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In vitro characterization of gamma oscillations in the hippocampal formation of the domestic chick

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Abstract

Avian and mammalian brains have evolved independently from each other for about 300 million years. During that time, the hippocampal formation (HF) has diverged in morphology and cytoarchitecture, but seems to have conserved much of its function. It is therefore an open question how seemingly different neural organizations can generate the same function. A prominent feature of the mammalian hippocampus is that it generates different neural oscillations, including the gamma rhythm, which plays an important role in memory processing. In this study, we investigate whether the avian hippocampus also generates gamma oscillations, and whether similar pharmacological mechanisms are involved in this function. We investigated the existence of gamma oscillations in avian HF using in vitro electrophysiology in P0–P12 domestic chick (Gallus gallus domesticus) HF brain slices. Persistent gamma frequency oscillations were induced by the bath application of the cholinergic agonist carbachol, but not by kainate, a glutamate receptor agonist. Similar to other species, carbachol-evoked gamma oscillations were sensitive to GABA, AMPA/kainate and muscarinic (M1) receptor antagonism. Therefore, similar to mammalian species, muscarinic receptor-activated avian HF gamma oscillations may arise via a pyramidal-interneuron gamma (PING)-based mechanism. Gamma oscillations are most prominent in the ventromedial area of the hippocampal slices, and gamma power is reduced more laterally and dorsally in the HF. We conclude that similar micro-circuitry may exist in the avian and mammalian hippocampal formation, and this is likely to relate to the shared function of the two structures.

Introduction

Avian and mammalian brains have diverged for over 300 million years since their last common ancestor. The telencephalon of these two lineages is very differently organized (Jarvis et al., 2005), and there is an ongoing debate about whether the avian dorsoventricular ridge (DVR) is homologous to the mammalian neocortex, or whether they derive from different developmental regions (Striedter et al., 2014; Karten, 2015). Functionally, however, they seem to exhibit convergent abilities (Emery, 2004, 2006; Clayton & Emery, 2015). Unlike for cortex, there is a general consensus that the hippocampal formations (HF) of both lineages are homologous to each other, as they both derive from the medial pallium (Striedter, 2016). Although hippocampal function seems conserved in the two lineages, with important roles in memory, spatial navigation and feedback onto the hypothalamic-pituitary-adrenal axis (Kahn & Bingman, 2009; Mayer et al., 2013; Herold et al., 2015; Smulders, 2017), the anatomical organization of the two structures differs dramatically. Whereas the mammalian hippocampal formation has clearly delineated subdivisions such as the entorhinal cortex, subiculum, Ammon’s horn regions and dentate gyrus (Andersen, 2007), the avian HF has no such clear cell fields (Gupta et al., 2012; Herold et al., 2014; Striedter, 2016). Different subdivisional homologies have been proposed between avian and mammalian hippocampal formations, but the debate is still ongoing (Striedter, 2016). The existence of such functionally similar, yet anatomically dissimilar structures between birds and mammals leads us to the question: How do such divergent structures perform such similar functions?

Much has been made about the macroscopic organization of the mammalian HF (and indeed the cortex) in discrete cell layers with regard to its function (Andersen, 2007), but the fact that the avian HF can carry out the same functions without the clearly delineated cell layers suggests that this is not the case. However, it is still possible that a similar connectional organization underlies the divergent macroscopic organizations to deliver a similar function (Striedter, 2016). Again, connections between avian hippocampal subdivisions have been studied and different hypotheses put forward about the
similarities and differences between the avian and mammalian HF (Szekely, 1999; Hough et al., 2002; Kahn et al., 2003; Atoji & Wild, 2004, 2006). However, to date, nobody has yet studied local circuit functionality in the avian HF and compared it to that in the mammalian hippocampus. The study we present here aims to make a start at better understanding local circuit properties in the avian HF and relate these to what we know about local circuit properties in the mammalian system.

One prominent feature of the mammalian hippocampus is its ability to generate discrete forms of neuronal oscillations. In particular, gamma rhythms (30–80 Hz) are a physiological feature of the mammalian hippocampus (Traub et al., 1996), which play an important role in memory processing. They are posited to provide a substrate for organization and binding of different perceptions (Singer, 1996). In vitro studies using pharmacologically driven models (e.g. application of carbachol or kainate) of gamma oscillations have revealed the circuitry responsible for this activity in animals. While pyramidal neurons fire sporadically during gamma oscillations (Fisahn et al., 1998; Gloveli et al., 2005), in contrast, fast-spiking parvalbumin-expressing interneurons fire on each gamma cycle and are strongly phase locked to the oscillation (Gloveli et al., 1997; Hajos et al., 2004). Further studies have demonstrated that these perisomatic targeting interneurons are critical for the generation of gamma oscillations in the rodent CA3 region (Oren et al., 2006, 2010; Mann & Mody, 2010). As pyramidal-interneuron network gamma oscillations require phasic excitatory synaptic input onto interneurons, it is unsurprising that the selective reduction in GluA1- or GluA4-containing AMPA receptors on hippocampal interneurons (Fuchs et al., 2001, 2007) has profound implications for network gamma oscillations. It is this negative feedback loop (Wang & Buzsáki, 1996) between pyramidal neurons and parvalbumin-expressing fast-spiking interneurons (Freund & Buzsáki, 1996) that generates the gamma oscillations in mammals. Gamma rhythm generation in the mammalian HF is most prominent in the CA3 region (Csicsvari et al., 2003) and is communicated to other regions of the HF and beyond from there.

The existence of gamma oscillations has not yet been explored in the avian HF. They have, however, been described in the avian optic tectum (OT) in the midbrain (Sridharan & Knudsen, 2015). In barn owls (Tyto alba), a brief visual stimulation that engages the bird’s attention triggers a brief gamma oscillation in the OT (Sridharan et al., 2011). This can be mimicked in vitro in OT slices taken from poultry chicks (Gallus gallus domesticus) by providing a brief electrical stimulation to the region that receives retinal inputs (Goddard et al., 2012). The local OT circuit generating these oscillations depends on GABA_A receptors (both to generate the rhythm and to set the peak frequency of this activity), NMDA glutamate receptors (to prolong the oscillation following termination of the stimulus) and on acetylcholine receptors. The GABAergic neurons involved in the gamma-generating circuitry seem to be parvalbumin+ inhibitory interneurons (Goddard et al., 2012), as is also the case for mammalian CA3 gamma oscillations (Cardin et al., 2009; Sohal et al., 2009).

It is clear that avian brains can generate gamma oscillations, just like mammalian brains can. In this study, the questions we aim to address through in vitro electrophysiology are whether (i) the avian HF also contains local circuits that generate and sustain gamma frequency oscillations, (ii) these HF micro-circuits exhibit pharmacological similarities to the gamma-generating circuits in the mammalian hippocampus, and (iii) there is a subregional differentiation in the generation of gamma rhythms in the avian HF.

Materials and methods

Animals

For the pharmacological characterization, Welsummer breed chicken eggs (Gallus gallus domesticus) were obtained from a local breeder and incubated in the Comparative Biology Centre, Newcastle University. A total of 17 birds were used for this study. Upon hatching, we housed them as a group in a heated cage, provided with food and water ad libitum, and subjected to a regular 12-h light/dark cycle (0800-2000 hours).

For the spatial mapping, we obtained 1-day-old broiler chicks (Ross 308) from a commercial hatchery in Thirsk, UK, which provided a more reliable supply of birds. A total of 18 birds were used in this study. They were transported to the laboratory by car, where they were housed in floor pens measuring 120 × 70 cm. The floor of the pens was covered with wood chips, and each pen contained a food hopper, a water hopper and a plastic shelter. Chicks were subjected to a 14L : 10D cycle using uncovered fluorescent lights, and the temperature of the laboratory was maintained at 26–30 °C using room heaters. Water was provided ad lib., as were chick starter crumbs. Chicks also received mealworms Tenebrio molitor once a day in their home cages. All subjects were marked with non-toxic ‘child-friendly’ coloured marker pens, and their weights were monitored daily for welfare purposes.

To obtain brains for slice preparation, we anaesthetized chicks aged P0–P12 days (of either sex) by the brief inhalation of isoflurane (0.05%) and euthanized them by cervical dislocation. We carried out all procedures in accordance with UK Home Office guidelines set out in the Animals (Scientific Procedures) Act 1986. The protocol was approved by Newcastle University’s Animal Welfare and Ethical Review Board (Project ID No 380).

Slice preparation

Following decapitation, the brain was immediately removed and placed in ice-cold (~4 °C) sucrose-based artificial cerebrospinal fluid (ACSF) solution composed of the following (in mM): sucrose, 25; D-glucose, 10; NaHCO_3, 24; KCl, 3; MgCl_2, 5; CaCl_2, 2; NaH_2PO_4, 1.25; MgSO_4, 2, continuously bubbled with carbogen (95% O_2, 5% CO_2). The forebrain was then mounted on a steel plate, and 400-μm thin coronal sections were cut in the cold, sucrose-based ACSF solution using a Leica VT1000 microtome (Leica Microsystems, Germany). The hippocampus was then dissected free from the surrounding brain regions and transferred to a holding chamber containing ACSF at room temperature. For recordings, slices were then transferred to an interface style chamber and maintained at the interface containing standard ACSF, composed of the following (in mM): NaCl, 126; KCl, 3; NaH_2PO_4, 1.25; NaHCO_3, 24; MgSO_4, 1; CaCl_2, 1.2; glucose, 10. The circulating ACSF solution was continuously bubbled with humidified carbogen (95% O_2, 5% CO_2) to maintain a pH of 7.4 and warmed to 34 ± 1 °C. Slices were allowed to equilibrate for at least 30 min prior to any recording.

Recording and data acquisition

Glass microelectrodes for extracellular local field potentials (LFP) recordings were pulled from thin-walled borosilicate tubing (1.2 mm O.D. × 0.94 I.D., Harvard Apparatus Ltd., Kent, UK) to a resistance of 2–5 MΩ using a PP-83 Narishige puller (Narishige Scientific Instrument Lab, Tokyo, Japan) and filled with standard ACSF. Up to three microelectrodes were then placed in pipette holders.
Electrophysiology data analysis

Power spectra of the recorded LFP were calculated from 60-s epochs of recorded field activity using a fast Fourier transform (FFT) in the AXOGRAPH X software package. We quantified it by examining frequency and amplitude corresponding to peak in the gamma band of the power spectrum, while area power was determined as the area within 20 and 80 Hz. This gave us peak power (μV²/Hz) at an oscillation frequency (Hz) as well as the area power (total power of the oscillation (μV²/Hz-kHz) in the gamma frequency band.

Pharmacological manipulations

All drugs were bath applied at known concentrations previously used in rodent brain slice experiments: Kainic acid [(2S,3S,4S)-3-carboxymethyl-4-(prop-1-en-2-yl)pyrrolidine-2-carboxylic acid], 50–800 nM; Carbachol [2-[(Aminocarboxyloxy)-N,N,N-trimethylethanaminium), 1–50 μM; D-AP5 [2-amino-5-phosphonopentanoic acid], 1–50 μM; Phenylpropanoate, 1 μM; Atropine [(RS)-(8-Methyl-8-azabicyclo[3.2.1]oct-3-yl) 3-hydroxy-2-phenylpropanoate], 1 μM; Phystostigmine [(3aR, 8aS)-1,3a,8-Trimethyl-1H,2H,3H,5aH,8H,8aH-pyrrolo[2,3-b]indol-5-yl N-methylcarbamate], 10 μM; NBQX [2,3-dioxo-6-nitro-1,2,4-tetrahydropyridine-5-carboxyl-phosphoric acid], 10 μM; Pirenzepine [(S)-N,N,N-trimethyl-4-(prop-1-en-2-yl)pyrrolidine-2-carboxylic acid], 50–100 μM; NBQX [2,3-dioxo-6-nitro-1,2,4-tetrahydropyridine-5-carboxyl-phosphoric acid], 10 μM; Vanadate [sodium vanadate], 1–50 μM; 11-[(4-methylpiperazin-1-yl)acetyl]-5,11-dihydro-6H-pyrrolo[2,3-b][1,4]benzodiazepin-6-one, 1 μM; D-AP5 [2-amino-5-phosphonopentanoic acid], 50 μM. All were obtained from Tocris Cookson (Bristol, UK). Salts for ACSF and sucrose-ACSF were from BDH-Merck.

Immunohistochemistry

Brains were removed using the same method as explained above, but then placed into 4% Paraformaldehyde solution immediately after dissection before being transferred to 30% sucrose solution after 2 days for cryoprotection. Brains were then submerged in optimal cutting temperature (OCT) compound and cut to 40 μm on a MICROM HM560 cryostat before being stored at 0–4 °C in PBS. Sections were exposed to primary mouse anti-parvalbumin antibody (Sigma p3088) at 1:2000 concentration in PBS+1% Triton X-100 and 3% Horse serum and left to incubate overnight at 4 °C. The secondary horse anti-mouse antibody was applied at room temperature for 2 h at 1:200 concentration in PBS. Sections were incubated in horseradish peroxidase (Ready-To-Use Streptavidin Horseradish Peroxidase, Vector Labs, UK) for 2 h at room temperature. The chromogenic reaction was then developed using DAB (SigmaFast™ DAB tablets, Sigma-Aldrich, UK) for 5 min. Sections were mounted onto gelatine-subbed glass slides (Fisher Scientific, UK). After air-drying overnight, slides were washed in distilled water and sections were then dehydrated by incubation in sequential ethanol concentration gradients; starting from 70, 95 and 100%; each for 5 min each, with a further 5 min in another 100% ethanol gradient. Sections were then bathed in Histoclear (National Diagnostics, USA) for 5 min twice before being coverslipped using Eukitt® (Sigma-Aldrich, UK).

Statistical analysis

Statistical analyses were performed in IBM SPSS v.23. Gamma oscillations are analysed as the Area Under the Curve power in the gamma range (20–80 Hz). Because of large variation in absolute values from experiment to experiment, the data were expressed as a percentage of the value before the treatment started and then log10-transformed. For the regional mapping, data were expressed as a percentage of the ventromedial region power and log10-transformed. Data are analysed with paired t tests when comparing two treatments, or repeated-measures ANOVA when more than two treatments are compared. For the regional mapping, we used the Generalized Estimating Equations function in SPSS, because it can run with missing data, which the repeated-measures ANOVA function cannot. The output from this analysis is reported as Wald’s χ². Descriptive statistics are expressed as Mean ± Standard Error of the Mean.

Results

Gamma induction in avian hippocampal slices

Low concentrations of the kainate receptor agonist kainic acid are known to induce robust oscillations in mammalian hippocampus and associated neocortical structures. Interestingly, 100 nM of kainic acid was unable to elicit gamma activity in the avian HF (on average 1.46 (±1.42) times the area power of baseline, t_{7} = 1.09, P = 0.31; without any change in peak frequency, t_{7} = 0.40, P = 0.70; n = 8 slices from three birds). In three of these slices (from one bird), the concentration of kainic acid was slowly increased to 800 nM, but no significant change in gamma power ensued (on average 3.11 (±1.57) times baseline area power, t_{7} = 1.54, P = 0.26; no change in peak frequency, t_{7} = 0.16, P = 0.89).

As we had no immediate success with kainic acid, we examined the impact of carbachol, a broad-spectrum cholinergic agonist, which also induces robust gamma oscillations in mammalian hippocampal slices (Fisahn et al., 1998). Unlike kainic acid, carbachol induced robust gamma oscillations in the avian hippocampal formation, in a dose-dependent manner (Fig. 1; F_{4,12} = 18.14, P < 0.001, n = 16 slices from six birds). On average, 10 μM carbachol led to
an area power that was 22.80 (±1.51) times the baseline area power, with a mean peak frequency of 36 ± 1.4 Hz. Once initiated, the network activity was stable for few hours (2–3 h tested).

To investigate whether muscarinic cholinergic receptors play a role in the rhythmogenesis of gamma oscillations, we tested the effect of atropine, a generic muscarinic antagonist. After inducing gamma oscillations using 10 μM of carbachol, bath application of 1 μM of atropine eliminated the oscillations (Fig. 2; 18.5% of carbachol-induced area power, \( t_6 = 3.27, P = 0.017 \); reduction in peak frequency from 40 ± 2.2 to 28 ± 2.8 Hz, \( t_6 = 3.03, P = 0.023 \); in seven slices from three animals in which the area power increased again after atropine washout). Further, we aimed to investigate the role of M1 subtype of the muscarinic receptors. Bath application of 10 μM of scopolamine, an M1-specific antagonist, inhibited the carbachol-induced gamma oscillations (15% of carbachol-induced area power, \( t_5 = 4.44, P = 0.007, n = 6 \) slices from animals in which area power increased again upon washout). There was an almost significant drop in peak frequency from 34 ± 3.8 to 26 ± 1.5 Hz (\( t_5 = 2.51, P = 0.054 \); Fig. 2). This was confirmed with bath application of 1 μM of pirenzepine, another M1-subtype-specific muscarinic antagonist. Area under the curve did not significantly decrease (on average 57% of carbachol-induced gamma power, \( t_8 = 1.76, P = 0.12, n = 9 \) slices from two animals in which area power increased again after washout).
power increased after washout of pirenzepine), but the peak frequency did drop below the gamma range (from 36 ± 1.2 to 28 ± 2.0 Hz, \( t_9 = 3.94, P = 0.004 \); Fig. 2).

Involvement of GABAergic inhibition and Glutamatergic excitation?

Gamma activity in the mammalian hippocampus is known to be critically dependent on the rhythmic output of populations of GABAergic interneurons (Whittington et al., 1995). To test whether this hypothesis held in the avian HF, we first induced gamma activity with 10 \( \mu \)M of carbachol. Addition of 1 \( \mu \)M of Gabazine, a specific GABA\(_A\) receptor antagonist, resulted in almost complete abolition of the gamma activity (18% of carbachol-induced area power, \( t_9 = 3.30, P = 0.023 \); n = 6 slices from four animals in which area power increased after washout), reducing the peak frequency from 37 ± 1.3 to 30 ± 2.8 Hz (\( t_9 = 2.58, P = 0.05 \); Fig. 2).

Fast phasic glutamatergic transmission has been demonstrated to influence persistent gamma activity in the mammalian hippocampus (Whittington et al., 2000; Bartos et al., 2007). Thus to investigate the role of AMPA receptors in the avian HF, we tested the effect of NBQX, an AMPA/kainate receptor antagonist. We again induced gamma oscillations by bath application of 10 \( \mu \)M of carbachol. Application of 20 \( \mu \)M of NBQX resulted in complete abolition of the gamma activity (16% of carbachol-induced area power, \( t_9 = 4.74, P = 0.001 \), n = 9 slices from four animals; Fig. 2). The peak frequency is also significantly reduced from 38 ± 2.6 to 25 ± 3.2 Hz (\( t_9 = 4.93, P = 0.001 \)). Finally, to investigate the role of NMDA receptors in the gamma rhythmogenesis, we tested the effect of D-AP5, an NMDA receptor antagonist, on the gamma activity induced by 10 \( \mu \)M of carbachol. After inducing gamma oscillations by a bath application of carbachol, we bath applied 50 \( \mu \)M of D-AP5. Interestingly, whereas D-AP5 slightly reduced the peak frequency from 40 ± 2.2 to 36 ± 1.4 Hz (\( t_9 = 2.53, P = 0.032 \)), the area power in the gamma range significantly increased (1.8 times the carbachol-induced area power, \( t_9 = 3.54, P = 0.006 \), 10 slices from five animals; Fig. 2).

Mapping of gamma power in the different subdivisions

We recorded the gamma power in six different areas in the slice and analysed these with a Generalized Estimating Equations approach. Gamma power was highest in the ventromedial region and became progressively weaker as the electrode was moved dorsally (\( \chi^2 = 35.32, P < 0.001 \)) and laterally (\( \chi^2 = 12.41, P < 0.001 \); nine slices from six animals; Fig. 3). There was no interaction between the two spatial axes (\( \chi^2 = 2.71, P = 0.26 \)). For the four slices where we were able to record in the ventral-most tip of the slice, the gamma power was also lower than in the ventromedial region (\( t_3 = 3.52, P = 0.039 \)).

Parvalbumin staining

The anti-parvalbumin antibody clearly stained a population of PV\(^+\) cells throughout the hippocampal formation (Fig. 3A inset).

Discussion

Our main finding is that, like the mammalian hippocampus, the avian hippocampal formation is capable of endogenously generating gamma oscillations when stimulated with a cholinergic agonist. In terms of peak frequency (c. 40 Hz), this cholinergic rhythm in avian slices is similar to the peak frequency of gamma observed in a number of pharmacologically induced (e.g. by kainate, acetylcholine receptor agonists, or metabotropic glutamate receptor agonists) gamma oscillations in the mammalian hippocampus (Fisahn et al., 1998, 2004; Palhalmi et al., 2004; Fisahn, 2005). Both fast inhibition mediated by GABA\(_A\) receptors and fast excitation via predominantly AMPA receptors are a requirement for generating and maintaining this oscillation. This suggests that similar functional micro-circuitry, with similar pharmacological properties, may be present in the avian HF as in its mammalian counterpart, despite a clearly different cytoarchitecture of the structures. Further studies looking at single cells and their connections will allow us to fully characterize this circuitry.

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**Similarities to mammalian gamma oscillations**

Gamma oscillations were robustly induced using a direct cholinergic agonist. This suggests that there must be a natural cholinergic input into the avian HF that may contribute to the generation of gamma oscillations in vivo. The cholinergic antagonists we used suggest a necessary involvement of M1 muscarinic receptors in this process. This finding mimics observations made in the mouse hippocampus in which gamma oscillations were absent in M1 receptor-deficient mice (Fisahn et al., 2002). It is different, however, from the brief gamma oscillations triggered in the chickmidbrain, which depend on nicotinic, but not muscarinic acetylcholine receptors (Bryant et al., 2015). Interestingly, in adult pigeons, the density of M1 receptors is very low throughout the hippocampus (Kohler et al., 1995; Herold et al., 2014). Nevertheless, blocking of M1 receptors by intramuscular injection of scopolamine interferes with short-distance homing in pigeons, a task known to rely on hippocampal-dependent spatial memory (Kohler et al., 1996). It is therefore possible that mAChRs are only expressed by a small, but crucial population of neurons [e.g. on excitatory neurons (Fisahn et al., 2002)] and that activation of these receptors would be sufficient to indirectly activate interneurons to produce gamma activity.

The fact that a cholinergic agonist can induce gamma oscillations in the avian HF suggests that there must be an innate cholinergic innervation of the HF. In mammals, there is a strong cholinergic innervation from the septum into the hippocampus (Hasselmo & Bower, 1993). In birds as well, the HF and the septal nuclei are reciprocally connected (Atoji & Wild, 2004; Montagnese et al., 2004), but the density of the cholinergic innervation of the HF is lower and originates mainly in the nucleus of the diagonal band (Krayniak & Siegel, 1978; Krebs et al., 1991; Montagnese et al., 2004).

The circuit that generates the gamma rhythms includes both excitatory and inhibitory synapses, as indicated by the interruption of the oscillations by both a GABA<sub>A</sub> receptor antagonist and an AMPA receptor antagonist. This suggests that a similar circuit to the proposed pyramidal-interneuron gamma (PING) oscillator may also exist in the avian hippocampal formation. The presence of parvalbumin<sup>+</sup> cells in the avian HF is consistent with this, although further work needs to be done to ascertain whether they are also fast spiking, as is the case in the mammalian hippocampus (Freund & Buzsáki, 1996; Gioveli et al., 1997; Hajos et al., 2004). Indeed, circuits of local inhibitory interneurons and pyramidal projection neurons have been described in both the chick and pigeon hippocampus using Golgi staining methods (Tombol et al., 2000a,b). A similar circuit also produces gamma oscillations in the chick midbrain (Goddard et al., 2012). However, in those oscillations, which were triggered by a brief electrical stimulus to the slice, D-AP5 reduced the duration of the gamma oscillations, implying that they are dependent on NMDA receptors for their maintenance (Goddard et al., 2012; Sridharan & Knudsen, 2015). The carbachol-induced gamma oscillations in the hippocampus were not reduced by D-AP5, however. To the contrary, the power in the gamma band slightly increased, even though the peak frequency was slightly decreased (but remained in the gamma band). This is consistent with carbachol-induced gamma oscillations in rodent hippocampus, which are also not dependent on NMDA receptors (Fisahn et al., 1998).

We do not know what may have caused the slight (but significant) increase in gamma power, but similar effects have been seen in hippocampal and cortical gamma oscillations in vivo (Pinault, 2008; Lazarewicz et al., 2010). The strongest gamma power by far was recorded in the most ventromedial region of the rostral HF, which is often referred to as Vm (Herold et al., 2014). This region has been compared by some to CA3, and by others to the DG of mammals, if any such equivalences can be made at all (Striedter, 2016). What is sure is that in terms of the distribution of different synaptic neurotransmitter receptors (glutamatergic, GABAergic, cholinergic, noradrenergic, serotonergic and dopaminergic), this region is most similar to CA3 (Herold et al., 2014). The distribution of receptor binding sites is probably a poor feature to use to derive evolutionary homologies between brain areas, as it is very sensitive to selection pressures that might change the function of a cell population. For that same reason, however, it is a good proxy for functional equivalences, and it is therefore not surprising that the area in the avian HF with the highest pharmacological similarity to CA3 is also the region that generates endogenous gamma rhythms, as does CA3 in mammals.

Of course, in mammals, other regions within the limbic system are also known to be capable of independently generating gamma activity. Subfields of the hippocampus such as the dentate gyrus (Poschel et al., 2002; Towers et al., 2002), CA1 (Traub et al., 2003; Middleton et al., 2008; Piersen et al., 2014; Craig & McBain, 2015) and the subiculum (Colling et al., 1998; Eller et al., 2015) have been demonstrated to be capable of acting as independent gamma generators. In the rodent parahippocampal region, the medial entorhinal cortex has also been shown to be a potent source of gamma frequency oscillations (Cunningham et al., 2003; Middleton et al., 2008; Colgin et al., 2009) capable of entraining network oscillations in CA1. Further work therefore remains to be done to pinpoint the areas in the avian HF that generate gamma oscillations, and those that follow them.

Given the ubiquity of gamma oscillation generators throughout the mammalian brain (Headley & Paré, 2017), and given the existence of gamma oscillations in the avian midbrain (Sridharan & Knudsen, 2015), it is very likely that gamma oscillations can be generated in other parts of the avian forebrain as well. This remains to be explored. Interestingly, however, within the hippocampal formation, data from the spatial studies (Fig. 3) would indicate a localized network in the ventral-medial avian HF capable of supporting gamma oscillations. Data are now emerging from rodent studies, which have demonstrated coupling of place cell firing to gamma oscillations (Lasztoczi & Klausberger, 2016). Previous studies in freely moving homing pigeons to determine the presence of spatially responsive neurons have only reported unstable firing fields (Siegel et al., 2002, 2005, 2006; Bingman et al., 2003; Hough & Bingman, 2004, 2008; Kahn et al., 2008). Interestingly, the recordings in these studies were conducted in more lateral areas of the HF. Therefore, it may be the case that medial structures of the avian HF, which we have shown to be potent gamma-generating regions, are the site of location for avian place cells. This hypothesis, of course, remains to be tested.

**Differences from mammalian gamma oscillations**

This present study did reveal differences as compared to the mammalian hippocampal slice preparation, however. Unlike in mammalian brain slices (Hajos et al., 2000; Fisahn et al., 2004; Cunningham et al., 2006), kainate was not able to induce gamma oscillations in the avian hippocampus. If this is indeed a feature of the avian HF in general, this may indicate a difference in the conditions that may trigger gamma oscillations in vivo. In mammals, kainate-induced gamma oscillations arise from the influence of kainate receptors (KARs) on interneuronal and excitatory cell function. Activation of KARs on both somatodendritic and axonal compartments of rodent hippocampal interneurons will lead to depolarization...
Implications for brain evolution

The discovery of robust gamma oscillations in the avian HF in vitro confirms that the avian and mammalian hippocampal formations may well share functional micro-circuitry. It also reinforces the idea that certain rhythmic patterns are important for proper functioning of brains, and the frequencies of these rhythms are found across a range of species (Buzsaki et al., 2013). It is difficult at this point in time to tell whether the similarity in pharmacology implies similar micro-circuits, and whether these micro-circuits are homologous or the result of convergent evolution, but they strongly suggest that in order for brain areas to carry out similar behavioural functions, they also need similar physiological functions (and their underlying circuitry). Our first descriptions of gamma oscillations in the avian HF open up a great number of new questions. On the one hand, they beg the question of whether avian hippocampal gamma is triggered under the same conditions in vivo as it is in mammals and, therefore, whether gamma oscillations play a similar role of synchronizing populations of neurons with each other for efficient communication (Colgin, 2016). On the other hand, we have only started to scratch the surface of the actual structure of the local circuits in the HF that generate these (and other) hippocampal rhythms. Further work combining pharmacology, patch-clamp recordings and detailed cellular anatomy will be required to be able to compare the mammalian and avian local circuits in detail. In the mammalian brain, gamma rhythms are generated by different circuitry in different brain areas. Comparing mammalian and avian hippocampal circuits will give us deeper insights into the evolution of these circuits and how they relate to the specific functions of the brain areas in which they occur.

Conclusion

We have shown that the avian hippocampal formation can generate gamma oscillations in vitro and that these share a number of features with similar oscillations in the mammalian hippocampus. The very existence of local circuits for generating gamma oscillations in the avian HF points to similarity in physiological processes and features that are required for similar information processing functions that exist in the avian and mammalian structures.

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Conflict of interest

The authors declare that there are no conflict of interests.

Author contributions

MOC and TVS designed the study and trained the others in data collection. PD, NML and JC collected the data. PD and TVS analysed the data. PD and TVS wrote the first draft of the manuscript, and the other authors commented on subsequent versions. All authors approved the submission of the final manuscript.

Data accessibility

Data used in the statistical analyses are uploaded with the manuscript. Raw data files are archived on the laboratory servers and are available upon request.

References

Abellan, A., Desflis, E. & Medina, L. (2014) Combinatorial expression of Lef1, Lhx2, Lhx5, Lhx9, Lmo3, Lmo4, and Prox1 helps to identify com-
Atoji, Y. & Wild, J.M. (2004) Fiber connections of the hippocampal forma-
tion and septum and subdivisions of the hippocampal formation in the pigeon as revealed by tract tracing and kainic acid lesions. J. Comp. Neu-
rol., 475, 426–461.
Bartos, M., Vida, I. & Jonas, P. (2007) Synaptic mechanisms of synchroni-

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European Journal of Neuroscience, 1–9.


