

## Pharmacological rescue of adult hippocampal neurogenesis in a mouse model of X-linked intellectual disability



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### ABSTRACT

Oligophrenin-1 (OPHN1) is a Rho GTPase activating protein whose mutations cause X-linked intellectual disability (XLID). How loss of function of Ophn1 affects neuronal development is only partly understood. Here we have exploited adult hippocampal neurogenesis to dissect the steps of neuronal differentiation that are affected by Ophn1 deletion. We found that mice lacking Ophn1 display a reduction in the number of newborn neurons in the dentate gyrus. A significant fraction of the Ophn1-deficient newly generated neurons failed to extend an axon towards CA3, and showed an altered density of dendritic protrusions. Since Ophn1-deficient mice display overactivation of Rho-associated protein kinase (ROCK) and protein kinase A (PKA) signaling, we administered a clinically approved ROCK/PKA inhibitor (fasudil) to correct the neurogenesis defects. While administration of fasudil was not effective in rescuing axon formation, the same treatment completely restored spine density to control levels, and enhanced the long-term survival of adult-born neurons in mice lacking Ophn1. These results identify specific neurodevelopmental steps that are impacted by Ophn1 deletion, and indicate that they may be at least partially corrected by pharmacological treatment.

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### 1. Introduction

Intellectual Disability (ID) is a complex disease of the central nervous system (CNS) that results in impaired cognitive abilities (Chelly et al., 2006; Van Bokhoven, 2011). A genetic contribution to the etiology of ID is well established and, among the genetic conditions, the most frequent are the X-linked intellectual disabilities (XLIDs) forms caused by single gene mutations on the X chromosome (Humeau et al., 2009; Ropers and Hamel, 2005). Among the XLID genes, oligophrenin-1 (Ophn1) encodes a RhoGTPase-activating protein (Rho GAP) which is expressed in the developing and adult nervous system (Billuart et al., 1998; Fauchereau et al., 2003; Khelifaoui et al., 2007; Valnegri et al., 2011). At the cellular level, the protein is present both in glial cells and neurons where it colocalizes with F-actin, notably at the tip of growing dendrites (Fauchereau et al., 2003). OPHN1 is detected on

both sides of the synapse, indicating that it participates to synaptic function (Nadif Kasri et al., 2009; Nakano-Kobayashi et al., 2014; Khelifaoui et al., 2009; Nakano-Kobayashi et al., 2009). Indeed, decreased or defective Ophn1 signaling prevents synapse maturation and causes loss of synaptic structure, function and plasticity (Govek et al., 2004; Khelifaoui et al., 2007; Powell et al., 2012; Powell et al., 2014). Ophn1 knock-out (KO) mice, the murine model of the disease, exhibit behavioral impairments in spatial memory, and an immature phenotype of dendritic spines in hippocampal pyramidal neurons associated with altered synaptic plasticity (Khelifaoui et al., 2007; Khelifaoui et al., 2014). However, how Ophn1 loss of function impacts on neuronal development and circuit formation, leading to impaired information processing remains still incompletely understood. This is mainly due to the technical difficulties to study embryonic/perinatal neurogenic processes (i.e. proliferation, migration, morphological differentiation and functional integration) that largely overlap in time.

Here, we have examined how lack of Ophn1 impacts on neuronal maturation by following the process of adult neurogenesis - i.e. the morphological and functional development of new dentate granule neurons that are normally and constantly added to hippocampus (Kempermann et al., 2015; Braun and Jessberger, 2014; Christian et al., 2014). In

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mammals, new neurons are produced in two specific neurogenic niches, the subventricular zone (SVZ) and the subgranular zone (SGZ) in the dentate gyrus (DG) of the hippocampus. Adult hippocampal neurogenesis recapitulates at least some neuronal developmental processes including proliferation of progenitors, morphological differentiation and integration of newborn neurons into brain networks (Kempermann et al., 2015). In the SGZ, stem/progenitor cells proliferate and give rise to neuroblasts that localize within the granule cell layer of the DG, elaborate their dendritic arbor into the molecular layer and extend axons to innervate target cells in the CA3 region. Adult newborn neurons finally receive functional excitatory and inhibitory synaptic connections in order to be integrated in the existing hippocampal circuitry (Kempermann et al., 2015; Braun and Jessberger, 2014; Christian et al., 2014; Toni and Schinder, 2016). Principal dentate granule cells are the only neuronal subtype that is generated, and newly generated neurons have enhanced synaptic plasticity properties that enable them to contribute to specific forms of hippocampal-dependent learning and memory (Deng et al., 2010; Christian et al., 2014; Schmidt-Hieber et al., 2004). Accordingly, impairments in adult hippocampal neurogenesis have been suggested as one of the processes underlying learning and memory impairments in neurodevelopmental disorders (Contestabile et al., 2013; Guo et al., 2011; Ishihara et al., 2010; Giacomini et al., 2015; reviewed in Pons-Espinal et al., 2013a).

In this manuscript, we used *Ophn1* KO mice to investigate how loss of *Ophn1* affects adult neurogenesis in the DG. We observed deficits in newborn neuron survival, while proliferation of stem/progenitor cells was not affected. We also found an altered morphological maturation of newly generated cells with a robust impairment of axonal extension and immature dendritic spines in mice lacking *Ophn1*. Moreover, we also tested a rescue strategy based on inhibition of two kinases, Rho-associated protein kinase (ROCK) and protein kinase A (PKA), whose activity is potently up-regulated by loss of *Ophn1* (Khelifaoui et al., 2014). In particular, we administered the ROCK/PKA inhibitor fasudil to *Ophn1* KO mice, and found that dendritic spine alterations and survival of newborn neurons in the granule cell layer of DG could be restored upon this pharmacological treatment.

## 2. Materials and methods

### 2.1. Animals and treatment

All procedures conformed to the EU Directive 2010/63/EU for animal experiments and to the ARRIVE guidelines. Experimental protocols were approved by the Italian Ministry of Health. All analyses were performed blind to genotype and treatment. Experiments were carried out using male KO (*Ophn1*<sup>-/-</sup>) and their control WT littermates (*Ophn1*<sup>+/+</sup>) (Khelifaoui et al., 2007; Meziane et al., 2016). Mice were housed in a room with a temperature of 21 °C, 12 h light/dark cycle, and food and water available ad libitum. All animals were 2–3 months old at the beginning of the experiments. No mice were excluded from the analyses. Heterozygous females were bred with C57/Bl6 wild type mice to maintain the genetically modified colony. The animals were genotyped using the polymerase chain reaction (PCR) with a set of 2 oligonucleotides previously validated (Khelifaoui et al., 2007).

For the rescue experiments, KO and control littermate mice were treated with the clinically approved ROCK and PKA inhibitor fasudil for 7 weeks. At 2 months of age, mice were randomly assigned to the treatment or control groups. The animals in the treatment group received fasudil in the drinking water (0.65 mg/mL; Khelifaoui et al., 2014), while the controls received only water. The fasudil experiments always included a subset of naïve WT mice to be used for comparison.

### 2.2. BrdU injections

For evaluation of adult hippocampal neurogenesis in *Ophn1* KO and WT mice, BrdU (Sigma-Aldrich) was administered intraperitoneally at

50 mg/kg body weight. For the experiments on neuronal proliferation (Fig. 1A, C), mice received 2 injections spaced by 2 h and were sacrificed after 24 h. For neuronal survival experiments, we performed 4 BrdU injections every 2 h and animals were sacrificed at 15 or 50 dpi (Fig. 1D, E; Kempermann et al., 2003). Because of the different doses of BrdU, and since independent cohorts of WT and KO mice were analyzed at each time point, the difference in BrdU-positive cells between genotypes was assessed by pairwise comparisons at 1, 15 and 50 dpi.

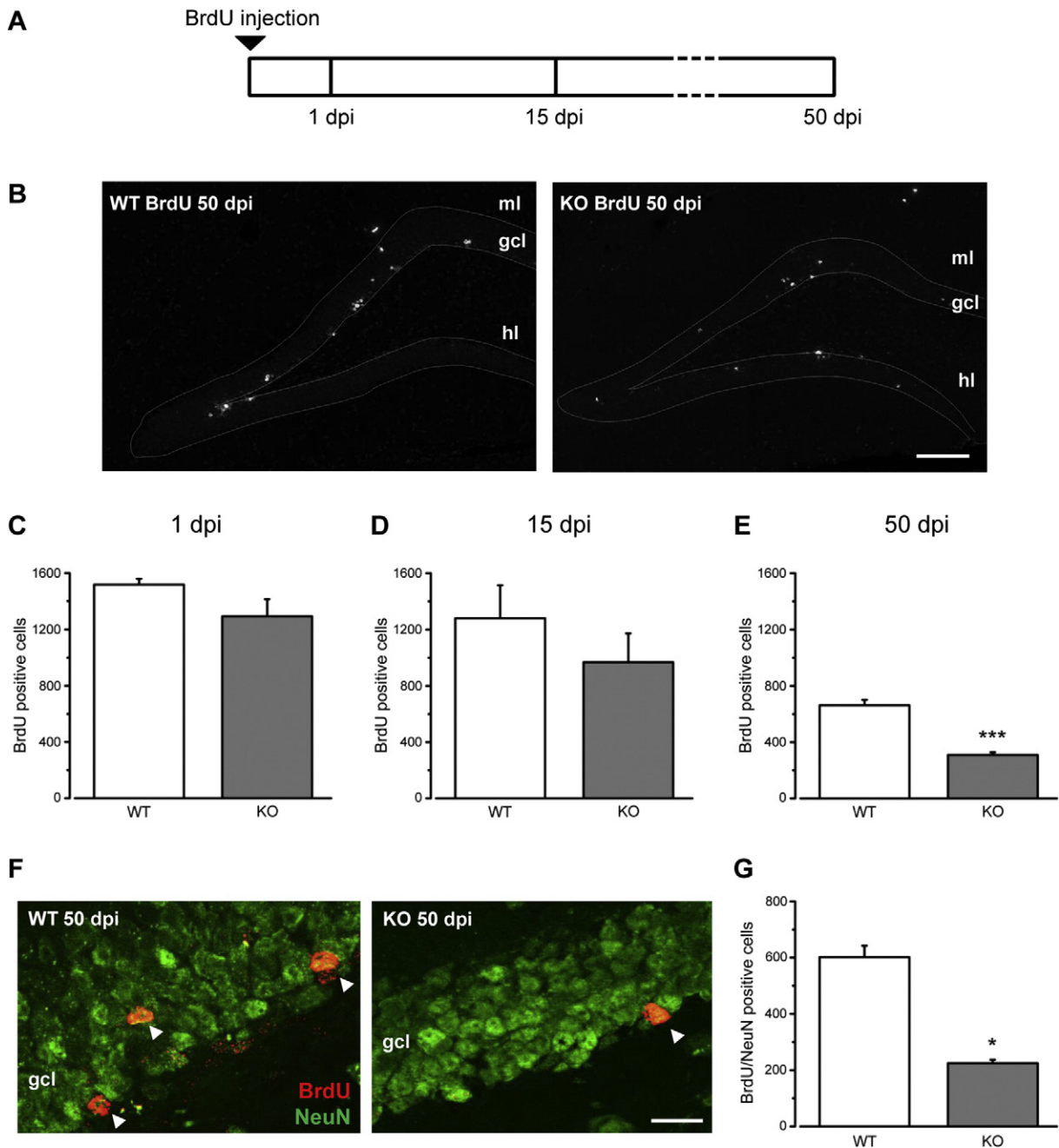
### 2.3. Immunohistochemistry and stereological cell counting

Animals were anesthetized with chloral hydrate and perfused through the heart with 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer (PB), pH 7.4. Brains were post fixed for 2 h at 4 °C, and cryoprotected in 30% sucrose in 0.1 M PB at 4 °C, and cut coronally at 40 µm with a freezing microtome. For each animal, serial sections (1 out of 6) spanning the entire rostro-caudal extension of hippocampus were processed for different stainings. For BrdU detection, free-floating sections were pretreated by denaturing DNA with 2 M HCl for 30 min at 37 °C (Rossi et al., 2006; Kempermann et al., 2003). Slices were blocked for 1 h at room temperature (RT) with 10% normal goat serum (NGS), 0.3% Triton X-100 in PBS, and then incubated overnight at 4 °C with a rat monoclonal anti-BrdU antibody (1:200, Abcam, AB6326) diluted in NGS 5% and Triton 0.1% in PBS. Signal was finally revealed with 2 h incubation at RT with Cy3 conjugated anti-rat IgG (1:500, Jackson Immuno Research). We also performed a double labeling for BrdU and neuronal nuclei (NeuN). Sections were incubated overnight with the rat anti-BrdU antibody (1:200) and guinea pig polyclonal anti-NeuN (1:1000, Millipore, ABN90P). Bound primary antibodies were revealed with Cy3 anti-rat IgG (1:500) and AlexaFluor488 anti-guinea pig IgG (1:400, Jackson Immuno Research). For Dcx staining, sections were blocked for 1 h at RT with 10% rabbit serum (RS), 0.3% Triton X-100 in PBS, and then incubated overnight at 4 °C with goat polyclonal anti-Dcx primary antibody (1:1000, Santa Cruz, sc8066), 0.2% RS, and 0.1% Triton X-100 in PBS. The primary antibody was revealed by incubation for 2 h at RT with Rhodamine Red-X conjugated anti-goat IgG (1:500, Jackson Immuno Research) diluted in 1% RS and 0.1% Triton in PBS. To detect early neuronal cells expressing Dcx and NeuN, we performed a double staining for both. Sections were incubated with the goat anti-Dcx (1:1000) and guinea pig anti-NeuN (1:1000) overnight at 4 °C. Signals were revealed with Rhodamine Red-X anti-goat IgG (1:500) and AlexaFluor488 anti-guinea pig IgG (1:400). All primary antibodies used in this report have been validated by several previous studies on hippocampal neurogenesis (Contestabile et al., 2013; Iggena et al., 2015; Vannini et al., 2016).

#### 2.3.1. Stereological counting

The number of BrdU, Dcx and Dcx/NeuN positive cells was estimated in serial coronal sections covering the complete rostro-caudal extension of the DG as previously described (Rossi et al., 2006). Cells were counted on a fluorescence microscope (Zeiss) using a 20× objective and the analysis was performed using Stereo Investigator software (MicroBrightfield). All cells in the granule cell layer and subgranular zone of every sixth section were counted, and the resulting number of cells was multiplied by six to give an estimate of the total number of labeled cells.

To investigate the fate of the newly generated cells, we performed the analysis of Dcx/NeuN and BrdU/NeuN double labeled neurons. For Dcx/NeuN staining, fluorescence images were captured with a microscope (Axio Imager. Z2, Zeiss) equipped with Apotome.2 (Zeiss) at resolutions of 1350 × 1024 pixels and a 63× EC-PLAN-NEOFLUAR oil objective (0.75 NA). For each experimental animal, 4 sections were analyzed, and for each section, z-stack images (1 µm z-step size) of 3 different regions of the dorsal granule cell layer were acquired with the ZENpro software (Zeiss). For each animal, Dcx positive cells were analyzed for the colocalization with NeuN with the cell counter plug in



**Fig. 1.** Impaired adult hippocampal neurogenesis in Ophn1 KO mice. **A**) Schematic of BrdU labeling experiments. To evaluate proliferation (i.e. number of BrdU-positive cells at 1 dpi), mice received 2 BrdU injections spaced by 2 h and were sacrificed after 24 h. For the analyses at 15 and 50 dpi, we performed 4 BrdU injections every 2 h at day 0. **B**) Representative images of BrdU labeling in the dentate gyrus of WT and KO mice at 50 dpi. Dorsal is up and medial is to the left. Scale bar: 100  $\mu$ m; ml, molecular layer; gcl, granule cell layer; hl, hilus. **C**) Stereological counting of BrdU-positive cells in the DG of WT and KO mice shows that proliferation is not affected by Ophn1 deficiency at 1 dpi (WT, 1517  $\pm$  40, n = 4; KO, 1294  $\pm$  118, n = 4; two-tailed Student's *t*-test, *P* = 0.12). **D**) The quantification of BrdU positive cells at 15 dpi indicates that the early stage of neurogenesis is similar between the two groups (WT, 1281  $\pm$  232, n = 4; KO, 968  $\pm$  205, n = 3; two-tailed Student's *t*-test, *P* = 0.37). **E**) Survival of newly generated cells is significantly reduced at 50 dpi (WT, 662  $\pm$  38, n = 11; KO, 309  $\pm$  18, n = 8; Mann-Whitney rank sum test, *P* < 0.0001). **F**) Representative images of the granule cell layer (gcl) stained for both BrdU (red) and NeuN (green) at 50 dpi in the DG of WT mice (left) and KO animals (right). Arrowheads indicate double stained cells. Dorsal is up and medial is to the left. Scale bar: 20  $\mu$ m. **G**) Stereological quantification of BrdU/NeuN double stained cells at 50 dpi shows reduced numbers of mature neuronal cells in KO mice respect to the control group (WT, 601  $\pm$  40, n = 4; KO, 224  $\pm$  12, n = 4; Mann-Whitney rank sum test, *P* = 0.028). Histograms represent mean  $\pm$  SEM. Statistical significance, \**P* < 0.05; \*\*\**P* < 0.001.

(ImageJ) and in particular we evaluated the percentage of Dcx/NeuN early neurons over the total number of Dcx-positive cells. To evaluate BrdU/NeuN double staining, for each animal, 50 BrdU positive cells were assessed for NeuN colocalization with a 40 $\times$  objective and analysis was performed using Stereo Investigator software.

To quantify the effect of fasudil treatment on early neuronal cells, the numbers of Dcx-positive and Dcx-NeuN double labeled cells of each group (naïve KO, KO treated with fasudil, WT treated with fasudil) were normalized to the mean of untreated WT mice of the same litter.

All experiments, data acquisition and quantification were performed blind to the genotype and treatment. The representative images shown in the Figures were obtained with a microscope (Axio Imager. Z2, Zeiss) equipped with Apotome.2 (Zeiss), using either a 20 $\times$  EC-PLAN-NEOFLUAR objective or a 63 $\times$  EC-PLAN-NEOFLUAR oil objective.

### 2.3.2. Stereological volume analysis

To determine the volume of the dentate gyrus and hilus, one-in-six series of sections was stained with Hoechst dye. In each section, the

DG and hilus areas were traced at 10× magnification. The total volume was determined by summing the areas contoured in each section and multiplying the result by the distance between sections sampled (300 μm).

#### 2.4. Stereotaxic injections of retroviral vectors

For virus delivery into dentate gyrus of the hippocampus, mice were anesthetized and a total volume of 1 μl of retrovirus transducing murine Moloney leukemia virus (MoMuLV)-based vector CAG-GFP (provided by Prof. F.H. Gage) was infused into each hemisphere (coordinates from Bregma: anteroposterior – 2 mm, lateral ± 1.6 mm, ventral + 1.8 mm) through the insertion of capillary glasses (WPI) connected to a manual syringe pump (Narishige). Mice were allowed to recover and housed in standard cages until the day of sacrifice.

#### 2.5. Analysis of newborn DG granule cells morphology

Confocal analysis was performed using a laser scanning motorized confocal system (Nikon A1) equipped with an Eclipse Ti-E inverted microscope.

##### 2.5.1. Analysis of dendrites

Coronal brain sections from injected brains (120 μm thick) were acquired with a 20× oil objective and digital 2 magnification. Z-series stacks were acquired with an interval of 1 μm. Maximum intensity projections were produced using ImageJ software and tracing and quantification of dendrites were obtained using NeuronJ Plugin (<http://imagescience.bigr.nl/meijering/software/neuronj>). Only neurons showing intact dendritic arborization were analyzed. Sholl analysis was performed by Sholl Analysis Plugin (Ghosh Lab Website, <http://biology.ucsd.edu/labs/ghosh/software/ShollAnalysis.class>) using a 10 μm interval between concentric circles.

##### 2.5.2. Analysis of spine density

Spines were visualized using a 60× oil objective and digital 4 magnification. Z-series stacks were acquired with an interval of 0.3 μm. The area of analysis was set on dendritic segment located in the medial region of the molecular layer of the upper blade at approximately the same distance from the cell body. The number of spines was counted manually and density was calculated dividing the total number of spines by the length of dendritic segment. Three dimensional reconstruction was performed using Nis Element Software (Nikon).

##### 2.5.3. Analysis of axons

To analyze axons of newborn neurons, cells were acquired with 60× oil objective with a z-series stack of 0.5 μm. Cells were classified for their orientation respect to the granular cell layer (horizontal or radial). Radially oriented cells were classified for presence of one single basal process extending to the hilus for a distance of at least 50 μm (single axon) or for the presence of multiple short basal processes terminating within 20–30 μm from the cell body (no axon). Three dimensional (3D) reconstructions of newborn neurons were performed using Nis Element Software (Nikon) by using pseudocolor z-depth coding. By knowing the 3D coordinates of any point on the reconstruction we were able to determine the axon's profile including the axon's depth map, an image that contains information relating to the ending distance of an axon in respect to the cell body. To provide the distance information needed to establish axogenesis we used pseudo-colors: surfaces closer to the cell body are blue; surfaces further away from the cell body are red.

Mossy fibers labeled with GFP at CA3 region were quantified measuring mean fluorescence intensity in regions of interest (ROIs).

#### 2.6. Statistical analysis

Statistical analysis was performed with SigmaPlot 12.0 software. Based on the preliminary analysis of statistical power, at least 3–6 mice were allocated to each experimental group. Pairwise comparisons of quantitative phenotypes between mice of different groups were assessed with a two-tailed Student's *t*-test when the observed treatment effects were normally distributed, or with the Mann–Whitney rank sum test when the samples were not drawn from normally distributed populations with the same variances. When more than two groups were analyzed, one way ANOVA followed by Holm–Sidak/Tukey/Dunn's post hoc test were used. The number of intersections in the Sholl analysis was examined with two-way repeated measures ANOVA, followed by Holm–Sidak test. Normality of distributions was assessed with Kolmogorov–Smirnov test. Level of significance was  $P < 0.05$ .

### 3. Results

#### 3.1. *Ophn1* deletion impairs survival of newborn cells in the hippocampal dentate gyrus (DG)

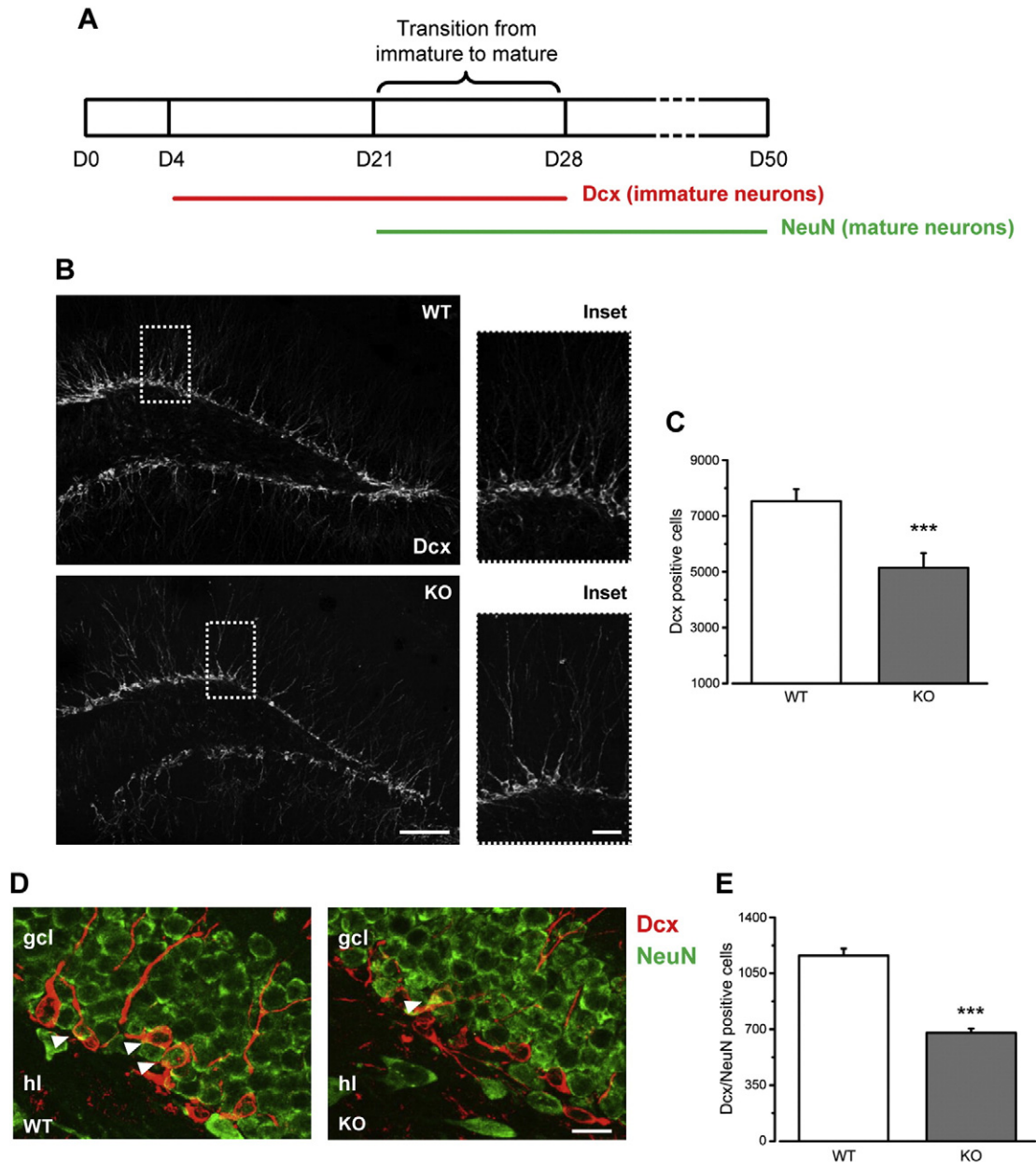
To investigate adult neurogenesis in *Ophn1*<sup>−/y</sup> mice, we performed experiments of bromo-deoxyuridine (BrdU) incorporation. In particular, we dated from birth (i.e. mitotic stage) the newly generated neurons by systemic injections of the thymidine analogue BrdU and assessed the total number of BrdU-positive cells 1, 15 and 50 days post-injection (dpi; Fig. 1A, B). This protocol allows visualizing proliferating cells and their progeny, i.e. those cells that incorporated the BrdU during the mitotic stage and then integrated in the DG circuitry.

The stereological analysis of adult hippocampal sections showed that, at 1 day after BrdU administration, the total number of newborn cells was similar in both groups, wild type (WT) and knock-out (KO) mice (Fig. 1C; Student's *t*-test,  $P > 0.05$ ). Next, we examined the survival of newly generated cells until 15 and 50 dpi. While at 15 dpi there were no significant differences between the two groups (Fig. 1D; *t*-test,  $P > 0.05$ ), we found a decrease of BrdU-positive cells at 50 dpi in KO compared to WT littermates (Fig. 1B, E; Mann–Whitney test,  $P < 0.0001$ ). Double labeling for BrdU and the neuronal marker NeuN was performed at 50 dpi to determine the phenotype of the BrdU-positive cells. We found a reduction of mature neuronal cells in KO mice with respect to the control group (Fig. 1F, G; Mann–Whitney rank sum test,  $P < 0.05$ ), pointing to impaired addition of new neurons. These results indicate that loss of *Ophn1* has no impact on cell proliferation while affecting the survival/integration of the newborn neurons in the DG hippocampal network.

Newly generated hippocampal cells committed to a neuronal fate begin expressing doublecortin (Dcx), a microtubule-associated protein that labels the newly generated neurons from the late precursor stage (about 4 days after birth) up to 4 weeks (Christie and Cameron, 2006; Fig. 2A). To quantify the total number of Dcx-positive cells, we performed the stereological counts in one hemisphere of brain sections of KO and WT animals and we found a substantial decrease of Dcx-positive newborn neurons in KO animals compared to controls (Fig. 2B, C; *t*-test,  $P \leq 0.001$ ). Between 21 and 28 days after birth, Dcx-positive newborn cells undergo a morphological and physiological development associated with the expression of neuronal nuclei (NeuN) upon final maturation (Fig. 2A). We performed a double staining with Dcx and NeuN (Fig. 2D) and we quantified the total number of early neuronal cells expressing both markers. The stereological results showed a significantly decreased number of total Dcx/NeuN positive cells in KO mice with respect to WT (Fig. 2E; *t*-test,  $P < 0.001$ ).

Altogether, these data demonstrate that *Ophn1* mainly affects the late stages of adult neurogenesis, when a postmitotic cell, already committed to the neuronal fate, is becoming a dentate granule cell.

Since previous studies have highlighted brain wide alterations in *Ophn1* KO mice (Khelifaoui et al., 2007), we performed a



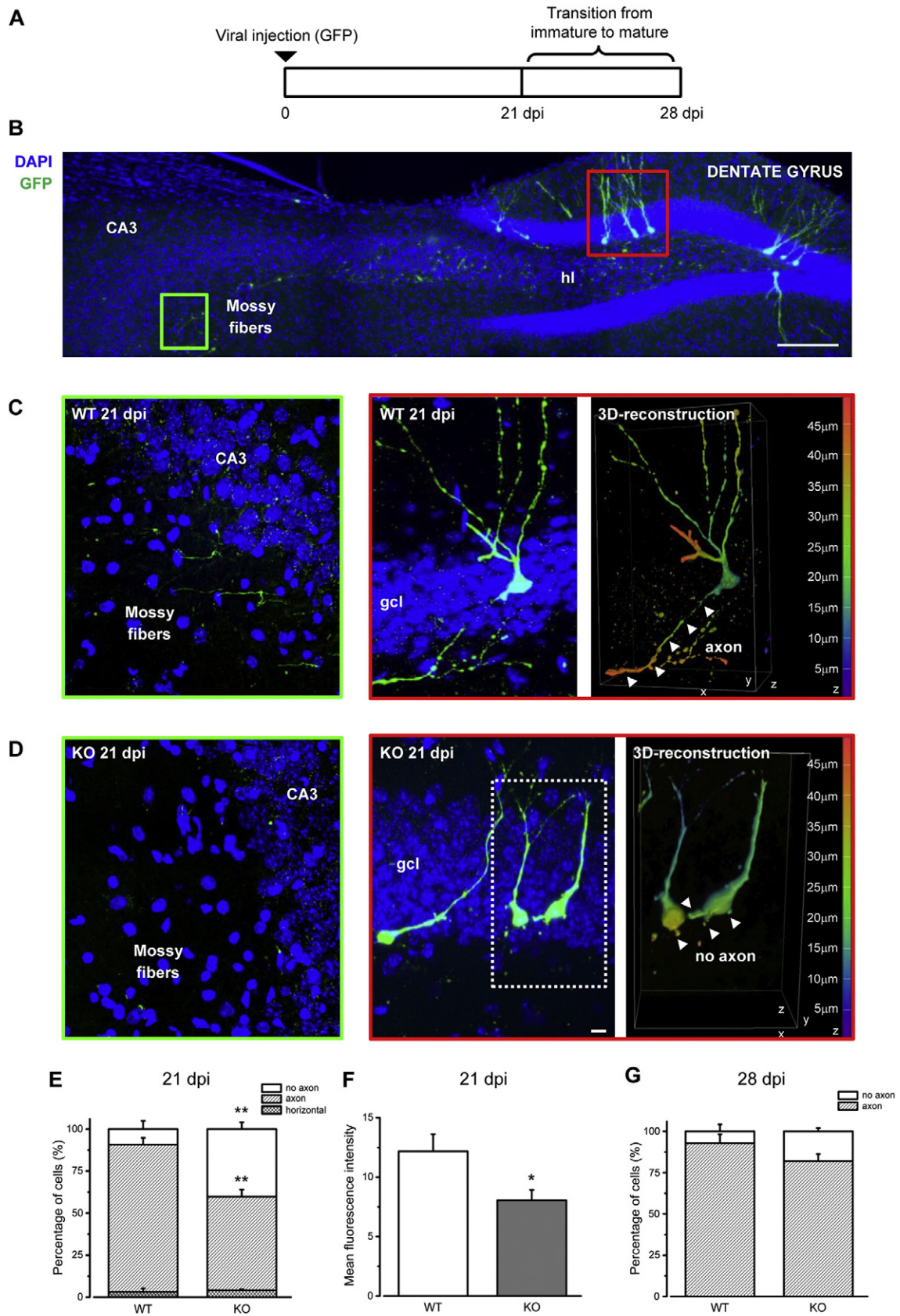
**Fig. 2.** Reduced number of Dcx-positive cells in Ophn1 KO mice. **A**) Schematic of newborn cell maturation. Neural progenitors express Dcx from day (D) 4 to D28. Dcx-positive cells undergo a morphological and physiological maturation with the expression of NeuN upon final maturation. **B**) Representative immunoreactivity for Dcx in the DG of WT (top) and KO (bottom) mice. Dorsal is up and medial is to the right; ml, molecular layer; gcl, granule cell layer; hl, hilus. Scale bar: 100  $\mu$ m (insets, 40  $\mu$ m). **C**) Stereological counting of the total number of Dcx-positive cells reveals a strong reduction in the DG of KO mice with respect to controls (WT,  $7531 \pm 428$ ,  $n = 8$ ; KO,  $5144 \pm 530$ ,  $n = 7$ ; two-tailed  $t$ -test,  $P = 0.001$ ). **D**) Representative immunolabeling for Dcx (red) and NeuN (green) in the hippocampus of WT and KO mice (arrowheads point to double stained cells). Scale bar = 20  $\mu$ m. **E**) Stereological quantification of early neuronal cells double labeled for both Dcx and NeuN in the hippocampal DG shows a significant reduction in KO (grey) mice in comparison to WT (white) animals (WT,  $1161 \pm 44$ ,  $n = 4$ ; KO,  $678 \pm 26$ ,  $n = 3$ ; two-tailed  $t$ -test,  $P = 0.0003$ ). Histograms represent mean  $\pm$  SEM. Statistical significance, \*\*\* $P \leq 0.001$ .

neuroanatomical, stereological analysis of the volume of the neurogenic hippocampal niche. We found no differences in the volume of either granule cell layer or the hilus between WT and KO mice ( $t$ -test,  $P > 0.05$ ; Suppl. Fig. 1).

### 3.2. Impaired morphological maturation of newborn neurons in Ophn1 KO mice

The decrease of Dcx/NeuN double positive cells suggests an impairment of neuron integration in the pre-existing circuits of the dentate gyrus (DG) of Ophn1-deficient mice. Accordingly, we investigated axonal and dendritic maturation of newly generated neurons in Ophn1 KO mice and controls. Specifically, we injected WT and KO hippocampi

with retroviral vectors expressing GFP to label proliferating cells (Bergami et al., 2008; Bergami et al., 2013; Zuccaro et al., 2014) and examined axonal formation at 21 and 28 dpi (Fig. 3A). We found that while about 88% of the newborn cells in WT animals have extended an axon at 21 days, this percentage is dramatically reduced in KO mice (about 60%; Fig. 3B–E;  $t$ -test,  $P < 0.05$ ). The majority of neurons from KO mice extended multiple small processes ending at a short distance from the cell body, as revealed by three-dimensional (3D) reconstruction images (Fig. 3C, D; right panels). Moreover, we quantified the presence of GFP-positive axons reaching the CA3 region by measuring fluorescence intensity in defined areas (Fig. 3B–D). In contrast to control mice, KO mice showed very little and thin axon fibers found in the CA3 region (Fig. 3C, D, F;  $t$ -test,  $P = 0.032$ ). Impaired axogenesis was no



longer significant at 28 dpi (Fig. 3G; *t*-test,  $P > 0.05$ ). Thus, loss of Ophn1 impairs axon formation at a specific developmental stage in adult-generated neurons.

Prompted by the axonal phenotype, we reconstructed GFP-stained neurons in the DG to examine the dendritic structure of newborn cells at 21 dpi (Fig. 4A). We found that overall dendritic length was not impacted by lack of Ophn1 (Fig. 4B; *t*-test,  $P > 0.05$ ). Dendrite complexity was examined by Sholl analysis. We found a slight increase in the number of intersections between dendrites and Sholl circles at a distance of 180–230  $\mu\text{m}$  from the cell soma in the KO animals (two-way repeated measures ANOVA followed by Holm-Sidak test,  $P < 0.0001$ ; Fig. 4C).

We also examined density of spines on the dendrites of newborn DG neurons. We found that Ophn1-deficient mice displayed an overall greater density of dendritic protrusions at 21 dpi (Fig. 4D, E; *t*-test,  $P < 0.001$ ). In WT animals, density of dendritic spines was dramatically increased from 21 to 28 dpi, while this rate of increase was much less in KO mice (Fig. 4E; *t*-test,  $P < 0.001$ ).

Overall, the morphological analysis reveals robust alterations in the structure of newborn neurons, in particular a reduction in the percentage of cells able to extend an axon towards CA3 and an alteration in the spine density in Ophn1-deficient mice.

### 3.3. Rescue of adult hippocampal neurogenesis via systemic fasudil administration in Ophn1 KO mice

We next investigated whether the hyperactivation of the ROCK/PKA cascade, set in motion by Ophn1 deficiency (Govek et al., 2004; Khelifaoui et al., 2014), mediates the impairments of adult neurogenesis in Ophn1<sup>-/-</sup> mice. We performed a rescue experiment by treatment with the clinically approved ROCK/PKA inhibitor fasudil (Khelifaoui et al., 2014; Meziane et al., 2016) and assessed whether this pharmacological strategy was able to restore the impairments in hippocampal neurogenesis.

As shown in Fig. 5A, B, seven weeks of fasudil administration were able to completely normalize the total number of Dcx-stained cells (*t*-test,  $P < 0.01$ ). The percentage of Dcx/NeuN double stained cells over the total sample of Dcx-positive cells was also rescued by fasudil treatment (Fig. 5C; *t*-test,  $P < 0.05$ ).

We next evaluated the impact of fasudil delivery on the morphology of adult-generated neurons, 21 days after birth (21 dpi; Fig. 6A). The axonal phenotype was not impacted by fasudil and indeed we detected an abnormally low proportion (about 60%) of granule cells with an axon extending towards CA3. This proportion was not different in fasudil-treated and untreated KO mice (*t*-test,  $P > 0.05$ ; Fig. 6B). In contrast, the spine alterations were rescued by fasudil treatment, with a significant decrease of the density of dendritic protrusions (*t*-test,  $P < 0.05$ ; Fig. 6C).

To ascertain whether fasudil treatment could normalize the number of mature newborn neurons, we evaluated the total number of BrdU-positive cells in the hippocampal DG, 50 days after injection. The data demonstrated that systemic fasudil administration partially rescues the impairment found at 50 dpi (Fig. 6D, E). Fasudil-treated KO mice displayed a higher number of BrdU-positive cells in comparison to the KO vehicle animals (Mann-Whitney test,  $P < 0.05$ ). Moreover, the quantification of BrdU and NeuN double stained cells demonstrated that

fasudil treatment is able to increase the number of mature newborn neurons at 50 dpi in KO mice (Mann-Whitney rank sum test,  $P \leq 0.01$ ; Fig. 6F, G). Overall, these data demonstrate that fasudil treatment for 7 weeks rescues a significant proportion of adult-generated neurons in Ophn1 KO mice.

As a control, we also checked whether fasudil treatment per se increases the neurogenic cellular population in the hippocampal DG of WT animals. As shown in Fig. 7, control and fasudil-treated WT mice displayed similar numbers of both Dcx-positive cells (Fig. 7A *t*-test,  $P > 0.05$ ) and BrdU-labeled mature newborn neurons at 50 dpi (Fig. 7B, *t*-test,  $P > 0.05$ ). Thus, fasudil appears to restore impaired generation of newborn cells in Ophn1-deficient mice but has no impact on basal neurogenesis in healthy mice.

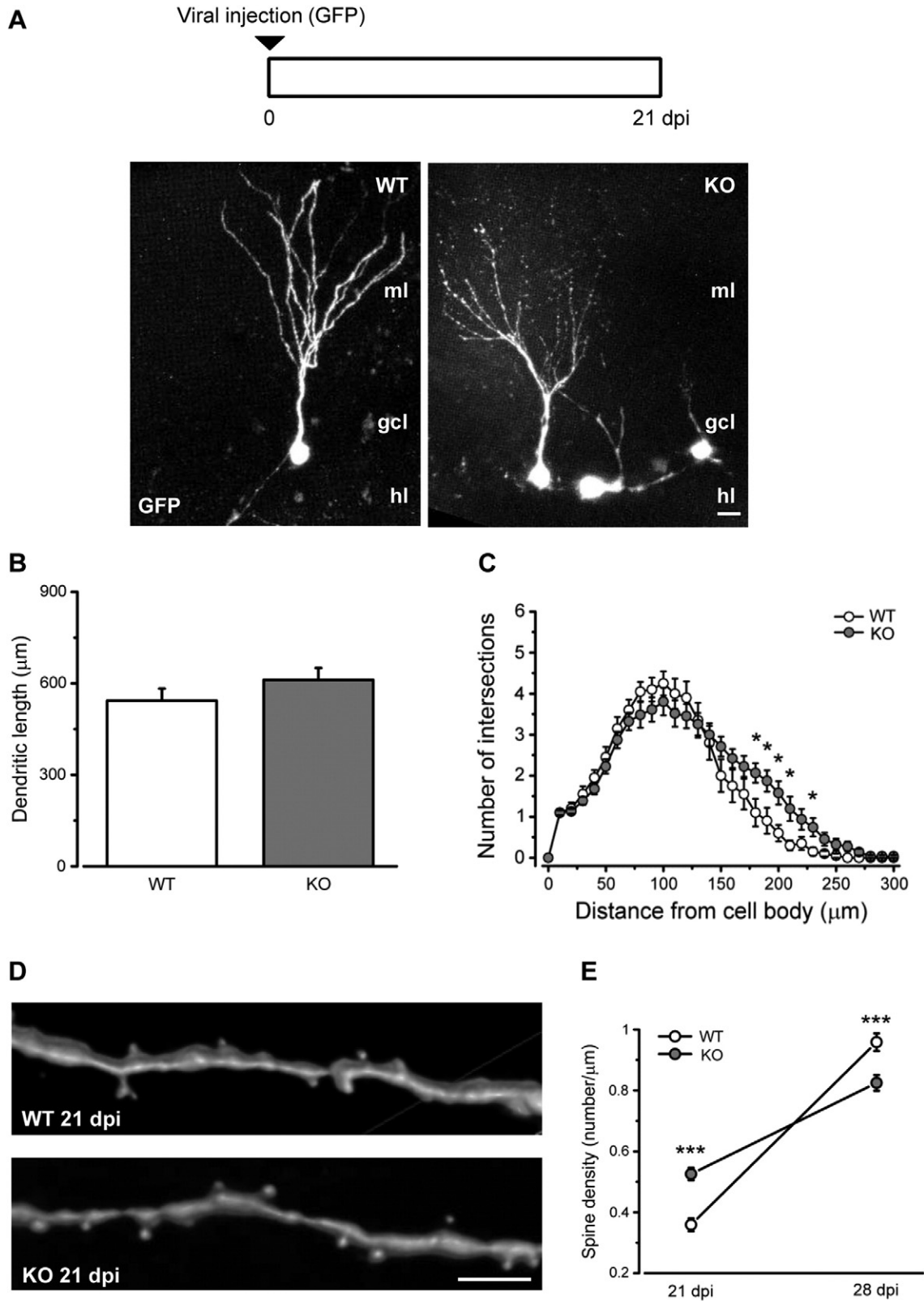
## 4. Discussion

Ophn1-dependent intellectual disability is a complex neurodevelopmental disorder involving several pathophysiological processes occurring in various brain areas. Studying adult hippocampal neurogenesis may be useful to shed light on the alterations in neuronal maturation that underlie cognitive impairment in XLID. Indeed, adult neurogenesis recapitulates at least some of the aspects of brain development and comprises a series of sequential events including proliferation of stem/progenitor cells, morphological maturation and functional integration of newly generated neurons into hippocampal circuitry. These steps have been characterized in detail (Kempermann et al., 2015; Braun and Jessberger, 2014; Christian et al., 2014; Bergami et al., 2015) and allow a window on brain development and hippocampal function (Toni and Schinder, 2016; Deng et al., 2010; Akers et al., 2014). Thus, adult neurogenesis represents an accessible model to dissect abnormalities in neuronal maturation/wiring, and test novel treatment options (Contestabile et al., 2013; Guo et al., 2011; Pons-Espinal et al., 2013b; Giacomini et al., 2015). Importantly, hippocampal neurogenesis deficits may contribute to the pathophysiology of intellectual disabilities, such as Rett, Fragile X and Down syndrome (Guo et al., 2011; Christian et al., 2014; Pons-Espinal et al., 2013a). The data presented here demonstrate that Ophn1 loss of function causes robust alterations in adult hippocampal neurogenesis, and suggest a possible therapeutic strategy for counteracting the impairments in neuronal maturation by inhibiting ROCK/PKA signaling pathways.

### 4.1. Impaired adult hippocampal neurogenesis in mice lacking Ophn1

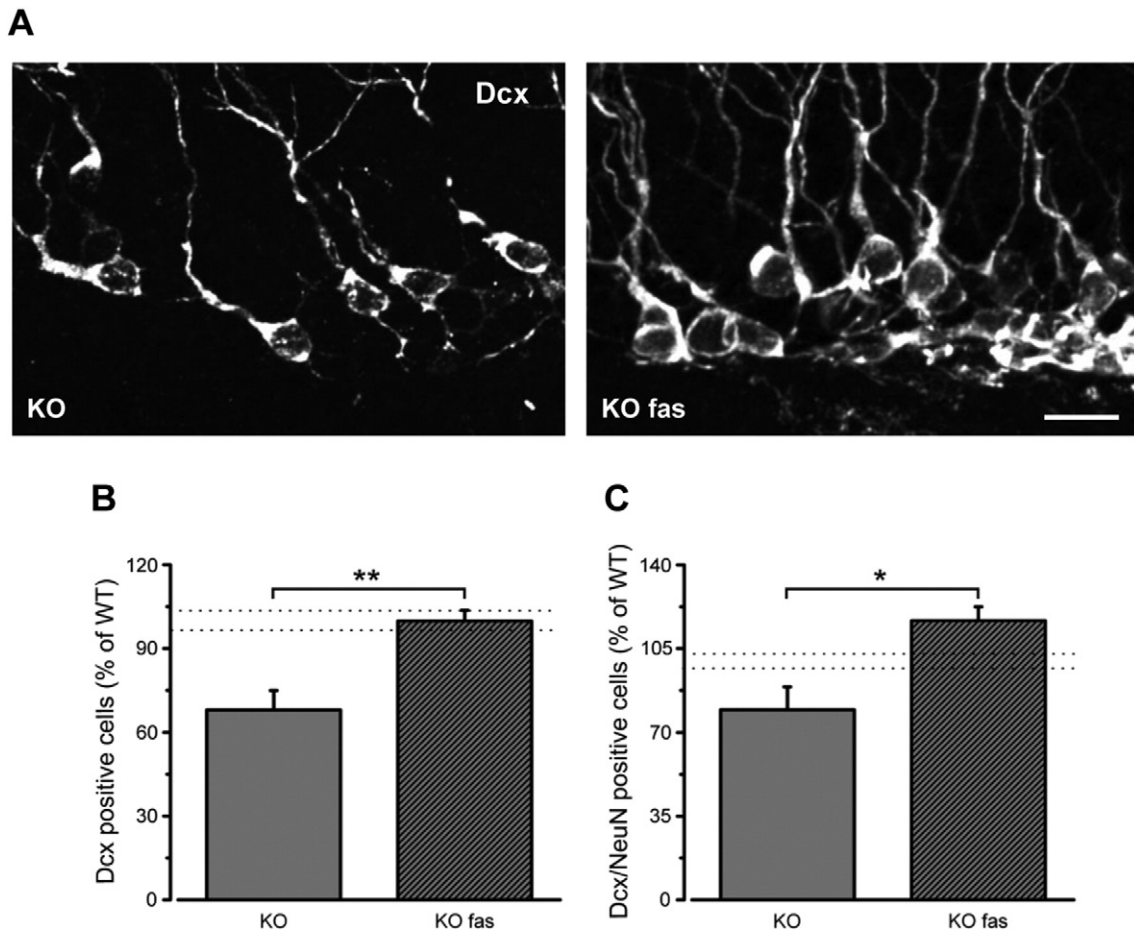
The number of BrdU-labeled cells at 1dpi was superimposable in Ophn1<sup>+/-</sup> and Ophn1<sup>-/-</sup> mice, indicating that proliferation of stem/progenitor cells was not impacted by Ophn1 deficiency. Similar data were obtained at 15 days. Analysis of Dcx-positive immature neurons, however, revealed a significant deficit in Ophn1 KO animals. Dcx expression extends from 4 days post birth up to a phase of postmitotic maturation lasting until about P28 (Christie and Cameron, 2006; Kempermann et al., 2015). Taken together, these data indicate that newly generated cells become particularly vulnerable to Ophn1 deficiency in a phase comprised between the third and fourth week post birth. Newborn neurons at 21 days also displayed significant alterations in neurite extension, with reduced axon formation, a phenotype that

**Fig. 3.** Ophn1 regulates axon formation in adult newborn neurons. A) Schematics of the experimental paradigm. Retroviral vectors expressing GFP were injected in the dentate gyrus of WT and KO animals, and mice perfused immediately before (21 dpi) and after (28 dpi) conversion from immature to mature neurons. B) Coronal section of the mouse hippocampus showing adult-generated neurons transduced with GFP-retrovirus in the granule cell layer (gcl), extending dendrites in the molecular layer (ml) of the dentate gyrus and axons through the hilus (hl) towards the CA3 area. Ventral is up and medial is to the right. Scale bar: 100  $\mu\text{m}$ . C) Left panel shows confocal image of mossy fibers from GFP-labeled newborn neuron from a WT mouse reaching the CA3 area (green box) at 21 dpi. Right panel shows confocal image (red box) of one axon extending from GFP-labeled newborn neurons. D) Confocal images from KO mice as in C. Three dimensional reconstructions display axons exceeding z axes (WT) or short processes not exceeding z axes (KO;  $z = 45 \mu\text{m}$ ); pseudocolor measures z distance. E) Histogram depicts percentage of newborn neurons with single axon, no axon or horizontally oriented at 21 dpi in WT and KO mice (WT with axon,  $90.6 \pm 4.1\%$  and WT without axon,  $6.25 \pm 4.7\%$ ,  $n = 3$  mice; KO with axon,  $59.8 \pm 4.15\%$  and KO without axon,  $36 \pm 3.9\%$ ,  $n = 4$  mice; two-tailed *t*-test,  $P = 0.003$  and  $P = 0.004$ , respectively). F) Histogram depicts quantification of mean fluorescence intensity of fibers reaching CA3 area at 21 dpi (WT  $12.18 \pm 1.43$ ,  $n = 6$  slices from 3 mice; KO  $8.05 \pm 0.87$ ,  $n = 6$  slices from 3 mice; two-tailed *t*-test,  $P = 0.032$ ). G) Histogram depicts percentage of newborn neurons with single axon or no axon at 28 dpi (WT with axon,  $92.8 \pm 5.3\%$ ,  $n = 3$  mice; KO with axon,  $82 \pm 4.1\%$ ,  $n = 4$  mice; two-tailed *t*-test,  $P = 0.16$ ). Histograms represent mean  $\pm$  SEM. Statistical significance, \* $P < 0.05$ , \*\* $P < 0.01$ .



**Fig. 4.** Dendritic complexity and spine density are regulated by Ophn1 in adult born neurons. **A)** Top, schematic of the experimental protocol. Bottom, representative confocal images depict newborn neurons morphology from the two different genotypes (ml: molecular layer; gcl: granule cell layer; hl: hilus). Scale bar: 10  $\mu\text{m}$ . **B)** Total dendritic length of newborn neurons is similar for both groups (WT, 543.3  $\pm$  39.2,  $n = 22$  cells; KO, 611.4  $\pm$  38.9,  $n = 31$  cells; two-tailed  $t$ -test,  $P = 0.23$ ). **C)** Sholl analysis of the dendritic arbors shows a slightly increased complexity 180–230  $\mu\text{m}$  from the cell soma in KO mice (WT,  $n = 20$  cells; KO,  $n = 31$  cells; Two-way RM ANOVA,  $P < 0.0001$ , followed by Holm-Sidak test,  $P < 0.05$ ). **D)** Representative images show 3D reconstruction of dendritic segments and protrusions of WT and KO newborn neurons at 21 dpi. Scale bar = 5  $\mu\text{m}$ . **E)** Quantification of spine density expressed as the number of protrusions per micrometer of dendritic segment length at 21 and 28 dpi in WT and KO mice (21 dpi: WT, 0.36  $\pm$  0.02,  $n = 32$  dendritic segments; KO, 0.53  $\pm$  0.02,  $n = 35$  dendritic segments; two-tailed  $t$ -test,  $P < 0.0001$ ; 28 dpi: WT, 0.96  $\pm$  0.03,  $n = 42$ ; KO, 0.82  $\pm$  0.03,  $n = 50$ ; two-tailed  $t$ -test,  $P < 0.001$ ). Graphs represent mean  $\pm$  SEM. Statistical significance, \* $P < 0.05$ , \*\*\* $P \leq 0.001$ .





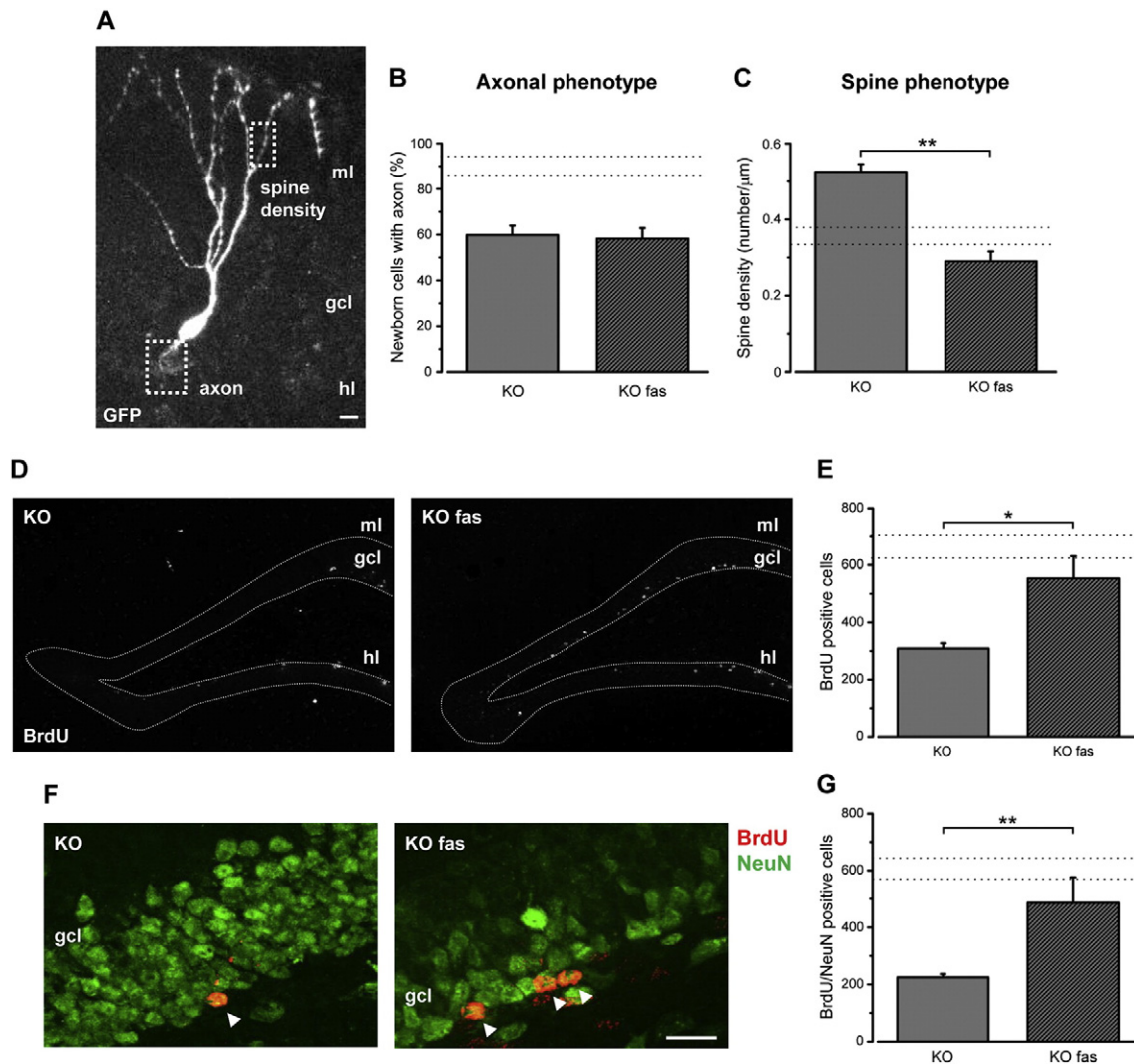
**Fig. 5.** Fasudil treatment rescues the population of Dcx-positive cells in Ophn1 KO mice. **A)** Representative images of hippocampal coronal sections (dorsal is up and medial is to the right) showing Dcx labeling in the DG of naive KO mice (left) and KO animals treated with fasudil for 7 weeks (KO fas, right). Scale bar: 20  $\mu$ m. **B)** Stereological counts show that fasudil administration rescues the total number of Dcx positive cells in the DG of KO mice (percentage of WT: KO, 67.9  $\pm$  6.9%, n = 7; KO + fasudil, 99.8  $\pm$  3.8%, n = 5; two-tailed *t*-test, *P* = 0.002). Data are expressed as percentage of the WT group (n = 13; the range values of WT mice is indicated by the dotted lines). **C)** Graph indicates the proportion of early neuronal cells (Dcx-NeuN double labeled) over the total sample of Dcx-positive cells in KO naive mice and upon fasudil treatment (KO, 79.3  $\pm$  9.6%, n = 3; KO + fasudil, 116.6  $\pm$  5.7%, n = 3; two-tailed *t*-test, *P* = 0.029). Data are expressed as percentage of the WT group (n = 8; the range values of WT mice is indicated by the dotted lines). Histograms represent mean  $\pm$  SEM. Statistical significance, \**P* < 0.05, \*\**P* < 0.01.

was not observed at 28 days (see Fig. 3). One possible explanation for this finding is a delay in axonal growth of neurons lacking Ophn1. It is also possible that Ophn1-deficient neurons unable to extend their axons towards CA3 undergo programmed cell death and are removed from the hippocampal network at the time of their integration. This is consistent with the robust drop of early neuronal cells (double labeled for Dcx and NeuN, corresponding to the time window 21–28 days post-birth) in the DG of Ophn1-deficient mice.

Maturation of newborn cells continues over the subsequent weeks and there is consensus in the literature that after a period of 7–8 weeks, the new neurons become indistinguishable from the resident granule cells (Kempermann et al., 2015; Toni and Schinder, 2016). We therefore assessed the number of mature, BrdU-labeled neurons at 50 dpi. We found a significant decline in the number of newborn cells in the DG of Ophn1 KO mice. Thus, decreased new neuron production and integration may contribute to dysfunction of the hippocampus in Ophn1-dependent XLID (Khelifaoui et al., 2007; Meziane et al., 2016). Specifically, recent data indicate that high levels of adult hippocampal neurogenesis contribute to removal of context fear memory, in particular removal of context fear memory (Akers et al., 2014; Saxe et al., 2006). Consistent with these data, Ophn1<sup>-/-</sup> mice display a deficit in fear extinction memory (Khelifaoui et al., 2014).

#### 4.2. Morphological abnormalities of newly generated granule cells in the DG of Ophn1 KO mice

To study the morphological maturation of adult-generated neurons we injected a retroviral vector expressing GFP into the DG (Zuccaro et al., 2014). We analyzed axon formation and dendritic structure by confocal microscopy. In the control mice, the vast majority (almost 90%) of 21-days-old neurons displayed one axon traveling through the hilus and reaching the CA3 region. Conversely, loss of Ophn1 caused a significant decrease in the proportion of neurons forming an axon. This was demonstrated by both the 3D analysis of axonal formation and by the assessment of fluorescence intensity in CA3 region. However, it is difficult to establish whether these axon terminals form functional synaptic boutons onto the target pyramidal neurons. A role for OPHN1 in axon extension is supported by its localization at the tip of growing neurites (Fauchereau et al., 2003). To our knowledge, this is the first report of aberrant axon formation in Ophn1-dependent XLID. It is likely that alterations in long-range axonal extension may result in abnormal network connectivity and consequent deficits in information processing in XLID. Indeed disruptions in functional connectivity are a typical hallmark of neurodevelopmental disorders (Van Der Molen et al., 2014; Plitt et al., 2015).



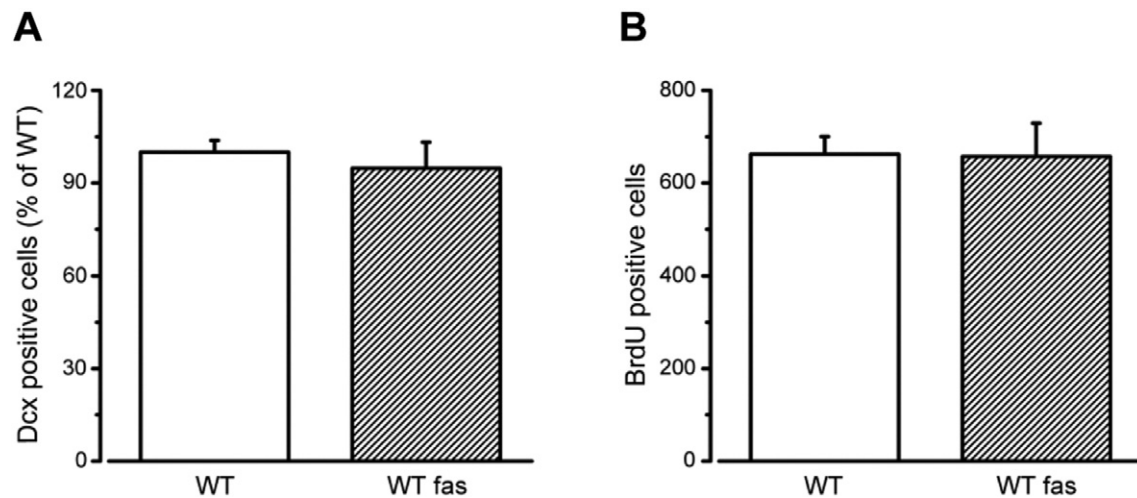
**Fig. 6.** Fasudil treatment rescues the dendritic phenotype and promotes long-term survival of newly generated neurons in Ophn1 KO mice. A) Confocal image depicts a representative GFP-labeled newborn neuron at 21 dpi, analyzed for axonal phenotype and spine density. B) Percentage of newborn neurons displaying an axon in vehicle- and fasudil-treated KO mice. No difference is detectable between the two groups (KO,  $59.8 \pm 4.1\%$ ,  $n = 4$  mice; KO + fasudil,  $58.2 \pm 4.6\%$ ,  $n = 3$ ; two-tailed *t*-test,  $P = 0.808$ ). C) Spine density of newborn neurons (21 dpi) is significantly reduced by fasudil administration (KO,  $0.53 \pm 0.02$ ,  $n = 35$  dendritic segments; KO + fasudil,  $0.29 \pm 0.02$ ,  $n = 15$  dendritic segments; two-tailed *t*-test,  $P = 0.005$ ). D) Representative low magnification images of hippocampal coronal sections stained for BrdU at 50 dpi in the DG of KO animals (left) and KO animals treated with fasudil (KO fas, right). Dorsal is up and medial is to the left. Scale bar: 100  $\mu\text{m}$ ; gcl, granule cell layer; hl, hilus; ml, molecular layer. E) Stereological counts show that 7 weeks of fasudil treatment significantly increases BrdU-positive newly generated cells in KO mice 50 dpi (KO,  $309 \pm 18$ ,  $n = 8$ ; KO + fasudil,  $554 \pm 77$ ,  $n = 6$ ; Mann-Whitney rank sum test,  $P = 0.029$ ). F) Representative images of the gcl stained for both BrdU (red) and NeuN (green) at 50 dpi in KO mice (left) and KO animals treated with fasudil for 7 weeks (KO fas, right). Arrowheads indicate double stained cells. Dorsal is up and medial is to the left. Scale bar: 20  $\mu\text{m}$ . G) The total number of BrdU/NeuN double labeled cells at 50 dpi is at least partially rescued in KO mice upon fasudil treatment (KO,  $224 \pm 12$ ,  $n = 4$ ; KO fas,  $485 \pm 91$ ,  $n = 5$ ; Mann-Whitney rank sum test,  $P = 0.015$ ). Histograms represent mean  $\pm$  SEM and the dotted lines indicate the range values of the WT group. Statistical significance, \* $P \leq 0.05$ , \*\* $P \leq 0.01$ .

We also examined the structure of the dendritic arbor. Newly generated neurons start to extend dendritic processes into the molecular layer at around 7 days after birth (Zhao et al., 2006), and to form afferent connections with the local neuronal network during the third week after birth (Deng et al., 2010). In particular, adult-born neurons begin to elaborate spines that form synapses with the afferent perforant fibers from the entorhinal cortex, and this phase of synaptic integration is characterized by increased synaptic plasticity (Ge et al., 2007; Ge et al., 2008). We found that while total dendritic length was unaltered, Ophn1 deficiency produced an enhancement of the density of spines 21 dpi. At this stage of maturation, most of the dendritic protrusions had an immature appearance. Interestingly, the overall density of dendritic spines increases more robustly from 21 to 28 dpi in WT than in KO animals. One possible explanation is that newly generated Ophn1 deficient neurons display an initial overproduction of immature spines

that is later followed by a reduced addition, ultimately resulting in a lower density of mature protrusions. These data are consistent with previous findings reporting a key role for Ophn1 in dendritic spine morphogenesis. Indeed, inactivation of Ophn1 function increases the density and proportion of immature spines in neuronal cultures and in vivo (Khelifaoui et al., 2007; Govak et al., 2004; Redolfi et al., 2016). In the Ophn1 KO mouse, mature dendritic spines are reduced and this is associated with impairments of spatial learning and alterations of social behavior (Khelifaoui et al., 2007).

#### 4.3. Restoration of adult hippocampal neurogenesis by inhibition of ROCK/PKA signaling

One of the best known downstream target of OPHN1 is the Rho kinase (ROCK) protein. Ophn1 loss of function results in persistent



**Fig. 7.** Fasudil treatment per se does not affect adult hippocampal neurogenesis in WT mice. A) Stereological counts in WT animals show that the percentage of Dcx-positive cells is unaffected upon 7 weeks of fasudil treatment (WT,  $100 \pm 3.8$ ,  $n = 5$ ; WT + fasudil,  $94.7 \pm 8.4$ ,  $n = 3$ ; two-tailed *t*-test,  $P = 0.539$ ). Data are expressed as percentage of the control group (WT). B) Quantification of the total number of mature newborn neurons (50 dpi) shows that vehicle- and fasudil-treated WT mice display a similar range of BrdU-positive cells (WT,  $662 \pm 38$ ,  $n = 11$ ; WT + fasudil,  $657 \pm 72$ ,  $n = 4$ ; two-tailed *t*-test,  $P = 0.946$ ). Histograms represent mean  $\pm$  SEM.

activation of RhoGTPases and consequent hyperstimulation of the RhoA/ROCK pathway (Govek et al., 2004; Khelifaoui et al., 2009). In addition, lack of OPHN1 leads to high levels of PKA activity (Khelifaoui et al., 2014). Presynaptic PKA-dependent LTP, and hippocampus- and amygdala-related learning (in the fear conditioning test) are compromised in Ophn1 KO animals. Some of the electrophysiological and behavioral deficits of Ophn1 KO mice can be rescued by treatment with ROCK/PKA inhibitor fasudil (Khelifaoui et al., 2014; Meziane et al., 2016), a drug already approved for clinical use in Japan and China. Based on these premises, we tested whether chronic fasudil treatment is effective in restoring adult hippocampal neurogenesis. We found that fasudil normalized the number of Dcx-positive early neuronal cells, indicating that pharmacological ROCK/PKA inhibition may overcome the deficits caused by Ophn1 mutation at an early stage of neuronal maturation. At the morphological level, fasudil corrected the dendritic spine deficit at 21 dpi. In contrast to the dendritic effects, fasudil administration was unable to restore a normal proportion of newborn neurons extending their axon towards CA3.

Altogether, the morphological analysis reveals two key processes impacted by Ophn1 deficiency: axonal extension and dendritic spine morphogenesis, both of which are critical for proper integration of newborn neurons (i.e. for establishing afferent and efferent connectivity). The fasudil rescue experiments indicate that these two processes proceed via at least partly independent pathways: dendritic spine density can be restored by downregulating abnormally high ROCK/PKA activity in Ophn1 KO mice, while the same approach is not effective on aberrant axonogenesis. Previous studies have shown that Ophn1 can act as a GAP for all three Rho GTPases: RhoA, Rac1 and Cdc42 (Govek et al., 2004). Thus, the unchecked activation of all these signaling cascades may potentially explain the pathological consequences of OPHN1 deficiency. Since the expression of constitutively active Rac1 in mouse Purkinje neurons inhibits axonal outgrowth (Luo et al., 1996), an excessive Rac1-mediated signaling (not counteracted by fasudil treatment, that targets only the pathway downstream of RhoA) may explain the persistence of impaired axon formation/extension in Ophn1-deficient newborn neurons treated with fasudil.

Importantly, we found the total number of mature dentate granule neurons at 50 dpi was significantly increased by fasudil administration in Ophn1-deficient mice. Still, we observed a 20% reduction in the number of mature neurons as compared to WT, healthy mice (see Fig. 6E).

These data indicate that only a proportion of the newly generated cells can be rescued by fasudil administration, and this is in line with the persistence of axonless newborn cells in fasudil-treated KO mice (see Fig. 6B). Obviously, adult-generated neurons unable to connect with their natural targets in the CA3 region would be eliminated from the network (despite the rescue of the dendritic spine phenotype), resulting in lower numbers of mature neurons at 50 dpi.

Overall, the present findings allow a precise dissection of alterations of neuronal maturation in XLID, and suggest a line along which possible therapeutic strategies for XLID may be developed. In particular, early fasudil treatment (i.e. in embryonic or early postnatal period) might be envisaged to correct neuronal maturation and prevent wiring deficits and consequent network dysfunction in Ophn1-dependent XLID. Given the complex pathology associated with XLID, it is likely that a combination of drugs aimed at downregulating Rho GTPase activity and administered during specific “critical periods” of brain development may be needed to successfully treat this pathology.

Supplementary data to this article can be found online at <http://dx.doi.org/10.1016/j.nbd.2017.01.003>.

### Competing interests

The authors declare no competing interests.

### Authors' contributions

Conceptualization: M.A., C.S., B.V., M. Can., M. Cal.; Formal analysis and investigation: M.A., C.S., B.V., S.A., I.B.; Writing - original draft preparation: M.A., M. Cal.; Writing - review and editing: all authors; Funding acquisition: M. Can., M. Cal.; Resources: P.B.

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