Hsp83/Hsp90 Physically Associates with Insulin Receptor to Promote Neural Stem Cell Reactivation

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SUMMARY

Neural stem cells (NSCs) have the ability to exit quiescence and reactivate in response to physiological stimuli. In the Drosophila brain, insulin receptor (InR)/phosphatidylinositol 3-kinase (PI3K)/Akt pathway triggers NSC reactivation. However, intrinsic mechanisms that control the InR/PI3K/Akt pathway during reactivation remain unknown. Here, we have identified heat shock protein 83 (Hsp83/Hsp90), a molecular chaperone, as an intrinsic regulator of NSC reactivation. Hsp83 is both necessary and sufficient for NSC reactivation by promoting the activation of InR pathway in larval brains in the presence of dietary amino acids. Both Hsp83 and its co-chaperone Cdc37 physically associate with InR. Finally, reactivation defects observed in brains depleted of Hsp83 were rescued by over-activation of the InR/PI3K/Akt pathway, suggesting that Hsp83 functions upstream of the InR/PI3K/Akt pathway during NSC reactivation. Given the conservation of Hsp83 and the InR pathway, our finding may provide insights into the molecular mechanisms underlying mammalian NSC reactivation.

INTRODUCTION

The balance between proliferation and quiescence of stem cells is crucial in maintaining tissue homeostasis. Neural stem cells (NSCs) in the brain have the ability to be reactivated from a reversible quiescent state to generate new neurons. In the mammalian adult brain, the majority of adult NSCs are quiescent and are not actively dividing (Doetsch et al., 1999; Morshead et al., 1994). Interestingly, these quiescent NSCs can be reactivated and participate in neurogenesis upon various extrinsic stimuli (Ahn and Joyner, 2005; Daynac et al., 2013, 2016; Faiz et al., 2015; Kawai et al., 2017; Lugert et al., 2010; Wang et al., 2011). Dysregulation of the balance between proliferation and quiescence of NSCs may contribute to neurodevelopmental disorders such as microcephaly (Baser et al., 2009; Cloetta et al., 2013). Reactivating quiescent NSCs may also provide therapeutic strategies for the treatment of brain injuries or neurodegenerative disorders.

Recently, Drosophila NSCs, neuroblasts, have emerged as an excellent in vivo model for the study of NSC quiescence and reactivation. NSCs exit the cell cycle, shrink in size, and enter quiescence at the end of embryogenesis under the control of the spatial Hox protein, temporal identity factors, and a homeodomain differentiation factor Prospero (Isshiki et al., 2001; Lai and Doe, 2014; Tsuji et al., 2008). At early larval stages, following 24 hr of quiescence, they are reactivated in response to feeding (Britton and Edgar, 1998; Ito and Hotta, 1992; Truman and Bate, 1988). Dietary amino acids present in the food are sensed by the fat body, which functions equivalent to the mammalian liver and adipose tissue (Colombani et al., 2003). The fat body generates mitogens, stimulating blood-brain barrier glial cells to secrete insulin-like peptides (dILPs). dILPs act locally by activating the insulin receptor (InR)/phosphatidylinositol 3-kinase (PI3K)/Akt pathway in underlying NSCs and promoting their reactivation (Britton and Edgar, 1998; Chell and Brand, 2010; Sousa-Nunes et al., 2011). Blood-brain barrier glial cells synchronize NSC reactivation via calcium oscillations through gap junctions (Speder and Brand, 2014), while cortex glial cells remodel to promote new-born neuron survival (Speder and Brand, 2018). Besides InR pathway, the spindle matrix complex intrinsically promotes NSC reactivation (Li et al., 2017), while the Hippo pathway maintains NSC quiescence (Ding et al., 2016; Poon et al., 2016). In mammalian brains, insulin-like growth factor-1 (IGF-1) produced by astroglial cells has a similar role in promoting NSC proliferation in response to brain injury (Mairet-Coello et al., 2009; Van et al., 2006; Ye et al., 2004). In addition, IGF-1 receptor (IGF-1R) promotes the proliferation of mammalian NSCs (Arsenijevic et al., 2001), and human IGF-1R mutations are associated with microcephaly (Juanes et al., 2015), suggesting that the insulin pathway is likely conserved from flies to humans in promoting NSC proliferation. However, how the insulin pathway is regulated during NSC reactivation is still poorly understood.

Here, we show that a highly conserved chaperone, Hsp83/Hsp90, is required for the activation of the InR/PI3K/Akt pathway during NSC reactivation. Hsp83 functions in a wide variety of biological processes, including
A) Dpn, Mira, Mira/Dpn
   hsp83 RNAi

B) EdU, EdU/Mira
   hsp83 RNAi

C) Dpn, Mira, Dpn/Mira/CD8
   hsp83-m4A

D) Dpn, EdU, Dpn/EdU/CD8
   hsp83-m4A

E) NSC% with cellular extensions
   Control, hsp83 RNAi, MARCM Control, hsp83-m4A
   NSC% without EdU incorporation
   Control, hsp83 RNAi, MARCM Control, hsp83-m4A

F) Mira/Dpn
   v47776
   v110727 BDSC#2875

H) Control
   EdU/Mira
   cdc37 RNAi
   EdU/Mira

I) Control
   O/E hsp83-HA
   EdU/Dpn

J) 6h ALH
   NSC% with EdU incorporation
   Control, O/E hsp83-HA

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cell polarity, DNA transcription, and chromatin remodeling (Andersen et al., 2012; Pratt and Toft, 1997; Sawarkar et al., 2012; Tariq et al., 2009). Unlike Hsp70, eukaryotic Hsp83/Hsp90 does not act in nascent protein folding, rather, it binds to substrate proteins that are in a near native state at a late stage of folding (Young et al., 2001). We show that Hsp83 is necessary and sufficient for NSC reactivation in the presence of dietary amino acids. Its co-chaperones Cdc37 and Hsp70 mediate the association between Hsp83 and its client protein kinases and steroid hormone receptors, respectively (Eckl and Richter, 2013; Young et al., 2001). We show that the activation of the InR/PI3K/Akt pathway is dependent on Hsp83 and Cdc37 function in larval brains. Hsp83 and Cdc37 physically associate with InR, but not Akt. Finally, the hsp83 knockdown phenotype is rescued by over-activation of the InR/PI3K/Akt pathway.

RESULTS

Hsp83 Is Required Intrinsically for NSC Reactivation

To identify regulators of NSC reactivation, we used an NSC-specific driver insc-GAL4 to perform an unbiased RNAi screening in NSCs. Our screen identified an RNAi line (VDRC no. 108568), which knocks down hsp83 expression, exhibiting defective NSC reactivation, with cellular extensions, a hallmark of quiescent NSCs, observed in 27.7% ± 3.0% (Figures 1A and 1E; n = 1,452, 41 brain lobes) of NSCs at 72 hr after larval hatching (ALH) at 29°C. In comparison, all control NSCs were reactivated and lost their cellular extensions at the same stage (Figures 1A and 1E; n = 1,084, 20 brain lobes). This finding suggests that these NSCs failed to be reactivated and remained in a quiescent state upon hsp83 knockdown. Next, in order to examine if NSCs with hsp83 knockdown are able to enter the cell cycle, we carried out 5-ethyl-2′-deoxyuridine (EdU) incorporation experiments. All control NSCs (Figures 1B and 1F; n = 756, 20 brain lobes) incorporated EdU at 72 hr ALH at 29°C. By contrast, at the same condition, 44.7% ± 6.3% (n = 824, 24 brain lobes) of NSCs failed to incorporate EdU with hsp83 knockdown (Figures 1B and 1F). Next, we examined the phenotype of an amorphic hsp83<sup>j5C2</sup> and hypomorphic hsp83<sup>e6A</sup> in mosaic analysis with a repressible cell marker (MARCM) clones (Lee and Luo, 2001). While all NSCs in wild-type control clones exhibited normal round shapes without any cellular extensions (n = 58), 43.0% (n = 114) of hsp83<sup>j5C2</sup> NSC clones and 36.4% (n = 266) of hsp83<sup>e6A</sup> NSC clones contained NSCs with cellular extensions (Figures 1C and 1E). These observations further show that NSCs remain quiescent after loss of hsp83. To examine the proliferative status of the NSCs, we performed EdU pulse-chase analysis. At 96 hr ALH, all NSCs in wild-type control clones were actively dividing evident by EdU incorporation (n = 101). Unlike the wild-type control clones, 82.8% (n = 82) and 77.9% (n = 87) of NSC clones with hsp83<sup>j5C2</sup> and hsp83<sup>e6A</sup>, respectively, failed to incorporate EdU (Figures 1D and 1F). Furthermore, the NSC quiescence phenotype of hsp83 mutants were fully rescued by introducing a 21.8-kb bacterial artificial chromosome (BAC) CH322-129N17 containing hsp83 (hsp83-BAC). A total of 39.7% (n = 126) and 36.4% (n = 107) NSCs in hsp83<sup>j5C2</sup> and hsp83<sup>e6A</sup> MARCM clones showed cellular extensions (Figure S1A). However, when hsp83-BAC was expressed, none of NSCs in hsp83<sup>j5C2</sup> (n = 117) and hsp83<sup>e6A</sup> (n = 107) MARCM clones exhibited...
cellular extensions (Figure S1A). Similarly, 78.3% (n = 46) and 70.8% (n = 72) NSCs in hsp83^{S2C} and hsp83^{M2A} MARCM clones failed to incorporate with EdU (Figure S1A), while all NSCs from hsp83^{S2C} with hsp83-BAC (n = 95) and hsp83^{M2A} with hsp83-BAC (n = 70) were incorporated with EdU (Figure S1A). Taken together, these results indicate that Hsp83 is required for NSC reactivation.

We carried out a time course experiment and examined larval brains at 24, 48, and 72 hr ALH. At 24 hr ALH, most control NSCs were reactivated, with only 4.9% ± 1.0% (n = 2,174, 37 brain lobes) showing cellular extensions, suggesting that they were still in quiescence (Figures S1C and S1E). With hsp83 knockdown under the insc-Gal4 driver, 15.4% ± 1.3% (n = 1,535, 28 brain lobes) of NSCs possess cellular extensions at 24 hr ALH (Figures S1C and S1E). Loss of cellular extensions of NSCs seemed to precede cell-cycle re-entry. At 24 hr ALH, 19.5% ± 3.8% (n = 1,205, 22 brain lobes) of control NSCs failed to incorporate EdU (Figures S1B and S1D), while 63.1% ± 5.0% (n = 1,193, 24 brain lobes) NSCs with hsp83 knockdown were EdU-negative (Figures S1B and S1D). This observation suggests that NSC reactivation is defective when hsp83 is knocked down at early larval stages. At 48 hr ALH, all control NSCs showed round cell morphology without cellular extension (Figures S1C and S1E; n = 1,625, 30 brain lobes) and nearly all of the NSCs incorporated EdU (Figures S1B and S1D; 98.2% ± 1.4%, n = 704, 21 brain lobes). In comparison, with hsp83 knockdown, 27.2% ± 2.9% (n = 1,254, 20 brain lobes) of NSCs still showed cellular extensions (Figures S1C and S1E), while 48.7% ± 6.2% (n = 526, 13 brain lobes) of NSCs failed to incorporate EdU (Figures S1B and S1D). At 72 hr ALH, all control NSCs had no cellular extensions (Figures S1C and S1E; n = 1,221, 13 brain lobes) and incorporated EdU (Figures S1B and S1D; n = 322, 6 brain lobes). By contrast, with hsp83 knockdown 28.2% ± 2.1% (n = 1,003, 11 brain lobes) of NSCs showed cellular extensions (Figures S1C and S1E) and 49.5% ± 7.4% (n = 290, 5 brain lobes) of NSCs lacked EdU incorporation (Figures S1B and S1D). Taken together, Hsp83 is intrinsically required for NSC reactivation.

Upon hsp83 knockdown at 72 hr ALH, the sizes of NSCs are variable, ranging from 4 to 11 μm, even when they still maintained cellular extensions, a hallmark of quiescent NSCs. At 24 hr ALH, the diameter of control NSCs was 8.7 ± 0.6 μm (n = 121), while NSC diameters were decreased to 6.8 ± 1.3 μm (n = 129) upon hsp83 knockdown under insc-Gal4 (data not shown). At 72 hr ALH, the average NSC diameter upon hsp83 RNAi knockdown is 8.1 ± 1.8 μm (n = 124), still smaller than control NSCs (10.6 ± 0.7 μm; data not shown). Similarly, at 72 hr ALH the NSC diameters of hsp83^{M2A} and hsp83^{S2C} were 8.1 ± 1.7 μm (n = 121) and 8.2 ± 1.6 μm (n = 118), respectively. These observations suggest that, although NSC reactivation is dramatically disrupted upon knocking down hsp83, cell growth is incompletely blocked, resulting in enlargement of some of the quiescent NSCs. Thus, cell division and cell growth seemed to be partially uncoupled upon failure of NSC reactivation in NSCs upon hsp83 depletion. Similar uncoupling of cell division and growth were also reported in other mutants defective in NSC reactivation including chromator, which appears to function downstream of the Inr signaling pathway (Li et al., 2017).

Mushroom body (MB) neuroblasts do not enter quiescence at the end stage of embryogenesis and are capable of dividing in the absence of dietary amino acids throughout larval stages (Ito and Hotta, 1992; Sousa-Nunes et al., 2011; Truman and Bate, 1988). We investigated whether Hsp83 was required for general proliferation of cells, such as MB neuroblasts. At 24 hr ALH, all MB neuroblasts (Dpn^+), which were surrounded by Dachshund-positive MB neurons, from both control (n = 224) and hsp83 RNAi (n = 200), were incorporated with EdU (Figure S1F). This result suggests that proliferation of MB neuroblasts is unaffected by hsp83 RNAi knockdown.

Co-chaperone Cdc37 Is Required for NSC Reactivation

Functioning as a co-chaperone, Hsp83 has two major classes of clients, steroid hormone receptors and protein kinases, and requires co-chaperones heat shock protein 70 (Hsp70) or cell division cycle 37 (Cdc37), respectively (Eckl and Richter, 2013). To investigate whether steroid hormone receptors or protein kinases are the potential clients of Hsp83 during NSC reactivation, we tested whether Hsp70 or Cdc37 plays any role during NSC reactivation. There are seven isoforms of hsp70 in Drosophila and we acquired 14 RNAi stocks for these isoforms (Table S1) and knocked down these genes in NSCs using insc-Gal4. However, knocking down these isoforms of hsp70 or Hsp70/Hsp90 organizing protein (hop), which coordinates the interaction between Hsp70 and Hsp83 by binding to both of them, did not result in any NSC reactivation defect at both 24 and 72 hr ALH (Table S1 and data not shown). By contrast, at 72 hr ALH at 29°C, in cdc37 RNAi knockdown (VDRC no. 47776) brains, 16.8% ± 2.2% (n = 947, 11 brain lobes) of NSCs showed cellular processes (Figure 1G), and only 1.5% ± 1.4% (n = 1,135, 14 brain lobes) of NSCs were incorporated with EdU (Figure 1H). In wild-type NSC MARCM clones, all NSCs were devoid of...
cellular extensions at 96 hr ALH (Figures S1G and S1H; n = 50). By contrast, in cdc37e4D MARCM clones, 28.8% (n = 198) of NSCs extend cellular processes (Figures S1G and S1H), suggesting that Cdc37 is required for NSC activation. The process phenotype of cdc37e4D was fully rescued by introducing a 22.1-kb BAC CH322-35F18 containing cdc37 (cdc37-BAC). A total of 25.9% (n = 85) NSCs in cdc37e4D MARCM clones showed cellular extensions, while none of the NSCs (n = 71) from cdc37-BAC showed cellular extensions (Figures S1G and S1H). Therefore, we conclude that Cdc37 is required for NSC reactivation.

Overexpression of Hsp83 Leads to Premature NSC Reactivation
To test whether Hsp83 overexpression is sufficient to trigger NSC reactivation, we overexpressed hsp83-hemagglutinin (HA) (Kuo et al., 2013) in NSCs using insc-Gal4 and examined its effect on NSC proliferation at 6 hr ALH. Hsp83-HA is functional, as it largely rescued the NSC quiescence phenotype in hsp83j5C2 and hsp83e6A mutants (data not shown). At 6 hr ALH, most wild-type control NSCs remained in quiescence, as only 18.8% ± 4.8% of NSCs (n = 1,907, 55 brain lobes) incorporated EdU (Figures 1I and 1J). However, upon hsp83 overexpression, 41.9% ± 5.5% (n = 2,211, 62 brain lobes) of NSCs incorporated EdU (Figures 1I and 1J). This result suggests that Hsp83 overexpression leads to premature NSC reactivation. However, in the absence of dietary amino acids, Hsp83 overexpressing in NSCs was unable to promote NSC reactivation (data not shown), suggesting that Hsp83 overexpression effect is nutrition dependent. Taken together, Hsp83 is both necessary and sufficient for NSC reactivation in the presence of dietary amino acids.

plnR and Akt Levels Are Decreased upon hsp83 Depletion
Activation of the InR/PI3K/Akt pathway promotes NSC reactivation (Chell and Brand, 2010; Sousa-Nunes et al., 2011). We investigated whether the InR/PI3K/Akt pathway is activated in NSCs with hsp83 depleted. To this end, we first analyzed the protein levels of Akt in NSCs, a major component of the InR/PI3K/Akt pathway and a protein kinase that triggers NSC reactivation (Shim et al., 2013). In the central brain of wild-type larvae at 72 hr ALH, Akt protein levels were comparable with its levels in the optic lobe (Figure 2A). However, upon hsp83 knockdown in central brain NSCs by insc-Gal4 at 72 hr ALH, Akt protein levels in the central brain decreased significantly, while its levels in the optic lobe remained similar to control (Figure 2A). Likewise, Akt protein levels in the central brain were reduced significantly upon cdc37 knockdown by RNAi (Figure 2A). These observations suggest that Hsp83 and Cdc37 may be required for the activation of the InR/PI3K/Akt pathway in NSCs. Next, we examined Akt protein levels in whole-brain protein extracts by western blot. hsp83 knockdown in larval brains was efficient, as Hsp83 protein levels in the whole brain were reduced to 47.3% ± 6.8% (n = 3) in hsp83 knockdown samples compared with controls (Figures 2B and 2C). Remarkably, Akt protein levels was only 61.7% ± 5.5% (n = 3) in hsp83 knockdowns compared with wild-type controls (Figures 2B and 2C). Likewise, upon cdc37 knockdown in NSCs Akt level was reduced to 67.1% ± 15.8% (n = 3), compared with wild-type control (Figures 2B and 2C). These results suggest that the InR/PI3K/Akt pathway is likely impaired upon depletion of either hsp83 or cdc37 in NSCs.

We then examined whether the protein levels of phosphorylated insulin receptor (plnR), the activated form of InR, is altered upon hsp83 or cdc37 depletion in larval brains. After hsp83 knockdown in NSCs under insc-Gal4 at 72 hr ALH, only 16.5% ± 9.7% (n = 3) of plnR remained compared with wild-type controls (Figures 2D and 2E). Similarly, knockdown of cdc37 in NSCs resulted in a dramatic reduction of plnR levels with 23.1% ± 0.6% (n = 3) compared with controls (Figures 2D and 2E). The mRNA levels of InR or akt, were not obviously altered with hsp83 or cdc37 RNAi knockdown (Figures S2A and S2B).

Taken together, we conclude that Hsp83 and Cdc37 are likely required for the activation of the InR/PI3K/Akt pathway.

Hsp83 Overexpression Results in Increase of plnR and Akt Level
Since Hsp83 overexpression triggers premature NSC reactivation, it is worthy to investigate whether Hsp83 overexpression can result in the activation of the InR/PI3K/Akt pathway. Due to insufficient amounts of protein that could be obtained due to the tiny size of larval brains at 6 hr ALH, we examined the protein levels of the whole larvae. Upon hsp83-HA overexpression in NSCs using insc-Gal4, the protein levels of Akt in the whole larvae were increased to 184.6% ± 25.9% (n = 4), compared with controls (Figures 2F and 2G). Similarly, hsp83 overexpression in NSCs resulted in a significant increase of plnR level in whole larvae, with 383.6% ± 88.2% (n = 4) compared with controls (Figures 2H and 2I). These results suggest that overexpression of Hsp83 is sufficient to drive the activation of the InR/PI3K/Akt pathway on the fed condition.

Hsp83, Cdc37, and InR Physically Interact
Given that Hsp83 and Cdc37 are required for the activation of the InR/PI3K/Akt pathway, we explored whether Hsp83 and Cdc37 physically interact with InR or Akt. To test the physical association between Hsp83, Cdc37, and InR, we performed a proximity ligation assay (PLA), a technique
enabling high specificity and sensitivity detection of protein-protein interactions (Figure 3A) (Fredriksson et al., 2002). We co-expressed various proteins tagged with Myc or HA in S2 cells and quantified PLA foci that indicated interactions in cells, which co-expressed both Myc and HA-tagged proteins (Figures 3B and 3C). In S2 cells co-expressing both HA and Myc constructs, the vast majority of cells (90.2%, n = 1,397) had no PLA signals and the rest of the cells displayed weak PLA fluorescence signal of less than 10 foci (Figures 3B and 3C). On average, these control cells only had 0.13 PLA foci per cell. We failed to generate full-length InR constructs, presumably due to its large size (>6 kb). As the intracellular domain of InR including the kinase domain (InR intra), an active form of InR, was previously used for demonstrating protein-protein interactions for InR (Almudi et al., 2013), we decided to use InR intra in PLA assays. Similarly, in the vast majority of cells co-expressing Myc-Hsp83 with control HA (n = 1,456) and HA-InR intra with control Myc (n = 1,767), no PLA signal was detected and, on average, there were 0.45 and 0.38 PLA foci per cell, respectively (Figures 3B and 3C). By contrast, cells co-expressing Myc-Hsp83 and HA-InR intra displayed 12.77 PLA foci per cell on average; 90.9% of cells exhibited PLA signal, with 41.8% of cells displaying moderate PLA signals (11–30 foci), 36.4% displaying weak PLA signals (<10 foci), and 12.7% displaying strong PLA signals (>30 foci; Figures 3B and 3C; n = 1,784). In a similar PLA assay, we tested whether Hsp83 interacts with Akt. The majority of S2 cells co-expressing Myc-Akt and HA-Hsp83 had no PLA signal (Figures S3A and S3B; 76.1%, n = 1,581), suggesting that Hsp83 did not interact with Akt. These results suggest that Hsp83 physically associated with InR intra, but not Akt.

Similarly, in controls expressing Myc-Cdc37 and HA, we observed that the majority of cells (88.2%) had no PLA signal (Figures 3B and 3C; n = 1,522). However, co-expressing Myc-Cdc37 and HA-InR intra resulted in 86.8% S2 cells showing PLA fluorescence; 51.3% of cells displayed weak PLA signals, 31.4% of cells showed moderate PLA signals, and 4.1% of cells showed strong PLA signals (Figures 3B and 3C; n = 1,144). In controls co-expressing HA-Hsp83 and Myc, 88.5% of S2 cells did not have any PLA signal (Figures 3B and 3C; n = 1,069). By contrast, in cells co-expressing Myc-Cdc37 and HA-Hsp83, 51.5%, 30.4%, and 4.2% of these cells displayed weak, moderate, and strong PLA signals, respectively (Figures 3B and 3C; n = 1,271).

To further validate the interactions between Hsp83, Cdc37, and InR, we employed a protein-protein interaction assay named biomolecular fluorescence complementation (BiFC), which is able to detect transient or weak interactions due to the irreversibility of the BiFC complex formation (Gohl et al., 2010; Shyu and Hu, 2008). We generated chimeric proteins InR intra-HA-CYFP (InR intra with C-terminal YFP tagged with HA) and Hsp83-Myc-NYFP (N-terminal YFP tagged with Myc was fused to the Hsp83). As expected, when transfecting either of these two chimeric constructs into S2 cells with their respective controls, Myc-NYFP and HA-CYFP, no YFP signal was detected in cells (Figure S4B). By contrast, strong YFP was detected when cells were expressing both Hsp83-myc-NYFP and InR intra-HA-CYFP, suggesting that Hsp83 and InR intra physically associate (Figure S4B). To further test the specificity of this interaction, we generated a C-terminal truncation of Hsp83 (Figure S4A; Hsp83C: amino acids 538–717), which presumably abolishes the dimerization of Hsp83 and its subsequent ability to form a pocket-like structure to bind to its client proteins. We tested the ability of this Hsp83 truncation to associate with InR intra. There was no YFP signal upon co-expression of Hsp83C-myc-NYFP with InR intra-HA-CYFP in S2 cells (Figure S4B), suggesting that the association between Hsp83 and InR intra was abolished with the N-terminal Hsp83 deletion. In addition, Hsp83 associated with its co-chaperone Cdc37, but not with an unrelated control WAVE in the BiFC assay (Figure S4B). Cdc37 also physically interacts with InR intra in the BiFC assay, as S2 cells expressing both InR intra-HA-CYFP and Cdc37-myc-NYFP, but not the controls, displayed YFP signal (Figure S4B).
Next, to examine their physical association in an in vivo BiFC assay, we generated transgenic flies expressing Hsp83-myc-NYFP, InR<sub>intra</sub>-HA-CYFP, Cdc37-myc-NYFP, and Cdc37-HA-CYFP in NSCs under the <i>insc</i>-Gal4 driver. At 72 hr ALH at 29°C, overexpressed Hsp83, Cdc37, and InR<sub>intra</sub> in NSCs were observed by their respective epitope tags. As expected, there was no detectable YFP when Hsp83-myc-NYFP, Cdc37-myc-NYFP, Cdc37-HA-CYFP Cdc37, or InR<sub>intra</sub>-HA-CYFP were co-expressed with their corresponding control half-YFP proteins (Figure 4). By contrast, when Hsp83-myc-NYFP and InR<sub>intra</sub>-HA-CYFP were co-expressed, strong YFP was detected in NSCs (Figure 4), suggesting that Hsp83 and InR<sub>intra</sub> physically associate in NSCs. Likewise, Cdc37 interacted with both InR<sub>intra</sub> and Hsp83 in NSCs in the BiFC assay (Figure 4). Taken together, Hsp83, Cdc37, and InR<sub>intra</sub> physically associate with one another both in vitro and in vivo in BiFC assays.

Hsp83 Functions Upstream of the InR/PI3K/Akt Pathway in NSC Reactivation
Next, we ascertained whether Hsp83 functions upstream or downstream of the InR/PI3K/Akt pathway during NSC reactivation. Over-activation of the InR/PI3K/Akt pathway by expressing various active components of this pathway over-rides the requirement of dietary amino acids for NSC reactivation (Chell and Brand, 2010; Sousa-Nunes et al., 2011). We explored whether depletion of <i>hsp83</i> was able to abrogate this effect of InR/PI3K/Akt pathway over-activation. Overexpression of an active form of InR (i.e., InR<sup>AD</sup>, BDSC no. 8440) under the <i>insc</i>-Gal4 drove NSC reactivation in the absence of dietary amino acid (sucrose-only food), as there were 32.8 ± 13.9 (n = 22) EdU<sup>+</sup> cells per brain hemisphere at 72 hr ALH (Figures 5A and 5B). Similarly, at the same growth condition, when <i>hsp83</i> was depleted in InR<sup>AD</sup>-overexpressing brains, there were 32.5 ± 14.6 (n = 25) EdU<sup>+</sup> cells per brain hemisphere (Figures 5A and 5B). Likewise, at 24 hr ALH in...
sucrose-only food, in InRAD-overexpressing brains with hsp83 RNAi knockdown, there were 17.1 ± 6.6 (n = 20) EdU+ cells per hemisphere (Figures S5A and S5B), similar to InRAD overexpression alone (Figures S5A and S5B; 21.3 ± 5.0 EdU+ cells per hemisphere, n = 32). These observations suggest that over-activation of InR was epistatic to hsp83 depletion in NSCs. Therefore, Hsp83 likely functions upstream of InR during NSC reactivation.

Rheb is a GTPase and a downstream effector of the InR/PI3K/Akt pathway in NSC reactivation (Shim et al., 2013). In sucrose-only food at 24 hr ALH, there were 15.2 ± 2.8 (n = 42) EdU+ cells per brain hemisphere upon Rhebwild-type expression (wild-type form, BDSC no. 9689) under the insc-Gal4 (Figures S5B and S5D). Under the same conditions, 13.8 ± 3.4 EdU+ cells (n = 48) were observed in larval brains overexpressing Rhebwild with hsp83 RNAi knockdown (Figures S5B and SSD). Under the same conditions, 13.8 ± 3.4 EdU+ cells (n = 48) were observed in larval brains overexpressing Rhebwild with hsp83 RNAi knockdown (Figures S5B and SSD). Likewise, at 48 hr ALH on sucrose-only food, the number of EdU+ cells was similar in Rhebwild-overexpressing brains with hsp83 RNAi knockdown (Figures S5B and SSD; 21.6 ± 4.0, n = 26) and without hsp83 depletion (Figures S5B and SSD; 18.5 ± 4.2, n = 27).

Under fed conditions, InRAD or Rhebwild overexpression also significantly rescued NSC quiescence phenotype caused by hsp83 knockdown. At 48 hr ALH, vast majority of control NSCs were reactivated, except for 1.62% ± 0.8% NSCs without EdU incorporation (Figures 5E and 5F; n = 2410, 24 brain lobes). Similarly, almost all NSCs with InRAD or Rhebwild overexpression were reactivated as well, with none (Figures 5E and 5F; n = 2,412, 24 brain lobes) or 0.3% ± 0.4% (Figures 5E and 5F; n = 1,757, 18 brain lobes) NSCs without EdU incorporation. Upon hsp83 RNAi knockdown in NSCs, 48.9% ± 8.2% (Figures 5E and 5F; n = 1,357, 17 brain lobes) lacked EdU incorporation. By contrast, at the same time point, in NSCs overexpressing InRAD or Rhebwild concomitant with hsp83 RNAi knockdown, the percentages of EdU+ NSCs were decreased significantly to 11.7% ± 1.5% (Figures 5E and 5F; n = 2,392, 26 brain lobes) and 10.4% ± 2.3% (Figures 5E and 5F; n = 1,827, 22 brain lobes).

Taken together, overexpression of InRAD or Rhebwild significantly rescued NSC reactivation defects observed upon hsp83 RNAi knockdown.

Amino Acids Deprivation Causes Significant Downregulation of hsp83 mRNA and Inactivation of InR/PI3K/Akt Pathway

To explore whether the expression of InR, hsp83, and cdc37 are nutrition dependent, we assessed the RNA level of InR, hsp83, and cdc37 in larvae upon amino acid deprivation. At 24 hr ALH in sucrose-only food, the mRNA levels of hsp83 were dramatically reduced to 13.4% ± 3.2% and 10.4% ± 3.1% measured by two different primer pairs (n = 4 each), compared with control larvae raised under fed conditions at the same time point (Figure S5E). RNA level of
cdc37 remains unchanged on the starvation condition (Figure S5E). This suggests that the expression of hsp83, but not cdc37, is dependent on the presence of dietary amino acids. Surprisingly, the mRNA level of InR and 4E-binding protein (4E-BP) on sucrose-only food was increased significantly to at least 5-fold (Figure S5E; n = 4). This result is in line with the previous finding that increased transcription of InR and 4E-BP was correlated with inactivation of InR pathway and growth inhibition when nutrients are limited due to a feedback control mechanism for the InR pathway (Puig et al., 2003). Since the upregulation of mRNA levels of InR in the entire larvae upon amino acid deprivation was not observed in larval brains upon knocking down of hsp83 alone, this effect likely involves additional factors that are altered by depletion of dietary amino acids or due to different experimental conditions. Therefore, amino acid deprivation causes dramatic downregulation of hsp83 and inactivation of InR/PI3K/Akt pathway.

DISCUSSION

How the InR/PI3K/Akt pathway is regulated during NSC reactivation is poorly understood. Here we show that molecular chaperone Hsp83/Hsp90, together with its co-chaperone Cdc37, play a role in the reactivation of Drosophila NSCs. Mechanistically, Hsp83 and Cdc37 physically associate with InR and are important for the activation of the InR/PI3K/Akt pathway in NSCs. Therefore, we demonstrate that Hsp83 serves as an intrinsic factor within NSCs that is necessary for the activation of the InR/PI3K/Akt pathway in NSCs. Our evidence suggests that Hsp83 and Cdc37 regulate the protein folding and activation of InR in the nervous system.

The role of Hsp83 in NSC reactivation at early larval stages is distinct from its known role in centrosomes or NSC polarity. Drosophila Hsp83 is a core centrosomal component required for proper mitotic spindle formation.
and chromosome segregation (Lange et al., 2000). In *Drosophila* larval CNS, Hsp83 and co-chaperone Sgt1 are required for the stabilization of Polo and centrosome organization in NSCs (Martins et al., 2009). Hsp83 and Sgt1 are also required for the establishment of NSC polarity via the LKB1/AMPK pathway in third-instar larvae (Andersen et al., 2012). However, Sgt1 RNAi (BDSC no. 34605) in NSCs did not display any phenotypes during NSC reactivation (data not shown), suggesting that Hsp83 interacts with different co-chaperones to control NSC reactivation and cortical polarity at different developmental stages. Consistent with this notion, we found that Cdc37, but not other co-chaperones of Hsp83, is required for NSC reactivation. We found that the proliferation of MB NSCs were unaffected by *hsp83* knockdown. Therefore, Hsp83 promotes NSC reactivation rather than general cell proliferation. Consistent with our observations, there is no significant difference in proliferation between *hsp83* mutant and wild-type eye imaginal discs in the proliferating zone (Bandura et al., 2013). Interestingly, in pupal eyes that undergo terminal differentiation, Hsp83 is required for cell-cycle exit by activating the anaphase-promoting complex/cyclosome (Bandura et al., 2013). We found cytokinesis defects in *cdc37*-depleted NSCs, but not in NSCs depleted of Hsp83 (data not shown). This observation is consistent with a known role of Cdc37 in cell division and cytokinesis in *Drosophila* (Lange et al., 2002).

Hsp90 plays a key role in signal transduction and appears to bind to its substrates in a near native state poised for activation by binding of ligand or other factors (Young et al., 2001). Since Hsp83 overexpression is sufficient to drive the activation of InR/Pi3K/Akt pathway and trigger premature NSC reactivation, Hsp83 likely plays an active role in promoting NSC reactivation by binding to InR at a late stage of folding poised for activation by dILP binding. Furthermore, in the absence of dietary amino acids, the expression of *hsp83* is downregulated, likely partially contributing to the inactivation of the InR pathway (Figure S5E). We propose that InR is a target of Hsp83 and Cdc37 during NSC reactivation. The physical association among Hsp83, Cdc37, and InR was strongly supported by PLA assays, and both *in vitro* and *in vivo* BiFC. Although tandem affinity purification-mass spectrometry in *Drosophila* S2 cells implied an interaction between Hsp83 and InR (Friedman et al., 2011), we failed to detect a consistent interaction between Hsp83 and InR *in vitro* in S2 cells in co-immunoprecipitation experiments, probably due to the transient nature of this interaction. In addition, our genetic interaction experiments indicate that Hsp83 activates the InR/Pi3K/Akt pathway to promote NSC reactivation. Taken together, InR is likely a client of Hsp83 in *Drosophila* NSCs. Consistent with our findings, in human fibroblasts, Hsp90

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**EXPERIMENTAL PROCEDURES**

**Fly Stocks and Genetics**
The fly strains used in this study were: UAS-CYFP-HA, UAS-NYFP-Myc, UAS-Hsp83-Myc-NYFP, UAS-InR<sup>intra</sup>-HA-CYFP, UAS-Cdc37-Myc-NYFP, UAS-Cdc37-HA-CYFP, UAS-Hsp83-HA (T. Wang), *hsp83*-BAC, and *cdc37*-BAC. The following stocks were obtained from Bloomington Drosophila Stock Center (BDSC): RNAi for *hsp83* (BDSC no. 28756), *cdc37<sup>2D</sup>* (BDSC no. 5693), UAS-InR<sup>A1325D</sup> (VDRC nos. 47776 and 108568) and *cdc37* RNAi (VDRC nos. 77776 and 110727) were obtained from Vienna Drosophila Resource Center (VDRC). *Hsp83<sup>36Cl2</sup>* (DGRC no. 111379) was obtained from Kyoto Drosophila Genetic Resource Center. All experiments were carried out at 25°C, except for RNAi knockdown or overexpression at 29°C.

**Immunohistochemistry**
Larval brains were dissected in PBS and fixed for 22 min in 0.3% PBS-Triton (PBT) with 4% electron microscopy (EM)-grade formaldehyde (methanol free). Fixed brains were processed for immunostaining as described previously (Koe et al., 2014). Further details and primary antibodies used can be found in Supplemental Experimental Procedures.

**MARCM Analysis**
To generate MARCM clones, late first-instar larvae were heat shocked for 2 hr at 37°C, and heat shocked for a second time 10–16 hr later after recovering at 25°C. Larvae were dissected at the third instar-larval stage and for the tissue processed by immunohistochemistry.
EdU Pulse-Chase Analysis
Larvae were fed with standard food supplemented with 0.2 mM EdU from Click-iT EdU Alexa Fluor 555 Imaging Kit (Invitrogen) for 4 hr prior to dissection. The dissected larval brains were then fixed with 4% EM-grade formaldehyde (in 0.3% PBT) for 22 min. The brains were then processed for as described previously (Li et al., 2017).

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, five figures, and three tables and can be found with this article online at https://doi.org/10.1016/j.stemcr.2018.08.014.

AUTHOR CONTRIBUTIONS
Conceptualization, H.W.; Methodology, H.J.; Writing – Original Draft, Review & Editing, H.W. and H.J.; Funding Acquisition, H.W.; Resources, H.W. and H.J.; Supervision, H.W.

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