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Enhanced synaptic plasticity in newly generated granule cells of the adult hippocampus

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Neural stem cells in various regions of the vertebrate brain continuously generate neurons throughout life¹⁻⁴. In the mammalian hippocampus, a region important for spatial and episodic memory^{5,6}, thousands of new granule cells are produced per day⁷, with the exact number depending on environmental conditions and physical exercise^{1,8}. The survival of these neurons is improved by learning and conversely learning may be promoted by neurogenesis⁸⁻¹⁰. Although it has been suggested that newly generated neurons may have specific properties to facilitate learning^{2,10,11}, the cellular and synaptic mechanisms of plasticity in these neurons are largely unknown. Here we show that young granule cells in the adult hippocampus differ substantially from mature granule cells in both active and passive membrane properties. In young neurons, T-type Ca²⁺ channels can generate isolated Ca²⁺ spikes and boost fast Na⁺ action potentials, contributing to the induction of synaptic plasticity. Associative longterm potentiation can be induced more easily in young neurons than in mature neurons under identical conditions. Thus, newly generated neurons express unique mechanisms to facilitate synaptic plasticity, which may be important for the formation of new memories.

To identify newly generated granule cells in the adult hippocampus, we applied electrophysiological, morphological and immunocytochemical criteria. We made whole-cell recordings from neurons in the granule cell layer and determined the input resistance (R_{in}), which is known to be characteristically different between young and mature neurons¹². Subsequently, we examined the immunoreactivity for polysialic acid neural cell adhesion molecule (PSA-NCAM), a marker for newly generated granule cells^{1,3,13,14}, and the morphology of the dendritic tree (Fig. 1).

On the basis of the distribution of R_{in} , most cells fell into two categories, with mean R_{in} values of 232 \pm 78 M Ω and 4.5 \pm 1.9 G Ω , respectively (mean \pm s.d.; Fig. 1a, c, e). Cells with high R_{in} were exclusively encountered in the inner granule cell layer adjacent to the subgranular zone, where neural stem cells are located¹. Cells with R_{in} below 400 M Ω were immunonegative for PSA-NCAM and showed a complex apical dendritic tree (total dendritic length $3,652 \pm 134 \,\mu\text{m}$; mean \pm s.e.m.; n = 14), indicating that they were mature granule cells. In contrast, all cells with R_{in} larger than $1.5 \,\mathrm{G}\Omega$ were immunopositive for PSA-NCAM (Fig. 1d, f; relative fluorescence intensity $76 \pm 6\%$) and had rudimentary apical and basal dendrites (Fig. 1f; total dendritic length 845 ± 111 µm, n = 14), indicative of immature granule cells^{15–17}. As PSA-NCAM is expressed transiently in granule cells 1-3 weeks after mitosis^{13,14}, these results suggest that the cells with high R_{in} are newly generated immature granule cells in the adult hippocampus. Cells with R_{in} larger than 1.5 G Ω were immunonegative for parvalbumin (n = 4), suggesting that they do not represent interneurons18,19.

We next studied the active and passive membrane properties of the two classes of neurons (Fig. 2). Both mature and young granule cells showed comparable resting potentials (-80.8± $0.9 \,\mathrm{mV}$ versus $-75.3 \pm 2.0 \,\mathrm{mV}$, respectively) and were able to generate fast action potentials with large amplitudes (Fig. 2a-d; $140 \pm 2 \text{ mV}, n = 9 \text{ versus } 115 \pm 2 \text{ mV}, n = 6$). Unlike mature granule cells, which generated trains of action potentials during long current pulses (Fig. 2a; mean action potential frequency $30 \pm 1 \,\text{Hz}$), young neurons fired only a few action potentials per stimulus: in 15 of 20 cells, a single action potential was evoked (Fig. 2b). However, the current threshold determined with 100-ms pulses was markedly lower in young neurons than in mature cells $(34 \pm 9 \text{ pA}, n = 6 \text{ versus } 141 \pm 12 \text{ pA}, n = 4, \text{ respectively};$ P < 0.01). In a subset of young neurons currents of less than 10 pA were sufficient to elicit an action potential. Finally, young neurons had a much slower membrane time constant, $\tau_{\rm m}$, than mature granule cells (Fig. 2e, f; $123 \pm 10 \text{ ms}$, n = 7 versus $54 \pm 4 \text{ ms}$, n = 13, respectively)²⁰. Thus, newly generated granule cells have membrane properties that favour action potential generation with very small current stimuli, such as the opening of less than ten glutamate receptor channels²¹.

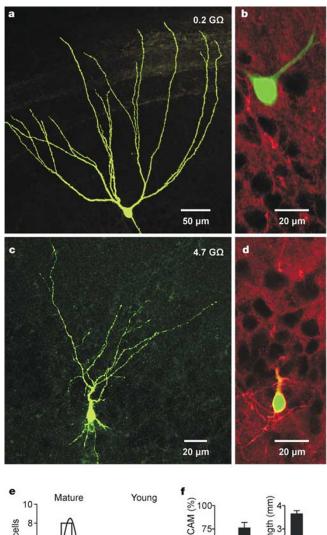
In contrast with mature neurons, young neurons generated transient low-threshold spikes when current injections below the threshold for Na⁺ action potential initiation were applied (Figs 2b and 3). Low-threshold spikes were insensitive to 1 μ M tetrodotoxin (TTX) but were blocked by 50 μ M Ni²⁺, showing that they were mediated by T-type Ca²⁺ channels²² (Fig. 3a–c). Ca²⁺ spikes in newly generated neurons had a voltage threshold of -56.0 ± 1.2 mV and an amplitude of 17.6 \pm 1.1 mV measured from the voltage at the end of the pulse (n = 16). Furthermore, they had a slow time course with a duration at half-maximal amplitude of 87 \pm 8 ms (n = 16). Thus, young granule cells generate low-threshold Ca²⁺ spikes under physiological conditions, whereas mature granule cells produce low-threshold Ca²⁺ spikes only in the presence of K⁺-channel blockers²³.

To examine whether the activation of T-type Ca²⁺ channels in young neurons contributed to the initiation of fast Na⁺ action potentials, we determined the probability of Na⁺ action potential initiation during current pulses of incremental amplitudes (Fig. 3d, e). A concentration of $50 \,\mu$ M Ni²⁺ markedly elevated the current threshold for the initiation of fast Na⁺ action potentials (Fig. 3e). On average, the current corresponding to the mid-point of the threshold curves increased to $176 \pm 56\%$ of the control value in young neurons (n = 6, P < 0.05). By contrast, the threshold remained unchanged in mature neurons ($98 \pm 3\%$, n = 4). In conclusion, newly generated immature granule cells show a unique

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pattern of excitability. As the stimulus intensity is increased, T-type Ca^{2+} channels first generate slow Ca^{2+} spikes and then boost the initiation of fast Na⁺ action potentials.

Postsynaptic action potentials are important associative signals for the induction of spike-timing-dependent plasticity at hippocampal glutamatergic synapses^{24–26}. We therefore tested the impact of postsynaptic excitability on synaptic plasticity in young granule cells. Excitatory postsynaptic potentials (EPSPs) evoked by stimulation of the perforant path were examined in the perforated-patch



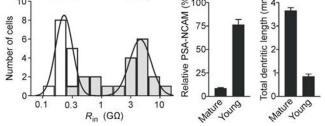


Figure 1 Newly generated granule cells in adult hippocampus are defined by high input resistance, PSA-NCAM immunoreactivity and immature dendritic morphology. **a**, **c**, Biocytin-filled mature (**a**) and young (**c**) granule cell. **b**, **d**, Confocal images of PSA-NCAM immunoreactivity (red) and biocytin staining (green) of the cells in **a** and **c**, respectively. Note the high PSA-NCAM expression of the immature cell (red dendrite and yellow rim at the soma). **e**, Distribution of input resistance of cells in the inner (grey bars) and outer part of the granule cell layer (open bars). Continuous curves represent the sum of two gaussian functions fitted to the histogram. **f**, Summary graph of PSA-NCAM immunoreactivity and total dendritic length.

configuration (Fig. 4). In vivo recordings suggest that dentate gyrus granule cells fire action potentials related to the location of the animal in the spatial environment and the phase of the hippocampal theta oscillation (5-8 Hz)^{27,28}. To mimic this situation, we applied stimulation paradigms corresponding to the physiological firing pattern of a granule cell outside (TBS_0) , at the border (TBS_1) , or in the centre of its place field (TBS₂; Fig. 4a, see Methods). In mature granule cells, TBS₀ and TBS₁ failed to induce long-lasting changes in synaptic strength (Fig. 4b, d, f; the EPSP amplitudes 25-30 min after the end of the induction paradigm were $88 \pm 13\%$ (n = 5) and $92 \pm 10\%$ (*n* = 6) of the control value, respectively; *P* > 0.4). However, if bursts of postsynaptic action potentials were evoked during TBS₂, a long-term potentiation (LTP) of the EPSP amplitude was induced (214 \pm 46% of control, n = 6, P < 0.05). Apparently, in mature granule cells a burst of postsynaptic action potentials was required for LTP induction, as reported previously for mature CA1 pyramidal neurons25,29.

Similar to mature granule cells, TBS_0 failed to induce significant changes in EPSP amplitude in young granule cells (123 ± 19%,

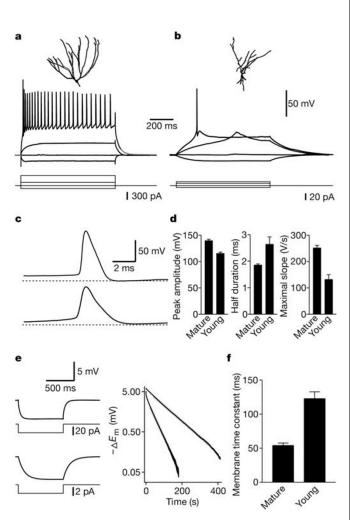


Figure 2 Young granule cells have distinct active and passive membrane properties. **a**, Train of action potentials in a mature neuron. **b**, Isolated low-threshold spike and single action potential in a young neuron. **c**, Single action potentials at higher time resolution in a mature (top trace) and a young neuron (bottom trace); dashed lines at -45 mV. **d**, Comparison of peak amplitude, half duration and maximal slope of action potentials. **e**, Voltage changes evoked by hyperpolarizing current pulses in a mature (top trace) and a young neuron (bottom trace). Right panel shows logarithmic plot of voltage changes (ΔE_m) after the pulse; regression lines (grey) are shown superimposed. **f**, Summary graph of membrane time constants.

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n = 5, P > 0.1). However, unlike in mature cells TBS₁ was sufficient to induce LTP (Fig. 4c, e, g; the EPSP amplitude was $154 \pm 9\%$ of control, n = 9, P < 0.01). Furthermore, in the young cells TBS₂ had only slightly larger effects than TBS₁ ($162 \pm 10\%, n = 5$). Statistical comparison between young and mature cells further indicates that TBS₁ has significantly larger effects in the young neurons (P < 0.001). Thus, our results demonstrate that excitatory synapses on newly generated granule cells show associative plasticity, and that the threshold for induction is lower than in mature neurons. Pairing of EPSPs with single postsynaptic action potentials is a highly efficient stimulus for the induction of LTP in young granule cells, whereas a burst of postsynaptic action potentials is required in mature neurons. Hence, different induction rules for LTP apply to young and mature granule cells.

Newly generated granule cells show specialized membrane properties and enhanced associative plasticity at their glutamatergic input synapses, suggesting a causal link. One scenario is that the high input resistance and the expression of T-type Ca^{2+} channels promote LTP because they facilitate the activation of neurons by sparse glutamatergic synaptic inputs. Furthermore, Ca^{2+} and Na⁺ action potentials as evoked by TBS₁ in young granule cells may efficiently induce LTP by generating both a global rise in postsynaptic Ca^{2+} concentration and a maximal relief of Mg²⁺ block of NMDA-type glutamate receptors.

Functional plasticity such as TBS₁-induced LTP may have profound structural consequences. It is well established that PSA-NCAM selectively labels newly generated neurons 1–3 weeks after

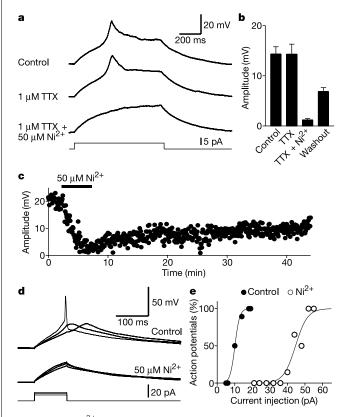


Figure 3 T-Type Ca²⁺ channels lead to enhanced excitability in young granule cells. **a**, **b**, The low-threshold spike in young neurons was insensitive to TTX, but was blocked by 50 μ M Ni²⁺, indicating that it is mediated by T-type Ca²⁺ channels. Spike amplitude was measured from the potential at the end of the current injection. **c**, Plot of Ca²⁺ spike amplitude against time in another experiment. The spike was reversibly blocked by Ni²⁺. **d**, **e**, The current threshold for the initiation of Na⁺ action potentials was increased when T-type Ca²⁺ channels were blocked by Ni²⁺. Continuous curves in **e** represent fitted Boltzmann functions.

mitosis^{13,14}. In this time period the fate of the newly generated neurons (survival versus cell death) is determined³⁰. Enriched environment and spatial learning increase the proportion of surviving neurons^{1,9}, suggesting the possibility that associative LTP promotes the survival and final integration of newly generated granule cells into the adult hippocampal circuit.

Both functional and structural plasticity in newly generated cells may be important for the formation of spatial memory. Single spikes during the theta cycle are observed *in vivo* when the animal approaches the border of the place field of a hippocampal granule cell^{27,28}. In the centre of the place field, however, granule cells fire brief bursts of action potentials. If the induction rules for synaptic plasticity as reported here also apply *in vivo*, our data would suggest that in young granule cells LTP will occur in a much larger 'induction area' as compared with mature neurons. This might be

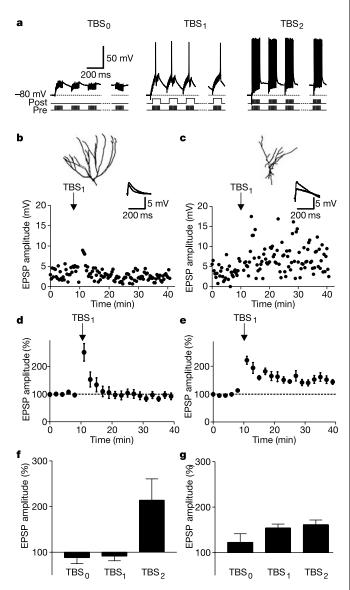


Figure 4 Different induction rules for long-term potentiation in young and mature neurons. **a**, Theta-burst induction paradigms, exemplified here for a young granule cell. **b**, **c**, Plot of EPSP amplitude against time in a mature (**b**) and a young (**c**) granule cell. **d**, **e**, Average plots for mature (**d**) and young (**e**) cells. In mature cells, TBS₁ had no effect, whereas it led to significant LTP in young cells. Data points represent means of six consecutive events. **f**, **g**, Summary of LTP experiments. TBS₁ led to LTP only in young neurons, whereas TBS₂ led to LTP in both mature and young cells. All data were from perforated-patch recordings.

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important for the formation of new place fields. As the dendritic tree grows and the cell becomes fully integrated into the hippocampal circuit, a burst of action potentials will be necessary for LTP induction. Thus, the induction area will be much more confined, finally leading to a high spatial selectivity and a precise neuronal encoding of new spatial environment⁶.

Methods

Slice preparation

Adult male Wistar rats (2-3 months old, approximately 300 g) were reared in large cages with running wheels and climbing frames. The animals were anaesthetized by isoflurane and killed by decapitation, in accordance with institutional guidelines. Transverse 350-µm-thick slices were cut from the hippocampus using a vibratome (DTK-1000; Dosaka). For the dissection and the storage of the slices, a solution containing 64 mM NaCl, 25 mM NaHCO₃, 10 mM glucose, 120 mM sucrose, 2.5 mM KCl, 1.25 mM NaH₂PO₄, 0.5 mM CaCl₂ and 7 mM MgCl₂ (equilibrated with 95% O₂/5% CO₂) was used. Slices were incubated at 35 °C for 30 min and subsequently held at room temperature.

Immunohistochemistry and morphology

Cells were filled with biocytin (1 mg ml⁻¹) during whole-cell recording and subsequently fixed in 4% paraformaldehyde. After washing, tissue sections were incubated overnight at 4 °C with the primary antibody for PSA-NCAM (mouse, Chemicon, 1:400) together with 5% goat serum and 0.3% Triton X-100. The secondary antibody (goat anti-mouse-Alexa546, 1:200, Molecular Probes) was applied together with fluorescein isothiocyanateconjugated avidin-D (2 $\mu l\,ml^{-1},$ Vector) and 0.3% Triton X-100 for 24 h at 4 °C. After washing, slices were embedded in Prolong Antifade (Molecular Probes). Parvalbumin immunoreactivity was tested using an antibody against parvalbumin (mouse, Swant, 1:5,000) and an Alexa568-conjugated secondary antibody. Fluorescence was analysed with a confocal laser-scanning microscope (LSM 510, Zeiss).

PSA-NCAM immunofluorescence was measured in 1- μ m optical sections and quantified as intensity per pixel along the somatic cell membrane. Background intensity was subtracted, and the signal was normalized by the mean intensity of the brightest 1% of pixels within the image $(325 \times 325 \,\mu\text{m})$. The morphology of the cells was reconstructed in three dimensions from a stack of 50-100 confocal images using NeuronTracer software (Bitplane) running on an O2 workstation (Silicon Graphics). The total dendritic length was calculated as the sum of the lengths of all dendritic segments. Only cells with intact dendrites were used.

Electrophysiology

Slices were superfused with a physiological extracellular solution containing 125 mM NaCl, 25 mM NaHCO₃, 25 mM glucose, 2.5 mM KCl, 1.25 mM NaH₂PO₄, 2 mM CaCl₂ and 1 mM MgCl₂ (equilibrated with 95% O₂/5% CO₂). A concentration of 10 µM bicuculline methiodide was added in plasticity experiments, and 10 µM bicuculline and 10 µM 6-cyano-7-nitroquinoxaline-2,3-dione (CNQX) in membrane time constant experiments. Recordings were performed under visual control. Patch pipettes (4–9 M Ω) were pulled from thick-walled borosilicate glass tubing. Voltage signals were measured with an Axopatch 200A amplifier (Axon Instruments) in the I-clamp fast mode, filtered at 5 kHz, and digitized at 10 or 20 kHz using a 1401plus interface (Cambridge Electronic Design). For whole-cell recordings, pipettes were filled with a solution containing 130 mM K-gluconate, 20 mM KCl, 2 mM MgCl₂, 4 mM K₂ATP, 0.3 mM NaGTP, 10 mM Na₂phosphocreatine and 10 mM HEPES (pH 7.2). For perforated-patch recordings, a pipette solution containing 140 mM K-gluconate, 20 mM KCl, 5 mM MgCl₂, 10 mM HEPES, 0.1 mM EGTA, 50–100 µg ml⁻¹ amphotericin B and 0.2–0.4% DMSO was used. Bridge balance was used to compensate the series resistance of 10–50 M\Omega (whole cell) or 60–90 M Ω (perforated patch). Resting potential was determined in a subset of recordings with seal resistances $>5R_{in}$. The holding potential was set to -80 mV. To stimulate the medial perforant path, 2–4-M Ω pipettes filled with HEPES-buffered Na⁺-rich solution were used. The pipettes were placed in the centre of the molecular layer close to the dendrites of the recorded cell, and 200-µs pulses were applied at a frequency of 0.05 Hz. EPSP amplitude measurements were started 10-15 min after a stable access resistance had developed. Recordings were performed at 21-23 °C.

LTP induction protocols

The LTP induction paradigms consisted of brief presynaptic bursts (ten stimuli at 100 Hz) repeated ten times at theta frequency (5 Hz). This presynaptic stimulation was applied alone (TBS₀), paired with a slow postsynaptic depolarization that typically evoked a single action potential (TBS1), or paired with a train of brief stimuli that evoked a burst of postsynaptic action potentials^{25,29} (TBS₂; 5-ms delay between pre- and postsynaptic train). Paradigms were applied four times at 0.1 Hz. Only one TBS paradigm was tested per cell. If no LTP occurred, a maximal high-frequency stimulation protocol was applied (100 Hz, 1 s, four times, paired to postsynaptic depolarizations to 0 mV), and experiments were rejected if no LTP could be induced (8 of 24 mature and 4 of 23 young cells).

Data analysis

The input resistance was determined from the voltage at the end of 500- or 800-ms current pulses leading to approximately 5 mV hyperpolarization. To determine the membrane time constant, 200-400 voltage traces were averaged, and the logarithmically transformed decay was analysed by linear regression. For the quantification of LTP, 15 or 30 consecutive

EPSPs were averaged 10 min before the onset and 25–30 min after the end of the induction protocol. Peak amplitudes were measured from holding potential and data are given as mean ± s.e.m. unless specified differently. A Wilcoxon signed rank test or a Mann-Whitney test were used to assess statistical significance.

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