

DR JEREMY LAMBERT (Orcid ID : 0000-0002-4204-3602)

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Corresponding Author Email Id : j.j.lambert@dundee.ac.uk

Endogenous neurosteroids influence synaptic GABA_A receptors during post-natal development.

Delia Belelli, Adam R. Brown, Scott J. Mitchell, Ben G. Gunn, Murray B. Herd, Grant Phillips, Mohsen Seifi, Jerome D. Swinny, Jeremy J. Lambert.

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Abstract

GABA plays a key role in both embryonic and neonatal brain development. For example, during early neonatal nervous system maturation, synaptic transmission, mediated by GABA_A receptors (GABA_ARs), undergoes a temporally specific form of synaptic plasticity, to accommodate the changing requirements of maturing neural networks. Specifically, the duration of miniature inhibitory postsynaptic currents (mIPSCs), resulting from vesicular GABA activating synaptic GABA_ARs, is reduced, permitting neurons to appropriately influence the window for postsynaptic excitation. Conventionally, programmed expression changes to the subtype of synaptic GABA_AR are primarily implicated in this plasticity. However, it is now evident that in developing thalamic and cortical principal- and inter-neurons an endogenous neurosteroid tone *e.g.* allopregnanolone, enhances synaptic GABA_AR function. Furthermore, a cessation of steroidogenesis, due to a lack of

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substrate, or a co-factor, appears primarily responsible for early neonatal changes to GABA-ergic synaptic transmission, followed by further refinement, which results from subsequent alterations of the GABA_AR subtype. The timing of this cessation of neurosteroid influence is neuron specific, occurring by postnatal day 10 (P10) in thalamus, but about a week later in cortex. Neurosteroid levels are not static, but change dynamically in a variety of physiological and pathophysiological scenarios. Given that GABA plays an important role in brain development, abnormal perturbations of neonatal GABA_AR-active neurosteroids, may have a considerable immediate, but also longer term impact, upon neural network activity. Here we review recent evidence that changes in neurosteroidogenesis substantially influence neonatal GABA-ergic synaptic transmission. We discuss the physiological relevance of these findings and how interference of neurosteroid-GABA_AR interaction early in life may contribute to psychiatric conditions later in life.

Introduction

From an early embryonic age, GABA-ergic signalling plays a fundamental role in neuronal development, influencing neuronal proliferation, migration and differentiation and is important in the establishment of neuronal networks¹⁻⁴. These effects of GABA are mediated by ionotropic GABA_A receptors (GABA_ARs), or by G-protein coupled GABA_B receptors. Certain steroids are potent, endogenous, positive allosteric modulators (PAMs) of the GABA_AR⁵⁻⁸ (neuroactive steroids). Furthermore, such steroids may be synthesised locally in the brain, or spinal cord and are classed as neurosteroids. Here we consider the role of neurosteroids and GABA in the developing neonatal central nervous system (CNS).

Neurosteroids: endogenous modulators of the GABA_AR.

The finding in 1966 by Craig and colleagues that the anticonvulsant properties of certain endogenously occurring steroids could be distinguished from classical hormonal actions⁹, paved the way nearly two decades later to the discovery that the synthetic general anaesthetic steroid alphaxalone¹⁰ and subsequently certain endogenous metabolites of progesterone (5 α -pregnan-3 α -ol-20-one; allopregnanolone) and deoxycorticosterone (5 α -pregnan-3 α ,21

diol-20-one; 5 α -THDOC) act as potent and selective positive allosteric modulators (PAMs) of the major inhibitory receptor in the mammalian brain, the GABA_AR^{5,10-14} (Figure 1).

GABA_ARs belong to the transmitter-gated ion channel superfamily and are composed of five subunits arranged around a central pore to form an anion-conducting ion channel^{15,16}. Low nM aqueous concentrations of these neuroactive steroids enhance the actions of GABA by promoting the anion-conducting state of the associated channel^{11,14,17,18}. Their activity at such low concentrations suggests the presence of a high affinity steroid binding site on the GABA_AR. However, these steroids are highly lipophilic, resulting in relatively high membrane concentrations in close proximity to the proposed transmembrane binding site on the GABA_AR^{19,20}. Therefore, the steroid may have a relatively low affinity for the receptor, but be optimally concentrated, locally in the membrane, leading to an increased probability that the steroid will occupy the receptor binding site^{19,20}. The GABA_AR is the target for clinically important drugs, *e.g.* benzodiazepines and certain general anaesthetics, which enhance the GABA_AR function⁵. In common with such drugs, administration of neuroactive steroids produce anxiolytic, analgesic, anticonvulsant and sedative effects, with higher doses capable of inducing a general anaesthetic state^{5,8}.

Initially, endocrine glands such as the adrenals and ovaries were considered the exclusive source of steroid, necessitating it to cross the blood brain barrier to influence neural activity. Subsequently, compelling evidence emerged for *de novo* brain synthesis *i.e.* neurosteroids²¹. Hence, these endogenous GABA_AR-modulators may potentially act in an endocrine, paracrine, or autocrine manner, to influence neuronal signalling in various physiological and pathological situations. Indeed, neurosteroid levels are dynamically changed in a variety of physiological (*e.g.* development, puberty, pregnancy, stress, ovarian cycle) and pathophysiological (*e.g.* major depression, postpartum depression, premenstrual tension, panic attacks and schizophrenia) scenarios⁵.

In the brain and spinal cord the GABA_AR-active steroids such as allopregnanolone and 5 α -THDOC may *a)* originate from peripheral sources *e.g.* placenta, ovaries and adrenals necessitating them crossing the blood brain barrier²²; *b)* be derived from their peripheral hormonal precursors, progesterone and deoxycorticosterone, or *c)* be synthesised *de novo* from

cholesterol *via* a series of multi-enzymatic steps²³. Diverse approaches including *in situ* hybridization, immunohistochemistry, gas chromatography mass spectrometry, electrophysiology and behaviour, purport that neurosteroids, synthesised by neurons and/or glia may achieve levels sufficient to influence in a paracrine and/or autocrine fashion GABA_AR signalling from the onset of embryonic GABAergic transmission into maturity^{5,24-29}. The suite of enzymes that synthesise neurosteroids exhibit regional and cellular-selective expression patterns, which change within discrete temporal windows and are susceptible to external challenges, *e.g.* stress^{23,30-33}. In the CNS the production of GABA_AR-active neurosteroids from cholesterol (Figure 1) first requires the translocation of the steroid across the mitochondrial membrane by translocator protein 18 kDa (TSPO)^{34,35}. Cholesterol is then metabolised to pregnenolone by the mitochondrial P450 side-chain cleavage enzyme CYP11A1, this metabolite is then exported across the mitochondrial membrane, where it may be converted to GABA_AR-active neurosteroids such as allopregnanolone following three sequential enzyme reactions catalysed by 3 β -hydroxysteroid dehydrogenase (3 β -HSD) to form progesterone, 5 α -reductase (5 α -R) to produce 5 α -dihydroprogesterone (5 α -DHP) and 3 α -hydroxysteroid dehydrogenase (3 α -HSD) to form allopregnanolone (Figure 1)⁷. Note both in the CNS and the periphery the enzymes 5 α -R and 3 α -HSD may sequentially participate in the conversion of deoxycorticosterone derived primarily from the adrenals into the GABA_AR-active 5 α -THDOC²². We have recently reviewed the expression of 5 α -R and 3 α -HSD in mammalian brain²⁴, (see also³⁰⁻³³). As GABA plays a crucial role in neurodevelopment, during this critical time of neonatal development it is conceivable that the changing levels of neurosteroids, from CNS-located paracrine, or autocrine sources, may influence the establishment of mature neuronal circuits and communication. Additionally, neonatal endocrine glands such as the adrenals may provide the brain and spinal cord with either the GABA_AR-active steroids *per se*, or their precursors^{22,36,37}.

GABA_ARs.

The GABA_AR subunits are drawn from a repertoire of 19 gene products belonging to distinct families including: α 1-6; β 1-3; γ 1-3, δ , ϵ , θ , π and ρ 1-3^{15,16}. In the adult mammalian brain this diversity underpins the expression of 20-30 GABA_AR subtypes, that are uniquely distributed and consequently influence particular behaviours³⁸⁻⁴⁰. Although neuroactive steroids are highly selective for GABA_ARs, they exhibit limited GABA_AR subtype selectivity⁵. However, the GABA_AR subunit composition does influence *a*) neuronal subcellular location *e.g.* synaptic vs extra/peri-synaptic expression⁴¹ *b*) the impact of enzymes (*e.g.* kinases) upon receptor function/location/expression⁴²⁻⁴⁴, *c*) the pharmacology of the receptor *e.g.* benzodiazepines and certain general anaesthetics^{15,40} and *d*) the physiological properties of the receptor, including the kinetics of receptor activity influenced by rates of deactivation, desensitization, and GABA affinity⁴².

The neuronal location of the receptor influences the GABA inhibitory signalling repertoire of the neuron. Receptors clustered in the post-synaptic domain primarily serve to mediate a fast, phasic form of inhibition in response to relatively high local concentrations of neurotransmitter, which occur in the synapse following the vesicular release of GABA⁴¹. By contrast, receptors in peri-synaptic and extra-synaptic locations are activated by ambient concentrations of GABA to mediate a sustained, or tonic form of neuronal inhibition⁴¹. However, in some neurons during physiological bursts of high frequency presynaptic stimulation the spill-over of GABA from the synapse is sufficient to additionally engage these extrasynaptic receptors to produce a greatly prolonged form of slow phasic inhibition⁴⁵⁻⁴⁷. The majority of GABA_ARs contain two α and two β subunits together with a single copy of the γ 2 subunit. Synaptic receptors often contain the γ 2 subunit, although receptors incorporating this subunit may be located out with the synapse. Receptors incorporating the δ subunit, in place of the γ 2 subunit are expressed extra- or peri-synaptically^{41,42,48,49}.

GABA_AR subunit expression and function during embryonic and post-natal development.

In the adult brain, activation of GABA_ARs usually causes an inhibitory effect on neuronal excitability, conferred both by a net inward directed flow of Cl⁻ ions and the shunting action of GABA⁵⁰. However, early in development (embryonic and first postnatal week), GABA_AR activation results in neuronal depolarization, due to a relatively high intracellular Cl⁻ concentration, resulting from limited expression of KCC2, the principal neuronal transporter for Cl⁻ ion extrusion⁵¹. Although the validity of this developmental ionic perturbation has been challenged, under certain experimental conditions⁵²⁻⁵⁴, the depolarizing nature of these early GABA-ergic signals may be sufficient to activate voltage-dependent calcium, or sodium channels and appear pivotal to the neurotrophic actions of the neurotransmitter⁵⁵⁻⁵⁷. Furthermore, during embryogenesis, such depolarizations are primarily mediated by activation of extra-synaptic GABA_ARs, which are expressed prior to and indeed facilitate the establishment of functionally relevant synaptic connections after birth^{2,3}.

During early development the subunit composition of GABA_ARs is highly plastic, consequently impacting upon the physiological and pharmacological properties of the receptors and influencing their subcellular localization^{15,42}. Preceding the generation, migration, and differentiation of most telencephalic and mesencephalic neurons^{58,59}, the first detection of GABA_AR subunits (including α 1-5, β 1-3 and γ 1-3), coincides with the appearance of GABA-positive neurons at approximately E13-14 in the marginal zone, sub-plate, and sub-ventricular zone^{60,61}. For rodent brain, the end of the first postnatal week generally signals a change to a hyperpolarizing GABA_AR response¹. Depending on the specific brain region, postnatal weeks two to three mark the establishment of a near-mature synaptic network, which reaches an adult stage by about two months of age following sexual maturation. From approximately the end of the first postnatal week synaptic GABA_AR signals, resulting from the vesicular release of GABA, begin to play an important role in shaping the development of neuronal circuits⁶².

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As the postnatal brain develops there is an increased requirement for temporal precision within neuronal networks and accordingly, in many neurons the duration of phasic events *e.g.* miniature inhibitory postsynaptic currents (mIPSCs) mediated by the activation of synaptic GABA_ARs by GABA released from a single vesicle, is progressively reduced^{42,63-69}. This critical development of GABA-ergic signalling is generally considered to result from a genetically programmed change of the subtype of synaptic GABA_AR. In this regard the α -subunit isoform exerts a profound role on the duration of GABA_AR-mediated phasic events. For example, electrophysiological studies with expressed recombinant GABA_ARs demonstrates the duration of GABA-evoked currents mediated by receptors incorporating either α 2-, or α 3-subunits are prolonged compared to equivalent receptors containing the α 1 subunit⁷⁰, although the β and γ subunit isoforms may also influence the mIPSC decay^{71,72}. Of physiological relevance, in different neurons (*e.g.* principal glutamatergic neurons and GABAergic interneurons) expression of different synaptic GABA_AR isoforms results in kinetically distinct synaptic events, underpinning the integration of neuronal signals and network activity and allowing the establishment of behaviourally relevant neuronal rhythms^{73,74}.

Many neurons express multiple GABA_AR isoforms, complicating evaluation of their role in neonatal development. To study this in detail we have used the ventrobasal (VB) neurons of the mouse thalamus. These neurons provide an ideal model as they exhibit a clear developmental transition from synaptic α 2-GABA_ARs from postnatal weeks 1-2, to α 1-GABA_ARs in weeks 2-3⁷⁵ (Figure 2). They are innervated by a band of GABA-ergic neurons, the nucleus reticularis (nRT neurons), which release GABA onto the VB neurons. Our studies, described below, reveal an important role for GABA-modulatory neurosteroids in determining the duration of mIPSCs during neonatal development.

During postnatal development of thalamic VB neurons neurosteroids, in concert with changes to GABA_AR isoforms, influence phasic and tonic inhibition.

We reported that the duration of thalamocortical VB neuron mIPSCs is greatly reduced over a critical, short period of 1-2 days between postnatal days (P) 9-11⁷⁵. Synaptic receptors incorporating the α 1-subunit are associated with brief mIPSCs^{63,75}. Implicating changes to synaptic GABA_ARs, our immunohistochemical and electrophysiological studies with an α 1 “knock-out” mouse and benzodiazepine-insensitive knock-in (α 1H101R; α 2H101R) mice revealed P20 neurons to exclusively express synaptic α 1 β γ 2 GABA_ARs, whilst younger mice expressed the α 2 β γ 2 isoform⁷⁵ (Figure 2). However, the major change to the mIPSC duration, occurred prior (P9-11) to the synaptic α 2- to α 1-GABA_AR transition⁷⁵ and were replicated in an α 1^{-/-} mouse⁷⁵. Clearly, this initial P9-11 synaptic plasticity does not result from expression of α 1-GABA_ARs⁷⁵. During postnatal development the levels of GABA_AR-modulatory neurosteroids change⁷⁶. Do neurosteroids influence neural inhibition during postnatal development (P7-24)?

Preventing neurosteroid synthesis in P7 thalamus with the 5 α -reductase inhibitor finasteride significantly reduced mIPSC duration⁷⁷. Intracellular delivery of the neurosteroid scavenger γ -cyclodextrin [γ -CD]⁷⁸⁻⁸⁰, produced the same effect⁷⁷, but by P10 was inert (Figure 2,3). Exogenous 5 α 3 α was equi-effective in prolonging mIPSCs of P7 and P20-24 neurons. Therefore, the developmental change in the mIPSCs is not a consequence of synaptic GABA_ARs steroid-insensitivity, but results from a loss of the neurosteroid⁷⁷. Note that the duration of both control and γ -CD-treated P20-24 mIPSCs are further reduced *c.f.* P10 mIPSCs, a change which corresponds temporally with the exchange of synaptic α 2-GABA_ARs by α 1-GABA_ARs⁷⁷ (Figure 2). Hence, prior to P10 the mIPSC duration is reduced, caused by decreased neurosteroid levels, with subsequent further kinetic refinement resulting from a change in the synaptic GABA_AR subtype (Figure 2). A recent study suggested that hippocampal expression of the GABA_AR α 2-subunit was increased by GABA_AR-active neurosteroids⁸¹. Therefore, it is conceivable that a waning neurosteroid tone heralds the programmed subsequent loss of thalamic synaptic α 2-GABA_ARs *post* P10.

The source of the neurosteroid influencing P7-8 neurons is not known. Studies using an antibody against the neurosteroid suggest that the VB neurons contain $5\alpha 3\alpha$ ⁸². Inclusion in the recording pipette of the membrane-impermeant γ -CD (0.5 mM), caused a time-dependent (> 6 min) decrease in the mIPSC duration of P7 VB neurons, mirroring the effect of pre-incubated γ -CD and of finasteride pre-treatment⁷⁷. These studies demonstrate that VB neurons contain GABA_AR-active neurosteroids, but do not elucidate whether their source is autocrine, paracrine, or indeed a consequence of prior endocrine release⁵.

We suggest that by P10 the local neurosteroid levels are insufficient to influence GABA-ergic neurotransmission. Can the more mature thalamus synthesise neuroactive steroids? Incubation of thalamic slices with 5α -DHP, the immediate precursor of allopregnanolone (Figure 1), caused a substantial prolongation of mIPSCs of both P20-24 and P7 neurons⁷⁷, which was prevented by indomethacin, an inhibitor of 3α -hydroxysteroid dehydrogenase (3α -HSD), the enzyme that converts 5α -DHP to the GABA_AR-active allopregnanolone (Figure 1). Although indomethacin is not a selective inhibitor of the 3α -HSD enzyme, this effect of 5α -DHP to prolong the mIPSCs was reversed by γ -CD treatment. Collectively, these results demonstrate that the more mature (P20-24) thalamic slice retains the ability to synthesise GABA_AR-active neurosteroids and that the change in mIPSC duration at P9-10 results from a loss of steroid substrate, or a co-factor *e.g.* NADPH, see⁸³.

P20-24 VB neurons express synaptic $\alpha 1\beta 2\gamma 2$ and extrasynaptic $\alpha 4\beta 2\delta$ GABA_ARs that mediate phasic and tonic inhibition respectively^{75,84}. Although the P20-24 mIPSCs were insensitive to γ -CD treatment, suggesting that local neurosteroid levels are insufficient to impact upon synaptic GABA_ARs, neurosteroids are particularly efficacious upon recombinant receptors incorporating the δ -subunit^{85,86}. However, the magnitude of the thalamic tonic current mediated by $\alpha 4\beta 2\delta$ -GABA_ARs was not influenced by γ -CD pre-incubation, suggesting that in P20-24 VB neurons the neurosteroid levels are not sufficient to influence either synaptic, or extrasynaptic GABA_ARs⁷⁷. Nevertheless, incubation with the precursor 5α -DHP, resulted in a large increase of the tonic current, illustrating that neurosteroid production can be reinitiated sufficiently to influence both phasic and tonic inhibition⁷⁷.

The mIPSC frequency of P10-24 VB neurons was much greater than that of P7 neurons. Unexpectedly, incubation with finasteride greatly increased the mIPSC frequency of P7, but not of P10 VB neurons. Furthermore, 5 α -DHP greatly reduced the mIPSC frequency of P20-24 VB neurons, an effect probably mediated by allopregnanolone, as indomethacin prevented this suppression of vesicular release by 5 α -DHP. Hence, vesicular GABA release from P7 nRT neurons is governed by a neurosteroid tone which, as it wanes, results in an increased mIPSC frequency for P10-24 neurons⁷⁷. These effects of the steroid metabolite appear unlikely to be mediated by presynaptic nRT GABA_ARs as other enhancers of GABA_AR function (pentobarbital, etomidate and zolpidem) had no effect on quantal GABA release⁷⁷ and therefore the steroid may act *via* alternative targets *e.g* voltage-gated T-type calcium channels⁸⁷.

A neurosteroid influence on GABA-ergic transmission of the developing mouse cortex.

Given the sudden loss of neurosteroid influence for P10 VB neurons we investigated whether GABA-ergic transmission in another brain region was similarly influenced and if so determine whether the temporal neurosteroid profile was similar to thalamus. These thalamic relay VB neurons send projections to cortical layer 4 (L4) neurons during P4-8 and subsequently (P10-14) L4 cortical neurons project to L2/3 neurons^{88,89}. We therefore investigated in L2/3 pyramidal neurons and GAD-67⁺ cortical interneurons the temporal profile of phasic inhibition and the influence of neurosteroid tone.

In L2/3 cortical pyramidal neurons the mIPSC profile changed considerably from neonatal (P7-15), to juvenile stages (P20-24)⁹⁰. Similar to VB neurons, the mIPSC frequency increased and their duration decreased between P7-8 and P20-24. In common with VB neurons, the mIPSC kinetic change did not result from neurosteroid insensitivity of synaptic GABA_AR, but resulted from a loss of neurosteroid production as the mIPSC decay time of P7, but not of P20-24 cortical neurons was substantially reduced by pre-incubation with γ -CD, or by finasteride pre-treatment⁹⁰. In contrast to the thalamic VB neurons, γ -CD influenced the mIPSC decay time at P10 and at P15, but as described above was inert by P20-24 (Figure 3). Hence, the neurosteroid profile is temporally distinct from that of the thalamocortical relay neurons. Mirroring the VB neurons, incubation of the cortical slice with 5 α -DHP greatly prolonged mIPSCs

of P20-24 pyramidal neurons. In common with thalamic VB neurons, this effect of 5 α -DHP was prevented by indomethacin and reversed by γ -CD treatment. Collectively, these results demonstrate that when supplied with the immediate steroid precursor, the cortical slice can metabolise 5 α -DHP into a GABA_AR-active neuroactive steroid⁹⁰. Indeed, phasic inhibition in hippocampal dentate gyrus granule cells and the medium spiny neurons of the nucleus accumbens is similarly influenced by 5 α -DHP (unpublished), suggesting this capacity to synthesise GABA_AR-active neurosteroids may be a common feature across different brain regions.

Even in the presence of γ -CD, the mIPSC decay time decreased from P7-8 to P20-24 suggesting synaptic GABA_ARs to be influenced by additional factor(s) during neonatal development. As described above, the GABA_AR subunit composition may influence the mIPSC duration, with particularly the α 1 subunit associated with brief events. We found the mIPSCs recorded from α 1^{-/-} cortical L2/3 pyramidal neurons to be prolonged *c.f.* their wild type counterparts at all ages studied (P7-8, P10, P15 and P20-24)⁹⁰. In common with WT neurons, treatment with γ -CD, or finasteride, reduced the mIPSC decay time of P7-15 α 1^{-/-} neurons, but with no effect on P20-24 neurons (Figure 3). Therefore, during cortical neuron development from P7, to at least P15, the duration of phasic inhibitory events is simultaneously influenced by a neurosteroid tone and by the subunit composition of the synaptic GABA_ARs, with a waning influence of the neurosteroid occurring between P15 and P20. By contrast, for thalamocortical neurons the neurosteroid influence dissipates by P10, but the mIPSC duration continues to decrease up to P20-24 due to the exchange of synaptic α 2-GABA_ARs by α 1-GABA_ARs (Figure 2,3). Finally, the frequency of mIPSCs increased during development in both cortical and thalamic neurons. However, whereas in thalamus the increased mIPSC frequency was due to a waning neurosteroid presynaptic influence, the frequency of cortical mIPSCs was not influenced by γ -CD pre-treatment⁹⁰ and therefore may reflect increased GABA-ergic innervation.

Immunohistochemical and *in situ* hybridisation studies reveal cortical principal neurons to express both the 5 α -reductase and the 3 α -HSD enzymes required for allopregnanolone synthesis, whereas interneurons do not³⁰. We therefore investigated whether phasic inhibition of cortical GABA-ergic interneurons is similarly influenced by neurosteroids during neonatal development.

Recordings were made from GAD67 GFP⁺ mice, engineered to co-express green fluorescent protein (GFP) with the GABA-synthesising γ -amino decarboxylase (GAD67) enzyme, which identifies, but does not distinguish between the three major interneuron classes in mouse cortex (*i.e.* somatostatin-positive (SS⁺), parvalbumin-positive (PV⁺), 5-HT₃R-positive (5-HT₃R⁺)] interneurons^{89,91-93}. Recordings from GFP⁺ neurons revealed P7-8 mIPSCs to be greatly prolonged *c.f.* equivalent recordings from P20-24 neurons. Intracellular γ -CD reduced the P7-8 mIPSC duration, with no effect on P20-24 neurons. Therefore, in common with cortical pyramidal neurons, interneuron phasic inhibition is influenced by neurosteroids early in postnatal development, an effect that waned by P20-24. As these interneurons do not appear to express the enzymes to synthesize allopregnanolone, these observations suggest such steroid may originate from paracrine, or endocrine sources and not emanate directly from autocrine synthesis³⁰.

Neurosteroid modulation of GABA_AR function during postnatal development:

Physiological relevance.

The findings presented above provide a compelling case for endogenous pregnane steroids enhancing GABA_AR-mediated phasic inhibition in both cortex and thalamus during the 2nd and 3rd post-natal weeks. During early postnatal maturation a programmed cessation of neurosteroidogenesis and changes to the expression of synaptic GABA_AR subtypes, act in concert to influence phasic inhibition. This interplay of neurosteroidogenesis with GABA_AR expression follows a neuron-specific temporal pattern. In somatosensory thalamus *i.e.* VB neurons, the neurosteroid tone dissipates by P10, prior to the subsequent exchange of synaptic α 2-GABA_ARs, by α 1-GABA_ARs, which further refines phasic inhibition. By contrast, in cortical L2/3 pyramidal neurons a neurosteroid influence on phasic inhibition persists for at least another week (Figure 3). The fading neurosteroid influence results from a loss of steroid substrate, or an essential co-factor, as incubation of P20-24 brain slices with 5 α -DHP prolongs both cortical and thalamic mIPSCs.

Neurosteroids enhance the function of both synaptic and extrasynaptic GABA_ARs^{5,48}, thereby potentially influencing both depolarising (embryonic and early post-natal) and hyperpolarising (later post-natal) actions of GABA. The mechanism(s) that dynamically regulate neonatal neurosteroid levels are not known, although NMDA receptor activation reportedly triggers their synthesis in CA1 neurons, providing an intriguing link between neural excitation and enhancement of neural inhibition^{94,95}.

What is the physiological significance of these neonatal changes to GABA-ergic transmission? As discussed, generally GABA-ergic signals mediate a depolarizing response up to the first post-natal week (but see below). Such an effect can cause activation of specific voltage-gated conductances, *e.g.* Ca²⁺, which act to initiate intracellular processes crucial to neuronal migration and maturation *e.g.* neurite growth and synapse formation, thus permitting the subsequent establishment and synchronization of neuronal networks^{2,96,97}. Neurosteroid levels are elevated during these early phases of neurodevelopment, *i.e.* during late gestation and early neonatal life, but have decreased by the third postnatal week, *e.g.* in the cortex⁷⁶ (See also Figure. 3 below).

The neurosteroid decline coincides temporally with the switch to a hyperpolarizing GABA signal¹. These events may be associated, as inhibition of neurosteroid synthesis, by finasteride, influences KCC2 expression in postnatal hippocampus, implying aberrant neuronal inhibition^{98,99}. Is the prolongation of the phasic GABA depolarisation by the neurosteroid action required for increased KCC2 expression? The differential neurosteroid profile of neonatal thalamic and cortical neurons (Figure 3), clearly warrants a comparison of the temporal expression of KCC2 in these neurons and the timing of their depolarising/hyperpolarising switch. Furthermore, certain cortical interneurons are of interest as, even for mature interneurons, GABA_AR activation may cause their depolarisation^{100,101}, an action important in the emergence of brain rhythms in the fast γ - frequency domain¹⁰²⁻¹⁰⁴ (Figure 3).

Enhancement by neurosteroids of the GABA-evoked depolarisation may influence the temporal window of crucial processes such as neuronal migration, morphological maturation and synapse formation^{3,57}. In support, allopregnanolone promotes cell proliferation both in human and rodent brain and regulates cell-cycle and gene expression^{105,106}. Although some of these

actions appear to be mediated by GABA_ARs, a putative involvement of G-protein-coupled membrane progesterone receptors requires consideration¹⁰⁷.

Our recordings from neonatal cortical L2/3 pyramidal and GABA-ergic interneurons reveal a role for neurosteroids in the normal maturation of the cortical network. The formation of functional GABAergic synapses on interneurons precedes the development of glutamatergic synapses in principal neurons^{97,108,109}, although there are exceptions^{110–112}. Whether neurosteroids influence the GABA_AR-mediated regulation of both GABAergic and glutamatergic transmission in these distinct neuronal types remains to be determined. We have not compared in detail the temporal pattern of neurosteroid regulation of GABAergic transmission of cortical interneurons, with that now established for pyramidal neurons. Furthermore, our recordings were from GAD-67⁺ neurons, therefore the neurosteroid influence may differ across distinct GABAergic, SS⁺, PV⁺, 5-HT₃R⁺ interneurons^{89,93,113}. The 5-HT₃R⁺ interneurons dominate in layers II-III of mature cortex⁹³. However, PV⁺ interneurons may be of particular interest as they exhibit a distinctive, extended developmental expression pattern *c.f.* other interneurons^{89,114} and a window of functional maturation exquisitely susceptible to extrinsic modulation during the second to third postnatal week⁸⁹. In particular, the overall homogenous distribution of PV⁺ interneurons across all cortical layers except for layer 1^{93,114} appears crucial for normal development of cortical connectivity. Importantly, in schizophrenics (see below) and in animal models of this condition, there is a layer selective reduction of the cortical PV and GAD67 mRNA signal¹¹³.

Suggesting a putative role for neurosteroids in the development of cortical connectivity, neonatal administration of allopregnanolone at (P5), a time which precedes the functional maturation of PV⁺ interneurons and when neurosteroids levels are relatively high (Figure 3), alters their distribution between superficial (II/III) and deep (V/VI) layers of the adult prefrontal cortex. Additionally, allopregnanolone reduces their abundance in the medial dorsal thalamus^{115–117}. These neurosteroid effects appear to be GABA_AR-mediated as they are mimicked by benzodiazepines¹¹⁸. Future genetic and pharmacological approaches to manipulate embryonic and postnatal neurosteroids may elucidate and differentiate their role in the three main classes of cortical

interneurons⁸⁹. Similar studies are required across different cortical layers and areas *e.g.* visual vs somatosensory cortex, as translaminar GABAergic inhibition is sub-served by different interneuron classes¹¹⁹.

A similar complexity may apply to somatosensory thalamic neurons (*e.g.* VB and nRT), where the neurosteroid influence wanes, approximately a week prior to that of cortical pyramidal neurons (Figure 2,3). The significance of this temporally distinct neurosteroidogenic profile is not known. However, unilateral lesion of neonatal thalamic lateral geniculate nucleus, or the VB, substantially alters the developmental expression of $\alpha 1$ - and $\alpha 5$ -GABA_AR subunits, across the layers of the corresponding cortical territories *i.e.* visual, V1 and somatosensory, S1, emphasising the profound influence of thalamic inputs on the development of cortical GABAergic circuits¹²⁰. Therefore, it is conceivable that thalamic neurosteroidogenesis may influence thalamo-cortical connectivity.

Neurosteroids, early life adversity and psychiatric disorders.

Given the proposed involvement of neurosteroids in brain development, experiences that perturb their levels during the establishment of neural circuitry, may subsequently influence juvenile and adult behaviour. In support, there are significant associations between abnormal levels of GABA_AR-active neurosteroids *e.g.* allopregnanolone and a variety of neuropsychiatric disorders³³ (see below). Moreover, strategies to restore neurosteroid levels in such conditions in humans, or equivalent animal models, have proved, beneficial^{33,121–125}. Note the majority of psychiatric disorders, ~2/3, are diagnosed by 24 years old, or earlier, consistent with a neurodevelopmental component^{126,127}.

Environmental events, grouped under the umbrella of early-life adversity or early life stress (ELS), are now recognised as a major preventable cause of future psychiatric disorders including anxiety, depression and substance abuse^{128–130}. Indeed, prior ELS may additionally be a risk factor for the future manifestation of some neurological disorders, *e.g.* certain forms of epilepsy¹³¹. Clinically, these terms describe a variety of negative experiences early in life, ranging from poverty, malnutrition, to physical and emotional trauma, or abuse¹³².

Patchev and colleagues (1997) first suggested that abnormal regulation of neurosteroid action as a consequence of negative experiences early in life could affect neurodevelopmental trajectories to contribute to adult psychopathology¹³³. In a rodent model of maternal separation the GABA_AR-active neurosteroid (5 α -THDOC), administered during early experience of reduced maternal care, prevented in adulthood, the development of a variety of neuroendocrine dysfunctions and behavioural abnormalities *i.e.* anxiety, associated with prior exposure to early-life adversity¹³³. Whether prior ELS subsequently influences adult levels of neurosteroids and/or the dynamic regulation of their levels under a variety of physiological conditions *e.g.* pregnancy is not known. However, such experiences greatly influence the functional effects of GABA_AR-active neurosteroids. Thus, using a naturalistic model of fragmented maternal care^{134,135}, we demonstrated that prior ELS produces a profound dysregulation of the neuronal circuits orchestrating the stress response in the paraventricular nucleus (PVN) of the hypothalamus¹³⁶. Specifically, prior ELS greatly increased the glutamatergic excitatory drive to the CRF-releasing PVN neurons, sufficient to prevent the normal suppression of PVN firing by physiological levels of allopregnanolone¹³⁶. Note that increased circulating allopregnanolone is purported to act as a part of a feedback circuit during acute stress, thereby limiting the duration of CRF release and consequently of glucocorticoids^{21,24}.

The nucleus accumbens is part of the reward pathway and is implicated in both depression and drug abuse¹³⁷⁻¹³⁹. In common with thalamus and cortex, it can synthesise GABA_AR-active neurosteroids (unpublished observations). Furthermore, we find that adult mice previously exposed to ELS, exhibit altered accumbal GABA_AR neurotransmission and an abnormal response to cocaine, *i.e.* an altered locomotor sensitization, a behaviour that recapitulates in rodents aspects of drug addiction¹⁴⁰.

Schizophrenia is negatively influenced by early-life adversity¹⁴¹⁻¹⁴³. The findings by Grobin and colleagues^{115,116}, discussed above, suggest a possible link between early neurosteroid dysregulation and an abnormal developmental pattern for prefrontal cortex, a region implicated in schizophrenia¹¹³. In support, a recent investigation revealed abnormal neurosteroid levels in a population of schizophrenic patients¹⁴⁴. Furthermore, Bortolato and co-workers have implicated altered neurosteroidogenesis in the accumbal-

mediated expression of behavioural deficits *i.e.* altered pre-pulse inhibition, which is typically altered in animal models of schizophrenia¹⁴⁵. Thus, the impact of abnormal neurosteroid levels upon neurodevelopment may extend beyond the cortex.

In conclusion, it is now evident that endogenous neurosteroids play a vital role in fine-tuning GABA-ergic transmission during neonatal development in a neuron specific manner. GABA is crucial to establishing and developing appropriate neural connections in the developing brain. Therefore perturbations of neurosteroid levels *e.g.* by stress, during this critical time of neonatal development may have a long term impact upon neuronal circuitry and plasticity. Further investigation of the influence of GABA_AR-active neurosteroids during development is now warranted. Such studies may allow a better understanding of the underlying neurobiology that results in the psychiatric disorders associated with early life adversity.

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Figure Legends:

Figure 1: The schematic illustrates the *de novo* neurosteroid synthesis. Initially cholesterol is translocated across the mitochondrial membrane by TSPO and accessory proteins (*e.g.* VDAC)^{34,35}. The mitochondrial P450scc (CYP11A1) converts cholesterol to pregnenolone, which diffuses into the cytosol. Cytosolic pregnenolone is subsequently converted to progesterone by 3 β -HSD. Progesterone, either newly synthesised locally, or of peripheral origin, is reduced to 5 α -dihydroprogesterone (5 α -DHP) and then to allopregnanolone (5 α -pregnan-3 α -ol-20-one, 5 α 3 α) by the enzymes 5 α -reductase (5 α -R) and 3 α -hydroxysteroid-dehydrogenase (3 α -HSD) respectively. Pharmacological agents such as finasteride and indomethacin inhibit the enzymes 5 α -R and 3 α -HSD respectively. The agent γ -CD can be employed to sequester the active neurosteroid. Although agents such as ethanol and certain antidepressants such as fluoxetine are known to enhance the levels of GABA_AR-active neurosteroids, their specific site(s) of action are not known. TSPO; translocator protein, VDAC; the 32-kDa voltage-dependent anion channel (required for benzodiazepine binding); 5 α -R; 5 α -reductase, 3 α -HSD; 3 α -hydroxysteroid dehydrogenase.

Figure 2: Illustrated is a diagrammatic representation of the changing temporal influence of an endogenous neurosteroid tone and the GABA_AR subunit composition upon the decay of mIPSCs recorded from VB neuron during development. During early developmental stages [postnatal day (P) 5-P14], the mIPSCs are exclusively mediated by α 2-GABA_AR (blue). By P16, the α 2 subunit is fully replaced by the α 1 subunit (green). During this developmental window, mIPSCs recorded from these neurons transition from slow to fast decaying phasic events, a trait often associated with a change in the receptor subunit composition. However, the developing α 2-GABA_AR containing VB synapses exhibit a significant endogenous neurosteroid tone that profoundly influences GABAergic neurotransmission, as indicated by the ability of γ -CD (grey lines) to shorten the mIPSC decay time course *c.f.* control (black line) up to P10. Importantly, this kinetic change precedes the switch from α 2-GABA_ARs to α 1-GABA_ARs. Thus, dissipation of an endogenous neurosteroid tone represents an alternative physiological mechanism to impart a rapid kinetic profile to GABAergic inhibitory signals. The steroid icon indicates 1) a postsynaptic action to enhance the actions of GABA interacting with GABA_ARs and 2) a presynaptic effect of the neurosteroid to suppress the quantal release of GABA mediated

by an unknown target. Note that mature $\alpha 1$ -GABA_AR containing synapses lack an endogenous neurosteroid tone.

Figure 3: A schematic representation illustrating the relationship between the development of cortical (layers 2 & 3) and the somatosensory thalamic (VB complex) connectivity and endogenous neurosteroid levels between P0 and P24. Left Y axis: A diagrammatic representation of the temporal profile of the development of cortical circuits between postnatal day (P) 0 – P24 describing: the oscillatory capacity of the network (green box), interneuron maturation (red box) and maturation of cortical neuronal projections for layers (L) 2,3 and 4 (black box). The Centre Panel illustrates the influence of endogenous neurosteroid tone upon GABA_AR-mediated phasic inhibition between P0 and P20-24, for thalamic VB neurons (blue), L2/3 cortical pyramidal neurons (green) and interneurons (red) suggesting a physiologically relevant role for endogenous neurosteroids in the processes of developmental maturation: representative normalised mIPSCs recorded from VB neurons (blue), L2/3 cortical pyramidal neurons (green) and interneurons (red) in the presence and absence of γ -CD (grey lines) during discrete temporal windows, namely P7, P10, P15 and P20-24. The presence of an endogenous neurosteroid tone is revealed by the effect of γ -CD (grey lines) on the decay time course (*i.e.* shortening) of the mIPSC (control black lines). This effect of γ -CD is evident at \leq P9 for VB neurons and \leq P15 for cortical pyramidal neurons. An endogenous neurosteroid tone is additionally revealed by γ -CD in cortical interneurons at P7, but it dissipates by P20-24. The endogenous neurosteroid tone profile between P7 and P20-24 for these interneurons remains to be determined. The Right Y axis illustrates developmental changes to neurosteroid levels (arbitrary units and deduced by their impact on GABA_AR phasic transmission) for VB neurons (blue lines), cortical neurons (green lines) and cortical interneurons (red lines). Dotted lines indicate that we do not have equivalent data on neurosteroid levels within these temporal windows. Note an arbitrary neurosteroid level of 0 here is used to infer a level of neurosteroid that is insufficient to influence phasic inhibition.



