Development

Role of NMDA Receptors in Adult Neurogenesis and Normal Development of the Dentate Gyrus

Ingrid Åmellem,1,2, Glen Yovianto,1,∗ Hai Tarng Chong,1,∗ Rajeevkumar Raveendran Nair,2,∗ Vanja Cnops,1,2 Ayesha Thanawalla,1 and Ayumu Tashiro (田代 歩)1,2

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1School of Biological Sciences, Nanyang Technological University, Singapore 308232 and 2Kavli Institute for Systems Neuroscience, Norwegian University of Science and Technology, Trondheim 7491, Norway

Abstract

The NMDA receptors are a type of glutamate receptors, which is involved in neuronal function, plasticity and development in the mammalian brain. However, how the NMDA receptors contribute to adult neurogenesis and development of the dentate gyrus is unclear. In this study, we investigate this question by examining a region-specific knock-out mouse line that lacks the NR1 gene, which encodes the essential subunit of the NMDA receptors, in granule cells of the dentate gyrus (DG-NR1KO mice). We found that the survival of newly-generated granule cells, cell proliferation and the size of the granule cell layer are significantly reduced in the dorsal dentate gyrus of adult DG-NR1KO mice. Our results also show a significant reduction in the number of immature neurons and in the volume of the granule cell layer, starting from three weeks of postnatal age. DG-NR1KO mice also showed impairment in the expression of an immediate early gene, Arc, and behavior during the novelty-suppressed feeding and open field test. These results suggest that the NMDA receptors in granule cells have a role in adult neurogenesis in the adult brain and contributes to the normal development of the dentate gyrus.

Key words: granule cell; hippocampus; neuronal survival; novelty suppressed feeding test; pattern separation

Significance Statement

A type of glutamate receptors, the NMDA receptors, are known to be generally important in brain development. However, a previous study, which genetically ablated the NMDA receptors in the developing dentate gyrus, did not detect malformation of the dentate gyrus. Here, we performed more detailed analyses of the knock-out mice and showed that the extent of neurogenesis and the size of the neuronal layer are reduced in the postnatal and adult dentate gyrus. We found that the knock-out mice exhibit behavioral abnormality during adulthood. These results indicate that the NMDA receptor is essential for normal development of the dentate gyrus and that a malfunction of the NMDA receptors during the postnatal period could lead to life-long disturbance in brain functions.

Introduction

A type of ionotropic glutamate receptors, the NMDA receptors, are involved in postnatal brain development (Haberny et al., 2002; Hansen et al., 2004; Ewald and Cline, 2009). It has been shown that the NMDA receptors support the survival of developing neurons (Ikonomidou et al., 1999) and regulate the formation of neuronal networks in several brain areas (Bear et al., 1987; Constantine-Paton et al., 1990), including the dentate gyrus (Gould et al., 1994). The majority of granule cells, a principal cell type in the dentate gyrus, are generated postnatally, predominantly during the first few weeks after birth in rodents (Cameron et al., 1993) and the structural

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development of the dentate gyrus occurs mainly during the first month of the postnatal period in rodents (Li and Pleasure, 2005). Early life experience, such as stress, has been found to regulate the expression of the NMDA receptors in the hippocampus (Roceri et al., 2002; Pickering et al., 2006) and to reduce the number of granule cells (Oreland et al., 2010).

The involvement of the NMDA receptors in neuronal survival and circuit formation continues in newly-generated neurons in the adult brain. The survival of newborn neurons in the adult brain is regulated by neuronal activity in the dentate gyrus, as evidenced by the finding that induction of NMDA receptor-dependent long-term potentiation facilitates the survival of newborn neurons (Bruel-Jungerman et al., 2006). A study using a cell-specific gene knock-out technique (Tashiro et al., 2006a) demonstrated that the NMDA receptors positively regulate the survival of immature granule cells in a cell-autonomous manner, within a few weeks of neuronal birth (Tashiro et al., 2006b). However, once new granule cells pass a certain maturational stage, the NMDA receptors are not required for their survival but continue to be required for the regulation of dendritic spine morphology and synaptic plasticity (Ge et al., 2007; Mu et al., 2015).

A previous study demonstrated that the NMDA receptors in granule cells of the dentate gyrus play a role in a context discrimination task using a region-specific gene knock-out mouse line. This mouse line lacks the NR1 gene, which encodes an essential subunit of the NMDA receptors, specifically in granule cells of the dentate gyrus (McHugh et al., 2007). The gene knock-out in granule cells starts between 1.5 and 4 weeks after the birth of mice. The study found that the gross structure of the dentate gyrus is intact but did not examine how adult neurogenesis is affected in the mouse line.

In this study, we have two aims. The first aim is to determine a role of NMDA receptors in adult neurogenesis of the dentate gyrus. We used the same knock-out mouse line created by McHugh et al. (2007; DG-NR1KO mice) and examined neurogenesis in the adult dentate gyrus. We used the same knock-out mouse line. This mouse line lacks the NR1 gene, which encodes an essential subunit of the NMDA receptors in granule cells of the dentate gyrus (McHugh et al., 2007). The gene knock-out in granule cells starts between 1.5 and 4 weeks after the birth of mice. The study found that the gross structure of the dentate gyrus is intact but did not examine how adult neurogenesis is affected in the mouse line.

In this study, we have two aims. The first aim is to determine a role of NMDA receptors in adult neurogenesis of the dentate gyrus. We used the same knock-out mouse line created by McHugh et al. (2007; DG-NR1KO mice) and examined neurogenesis in the adult dentate gyrus. The second aim is to reveal the role of NMDA receptors in normal formation of the dentate gyrus. As described above, the NMDA receptors have been shown to contribute to the development of granule cells in the postnatal and adult dentate gyrus. However, McHugh et al. (2007) did not detect that the NR1 gene knock-out starting from the postnatal development affects the development of the dentate gyrus. To reconcile this discrepancy in the contribution of the NMDA receptors to the development of the dentate gyrus, we examined the structure and neurogenesis in the dentate gyrus of the postnatal and adult brain.

Materials and Methods

Mice

All animal procedures were approved by the Norwegian Animal Research Authority and/or Institutional animal care and use committee at A*STAR and Nanyang Technological University. We re-generated DG-NR1KO (Pomc-cre+/−, fNR1+/+) mice using the same transgenic mouse lines used by McHugh et al. (2007), B6.FVB-Tg (Pomc-cre)1Stl/J and B6.129S4-Grin1Ptm2StlP/J (fNR1; both from The Jackson Laboratory). Cre recombinase is under transcriptional control by the proopiomelanocortin (Pomc) promoter, which initiates gene expression during a short time window in the maturation of newborn granule cells of the dentate gyrus (Overstreet et al., 2004). Specificity of Cre expression in Pomc-cre+/− mice was equivalent with McHugh et al. (2007) when we evaluated it by crossing them with a cre-dependent reporter line (B6;129S-Gt(Rosa)26Sortm32(CAG-COP4*H134R/EYFP)Hze/J, from The Jackson Laboratory; Fig. 1). Cre recombines a floxed part of the NR1 gene (fNR1) to ablate the functional NR1 gene. These mice were bred with Pomc-cre−/−, fNR1+/+ mice, and their Cre−/+ offspring were used as DG-NR1KO mice and the Cre-negative littermates were used as controls. Both female and male mice were used except Figure 6, where we used female mice only. For Figures 3, 4, 8G–O, we used young adult mice (42–51 d old at the start of experiments). For Figure 5, we used two-, three-, or four-week-old mice. For Figures 6, 7, 8A–F, we used young adult to mature adult mice (42–83 d old). Mice were housed with ad libitum access to food and water under 12/12 h light/dark cycle conditions, unless otherwise specified below.

Genotyping

The genotypes of mice were determined by PCR analyses. The primer sequences for Cre are 5′-CCGTGACCTGCA AACAGGCTCTA-3′ and 5′-GATTACATTCTCCACCG TCAGT-3′ and those for the fNR1 allele are 5′-CTTGGGTG GAGAGGCTATTC-3′ and 5′-AGGTGAGATGACAGAGA GC-3′. As previously described (McHugh et al., 2007), germ-line recombination occurs in this mouse line. To detect recombined alleles in control mice, nested PCR was performed. The primer sequences for the initial amplification were 5′-AATGCTGAGGTGGTAGGA-3′ and 5′-AGGTGAGA TGACAGAGATC-3′. The primer sequences for the second step are 5′-GCTACAAGGCAAAGATACAAGACC-3′ and 5′- ACCGTGCAGGAATTTCCGGATCAT-3′. When a recombined allele was detected in control mice, they were excluded from the experiments.

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Correspondence should be addressed to Ayumu Tashiro at atashiro@ntu.edu.sg.

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5-Bromo-2'-deoxyuridine (BrdU) injections
BrdU (catalog #B5002, Sigma) was dissolved in 0.9% saline at a concentration of 10 mg/ml. After filter sterilization (pore size: 0.2 μm), mice were intraperitoneally injected with a dose of 100 μg/g body weight per day.

Enriched environment
Female mice were injected with BrdU on days −1 and 0. Between days 7 and 14, they were either housed in an enriched cage (48 x 48 x 48 cm, eight mice per cage) which contained eight pieces of plastic blocks, one cardboard shelter, one plastic shelter, four tunnels, two sheets of paper for nesting, four pieces of small wood, and one running wheel (Fig. 2), or they stayed in the standard housing (26.7 x 20.7 x 14 cm, four mice per cage) with bedding only. On days 0–7 and days 14–28, all the mice were in the standard housing. The mice were perfusion-fixed on day 28 as described below.

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Novel environment exploration
The mice were exposed to a novel environment, which was a 32 x 28 x 28 cm open field with plastic walls with black and white stripes. They were allowed to explore the novel environment for 10 min. Then, the mice were moved back to their home cages in the experimental room. After 90 min, they were perfusion-fixated as described below.

Novelty suppressed feeding test
The mice were food deprived for 24 h before the test. A food pellet was placed on a white paper platform (12.5 cm) at the center of a square open field (60 x 60 cm, with white colored walls), which the mice had never been exposed to before. The room was lit with regular room lighting (around 700 lux at the center of the box) and the floor of the open field was covered with the same type of bedding as used in their home cages. At the start of a test, a mouse was placed at a corner of the open field. When the mouse started eating the food pellet or 10 min passed after the test started, the test was terminated. The test was recorded by an overhead camera and analyzed using EthoVision XT13 software (Noldus Information Technology BV). The area in the open field was divided into 5 x 5 square bins. A total of 16 outer bins were defined as the peripheral zone. The area on the white paper was defined as the center zone. The remaining area was defined as the inner zone.

Open field test
A square open field (60 x 60 cm, with white-colored walls and floor) was placed in a room that was lit with ceiling lights (~700 lux at the center of the box). At the start of a test, a mouse was placed at a corner of the open field and allowed to explore it. After 10 min, the test was terminated. The mice were removed from the open field and fecal boli were counted. The test was recorded by an overhead camera and analyzed using EthoVision XT13 software. The area in the open field was divided into 5 x 5 square bins. A total of 16 outer bins were defined as the peripheral zone. The remaining area was defined as the inner zone.

Elevated plus maze
A plus shaped maze was elevated 60 cm above the ground. Two arms opposite to each other were surrounded with 15-cm-high walls at the side and end of the
arm. These are called “closed arm” while the other two arms without walls were called “open arm.” The lengths of closed and open arms were 35 and 36.5 cm, respectively. The width of each arm was 5 cm. The room was lit dimly (~100 lux at the center of the maze). At the start of a test, a mouse was placed at the center of the maze, with its face facing one of the closed arms. The mouse was allowed to explore the maze. After 10 min, the test was terminated. The test was recorded by an overhead camera and analyzed using Anymaze 6.34 (Stoelting Co). The entry to and exit from an arm were registered when 40% of body entered and 35% exited the arm, respectively.

**Histology**

Perfusion fixation, preparation of brain sections (35- or 40-µm thickness) and immunostaining were conducted as previously described (Tashiro et al., 2015). We prepared coronal brain sections containing the rostral two thirds of the dentate gyrus; therefore, the analysis presented in this study is mostly from the dorsal, but not ventral, part of the dentate gyrus. The primary antibodies used were goat anti-DCX (1:500 dilution, catalog #sc-8066, Santa Cruz Biotechnology) and mouse anti-activity-regulated cytoskeleton-associated protein (Arc; 1:100, catalog #sc-17839, Santa Cruz Biotechnology), rat anti-GFP (1:1000, catalog #04404-84, Nacalai Tesque), rat anti-BrdU (1:400, catalog #OBT0030G, AbD Serotec), rabbit anti-Prospéro homeobox protein 1 (Prox1; 1:1000, catalog #PRB-238C, Covance), goat anti-Prox1 (1:40, catalog #AF 2727, R&D Systems), and rabbit anti-Ki67 (1:500, catalog #NCL-Ki67p, Leica Biosystems). The secondary antibodies used were donkey anti-rat-Alexa Fluor 488 (catalog #712-545-153), anti-rabbit-Cy3 (catalog #712-165-152), anti-mouse-DyLight549 (catalog #715-505-151), anti-goat-Alexa Fluor 488 (catalog #705-545-147), and anti-goat-Alexa Fluor 647 (catalog #705-605-147). All secondary antibodies were purchased from Jackson ImmunoResearch and used at 1:600 dilutions. Nuclear staining with 4′,6-diamidino-2-phenylindole-dihydrochloride (DAPI; 5 µg/ml; Merck) was done on some of the sections. The brain sections were mounted on glass slides with mounting medium containing PVA-DABCO.

**Microscopy**

Axio Scope A1 or Axio Imager M1 microscopes (Zeiss, Germany) were used to count BrdU-positive (+), Ki67+, and DCX+ cells and image the granule cell layer of the dentate gyrus for size measurements. Counting was done using either 20× or 40× objective lens from the granule cell layer and subgranular zone in both hemispheres in six brain sections per mouse for BrdU (every 6th section; Figs. 3, 6), three brain sections for Ki67 and DCX (every 12th section; Fig. 3) in adult mice, six brain sections from the anterior part of the dentate gyrus for Ki67 and granule cell layer volume measurements in two- to four-week-old mice (every 6th section; Fig. 5), and two brain sections from similar anterior part for DCX in two- to four-week-old mice (Fig. 5). The DAPI or Prox1 fluorescence were used to image the area of the granule cell layer using a 10× objective lens. The volume of the granule cell layer was calculated by multiplying the area by the thickness of the sections, which was 35 or 40 µm. The volume values shown in Figures 4, 5 are summed volume from three and six analyzed sections, respectively.

Confocal imaging was conducted using LSM710 confocal microscope (Zeiss, Germany) equipped with 488-, 543-, and 633-nm laser lines and the ZEN image-acquisition software. For co-localization analysis (BrdU+/Prox1+ and Arc+/Prox1+), a 40× objective lens with oil (NA 1.3) was used. Three images from the granule cell layer from random positions in three different hemispheres/mice were used for BrdU, Prox1 colocalization analysis in Figure 3 and four images from the granule cell layer from random positions in four different mice were used for BrdU, Prox1 colocalization analysis in Figure 6, and Arc, Prox1 colocalization analysis in Figure 7. The density of BrdU+/Prox1+ and BrdU–/Prox1+ were estimated by multiplying the density of BrdU+ cells counted under Axio Scope A1 or Axio Imager M1 microscopes by % of BrdU+ cells which are Prox1+ and Prox1–, measured with the confocal microscope, respectively. For the measurement of the distribution of Arc+/Prox1+ cells in the granule cell layer, only the area that showed the entire depth of the granule cell layer was used from the same confocal images as used for the Arc/Prox1 co-localization counting. Measurements of the granule cell layer area from epifluorescence images and counting of cells from confocal images were performed using ImageJ software (National Institutes of Health).

The section was selected in a way that their anteroposterior level is comparable between mice used in individual analyses. The density was calculated by dividing the number of cells by the volume of the granule cell layer. Our analyses focused on the area covering approximately anterior two thirds of the dentate gyrus, so that the results were from the dorsal dentate gyrus.

**Statistics**

Statistical analyses were done using SPSS Statistics (IBM Corp.) and Statistica (StatSoft) software. For independent-sample t tests, Levene’s test for equal variance was performed, and depending on the results, a t test with (Student’s t test) or without (Welch’s t test) equal variance assumed was done. All data are presented as mean ± SEM. When we detected significant interaction in two-way ANOVAs or three-factor interaction in three-way ANOVAs, we performed Tukey’s HSD tests. When we detected significant interaction of two factors but not of three factors, we performed simple effects tests using two-way ANOVAs.

**Results**

**Reduced neurogenesis and granule cell layer size in the dentate gyrus of adult DG-NR1KO mice**

To evaluate neurogenesis in the dentate gyrus of adult control and DG-NR1KO mice, we examined the density of cells expressing Ki67 (a marker for proliferating cells) and DCX (a marker for neuronal progenitor cells and immature neurons). We found that the density of Ki67+ cells is
significantly reduced in the subgranular zone of DG-NR1KO mice compared with control (p = 0.023, \( t_{(11)} = 2.641 \), n = 6 control and 7 DG-NR1KO mice, independent-sample \( t \) test, two-tailed; Fig. 3A, B), indicating a reduction in proliferation in DG-NR1KO mice. We also found a significant reduction in the density of DCX\(^1\) cells in DG-NR1KO mice compared with control (p = 5.6 \( \times \) \( 10^{-7} \), \( t_{(11)} = -10.286 \), n = 6 control and 7 DG-NR1KO mice, independent-sample \( t \) test, two-tailed; Fig. 3C, D), indicating a decrease of neuronal progenitor cells and/or immature neurons in the dentate gyrus.

Reduction of DCX\(^1\) cells may be exclusively because of reduced proliferation or be partly because of reduced survival of newborn neurons. To examine the latter possibility, we injected BrdU into adult DG-NR1KO and control mice and measured the density of BrdU\(^+\)/Prox1\(^+\) cells 7 or 28 d after injection (Fig. 3E). The granule cell marker Prox1 was used to determine the identity of BrdU\(^+\) cells as granule cells. Seven days after BrdU injection, the density of BrdU\(^+\) cells was significantly lower in DG-NR1KO mice than in controls (p = 0.0099, \( t_{(14)} = 2.983 \), n = 7 control and 9 DG-NR1KO mice, respectively, independent-sample \( t \) test, two-tailed; Fig. 3F). Proportion of BrdU\(^+\) cells expressing Prox1 was not significantly different between control and DG-NR1KO mice (p = 0.657, \( t_{(14)} = 0.454 \), n = 7 and 9 mice, respectively, independent-sample \( t \) test, two-tailed; Fig. 3G), suggesting that neuronal differentiation was not affected. The density of BrdU\(^+\)/Prox1\(^+\) cells showed a tendency of reduction in DG-NR1KO mice, but it was not significant (p = 0.070, \( t_{(8.385)} = 2.079 \), n = 7 control and 9 DG-NR1KO mice, respectively, independent-sample \( t \) test, two-tailed; Fig. 3H); 28 d after BrdU injection, both BrdU\(^+\) and BrdU\(^+\)/Prox1\(^+\) cell densities were significantly lower in DG-NR1KO mice than in controls (BrdU\(^+\): p = 3.6 \( \times \) \( 10^{-4} \), \( t_{(7.527)} = 6.116P \); BrdU\(^+\)/Prox1\(^+\): p = 3.3 \( \times \) \( 10^{-4} \), \( t_{(7.367)} = 6.298 \), n = 8 control and 9

**Figure 3.** Cell proliferation and the survival of new neurons were reduced in the dentate gyrus of adult DG-NR1KO mice. A, Representative images visualizing Ki67\(^+\) (green) and DAPI-labeled (red) cells in the dentate gyrus of adult control and DG-NR1KO mice. Scale bar: 75 \( \mu \)m. B, Density of Ki67\(^+\) cells in the subgranular zone. C, Representative images showing DCX\(^+\) (green) and DAPI-labeled (red) cells in the dentate gyrus of adult control and DG-NR1KO mice. The images were maximum intensity projections of confocal Z stacks and formed by joining two overlapping images containing adjacent areas. Scale bar: 25 \( \mu \)m. D, Density of DCX\(^+\) cells. E, Representative images showing BrdU\(^+\) (green) and Prox1\(^+\) (red) cells in adult control and DG-NR1KO mice on 7 and 28 d after BrdU injections. Scale bar: 75 \( \mu \)m. F, Density of BrdU\(^+\) cells in the granule cell layer and subgranular zone. G, Proportion of BrdU\(^+\) cells expressing Prox1. H, Density of BrdU\(^+\)/Prox1\(^+\) cells in the granule cell layer. I, Density of BrdU\(^+\)/Prox1\(^+\) cells in the granule cell layer. J, Survival rate of BrdU\(^+\)/Prox1\(^+\) cells from 7 to 28 dpi. Density at 28 d after BrdU injection was divided by mean values of density at 7 d; *p < 0.05, **p < 0.01, ***p < 0.005, independent-sample \( t \) test, two-tailed.
DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 3F, H). Proportion of BrdU+ cells expressing Prox1 was significantly lower in DG-NR1KO mice than in controls ($p = 5.3 \times 10^{-5}$, $t(9.687) = 6.834$, $n = 8$ control and 9 DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 3G). Therefore, a proportion of BrdU+ cells that differentiated to neurons were lower in DG-NR1KO mice at 28 d. The density of other cell types (BrdU+/Prox1−) was

![Figure 4.](image)

**Figure 4.** Reduced size of the granule cell layer in adult DG-NR1KO mice. A, Representative images showing the granule cell layer visualized by DAPI staining in adult control and DG-NR1KO mice. Scale bar: 150 μm. B, Volume of the granule cell layer (GCL) in adult control and DG-NR1KO mice. Summed volume from three analyzed sections is presented. C, Thickness of the upper and lower blade of granule cell layer in adult control and DG-NR1KO mice. D, Length of the subgranular zone (SGZ) in adult control and DG-NR1KO mice; *$p < 0.05$, **$p < 0.005$, independent-sample t test, two-tailed.

DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 3F, H). Proportion of BrdU+ cells expressing Prox1 was significantly lower in DG-NR1KO mice than in controls ($p = 5.3 \times 10^{-5}$, $t(9.687) = 6.834$, $n = 8$ control and 9 DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 3G). Therefore, a proportion of BrdU+ cells that differentiated to neurons were lower in DG-NR1KO mice at 28 d. The density of other cell types (BrdU+/Prox1−) was

![Figure 5.](image)

**Figure 5.** Neurogenesis was impaired in the dentate gyrus of postnatal, developing DG-NR1KO mice. A, Representative images showing Ki67+ (green) and DAPI-labeled (red) cells in the dentate gyrus of four-week-old control and DG-NR1KO mice. Scale bar: 60 μm. B, Density of Ki67+ cells in the subgranular zone in two-, three-, and four-week-old control and DG-NR1KO mice. C, Representative images showing DCX+ (green) and DAPI-labeled (red) cells in the dentate gyrus of two-, three-, and four-week-old control and DG-NR1KO mice. The images were maximum intensity projections of confocal Z stacks. Scale bar: 25 μm. D, Density of DCX+ cells in the granule cell layer of two-, three-, and four-week-old control and DG-NR1KO mice. E, Representative images showing the granule cell layer visualized by DAPI staining in two-, three-, and four-week-old control and DG-NR1KO mice. F, Volume of the granule cell layer (GCL) in two-, three-, and four-week-old control and DG-NR1KO mice. Summed volume from six analyzed sections is presented; *$p < 0.05$, **$p < 0.01$, ***$p < 0.005$, independent-sample t test, two-tailed.
Figure 6. Exposure to an enriched environment increases the survival of newborn neurons in DG-NR1KO mice. A, Experimental timeline. B, Representative images showing BrdU+ (green) and Prox1+ (red) cells in the dentate gyrus of adult control and DG-NR1KO mice with or without exposure to the enriched environment. Scale bar: 60 μm. C, Density of BrdU+ cells in the dentate gyrus of adult control and DG-NR1KO mice with or without exposure to the enriched environment. D, Proportion of BrdU+ cells expressing Prox1. E, Density of BrdU+/Prox1+ cells in the granule cell layer of adult control and DG-NR1KO mice with or without exposure to the enriched environment. F, Normalized density of BrdU+/Prox1+ cells in the granule cell layer of adult control and DG-NR1KO mice with or without exposure to the enriched environment. Density of each mouse was divided by a mean value of mice without exposure to the enriched environment in the same genotype; *p < 0.05, ***p < 0.005, Tukey’s HSD test (performed because of significant genotype × housing interaction in two-way ANOVA); ###p < 0.005, the main effect of genotype or housing (without significant interaction).

Figure 7. Impairment in novelty-induced Arc gene expression in granule cells of DG-NR1KO mice. A, Representative images showing Arc+ (green) and Prox1+ (red) expressing cells in the dentate gyrus of adult control and DG-NR1KO mice, which stayed in their home cage or were exposed to a novel environment. Scale bar: 75 μm. B, Density of Arc+/Prox1+ cells in the granule cell layer of adult control and DG-NR1KO mice with or without exposure to the novel environment. C, Normalized density of Arc+/Prox1+ cells in the granule cell layer of adult control and DG-NR1KO mice with or without exposure to the novel environment. Density of each mouse was divided by a mean value of mice without exposure to the novel environment in the same genotype. D, Depth categories divided along the thickness of the granule cell layer. The granule cell layer (GCL) was divided into five 20%-thickness segments, with 0% starting at the border to the hilus and 100% at the border to the molecular layer (ML). E, Density distribution of Arc+/Prox1+ cells along depth; *p < 0.05, **p < 0.01, ***p < 0.005, Tukey’s HSD test.
not significantly different between control and DG-NR1KO mice at either time point (7 d: \( p = 0.366, t_{(14)} = 0.934, n = 7 \) and 9 mice, respectively; 28 d: \( p = 0.367, t_{(15)} = -0.930, n = 8 \) and 9 mice, respectively, independent-sample t tests, two-tailed; Fig. 3I). By comparing the density at 7 and 28 d, we calculated the survival rate of new neurons from 7 to 28 d. The survival rate was significantly reduced in DG-NR1KO mice compared with control mice (\( p = 5.0 \times 10^{-4}, t_{(7,68)} = 5.757, n = 8 \) control and 9 DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 3J).

We also noted that the size of the granule cell layer was smaller in DG-NR1KO mice (Fig. 4A). The volume of the granule cell layer (\( p = 4.2 \times 10^{-4}, t_{(11)} = -4.977, n = 6 \) control and 7 DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 4B), the thickness of the upper and lower blade (\( p = 1.3 \times 10^{-4} \) and 0.0014, \( t_{(11)} = -5.760 \) and \(-4.243\), respectively, \( n = 6 \) control and 7 DG-NR1KO mice, independent-sample t tests, two-tailed; Fig. 4C), and the length of the subgranular zone (\( p = 0.016, t_{(11)} = -2.857, n = 6 \) control and 7 DG-NR1KO mice, independent-sample t test, two-tailed; Fig. 4D) were reduced in DG-NR1KO mice compared with control.

**Reduced neurogenesis and granule cell layer size in juvenile DG-NR1KO mice**

It is known that the majority of granule cells in the dentate gyrus are generated during postnatal development (Li and Pleasure, 2005). Thus, the reduced size of the granule cell layer may be caused by reduced neurogenesis during the postnatal development. We therefore examined neurogenesis in postnatal, developing mice. We analyzed Ki67+ cells in the subgranular zone of two-, three-, and four-week-old mice (Fig. 5A) and found that the densities of Ki67+ cells were not significantly different between control and DG-NR1KO mice (two weeks: \( p = 0.482, t_{(11)} = -0.728, n = 5 \) and 8 mice, respectively, three weeks: \( p = 0.458, t_{(13)} = 0.764, n = 7 \) and 8 mice, respectively, four weeks: \( p = 0.610, t_{(13)} = 0.522, n = 7 \) and 8 mice, respectively, independent-sample t tests, two-tailed; Fig. 5B).

The densities of DCX+ cells were not significantly different in the dentate gyrus between control and DG-NR1KO mice at the age of two weeks (\( p = 0.981, t_{(10)} = 0.024, n = 5 \) and 7 mice, respectively, independent-sample t test, two-tailed; Fig. 5C,D). But in the three- and four-week-old mice the densities were significantly reduced in the DG-NR1KO mice (three weeks: \( p = 0.0075, t_{(13)} = 3.159, n = 7 \) control and 8 DG-NR1KO mice, four weeks: \( p = 0.0048, t_{(13)} = 3.394, n = 7 \) control and 8 DG-NR1KO mice, independent-sample t tests, two-tailed). Thus, while neurogenesis seems to occur normally initially at the postnatal age of two weeks in DG-NR1KO mice, the number of immature neurons/neuronal progenitor cells starts decreasing between two and three weeks of age. This reduction continues to adulthood in DG-NR1KO mice as shown in Figure 3.

The same developmental pattern of effects was found for the volume of the granule cell layer (Fig. 5E, F), with no significant difference in two-week-old mice (\( p = 0.500, t_{(4,515)} = 0.733, n = 5 \) control and 8 DG-NR1KO mice, independent-sample t tests, two-tailed), but significant decreases in three- and four-week-old DG-NR1KO mice compared with control (three weeks: \( p = 0.0070, t_{(13)} = 3.197, n = 7 \) control and 8 DG-NR1KO mice, four weeks: \( p = 0.014, t_{(13)} = 2.852, n = 7 \) control and 8 DG-NR1KO mice, independent-sample t tests, two-tailed).

**Enriched environment exposure increased the survival of newborn neurons in DG-NR1KO mice**

Previously it has been shown that exposure to an enriched environment increases the survival of newborn neurons in the dentate gyrus of adult rodents (Kempermann et al., 1997; Tashiro et al., 2007). Exposure to an enriched environment for one week during the second week after neuronal birth has been found to be the most effective time point for this increased survival (Tashiro et al., 2007). To examine whether the NMDA receptors in the dentate gyrus are involved in the effect of enriched environment on the survival of newborn neurons, we injected BrdU on days –1 and 0 and then exposed DG-NR1KO and control mice to an enriched environment or a standard cage with bedding only from day 7 to day 14 (Fig. 6A). The enriched environment contained several toys, shelters, tunnels and a running wheel (Fig. 2). Except on days 7–14, the mice were housed in the standard cage. On day 28, we euthanized the mice to quantify the number of BrdU+ and BrdU+/Prox1+ cells (\( n = 7 \) control-Enriched, 7 control-Standard, 8 DG-NR1KO-Enriched, and 8 DG-NR1KO-Standard mice; Fig. 6B). DG-NR1KO mice had significantly lower BrdU+ cell density than control mice while enrichment increases the density similarly in both genotypes (housing: \( p = 0.002, F_{(1,26)} = 11.362, \) genotype: \( p = 4 \times 10^{-6}, F_{(1,26)} = 33.282, \) housing \times genotype: \( p = 0.072, F_{(1,26)} = 3.513, \) two-way ANOVA; Fig. 6C). Proportion of BrdU+ cells expressing Prox1 was significantly lower in DG-NR1KO mice than control mice similarly in standard and enriched cages while enrichment did not affect the proportion (housing: \( p = 0.344, F_{(1,26)} = 0.928, \) genotype: \( p = 4.7 \times 10^{-4}, F_{(1,26)} = 16.004, \) housing \times genotype: \( p = 0.967, F_{(1,26)} = 0.002, \) two-way ANOVA; Fig. 6D). For BrdU+/Prox1+ cell density (Fig. 6E), two-way ANOVA detected significance in the main effect of housing conditions (\( p = 0.002, F_{(1,26)} = 11.425, \) and genotype (\( p = 1 \times 10^{-6}, F_{(1,26)} = 38.776, \) and interaction between them (\( p = 0.046, F_{(1,26)} = 4.410, \)) Exposure to the enriched environment increased the densities in control but not in DG-NR1KO mice (\( p = 0.0047 \) and 0.785, respectively, \( df = 26, \) Tukey’s HSD test) while the densities were lower in DG-NR1KO mice than controls regardless of whether housed in enriched or standard environments (\( p = 1.8 \times 10^{-4}, \) and \( p = 0.034, \) respectively, \( df = 26, \) Tukey’s HSD test, two-tailed). However, the percentage increase after exposure to the enriched environment was not significantly different between control and DG-NR1KO mice (housing: \( p = 0.002, F_{(1,26)} = 12.336, \) genotype: \( p = 0.530, F_{(1,26)} = 0.404, \) housing \times genotype: \( p = 0.589, F_{(1,26)} = 0.299, \) two-way ANOVA; Fig. 6F). Thus, despite overall reduction of neurogenesis in DG-
NR1KO mice, exposure to the enriched environment still increased the survival of new neurons in DG-NR1KO mice.

Abnormalities in novelty-induced Arc gene expression of granule cells in the dentate gyrus of DG-NR1KO mice

Immediate early genes are a group of genes that are expressed in response to neuronal activation. Arc is one of such genes and known to be expressed in rodents after novel environment exploration and learning (Steward et al., 1998; Guzowski et al., 1999, 2001; Czerniawski et al., 2011). The novelty-induced expression of Arc is blocked with an NMDA receptor antagonist (Czerniawski et al., 2011). We examined whether novelty-induced Arc expression requires the NMDA receptors in the dentate gyrus.

For this purpose, we exposed control and DG-NR1KO mice to a novel environment for 10 min and perfusion-fixed them 90 min later. The non-exposed control and DG-NR1KO mice were perfusion-fixed directly from home cages. We then immunostained the brain sections with anti-Arc and -Prox1 antibodies to detect Arc expressing granule cells and quantified the density of Arc+/Prox1+ cells (n = 9 novel environment control, 6 home-cage control, 10 novel environment DG-NR1KO, 10 home-cage DG-NR1KO mice; Fig. 7A, B). Two-way ANOVA detected significance in the main effect of exposure (home cage vs novel environment, \( p = 5.8 \times 10^{-8} \), \( F_{(1,31)} = 63.087 \) and genotype (\( p = 9.3 \times 10^{-13} \), \( F_{(1,31)} = 133.291 \)) and interaction between them (\( p = 0.0017 \), \( F_{(1,31)} = 11.797 \)). Exposure to the novel environment increased the densities in both control and DG-NR1KO mice (\( p = 1.7 \times 10^{-4} \) and 0.008, respectively, df = 31, Tukey’s HSD test) while the densities were lower in DG-NR1KO mice than controls regardless of whether they were exposed to novel environment or not (\( p = 1.7 \times 10^{-4} \) and 2.0 \( \times 10^{-4} \), respectively, df = 31, Tukey’s HSD test). The percentage increase after exposure to the novel environment was significantly higher in DG-NR1KO mice than in control (exposure: \( p = 7.4 \times 10^{-8} \), \( F_{(1,31)} = 49.031 \) and genotype: \( p = 0.020 \), \( F_{(1,31)} = 6.028 \), exposure \& genotype: \( p = 0.020 \), \( F_{(1,31)} = 6.028 \), two-way ANOVA; \( p = 0.0046 \), df = 31, Tukey’s HSD test; Fig. 7C), while the percentage was significantly increased in both control and DG-NR1KO mice (\( p = 0.027 \) and 1.7 \( \times 10^{-4} \), df = 31, Tukey’s HSD test).

We divided the granule cell layer into parallel segments of 20% of the total thickness, and quantified Arc+/Prox1+ cells in each segment (Fig. 7D, E). Three-way mixed ANOVA detected a significant interaction between depth and genotype (\( p = 8.1 \times 10^{-5} \), \( F_{(3,656,113,325)} = 6.972 \)) but not among depth, genotype and exposure (home cage vs novel environment; \( p = 0.077 \), \( F_{(3,656,113,325)} = 2.224 \)). These results indicate that, regardless of whether exposed to the novel environment or not, Arc+/Prox1+ cell density showed significantly different changes along depth between control and DG-NR1KO mice. From the subgranular zone (0%) to the molecular layer (100%), control mice show a steady increase in Arc+/Prox1+ cell density up to the middle in the granule cell layer (60–80%), where it reaches a plateau level. In contrast, in DG-NR1KO mice, the density stays low up to the middle of the layer and jumps up at the depth close to the molecular layer (80–100%). Thus, DG-NR1KO mice show an impairment in relative amount of Arc expression along the depth of granule cell layer.

In sum, although DG-NR1KO mice showed a novelty-induced increase in Arc expressing granule cells, we detected impairments in three aspects: (1) overall reduction in Arc+ granule cells regardless of exposure to the novel environment or not; (2) higher percentage increase of Arc+ granule cells after exposure to the novel environment; and (3) abnormal distribution of Arc+ granule cells along the depth of granule cell layer.

The DG-NR1KO mice display higher tendency to explore the center of an open field during a novelty-suppressed feeding test and show higher fecal counts in an open field test

DG-NR1KO mice have been previously shown to have a deficit in a memory-based context discrimination task (McHugh et al., 2007). Previous studies suggested that adult neurogenesis has a role in anxiety-related behaviors (Jacobs et al., 2000; Santarelli et al., 2003; Surget et al., 2011). Therefore, we examined the behavior of adult, control and DG-NR1KO mice (\( n = 22 \) and 21 mice, respectively) in the novelty-suppressed feeding test, open field test and elevated plus maze (Santarelli et al., 2003; Walf and Frye, 2007; Seibenhener and Wooten, 2015).

After 1-d food deprivation, the mice were exposed to an open field (Fig. 8A), which was novel to the mice, with a food pellet located in the center. While mice naturally have a tendency to avoid open unprotected places, hunger drives them to move into the center of the open field to consume the food. The latency to consume the food is often considered to reflect the anxiety level of mice. We did not find a significant difference either in the latency to food consumption (\( p = 0.587 \), \( \chi^2(1) = 0.295 \), Kaplan–Meier survival analysis, log rank test; Fig. 8B) or overall speed (\( p = 0.304 \), \( t_{(41)} = -1.041 \), independent-sample \( t \) tests, two-tailed; Fig. 8C). We performed additional analyses on detailed behavioral performance as described before (Åmellén et al., 2017). The DG-NR1KO mice spent significantly smaller and larger percentages of time in the peripheral and center zones, respectively (\( p = 0.020 \) and 0.018, \( t_{(41)} = 2.411 \) and -2.464, respectively, independent-sample \( t \) tests, two-tailed; Fig. 8D), while the difference in the percentage time spent in the inner zone was not significant (\( p = 0.094 \), \( t_{(41)} = -1.715 \), independent-sample \( t \) tests, two-tailed). The number of entries into the center zone per 100 s was significantly higher in the DG-NR1KO mice (\( p = 0.008 \), \( t_{(41)} = -2.906 \), independent-sample \( t \) tests, two-tailed; Fig. 8E), while time spent in the center zone per visit was not (\( p = 0.887 \), \( t_{(41)} = 0.144 \), independent-sample \( t \) tests, two-tailed; Fig. 8F). These results indicate that the DG-NR1KO mice explore the center of open fields more extensively in the novelty-suppressed feeding test than control, suggesting that the removal of NR1 gene results in this behavioral impairment.

In the open field test (Fig. 8G), we quantified three measures which have been related to the anxiety level of mice, traveled distance, time in peripheral versus inner
zone and fecal counts. Larger traveled distance, longer time in peripheral zone and higher fecal counts are considered to be indices for higher anxiety level. We did not find significant difference in traveled distance and time in peripheral and inner zones (\(p = 0.157, 0.996, 0.996, t_{(34)} = 0.005, 0.005, \) independent-sample \(t\) test, two-tailed; Fig. 8H, I). On the other hand, fecal counts are significantly lower in DG-NR1KO mice than control (\(p = 0.028, t_{(34)} = 2.296,\) independent-sample \(t\) test, two-tailed; Fig. 8J). In the elevated plus maze (Fig. 8K), we quantified percentage of time in open arms, percentage of entries to open arms, total time in all arms and total number of entries to all arms. None of them show significant difference between control and DG-NR1KO mice (\(p = 0.728, 0.798, 0.581, 0.679, t_{(34)} = -0.352, 0.259, 0.560, 0.419, \) respectively, independent-sample \(t\) test, two-tailed; Fig. 8L–O).

The effect of NR1 gene knock-out did not show clear sex difference

Our original aim did not include the investigation of sex difference. Therefore, although we included both sexes in most experiments except Figure 6 (females only), we did not have a plan to analyze females and males separately. However, in response to reviewers’ requests, we here show the data for females and males separately (Fig. 9). A caveat is that sample size is relatively low for statistical analyses when we divided the data into two sexes, although we believe that the results would still give useful information for future follow-up study. From the graphs in Figure 9, the effect of NR1 gene knock-out seems to be generally consistent between females and males. To support this observation, we performed factorial ANOVA including sex as a between-subject factor (Table 1–Table 14). In most comparisons except six (see the next paragraph for more detail), no significant sex difference was detected as supported by no significance in the main effect of sex or interactions involving sex.

Four of such exceptions are results corresponding to 7 and 28 d for Figure 3F and 28 d for Figure 3H, J. For these exceptions, the main effects of sex and genotype are significant while their interactions are not significant. Thus, the effects of NR1 gene knock-out occurring similarly in both females and males, although the densities of BrdU+ and BrdU+/Prox1+ cells and survival rate are higher in females regardless of genotype. Another exception is a result corresponding to 7 d in Figure 3H. Two-way

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**Figure 8.** The DG-NR1KO mice display higher tendency to explore the center of an open field during a novelty-suppressed feeding test and show higher fecal counts in an open field test. A, A square open field used in a novelty-suppressed feeding test. Division of the open field into the peripheral, inner and center zone. B, Survival graph for latency to consume food for adult control and DG-NR1KO mice. C–F, Behavioral measurements in the novelty-suppressed feeding test for adult control and DG-NR1KO mice. G, A square open field used in an open field test. Division of the square open field into the peripheral and inner zone. H–J, Behavioral measurements in the open field test for adult control and DG-NR1KO mice. K, An elevated plus maze. L–O, Behavioral measurements in the elevated plus maze for adult control and DG-NR1KO mice; *\(p < 0.05\), **\(p < 0.01\), independent-sample \(t\) test, two-tailed.
Table 1: Two-way ANOVA results for Figure 9A (sex analysis for Fig. 3)

| Genotype | Sex | p | Genotype | Interaction |
|----------|-----|---|----------|-------------|
| 3B       | p   | 0.932 | 0.027 | 0.272 |
| 3D       | p   | 0.937 | 6 × 10^{-6} | 0.645 |
| 3F (7 d) | p   | 0.010 | 0.002 | 0.465 |
| 3F (28 d)| p   | 0.015 | 5 × 10^{-6} | 0.069 |
| 3G (7 d) | p   | 0.314 | 0.567 | 0.448 |
| 3G (28 d)| p   | 6 × 10^{-5} | 0.798 |
| 3H (7 d) | p   | 0.130 | 0.028 | 0.829 |
| 3H (28 d)| p   | 6.708 | 57.234 | 4.113 |
| 3I (7 d) | p   | 0.171 | 0.276 | 0.432 |
| 3J       | p   | 0.889 | 0.790 | 0.112 |
| p        | p   | 0.020 | 12 × 10^{-5} | 0.081 |
| F (7 d)  | p   | 6.98 | 46.673 | 3.575 |

Table 2: Sample size (unit: mice) for Figure 9A (sex analysis for Fig. 3)

| Genotype | Sex | Female | Male |
|----------|-----|--------|------|
| 3B, D    | Control | 3 | 3 |
| 3F–/– (7 d) | Control | 3 | 4 |
| 3F–/– (28 d) | Control | 5 | 4 |
| 3J       | Control | 5 | 3 |

Table 3: Two-way ANOVA results for Figure 9B (sex analysis for Fig. 4)

| Sex | Genotype | Interaction |
|-----|----------|-------------|
| p   | 3.7 × 10^{-4} | 0.524 |
| p   | 5.014 | 30.588 | 0.440 |
| p   | 1.9 × 10^{-4} | 0.582 |
| p   | 3.53 | 36.882 | 0.326 |
| p   | 0.792 | 0.004 | 0.928 |
| p   | 0.074 | 14.933 | 0.009 |
| p   | 0.523 | 0.030 | 0.894 |
| p   | 0.441 | 6.667 | 0.019 |

ANOVA (sex × genotype) showed the main effect of genotype is significant (p = 0.028; Tables 1, 2) although the test for the original analysis did not detect this difference as significant (p = 0.070). This discrepancy may be because there are variations between females and males (although not significant), which may prevent between-genotype difference in the original analysis from reaching statistical significance in the t test when the varied data from two sexes were analyzed without being sorted. The last exception is the analysis corresponding to two weeks old in Figure 5D; sex × genotype interaction is significant (Tables 5, 7). However, Tukey’s HSD tests did not reach significance for either female or male (Tables 6, 7). Thus, we did not detect significant difference between control and DG-NR1KO mice in either sex, similarly to the original analysis in Figure 5D.

For survival analysis in Figure 8B, we performed survival analysis stratified with sex. We did not detect significant difference between control and DG-NR1KO mice (Fig. 9E; Table 12).

In sum, we did not detect a clear sex difference in the effect of NR1 gene knock-out in any of the analyses in Figure 9.
Age of mice may affect the effect of NR1 gene knock-out in enrichment-induced survival of new neurons

In most experiments, we used young adult mice from 42 to 60 d old. However, for experiments described in Figures 6, 7, 8A–F, we also used more mature adult mice up to 86 d old. Although our original plan did not include the investigation of age difference, we have divided the mice into two groups of <60 d old and >60 d old and analyzed age effect in response to a reviewer’s request. A caveat is that sample size is relatively low for statistical analyses when we divided the data into the two age groups, although we believe that the results would still give useful information for future follow-up study.

The extent of adult neurogenesis is known to be reduced along age. Therefore, as expected, we found that <60-d-old mice have higher densities of BrdU+/and BrdU+/Prox1+ cells (Fig. 10A; Table 15—Table 18). Notably, these densities showed significant age x genotype x housing interaction (Tables 15, 18). Then, we performed preplanned comparisons (standard vs enrichment) in four groups (<60-d-old control, ≥60-d-old control, <60-d-old DG-NR1KO, ≥60-d-old DG-NR1KO). The densities of BrdU+ and BrdU+/Prox1+ mice are comparable between standard and enriched conditions in <60-d-old DG-NR1KO mice but the other three group showed significant increase or clear trend of non-significant increase in the densities under enriched conditions (Fig. 10A; Tables 16, 18). To further look into this age-dependent differences (Fig. 10B), we examined correlation between age and the densities. Enriched control mice show significant, negative correlations between age and BrdU+ or BrdU+/Prox1+ cell density. Standard control and standard DG-NR1KO mice also showed negative correlations, which were not significant probably because of small sample size. On the other hand, enriched DG-NR1KO mice seems to have completely lost the correlations. Because of the limited sample size, this potential age modulation in the effect of NR1 gene knock-out on the densities of BrdU+ and BrdU+/Prox1+ is not fully demonstrated. However, it would be of interest to follow-up this potential difference and elucidate a mechanism behind it.

None of other measurements corresponding to Figures 6D,F, 7, 8B–F (Fig. 10A,C,D) showed any notable age effect (main effects of age or interactions involving age; Table 15—Table 26).

Discussion

In this study, we have made three major findings. First, DG-NR1KO mice show impairment in neurogenesis during postnatal development and adulthood (Figs. 3, 5A–D). Second, DG-NR1KO mice have smaller size of the granule cell layer (Fig. 4), which becomes apparent during postnatal development (Fig. 5E,F). Third, DG-NR1KO mice have a functional deficit, which appears as increased...
NR1 gene is essential for the formation of functional NMDA receptors, these findings suggest that the NMDA receptors are essential for normal development and function of the dentate gyrus. McHugh et al. (2007) showed that the Cre-mediated removal of NR1 genes is specific to granule cells in the dentate gyrus. Based on this finding, our following discussion focuses on a role of the NMDA receptors in granule cells. However, a technical caveat for region-specific or cell type-specific knock-out mice is that one cannot completely exclude the possibility that the observed phenotype may be caused by gene knock-out in other regions or cell types than the regions or cell type of interest. We suggest readers to read our discussion while keeping this caveat in mind.

The NMDA receptors in granule cells support the normal development of the dentate gyrus

One of our major findings is that the size of the granule cell layer was smaller in the dorsal dentate gyrus of DG-NR1KO mice compared with control mice (Fig. 4), suggesting that the NR1 gene in granule cells is required for the normal development of the granule cell layer. This finding is consistent with a study by Bannerman and colleagues, which reported a reduced size of the dentate gyrus in mice lacking the NR1 gene in the hippocampus but not specifically in granule cells (Bannerman et al., 2012). In contrast to our finding, McHugh et al. (2007) did not report abnormality in the size of granule cell layer. However, considering high variability of granule cell layer size among coronal sections along the anteroposterior axis, relatively small reduction in size (~20%; Fig. 4) can be easily missed unless one systematically examines many sections and quantifies the volume as we did. We further found that this reduced size becomes apparent as early as at the postnatal age of three weeks (Fig. 5E,F), which demonstrates the developmental origin of this deficit.

We examined neurogenesis during the first few weeks of postnatal period, when the majority of granule cells in the dentate gyrus are generated (Encinas et al., 2013). We did not find any change in the number of Ki67 cells up to the postnatal age of four weeks (Fig. 5B), which suggests that cell proliferation is intact during postnatal development. In contrast, we detected a reduction of DCX+ cells starting at the postnatal age of three weeks (Fig. 5D), indicating that neurogenesis is impaired in the postnatal, developing dentate gyrus of DG-NR1KO mice. Because cell proliferation is not affected, this reduction is likely through a later process of neurogenesis such as neuronal survival. The reduction appears at the same timing as reduced size of granule cell layer (Fig. 5E,F), suggesting that impaired neurogenesis contributes to the reduction in the size of granule cell layer.

DG-NR1KO mice show impaired adult neurogenesis in the dentate gyrus

In DG-NR1KO mice, we found impairment in adult neurogenesis in the dorsal dentate gyrus. This impairment was shown as reduction in Ki67+, DCX+, BrdU+, and BrdU+/Prox1+ cells (Figs. 3A–F,H, 9A). Overall, the

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**Table 11: Sample size (unit: mice) for Figure 9D (sex analysis for Fig. 7)**

| Genotype | Exposure | Female | Male |
|----------|----------|--------|------|
| Control  | Home cage | 4      | 2    |
| KO       | Novel    | 5      | 4    |

**Table 12: Log rank test result (stratified with sex) for Figure 9E (sex analysis for Fig. 8)**

Fig. 8B \( \chi^2(1) = 0.380 \)

**Table 13: Two-way ANOVA results for Figure 9E (sex analysis for Fig. 8)**

| Genotype | Sex | Genotype | Interaction |
|----------|-----|----------|-------------|
| Control  | \( p \) | 0.303    | 0.188 0.183 |
| KO       | \( F(1,29) \) | 1.090 | 1.799 1.836 |
| Fig. 8D (peripheral) | \( p \) | 0.460 | 0.034 0.619 |
| Fig. 8D (inner) | \( p \) | 0.557 | 4.813 0.251 |
| Fig. 8D (left) | \( p \) | 0.776 | 0.151 0.397 |
| Fig. 8E | \( F(1,29) \) | 0.082 | 2.141 0.734 |
| Fig. 8F | \( p \) | 0.247 | 0.020 0.758 |
| Fig. 8H | \( F(1,32) \) | 1.381 | 5.880 0.096 |
| Fig. 8I | \( p \) | 0.926 | 0.010 0.887 |
| Fig. 8J | \( F(1,29) \) | 0.009 | 7.322 0.021 |
| Fig. 8K | \( p \) | 0.593 | 0.894 0.930 |
| Fig. 8L | \( F(1,29) \) | 0.291 | 0.018 0.008 |
| Fig. 8M | \( F(1,32) \) | 0.432 | 0.194 0.815 |
| Fig. 8N | \( p \) | 0.633 | 1.761 0.012 |
| Fig. 8O | \( F(1,32) \) | 0.380 | 0.983 0.693 |
| Fig. 8P | \( F(1,29) \) | 0.792 | 4.7 \times 10^{-4} 0.158 |
| Fig. 8Q | \( F(1,32) \) | 0.380 | 0.983 0.693 |
| Fig. 8R | \( F(1,32) \) | 0.792 | 4.7 \times 10^{-4} 0.158 |
| Fig. 8S | \( p \) | 0.731 | 0.030 0.770 |
| Fig. 8T | \( F(1,32) \) | 0.120 | 5.187 0.087 |
| Fig. 8U | \( p \) | 0.447 | 0.777 0.410 |
| Fig. 8V | \( F(1,22) \) | 0.599 | 0.082 0.704 |
| Fig. 8W | \( p \) | 0.431 | 0.755 0.457 |
| Fig. 8X | \( F(1,22) \) | 0.644 | 0.100 0.574 |
| Fig. 8Y | \( p \) | 0.288 | 0.646 0.910 |
| Fig. 8Z | \( F(1,22) \) | 1.186 | 0.216 0.013 |

**Table 14: Sample size (unit: mice) for Figure 9E (sex analysis for Fig. 8)**

| Genotype | Sex | Female | Male |
|----------|-----|--------|------|
| Control  | \( 8C–F \) | 9      | 13   |
| KO       | \( 8G–J \) | 8      | 13   |
| Control  | \( 8L–O \) | 7      | 6    |

exploration of the center of the open field during the novelty-suppressed feeding test and reduced fecal counts in the open field test (Fig. 8). Considering the fact that the NR1 gene is essential for the formation of functional neural activity, the impairment of adult neurogenesis in DG-NR1KO mice may underlie the behavioral deficits observed in these mice. Overall, our findings provide new insights into the role of the NR1 gene in the hippocampus and highlight the importance of investigating region-specific and cell type-specific effects of gene knock-outs.
number of approximately four-weeks-old, newly-generated neurons are reduced by >80% in DG-NR1KO mice (Fig. 3H). The reduced ratio of BrdU+/Prox1+ cells at 28 d to 7 d (Fig. 3J) indicates that the survival of newly-generated neurons is compromised. The result suggests that the NMDA receptors in the granule cells support the survival of new neurons in the adult dentate gyrus. The finding is consistent with previous studies which showed that the NMDA receptor-dependent long-term potentiation increases the survival of newborn neurons in the dentate gyrus (Bruel-Jungerman et al., 2006) and that the NMDA receptors on immature neurons regulate their own survival (Tashiro et al., 2006b).

A reduction of Ki67+ cells indicates a deficit in cell proliferation in adult DG-NR1KO mice, which was not observed in the postnatal, developing dentate gyrus. Considering that the impairment in neurogenesis (a reduction of DCX+ cells) and granule cell layer size precedes the reduction of Ki67+ cells, the effect on cell proliferation may be secondary to the abnormal development of the dentate gyrus. In addition, this possibility is also supported by the previous finding that the Pomc promoter initiates transcription in newborn neurons at some
Table 15: Three-way ANOVA results for Figure 10A,B (age analysis for Fig. 6)

| Genotype   | Age × genotype | Genotype × Age × housing |
|------------|----------------|--------------------------|
|            | Age            | Genotype                 | Housing | Age × genotype | Genotype × Age × housing |
| C          | 0.0016         | 4.4 × 10^{-8}            | 1.5 × 10^{-5} | 5.7 × 10^{-4} | 0.0015 | 0.431 | 0.035 |
|            | 130.003        | 66.284                   | 30.354    | 16.203        | 13.190 | 0.643 | 5.039 |
| E          | 0.413          | 0.0021                   | 0.306     | 0.865         | 0.906  | 0.684 | 0.939 |
|            | 0.696          | 12.100                   | 1.097     | 0.030         | 0.014  | 0.170 | 0.006 |
| F          | 0.0011         | 1.1 × 10^{-8}            | 1.3 × 10^{-5} | 6.5 × 10^{-4} | 7.0 × 10^{-4} | 0.322 | 0.037 |
|            | 14.025         | 78.211                   | 31.159    | 15.720        | 15.538 | 1.028 | 4.920 |
| C          | 0.279          | 0.282                    | 4.0 × 10^{-4} | 0.185         | 0.282  | 0.279 | 0.185 |
|            | 1.234          | 1.216                    | 17.361    | 1.871         | 1.216  | 1.234 | 1.871 |

Table 16: Tukey’s HSD test results (standard vs enrichment, two-tailed, df = 22) after significant interaction for Figure 10A,B (age analysis for Fig. 6)

| Genotype   | Age <60 d | Age >60 d |
|------------|-----------|-----------|
| C          | p = 1.9 × 10^{-4} | p = 0.182 |
| E          | p = 0.000 | p = 0.575 |

Table 17: Age (mean ± SD) for Figure 10A,B (age analysis for Fig. 6)

| Housing   | Age <60 d | Age >60 d |
|-----------|-----------|-----------|
| Control   | 53.2 ± 4.1 | 67.0 ± 5.7 |
| KO        | 51.0 ± 1.7 | 65.0 ± 3.8 |
| Standard  | 53.8 ± 4.7 | 65.5 ± 3.7 |
| Enrichment| 51.0 ± 1.7 | 65.6 ± 3.4 |

Table 18: Sample size (unit: mice) for Figure 10A,B (age analysis for Fig. 6)

| Housing   | Age <60 d | Age >60 d |
|-----------|-----------|-----------|
| Control   | 5         | 2         |
| KO        | 4         | 4         |
| Standard  | 3         | 4         |
| Enrichment| 3         | 5         |

Table 19: Pearson correlation between age and density for Figure 10A,B (age analysis for Fig. 6)

| BrdU+     | BrdU+/Prox1+ |
|-----------|--------------|
| Standard  | Enriched     | Standard  | Enriched     |
| p = 0.202 | p = 0.015    | p = 0.250 | p = 0.017    |
| r = -0.548| r = -0.851   | r = -0.503| r = -0.844   |
| p = 0.188 | p = 0.199    | p = 0.118 | p = 0.323    |
| r = -0.518| r = 0.507    | r = -0.597| r = 0.403    |

Table 20: Three-way ANOVA results for Figure 10C (age analysis for Fig. 7)

| Age | Genotype | Exposure | Age × genotype | Genotype × exposure | Age × exposure | Age × genotype × exposure |
|-----|----------|----------|----------------|---------------------|---------------|--------------------------|
|     |          |          |                |                     |               |                          |
| C   | p = 0.197| 1.1 × 10^{-11}| 4.0 × 10^{-8} | 0.536              | 0.0017        | 0.541                    | 0.515 |
|     | 1.752    | 125.968  | 57.115         | 0.394              | 12.081        | 0.383                    | 0.435 |
| C   | p = 0.519| 0.0072   | 1.0 × 10^{-9}  | 0.367              | 0.0072        | 0.519                    | 0.367 |
|     | 0.428    | 8.444    | 82.771         | 0.841              | 8.444         | 0.428                    | 0.841 |

This reduction of cell proliferation in the dentate gyrus of adult DG-NR1KO mice is in contrast to the previous studies which suggested NMDA receptor activation inhibits cell proliferation (Cameron et al., 1995; Gould and Cameron, 1997). These studies found that the systemic injection of NMDA receptor agonists and antagonists, respectively, decreases and increases cell proliferation in the adult dentate gyrus. This discrepancy may be because systemic injection blocks NMDA receptor function in the brain regions other than the dentate gyrus while NR1 gene removal in DG-NR1KO mice is specific to the dentate gyrus. Alternatively, it may be because of acute activation/blockade by the injection of an agonist/antagonist compared with chronic blockade starting from postnatal development in DG-NR1KO mice.

We found a reduced proportion of BrdU+ cells expressing Prox1 in DG-NR1KO mice at 28 d, but not 7 d, after BrdU injection (Fig. 3G). The proportion reached ~90% 7 d after BrdU injection and did not show difference between control and DG-NR1KO mice. Until 28 d, the proportion stayed at ~90% in control mice but reduced to ~60% in DG-NR1KO mice. The reduced proportion of BrdU+ cells expressing a neuronal marker is often interpreted as reduced tendency to neuronal differentiation. However, we believe that this reduction was caused by selective reduction in the survival of BrdU+/Prox1+ cells in DG-NR1KO mice in combination with the intact survival of BrdU+/Prox1− cells. However, the reviewer believes that this is partly caused by reduced neuronal differentiation. We believe that this is unlikely or, if any, makes only a small contribution because of the following reasons. First, by 7 d after BrdU injection, ~90% of BrdU+ cells had already committed to neuronal fate (Prox1+) and no further increase happened from 7 to 28 d after BrdU injection in control mice, suggesting that no (or only small) further neuronal differentiation normally occurs.
after 7 d. Second, the proportion of BrdU+ cells expressing Prox1+ reduced from ~90% to ~60%. If neuronal differentiation is a major cause of this reduction, de-differentiation from neuronal fate (losing Prox1 expression) needs to occur. Although we believe that reduced neuronal differentiation in DG-NR1KO mice is unlikely, we agree with the reviewer that our result does not completely exclude this possibility and that reduced neuronal differentiation or increased de-differentiation may have contributed to reduced neurogenesis in DG-NR1KO mice.

Exposure to an enriched environment has previously been shown to increase the survival of newborn neurons in the adult dentate gyrus (Kempermann et al., 1997; Tashiro et al., 2007), and our results confirm these findings (Fig. 6). Although the overall level of BrdU+/Prox1+ cells were lower in DG-NR1KO mice compared with control (Fig. 7E), the percentage increase of BrdU+/Prox1+ cells after exposure to the enriched environment did not differ significantly between the two genotypes (Fig. 6F). Therefore, the NMDA receptors in granule cells are not essential for enrichment-induced increase in survival of new neurons, although we cannot exclude the possibility that there is a sub-component of a survival-inducing effect which is dependent on the NMDA receptors.

The NMDA receptors in granule cells are required for normal function of the dentate gyrus

The immediate early gene Arc has been shown to be involved in synaptic plasticity in the hippocampus and in the consolidation of long-term memory (Guzowski et al., 2000; Plath et al., 2006), and it is known that Arc expression is induced by novel environment exploration, learning, and high-frequency stimulation (Steward et al., 1998; Guzowski et al., 1999, 2001), and is dependent on NMDA receptor function (Czernecki et al., 2011).

We found a significantly lower number of Arc+/Prox1+ cells in the DG-NR1KO mice compared with control mice (Fig. 6E, Fig. 7B), which confirms that Arc expression is dependent on functional NMDA receptors in granule cells. This reduction is regardless of whether the mice were exposed to the novel environment or not. In addition, we found that, on exposure to the novel environment, Arc+/Prox1+ cells still increase in DG-NR1KO mice, and in terms of percentage changes, the novelty-induced increase is higher in DG-NR1KO mice than control (Fig. 7C). These findings indicate that, although Arc expression is dependent on the NMDA receptors, at least some components of a novelty-induced increase in Arc expression are not dependent on the NMDA receptors; there may co-exist:

Table 21: Four-way mixed ANOVA results, between-subject effects, for Figure 10C (age analysis for Fig. 7)

|                      | Age | Genotype | Exposure | Age × genotype | Age × exposure | Genotype × exposure | Age × genotype × exposure |
|----------------------|-----|----------|----------|----------------|---------------|--------------------|--------------------------|
| Fig. 7E (between-subjects effects) | p   | 0.295    | 1.9 × 10⁻¹⁰ | 3.8 × 10⁻⁷  | 0.384         | 0.259              | 0.246                   | 0.847                     |
|                      | F(1,27) | 1.141 | 97.378   | 44.365         | 0.784         | 1.332              | 1.406                   | 0.038                     |

Table 22: Four-way mixed ANOVA results, within-subject effects, for Figure 10C (age analysis for Fig. 7)

|                      | Depth | Depth × age | Depth × genotype | Depth × exposure | Depth × age × genotype | Depth × age × exposure | Depth × genotype × exposure | Depth × age × genotype × exposure |
|----------------------|-------|------------|-----------------|-----------------|------------------------|------------------------|-----------------------------|----------------------------------|
| Fig. 7E (within-subjects effects) | p   | 3.4 × 10⁻²⁰ | 0.735           | 2.0 × 10⁻⁴      | 1.0 × 10⁻⁵             | 0.295                  | 0.697                       | 0.101                           | 0.822                           |
|                      | F(4,108) | 39.068 | 0.501           | 6.039           | 8.041                  | 1.248                  | 0.553                       | 1.992                           | 0.380                           |

Table 23: Sample size (unit: mice) for Figure 10C (age analysis for Fig. 7)

| Exposure | <60 d | >60 d |
|----------|-------|-------|
| Control  | Home cage | 4    | 2    |
| KO       | Home cage | 6    | 4    |
|          | Novel    | 6    | 4    |

Table 24: Log rank test results (stratified with age) for Figure 10D (age analysis for Fig. 8B–F)

Fig. 8B

\[ p = 0.660\]

\[ \chi^2(1) = 0.194\]

Table 25: Two-way ANOVA results of Figure 10D (age analysis for Fig. 8B–F)

|                      | Age | Genotype | Interaction |
|----------------------|-----|----------|-------------|
| Fig. 8C              | p   | 0.052    | 0.582       | 0.489       |
|                      | F(1,39) | 4.022 | 0.308       | 0.487       |
| Fig. 8D (peripheral) | p   | 0.862    | 0.058       | 0.951       |
|                      | F(1,39) | 0.031 | 3.815       | 0.004       |
| Fig. 8D (inner)      | p   | 0.669    | 0.186       | 0.900       |
|                      | F(1,39) | 0.185 | 1.808       | 0.016       |
| Fig. 8D (left)       | p   | 0.750    | 0.044       | 0.941       |
|                      | F(1,39) | 0.103 | 4.312       | 0.005       |
| Fig. 8E              | p   | 0.495    | 0.010       | 0.526       |
|                      | F(1,39) | 0.474 | 7.23        | 0.410       |
| Fig. 8F              | p   | 0.940    | 0.770       | 0.742       |
|                      | F(1,39) | 0.006 | 0.087       | 0.110       |

Table 26: Sample size (unit: mice) for Figure 10D (age analysis for Fig. 8B–F)

|                      | <60 d | >60 d |
|----------------------|-------|-------|
| Control              | 17    | 5     |
| KO                   | 16    | 5     |
mechanisms dependent on and independent of the NMDA receptors.

Another observation we made was a between-genotype difference in the distribution of Arc+/Prox1+ cells along the depth of the granule cell layer (Fig. 7E). Both control and DG-NR1KO mice had higher numbers of Arc+/Prox1+ cells toward the outer part of the granule cell layer. However, with or without exposure to the novel environment, control mice increased the number of Arc+/Prox1+ cells starting from the inner half of the layer. In contrast, DG-NR1KO mice kept the numbers of Arc+/Prox1+ cells low in the inner half and showed higher number of Arc+/Prox1+ cells mainly in the most outer part (80–100%) of the layer. This finding suggests that functioning of the dentate gyrus is compromised in DG-NR1KO mice.

Overall, these three observed abnormalities in Arc expression, (1) overall reduction in Arc+ granule cells; (2) an enhanced, novelty-induced increase in terms of percentage changes; and (3) abnormal distribution across the depth of granule cell layer, suggest that the function of the dentate gyrus, which is mediated by the Arc gene, may be compromised.

The novelty-suppressed feeding test is a conflict paradigm in which a food-deprived mouse has to choose between going into an open space to get food or stay close to the walls without access to the food. The latency until the mouse starts to eat the food has been considered as a measure of anxiety level of the mouse. Santarelli and colleagues found that the blockade of neurogenesis abolished the behavioral effect of fluoxetine in the novelty-suppressed feeding test (Santarelli et al., 2003). Adult neurogenesis has been linked to anxiety and mood disorders (Gould et al., 1997; Jacobs et al., 2000). Studies have found increased anxiety after a mild stressor (Snyder et al., 2011) or increased anxiety without a stressor (Revest et al., 2009) after ablation of adult neurogenesis. While we did not find a significant difference in the latency to consume food, we found multiple measurements pointing to an increased tendency for DG-NR1KO mice to explore the center of the open field, which is also often considered as a measure reflecting animal’s anxiety level, for example, in an open field test (Seibenhener and Wooten, 2015). In the open field test, we did not find similar tendency to spend more time in the inner part of the open field. However, we detected a significant reduction in fecal counts during the open field test, which has been considered as an index of lower anxiety level (Seibenhener and Wooten, 2015). From these mixed results, one may be inclined to interpret it as an indication that DG-NR1KO mice exhibit lower anxiety level. However, we are hesitant to conclude that the loss of the NMDA receptors in granule cells results in an anxiety-related phenotype. On the other hand, it is safe to say that DG-NR1KO mice have behavioral abnormality which is not necessarily based on compromised memory mechanisms which has been previously characterized (McHugh et al., 2007). This knowledge would be important to interpret their behavioral phenotypes involving spatial exploration and/or emotion.

Our work has contributed to the understanding of the role of the NMDA receptors in the normal development and adult neurogenesis of the dentate gyrus, and that the loss of the NMDA receptors during brain maturation can contribute to permanent alterations in brain function. Future work is required to reveal the complex contribution of the NMDA receptors to brain development and function in different brain areas.

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