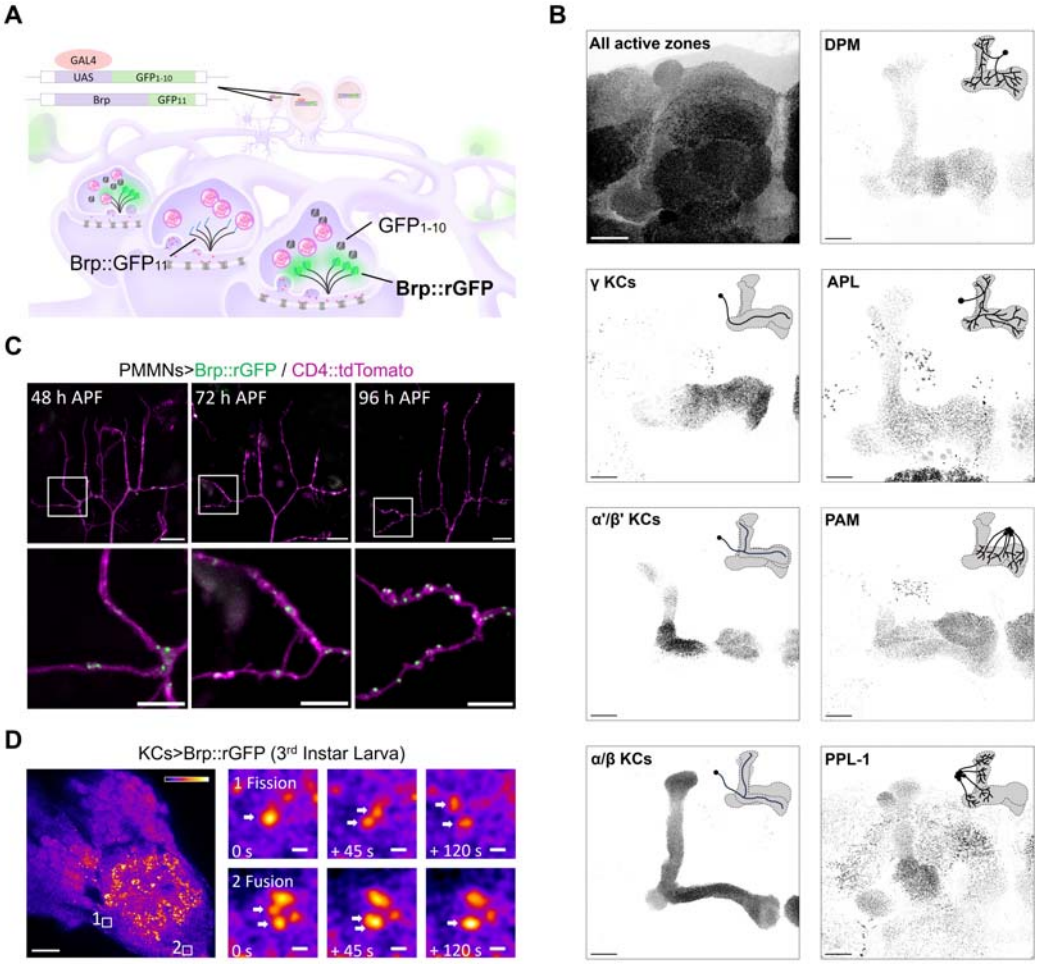

Figure 1. Cell-type specific dissection of the endogenous active zone scaffold

(A) Schematic of split-GFP tagging system. GFP_{11} was inserted prior to the stop codon of *brp*, while GFP_{1-10} was expressed in specific cell types using the GAL4-UAS system. GFP_{11} and GFP_{1-10} reconstitute to emit fluorescence.

(B) Visualization of Brp::rGFP in a cell-type specific manner. Immunostaining using antibody nc82 which labels all Brp in the brain, as a comparison to Brp::rGFP. Diagrams in each panel show the innervation pattern of neurons in the MB. GAL4 lines used: γ KCs (MB009B), α/β KCs (MB008B), α'/β' KCs (MB370B), PPL-1 (TH-GAL4), PAM (R58E02), DPM (VT64246), APL (GH146). Scale bar, 20 μm.

(C) Long-term live-imaging of Brp::rGFP (green) in a growing motor neuron (magenta, CD4::tdTomato) from 48 to 96 hours APF. OK371-GAL4 was used to express GFP_{1-10} and CD4::tdTomato. White boxes indicate zoomed-in areas shown in the lower panels. Scale bars: upper panels, 10 μm; lower panels, 5 μm.

(D) Ex vivo live-imaging of Brp::rGFP in 3rd instar larval KCs. GFP_{1-10} was expressed using R13F02-GAL4. The left panel shows the MB calyx region. White boxes indicate zoomed-in areas shown on the right panels. Box 1 and 2 demonstrate Brp::rGFP fission and fusion event. Whites arrows indicate Brp::rGFP clusters undergoing fission or fusion. Scale bars: left panel, 10 μm; right panels, 200 nm.

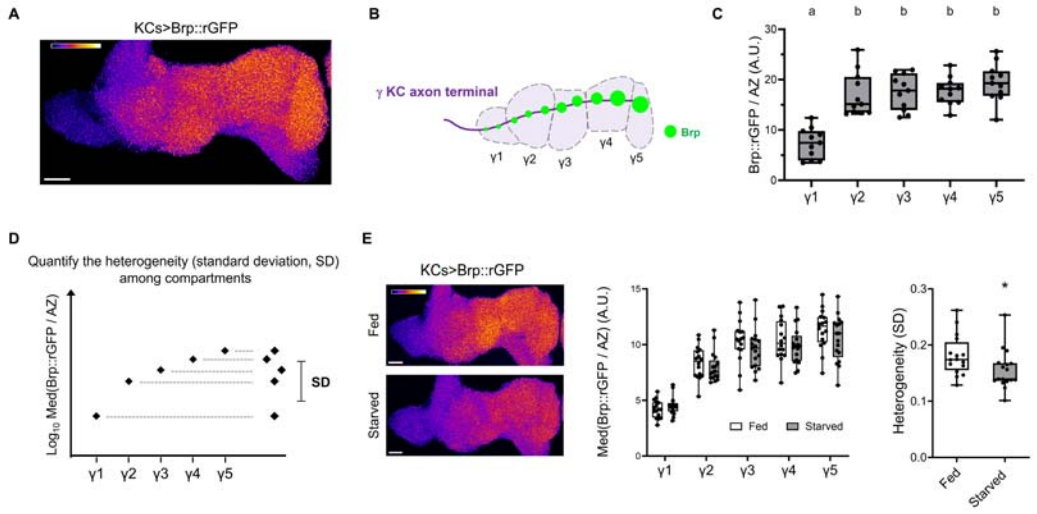

Figure 2. Compartmental presynaptic heterogeneity

(A) Brp::rGFP visualized in KCs. The horizontal lobe of the MB was imaged. Brp::rGFP was visualized using R13F02-GAL4.

(B) Diagram showing compartments in the MB γ lobe and the intracellular Brp localization in a single γ KC.

(C) Median intensity of Brp::rGFP clusters in each γ lobe compartment. Kruskal-Wallis test. n = 11. γ1 vs. γ2: P = 0.0012; γ1 vs. γ3: P = 0.0004; γ1 vs. γ4: P = 0.0004; γ1 vs. γ5: P < 0.0001.

(D) Standard deviation (SD) as an indicative of Brp::rGFP heterogeneity among compartments. Median Brp::rGFP intensity values of compartments were log-transformed for analysis. SD is calculated from five transformed medians.

(E) Starvation for 48 hours reduced The Brp compartmental heterogeneity in KCs. The middle panel shows the raw median Brp::rGFP intensities in each γ compartment for the control and starvation group. The heterogeneity (SD) was quantified for individual brain samples. Mann-Whitney U-test. Fed (n = 16) vs. Starved (n = 17): p = 0.0187; Scale bar, 10 μm.

Data are represented as box plots showing center (median), whiskers (Min. to Max.). Significant differences (P < 0.05) are indicated by distinct letters or *. Kruskal-Wallis test (C) and Mann-Whitney U-test (E).

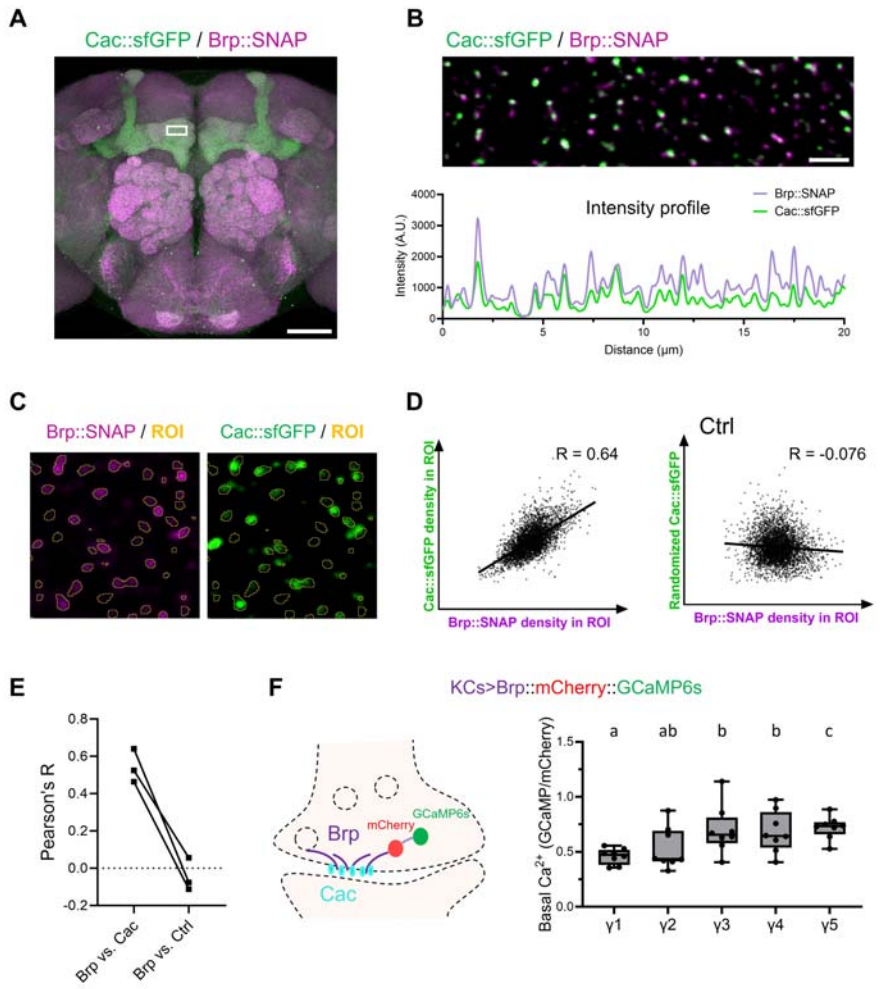

Figure 3. Brp is associated with Ca^{2+} channels in the fly brain

(A) Co-labeling of Cac::sfGFP (green) and Brp::SNAP (magenta) in an adult brain. The write box indicates the zoomed-in areas shown in (B**)**. Scale bar, 50 μm.

(B) Co-localization of Brp::SNAP and Cac::sfGFP signals. Signal intensity profiles of both Brp::SNAP (green) and Cac::sfGFP (magenta) in the image were plotted below. Scale bar, 10 μm.

(C) Correlation between Brp::SNAP (green) and Cac::sfGFP (magenta) signal intensities. The image is a selected area from γ5. Yellow circles show the 3D ROIs generated by segmenting Brp::SNAP signals. A loose setting was applied to include surrounding pixels. The same ROI set was used to quantify the signal density (total grey value divided by the ROI volume) for both Brp::SNAP and Cac::sfGFP.

(D) Scatter plot showing the correlation between Brp::SNAP and Cac::sfGFP signal intensities in an image sample. A 180˚ rotated Cac::sfGFP image was used as a control (see also Figure S4). Pearson's correlation coefficient (R) is shown.

(E) Pearson correlation coefficient (R) from three individual γ lobes showing the correlation between Brp::SNAP and Cac::sfGFP signal intensities.

(F) Basal Ca^{2+} concentration varies by compartment. The left panel shows the schematic of the Brp::mCherry::GCaMP6s ratio-matric sensor. GCaMP6s sensor is fused to

mCherry and Brp^{short}. The sensor was expressed by R13F02-GAL4 and GCaMP6s signal is normalized by mCherry for analysis. Kruskal-Wallis test. n = 8. γ1 vs. γ3: P = 0.0494; γ1 vs. γ4: P = 0.0494; γ1 vs. γ5: P = 0.0155; γ2 vs. γ5: P = 0.0494;
Data are represented as box plots showing center (median), whiskers (Min. to Max.). Significant differences (P < 0.05) are indicated by distinct letters. Kruskal-Wallis test.

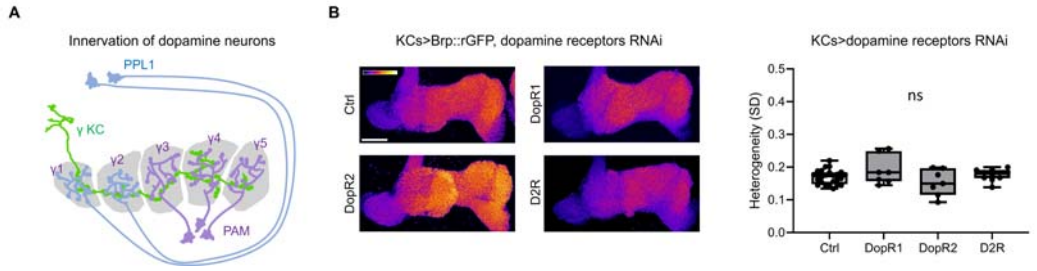

Figure 4. Dopamine receptors knockdown does not affect The Brp compartmental heterogeneity

(A) Schematic showing the innervation patterns of dopamine neurons. The γ lobe is innervated by dopamine neurons from the PPL1 and PAM clusters.
(B) Knockdown of dopamine receptors does not significantly alter the Brp compartmental heterogeneity. Three types of DA receptors, DopR1, DopR2 and D2R were knocked down using RNAi in KCs specifically with R13F02-GAL4. Representative images of Brp::rGFP from each group are shown. Scale bar: 20 μm. Ctrl (n = 24) vs. DopR1 (n = 7), DopR2 (n = 7) and D2R (n = 10): P > 0.05.
Data are represented as box plots showing center (median), whiskers (Min. to Max.). ns = not significant by Kruskal-Wallis test.

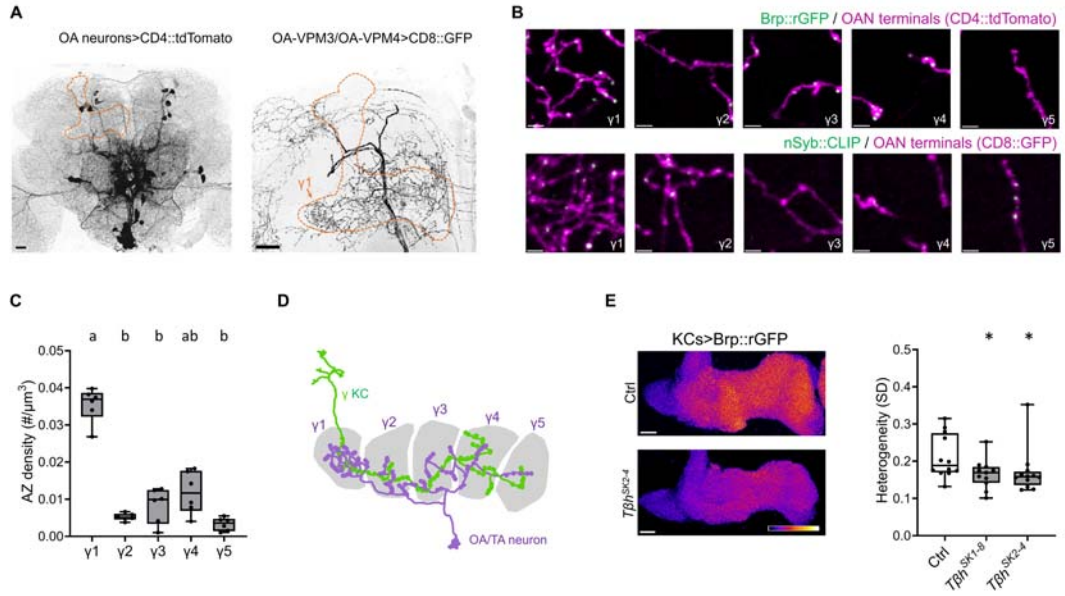

Figure 5. Octopamine controls the AZ structure in KCs

(A) OANs in the adult brain. OANs were labeled by Tdc2-GAL4 driven CD4::tdTomato in the left panel. OA-VPM3/4 neurons were labeled with MB022B-GAL4 [9] driven CD8::GFP (right panel). Dashed lines indicate the MB. Scale bars, 20

µm.
(B) AZ number and synaptic vesicle localization of OANs along the γ lobe. GFP_{1-10} and CD4::tdTomato were expressed using Tdc2-GAL4. nSyb::CLIP labeling synaptic vesicles and CD8::GFP were expressed using MB022B-GAL4. Scale bars, 2 µm. Images were cropped from the same MB and are identical in size.
(C) AZ density of OANs across γ compartment. Brp::rGFP was visualized using Tdc2-GAL4, and cluster density was quantified. n = 6 brains. Kruskal-Wallis test. γ1 vs. γ2: P = 0.0066; γ1 vs. γ3: P =0.0319; γ1 vs. γ5: P =0.0004.
(D) Schematic showing the innervation pattern of OANs along the γ lobe.
(E) Reduced Brp compartmental heterogeneity in *Tβh* mutants. Representative images of Brp::rGFP in control and $T\beta h^{SK2\text{-}4}$ background are shown. Scale bars, 10 µm. Kruskal-Wallis test. Ctrl (n = 12) vs. $T\beta h^{SK1\text{-}8}$ (n = 11) and $T\beta h^{SK2\text{-}4}$ (n = 12). Ctrl vs. $T\beta h^{SK1\text{-}8}$: P = 0.0442; Ctrl vs. $T\beta h^{SK2\text{-}4}$: P = 0.0228. *P < 0.05.
Data are represented as box plots showing center (median), whiskers (Min. to Max.). Significant differences (P < 0.05) are indicated by distinct letters or *. Kruskal-Wallis test.

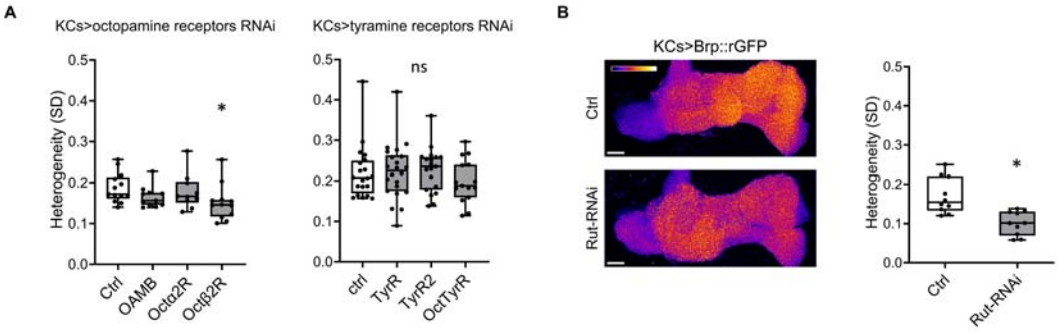

Figure 6. Octβ2R and cAMP underly the Brp compartmental heterogeneity
(A) Knockdown of Octβ2R reduced the Brp compartmental heterogeneity in γ KCs. Receptors are knocked-down in γ KCs specifically using RNAi with R13F02-GAL4. Kruskal-Wallis test. Ctrl (n = 13) vs. OAMB (n = 11), Octα2R (n = 10) and Octβ2R (n = 12): P= 0.0228 for Ctrl vs. Octβ2R. Ctrl (n = 20) vs. TyrR (n = 22), TyrR2 (n = 22) and OctTyrR (n = 15). *P < 0.05; ns = not significant.
(B) Knockdown of Rut significantly decreased the Brp compartmental heterogeneity in γ KCs. Representative images showing Brp::rGFP in WT and in Rut knockdown group. Mann-Whitney U-test. Ctrl (n = 10) vs. Rut-RNAi (n = 10): P = 0.0011.
Data are represented as box plots showing center (median), whiskers (Min. to Max.). *P < 0.05 and ns = not significant. Kruskal-Wallis test (A) and Mann-Whitney U-test (B).

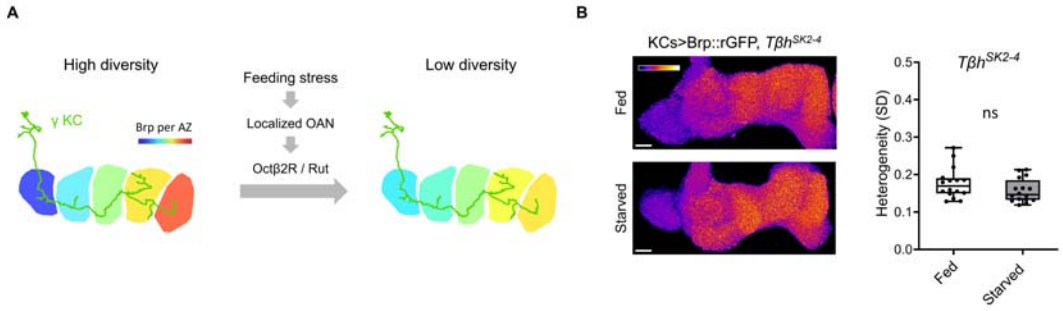

Figure 7. Nutritional states adjust the Brp compartmental heterogeneity through octopamine

(A) Graphical summary of the working model. Localized innervation of OANs onto γ KCs regulates the Brp accumulation and forms the Brp compartmental heterogeneity via Octβ2R and cAMP pathway. Nutritional states may modulate OAN activity, thereby adjusting the Brp heterogeneity.

(B) Starvation for 48 hours did not affect the Brp::rGFP compartmental heterogeneity in *Tβh*$^{SK2-4}$ mutant. GFP_{1-10} was expressed in KCs using R13F02-GAL4 in *Tβh*$^{SK2-4}$ background flies. Fed (n = 16) vs. starved (n = 17): $P > 0.05$.

Data are represented as box plots showing center (median), whiskers (Min. to Max.). ns = not significant by Mann-Whitney U-test.

START METHODS

KEY RESOURCE TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-Brp	DSHB	Cat#nc82; RRID: AB_2314866
AlexaFluor-568 goat anti-mouse	Invitrogen	Cat#A11004; RRID: AB_2534072
Chemicals, peptides, and recombinant proteins		
SNAP-Surface Alexa Fluor 647	New England Biolabs	Cat#S9136S
CLIP-Surface Alexa Fluor 647	New England Biolabs	Cat#S9234S
OMNIPAQUE 350	GE Healthcare	Cat#22000AMX02447
Histodenz	Sigma	Cat#D2158
Tetraspeck Microspheres 0.1 mm	Thermo Fisher Scientific	Cat#T7279
NaCl	Nacalai Tesque	Cat#31320-05
KCl	Nacalai Tesque	Cat#28514-75
TES	Nacalai Tesque	Cat#329810-55
Trehalose dihydrate	Tokyo Kasei Kogyo	Cat#T0331
D-glucose	Nacalai Tesque	Cat#16806-25
$NaHCO_3$	Nacalai Tesque	Cat#31213-15
NaH_2PO_4	Merck KGaA	Cat#106346
$CaCl_2$	Sigma-Aldrich	Cat#C5080
$MgCl_2$	Sigma-Aldrich	Cat#M2670
Experimental models: Organisms/strains		
Brp::SNAP	Bloomington Drosophila stock center (BDSC)	BDSC 58397
MB008B-GAL4	BDSC	BDSC 68291
MB009B-GAL4	BDSC	BDSC 68292
MB370B-GAL4	BDSC	BDSC 68319
VT64246-GAL4	VDRC	VDRC 204311
GH146-GAL4	BDSC	BDSC 30026
TH-GAL4	BDSC	BDSC 8848
R58E02-GAL4	BDSC	BDSC 41347
R13F02-GAL4	BDSC	BDSC 48571
OK371-GAL4	BDSC	BDSC 26160
UAS-Brp::mCherry::GCaMP6s	BDSC	BDSC 77131
Cac::sfGFP	Gratz et al. [6]	N/A
Tdc2-GAL4	BDSC	BDSC 9313
MB022B-GAL4	BDSC	BDSC 68298
UAS-nSyn::CLIP	BDSC	BDSC 58385
UAS-DopR1-RNAi	BDSC	BDSC 55239
UAS-Dop2R-RNAi	BDSC	BDSC 50621

UAS-DopR2-RNAi	BDSC	BDSC 51423
UAS-OAMB-RNAi	VDRC	VDRC 2861
UAS-Octα2R-RNAi	BDSC	BDSC 50678
UAS-Octβ1R-RNAi	BDSC	BDSC 58179
UAS-Octβ2R-RNAi	BDSC	BDSC 34673
UAS-Octβ3R-RNAi	BDSC	BDSC 62283
UAS-TyrR-RNAi	BDSC	BDSC 25857
UAS-TyrR2-RNAi	BDSC	BDSC 64964
UAS-OctTyrR-RNAi	BDSC	BDSC 28332
UAS-Rut-RNAi	BDSC	BDSC 80468
UAS-CD4::tdTomato	BDSC	BDSC 35841
UAS-CD8::GFP	BDSC	BDSC 32194
$T\beta h^{SK1-8}$	This study	N/A
$T\beta h^{SK2-4}$	This study	N/A
brp::GFP$_{11}$	This study	N/A
UAS-GFP$_{1-10}$	Kondo et al. [26]	N/A
Software and algorithms		
Python	N/A	N/A
Graphpad PRISM	Graphpad software	https://www.graphpad.com/
Fiji/Image J	NIH	https://imagej.net/software/fiji/downloads
Amira Software	Thermo Fisher Scientific	thermofisher.com/amira-avizo
DeconvolutionLab2	Sage et al. [81]	https://bigwww.epfl.ch/deconvolution/deconvolutionlab2/
CLIJ2	Haase et al. [82]	https://clij.github.io/
3D ImageJ Suite	Ollion et al. [83]	https://imagej.net/plugins/3d-imagej-suite/

RESOURCE AVAILBILITY

Lead contact

Further information and requests on resources and reagents should be directed to and will be fulfilled by the Lead Contact, Dr. Hiromu Tanimoto (hiromut@m.tohoku.ac.jp).

Material availability

All reagents generated in this study are available upon request from the Lead Contact.

Data and code availability

This study did not generate any unique code. All imaging data and quantification results reported in this paper are available from the Lead Contact upon request.

EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS

Animals

Flies were maintained on standard cornmeal food at 25 °C under a 12:12 hours light-

dark cycle for all experiments. All flies used for experiments are 3-7 days old adult males. After hatching, adult flies were transferred to fresh food vials and flip every 2-3 days before experiments. In experiments where flies were fasted, the same population of flies was separated before the experiment to ensure the homogeneity. All flies were food-deprived for 48 hours for the experimental group, while provide normal food supply for the control for the same time. The GAL4-UAS system was used to express transgenes. For each transgenic control in RNAi experiments, crossing respective transgenic line with Canton S strain, w1118, or y1w1118 to replace the balancer chromosomes. Flies for wild-type characterization were also crossed to replace balancer chromosomes. Strains used in this study are as indicated in the key resource table.

METHOD DETAILS:

Transgenic lines and mutants

Two independent null alleles of the *Tβh* gene were obtained by inducing frameshift indels by the transgenic CRISPR/Cas9 system as previously described [84]. The following 20-bp gRNA sequences were used:

gRNA#1: GATACGTACACCAGTCCGGA

gRNA#2: GGTGAGACGGGACTACCAGC

$T\beta h^{SK1\text{-}8}$ was derived from mutagenesis by gRNA#1 and carries the following frameshift insertion: CGTACACCAGgatggacactGGATGGACAG (the inserted sequence is indicated by lower-case characters). $T\beta h^{SK2\text{-}4\text{-}4}$ was derived from mutagenesis by gRNA#2 and carries the following frameshift deletion: TCCGGATGGA----------CTGTGAGGTC (the gap is indicated as hyphens). The *brp::GFP*$_{11}$ line was generated by targeted insertion of a GFP$_{11}$ cassette into the endogenous *brp* locus just prior to the stop codon as previously described [26].

Sample preparation

Animals used in experiments involving comparisons between control and experimental groups were dissected on the same day. Data were collected from multiple batches of experiments performed on different days. All steps were performed at room temperature otherwise specified. Flies were anesthetized on ice and placed on ice before dissection. The dissection was performed in ice PBS solution to collect the brain without the ventral nerve cord. Brains are subsequently fixed by 2% paraformaldehyde for 1 hour then washed 3×20 min with 0.1% PBT (0.1% Triton X-100 in PBS) in a PCR tube, typically containing 5-6 brains.

For SNAP or CLIP chemical tagging, brains were incubated in SNAP-Surface 647 (1:1000; NEB; S9136S) or CLIP-Surface 647 (1:1000; NEB; S9234S) in 0.3% PBT for 15 min and washed with 3×20 min with 0.1% PBT. Samples are mounted on microscope slides using the appropriate mounting medium (SeeDB2S, SeeDB2G [85] or 86% glycerol according to the imaging condition). Brp::SNAP, Brp::rGFP, Cac::sfGFP, CD4::tdTomato and CD8::GFP samples were imaged without immunohistochemistry.

Immunohistochemistry procedures (for anti-Brp staining in Figure 1B) were carried out as described [86]. In brief, fixed brains were washed 3×20 min with 0.1% PBT and incubated in 3% normal goat serum (NGS; Sigma-Aldrich; G9023) for 1 hour at room temperature. Antibodies used were as indicated in key resource table. Brains were incubated in antibody solutions at 4°C for 48 hours for both primary and secondary antibodies. Sub-resolution fluorescent beads (Tetraspeck Microspheres 0.1 mm, Thermo

Fisher Scientific, T7279) were imaged for generating experimental point spread function (PSF). The beads solution was diluted with distilled water and sonicated multiple times to eliminate aggregation.

For *in vivo* imaging of PM-Mns in developing pupae, samples were prepared as previously described [30]. In brief, white prepupal stage pupae (~ 0 hour APF) were collected and incubated at 25 °C until the experiment. Before imaging, the puparium case was carefully removed with forceps, and the pupa was transferred to an imaging chamber sealed with a cover slip to create an imaging window on the abdomen. For ex vivo imaging of 3rd instar pupal CNS, the CNS was dissected directly from the animal in PBS at room temperature. The CNS sample was then mounted with a spacer between the microscopic glass and cover slip with PBS, The entire process, from dissection to imaging, was performed within 5 minutes.

Image acquisition

For fixed samples and *ex vivo* imaging, images are acquired using the Olympus FV1200 confocal microscope platform equipped with GaAsP high sensitivity detectors and a 60×/1.42 NA oil immersion objective (PLAPON60XO, Olympus) and a 30×/1.05 NA silicone immersion objective (UPLSAPO30XS, Olympus). For *in vivo* imaging on pupae, images were acquired using the Zeiss LSM 800 series confocal microscopy equipped with a 40×/1.3 NA objective (Plan-Apochromat 40×/1.3 Oil DIC (UV) VIS-IR M27).

For Brp::SNAP chemical tagging, 30×/1.05 NA objective is used with a voxel size of 0.53 µm × 0.53 µm × 0.84 µm (lateral × lateral × axial) to image the whole brain except for the optical lobes. For Brp::rGFP and UAS-CD4::tdTomato samples in all GAL4 types except Tdc2-GAL4, 60×/1.42 NA oil immersion objective was used and 473 nm laser power: 0.1%; 559 nm laser power: 0.1%; scanning speed: 2 µs/pixel with a voxel size of lateral 0.079 × 0.079 × axial 0.370 µm. For Brp::SNAP/Cac::sfGFP images, 60×/1.42 NA objective is used with a voxel size of 0.079 × 0.079 × 0.370 µm. For Tdc2-GAL4 driven UAS4-CD8::GFP and UAS-nSyb::CLIP samples, 30×/1.05 NA objective is used with a voxel size of 0.414 × 0.414 × 0.84 µm. For MB022B-GAL driven UAS-CD8::GFP and UAS-nSyb-CLIP samples, 30×/1.05 NA objective is used with a voxel size of 0.276 × 0.276 × 0.87 µm. For Tdc2-GAL4 driven Brp::rGFP and UAS-CD4::tdTomato, 60×/1.42 NA objective is used with a scanning voxel size of 0.132 × 0.132 × 0.34 µm. For Brp::rGFP *in vivo* imaging of PM-Mns, a 40×/1.3 NA objective is used with a voxel size of 0.0725 × 0.0725 × 0.460 µm. For Brp::rGFP ex vivo imaging, 60×/1.42 NA objective is used with a voxel size of 0.079 × 0.079 × 0.370 µm and 473 nm laser power: 2.0%. For experimental PSF imaging, SeeDB2S immerged beads were scanned in a setting that is 60×/1.42 NA oil immersion objective; 473 nm laser power: 2.0%; 559 nm laser power: 1.5%; voxel size: 0.079 µm × 0.079 µm × 0.370 µm; scanning speed: 4 µs/pixel, to produce multiple bead images.

***In vivo* calcium imaging**

In vivo calcium imaging was performed following a previously described protocol [86]. Briefly, flies were anesthetized on ice for 3 minutes and placed in a custom-made holding dish on a Peltier plate (CP-085, Sinics) maintained at 5°C. The head capsule was fixed to the dish using UV curing optical adhesive (NOA68, Thorlabs). To minimize brain movement, the proboscis was glued to the capsule A small window was

opened on the top of the head capsule, and the exposed area was filled with *Drosophila* saline solution (final concentration: 103 mM NaCl, 3 mM KCl, 5 mM TES, 8 mM Trehalose dihydrate, 10 mM D-glucose, 26 mM $NaHCO_3$, 1 mM NaH_2PO_4, 1.5 mM $CaCl_2$, 4 mM $MgCl_2$, adjust to pH ~7.2). Air sacs and fat bodies covering the brain surface were carefully removed. Live imaging was conducted using a laser scanning confocal microscope equipped with GaAsP detectors (A1R, Nikon) and a 30×/1.1 NA water immersion objective (Apo LWD 25×, Nikon). GCaMP6s and mCherry were sequentially excited at 488 and 561 nm, respectively. Emission light was collected using dichroic mirrors and emission filters (BP500–550 and BP570-620) onto GaAsP detectors. The MB γ lobe was scanned at a pixel size of 0.5μm/pixel (512 × 128 pixels) at a frame rate of 1 Hz using line scans with 16× averages in the resonant scanning mode. The pinhole was set to 2.5 AU (561 nm). For recording, images were sequentially acquired over 40 s and saved for later image processing.

QUANTIFICATION AND STAISTICAL ANALYSIS

Data processing and analysis

PSF images were processed in the Amira software (Thermo Fisher Scientific). Using the Extract Point Spread Function module, PSFs extracted from each image were averaged into a single PSF, which was later used for image deconvolution (resized voxel size: 0.079 µm × 0.079 µm × 0.370 µm; image size: 32 pixels × 32 pixels × 21 slices). Image deconvolution was performed using the Richardson-Lucy iterative non-blind algorithm in the Fiji plugin DeconvolutionLab2 [81], or with CLIJ2 GPU-based Richardson-Lucy deconvolution [82].

AZ detection was performed as described in a compartmental manner. In brief, a sub-image stacks were cropped from every MB compartment in the original image. Compartments were manually identified referencing the CD4::tdTomato membrane signal. An intensity threshold was applied to reduce background noise. Sub-stacks were processed by the 3D maxima finder and the 3D spot segmentation function in the 3D suite plugin [83] in Fiji. 3D maxima were detected for each cluster and used as start points for pixel clustering using the 3D spot segmentation function. This process created 3D ROIs enclosing individual Brp clusters. ROIs were then used to extract signal intensities and cluster coordinates. Detection precision was optimized by comparing detection results with manually defined ground truths as previously described.

For calcium imaging, ROIs were manually drawn for each compartment in time-series image stacks. GCaMP6s signal was normalized by mCherry in each time-frame using the image calculator in Fiji. The mean intensity was calculated in each compartment by averaging 7 frames without any odor stimulation as the basal level.

Statistical analysis

Statistical analyses were performed using GraphPad Prism version 8, 9 and 10. Summarized data are represented as box plots showing center (median), whiskers (Min. to Max.). For comparison between more than two groups, Kruskal-Wallis test with original False Discovery Rate (FDR) method of Benjamini and Hochberg correction was applied. Desired false discovery rate was set to 0.05. For comparison between two groups, unpaired nonparametric two-sided Mann-Whitney U-test was used. Pearson's correlation coefficient (R) was calculated in Python.

SUPPLEMENTAL FIGURE TITLES AND LEGENDS

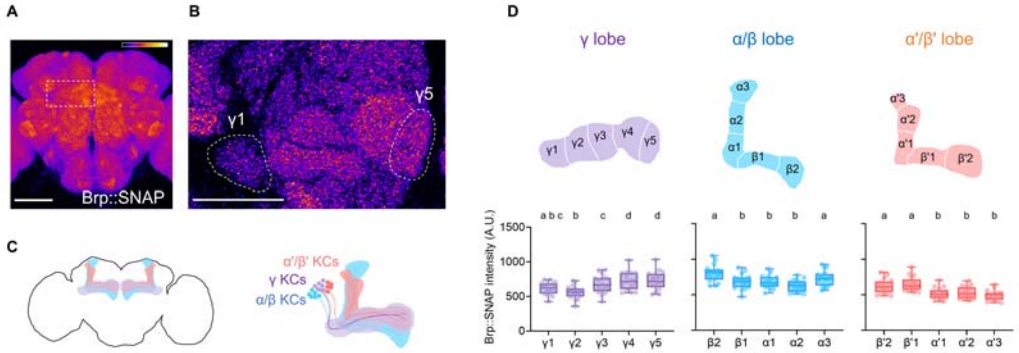

Figure S1. Heterogeneous Brp enrichment in the MB

(A) SNAP chemical tagging labeled endogenous Brp in the brain. Scale bar, 100 μm. The dash line area in indicates the zoomed-in area showed in (B**)**.

(B) Intensity difference of Brp::SNAP between γ1 and γ5 compartment on the same imaging plate. Scale bar, 50 μm.

(C) Schematic of the MB. The MB comprise three lobes based on the projection patterns of three KC subtypes: γ KCs, α'/β' KCs and α/β KCs.

(D) Brp::SNAP intensity difference across compartments. Schematic drawings above indicate compartments of each lobe.

Data are represented as box plots showing center (median), whiskers (Min. to Max.). Significant differences ($P < 0.05$) are indicated by distinct letters. Kruskal-Wallis test.

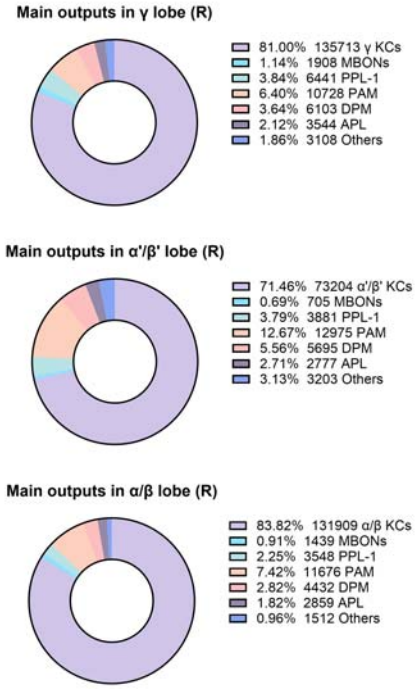

Figure S2. Pre-synapses in the MB are predominantly from KCs.

Pie charts showing the composition of pre-synapses in each of the three MB lobes. Data

is from the hemibrain online data but not all the cell types are listed. The number of pre-synapse and the percentage are indicated for each cell type, including KCs, MBONs PPL-1 neurons, PAM neurons, dorsal paired medial (DPM) neuron and anterior paired lateral (APL) neuron.

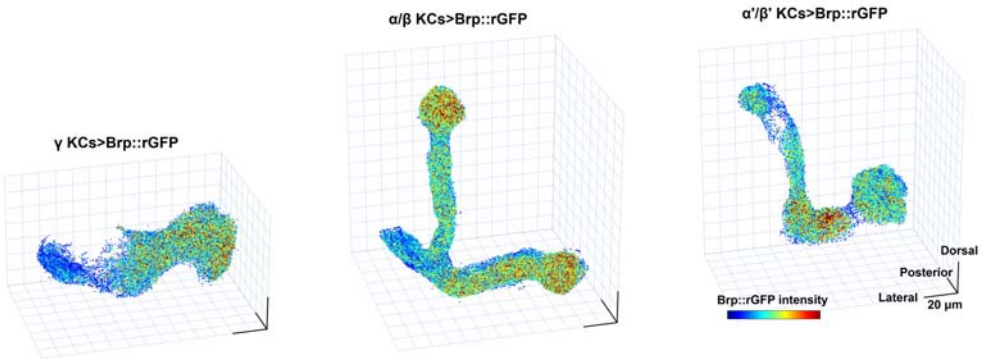

Figure S3. Brp compartmental heterogeneity in different KC subtypes
The 3D reconstruction of Brp::rGFP clusters, colored by Brp::rGFP intensity. Brp::rGFP is visualized in γ KCs using MB009B-GAL4, in α/β KCs using MB008B-GAL4 and in α'/β' KCs using MB0370B-GAL4. Scale bars, 20 μm.

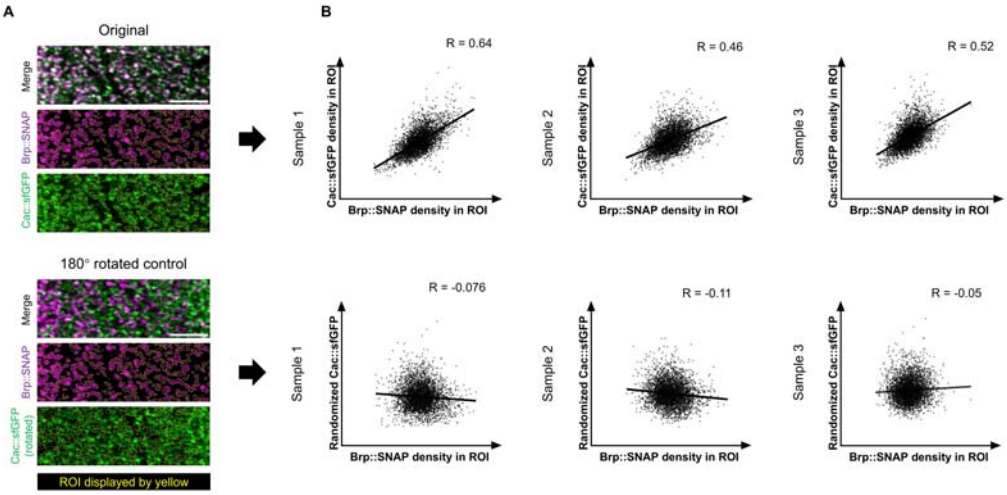

Figure S4. Correlation between Brp::SNAP and Cac::sfGFP signals intensities
(A) Correlation analysis of Brp::SNAP and Cac::sfGFP signal intensities. ROIs are generated using 3D spot segmentation method without a watershed process. Relatively loose setting was applied on Brp::SNAP images to generate wide ROIs. The same ROI set was used to calculated signal intensities in both Brp::SNAP and Cac::sfGFP channels. The 180˚ rotated version of Cac::sfGFP image was used as the control. A total of 4000 ROIs were analyzed, and Pearson correlation coefficient (R) values were calculated for each sample. Scale bars, 5 μm.
(B) Scatter plots showing the correlation between Brp::SNAP and Cac::sfGFP signal intensities (left) and control (right) in different brain samples. R value is indicated for each sample.

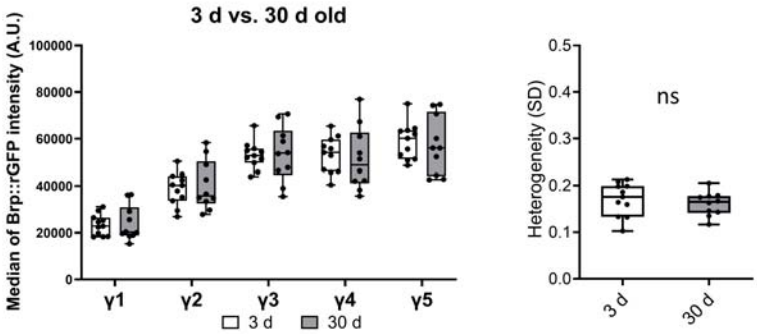

Figure S5. Brp compartmental heterogeneity in aging flies

Median intensity of Brp::rGFP clusters and the Brp::rGFP compartmental heterogeneity were compared between 3 days old and 30 days old flies. Mann-Whitney U-test, 3 d (n = 11) vs. 30 d (n = 10): $p > 0.05$.

Data are represented as box plots showing center (median), whiskers (Min. to Max.). ns = not significant by Mann-Whitney U-test.

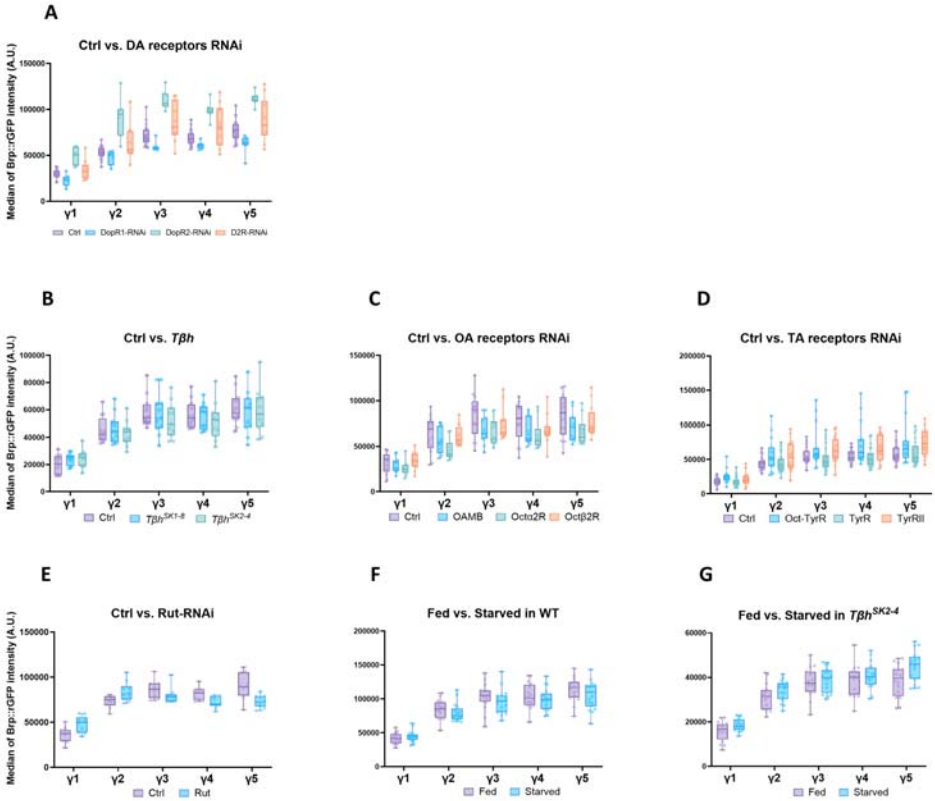

Figure S6. Median intensities of Brp::rGFP clusters in each experiment

(A) Dopamine receptor knockdown experiment.

(B) *Tβh* mutant's experiment.

(C) Octopamine receptor knockdown experiment.

(D) Tyramine receptor knockdown experiment.

(E) Rut knockdown experiment.

(F) Starvation experiment.
(G) Starvation experiment of $T\beta h^{SK2\text{-}4}$ mutants.
Data are represented as box plots showing center (median), whiskers (Min. to Max.).

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