Sex specific effects of pre-pubertal stress on hippocampal neurogenesis and behaviour

Running title: pre-pubertal stress, hippocampus and neurogenesis

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Abstract

Experience of traumatic events in childhood is linked to an elevated risk of developing psychiatric disorders in adulthood. The neurobiological mechanisms underlying this phenomenon are not fully understood. The limbic system, particularly the hippocampus, is significantly impacted by childhood trauma. In particular it has been hypothesised that childhood stress may impact adult hippocampal neurogenesis (AHN) and related behaviours, conferring increased risk for later mental illness. Stress in utero can lead to impaired hippocampal synaptic plasticity, and stress in the first 2-3 weeks of life reduces AHN in animal models. Less is known about the effects of stress in the post-weaning, pre-pubertal phase, a developmental time-point more akin to human childhood. Therefore, we investigated persistent effects of pre-pubertal stress (PPS) on functional and molecular aspects of the hippocampus. AHN was altered following PPS in male rats only. Specifically males showed reduced production of new neurons following PPS, but increased survival in the ventral dentate gyrus. In adult males, but not females, pattern separation and trace fear conditioning, behaviours which rely heavily on AHN, were also impaired after PPS. PPS also increased expression of parvalbumin positive GABAergic interneurons in the ventral dentate gyrus and increased GAD67 expression in the ventral hilus, in males only. Our results demonstrate the lasting effects of pre-pubertal stress on the hippocampus in a sex- and time-dependent manner, provide a potential mechanistic link between PPS and later behavioural impairments, and highlight sex differences in vulnerability to neuropsychiatric conditions after early life stress.
Introduction

Childhood adversity has repeatedly been identified as a risk factor for mental and physical illness. However, neurobiological mechanisms underpinning this relationship are not fully understood. Stressful experiences activate the hypothalamic-pituitary-adrenal (HPA) axis, resulting in release of corticosteroid hormones. These hormones traverse the blood brain barrier, providing a pathway through which stress can alter brain development and leave individuals susceptible to a range of psychiatric disorders later in life. The limbic system, especially the hippocampus, is predicted to display considerable sensitivity to stress during early life; it contains high levels of glucocorticoid receptors, and is undergoing significant structural and functional changes as it matures.

Accordingly, childhood maltreatment is consistently associated in humans with reduced hippocampal volume, particularly in dentate gyrus and Cornu Ammonis 3 (CA3) subfields. Hippocampal-dependent behaviours, such as pattern recognition, spatial memory and verbal declarative memory are also impaired in human sufferers of childhood maltreatment. Although informative, studies in humans do not allow extensive investigation into the underlying mechanisms of the impact of early life stress on the hippocampus to be explored.

In rodents, stress early in life can produce significant alterations in the adult hippocampus. Stress in utero can lead to impaired hippocampal synaptic plasticity, and stress in the first 2-3 weeks of life reduces adult hippocampal neurogenesis (AHN) in the dentate gyrus and BDNF expression. Less is known about the effects of stress in the post-weaning, pre-pubertal phase, a developmental time-point more akin to human childhood. Studies to date have revealed that pre-pubertal stress (PPS) impairs synaptic plasticity in the dorsal hippocampus, alters the expression of corticosteroid receptors and impairs performance in a hippocampal-dependent contextual fear task in males. Adult hippocampal neurogenesis (AHN) is believed to play a crucial role in a number of hippocampal-dependent cognitive processes, including contextual fear responses, trace fear conditioning and pattern separation. Pattern separation is a computational process that keeps similar input patterns distinct, allowing similar memories to remain separate. The more comparable the memories, the harder it is to 'pattern separate' them. Experimentally, pattern separation can be assessed through analysing the ability to spatially discriminate between objects placed relatively far apart (easy condition, spatial locations do not overlap greatly) or close together (difficult condition, spatial overlap and potential for interference greater). This function is attributed to the dentate gyrus, through the unique action of young adult born neurons. These neurons also play a specific role in fear conditioning – reductions in AHN using either.
antimitotic agents or transgenic models specifically impairs fear learning only when a delay is
interposed between the fear–associated and aversive stimuli, known as trace conditioning26-28. AHN
is exquisitely sensitive to a range of environmental factors, notably stress, physical exercise and
environmental enrichment19. Therefore, the first aim of this study was to determine whether PPS
would decrease adult hippocampal neurogenesis and impair behaviours which rely on AHN,
specifically, pattern separation and trace conditioning.

A number of intracellular pathways and extrinsic factors regulate AHN29,30. GABA signalling
plays a particularly important role in AHN, regulating division of neural stems cells and modulating
survival of new born neurons31,32. Therefore, the second aim of this study was to investigate the
consequences of PPS on PV+ GABAergic interneurons and GAD67 expression to elucidate a
mechanistic link between PPS and adult hippocampal function.

Methods and Materials

Animals

Animal experiments adhered to the European regulations on animal experimentation (Directive
2010/63/EU) and the UK Home Office Animals (Scientific Procedures) Act 1986. Male and female
Lister Hooded rats were bred in house at Cardiff University from adult pairs (Charles River) for this
study. Litters were weaned on postnatal day (PND) 21, and housed in same sex cages (32cm x 50cm
x 21cm) with littermates. Males and females were housed in the same room, food and water were
provided ad libitum unless stated otherwise, cages were lined with wood shavings, a cardboard tube
and wooden stick were provided as enrichment and light was maintained on a 12:12-h light/dark
cycle (not reverse). 9-13 animals per group and sex were used in each experiment, and separate
cohorts of animals were used for each behavioural experiment and for immunohistochemistry.

Sample sizes were based on power calculations from previous studies using the same animal model
and similar behavioural analyses16,17.

Pre-pubertal stress

On PND 25-27, half of the litters experienced a short-term PPS protocol that has been described
previously16,33,34, and was originally described by Jacobson-Pick and Richter-Levin35. This took place
in a designated room, separate from the holding room. On PND 25, between 9am and 13.00pm,
animals were given a 10 min swim stress in an opaque swimming tank (25cm high, 34cm diameter), 12 L capacity filled with 6L of 25±1°C water. PND 26 involved restraint stress: between 9am and 13.00pm animals were placed into plastic restraint tubes (15cm length, 5cm diameter) for 3 sessions of 30 minutes, separated by 30 minute breaks in the home cages. The final stressor on PND 27 involved placing animals onto an elevated platform (15x15cm, 115cm high) for three 30 minute sessions, separated by 60 minutes in the home cage between 9am and 5pm. Animals were then returned to their home cages and holding rooms and left undisturbed (aside from cage cleaning) until early adulthood (PND 60).

Litters were alternately allocated to experimental groups (PPS or control) by order of birth. A maximum of 3 animals per litter were used in each experiment, and a minimum of 5 litters per group (PPS or control) were generated to minimise effects of pseudoreplication 36. The litter of origin was accounted for in all statistical analyses.

**BrdU administration**

Forty four rats (Male: 10 control, 13 PPS; Female: 10 control, 11 PPS) were injected intraperitoneally on PND 60-65 with bromodeoxyuridine (BrdU, 200mg/kg in 0.9% sterile saline solution) and killed by transcardial perfusion with 0.01M PBS and 4% paraformaldehyde (PFA) under anaesthesia 24 hours later to assess baseline rates of cell proliferation and neurogenesis in the dentate gyrus. Brains were left in PFA overnight (4°C), transferred to 30% sucrose solution for cryoprotection. Coronal 30µm sections were cut through the entire hippocampal extent on a freezing microtome (Leica RM2245) and placed into a solution of cryoprotectant for storage at -20°C until immunohistochemical analysis.

**Immunohistochemistry**

Sections were stained for either: i) BrdU & doublecortin, ii) parvalbumin & doublecortin or iii) GAD67.

**BrdU & Doublecortin:** Sections were washed between each step for 3 x 5 minutes in 0.01M Tris-buffered saline (TBS, pH 7.4) and all steps were carried out at room temperature unless otherwise specified. One in every 12 sections was denatured in 45°C, 1M HCl for 30 minutes, followed by incubation in blocking solution (0.3% Triton-X in 0.01M TBS (TTBS), 2% donkey serum) for 60 minutes, rat anti-BrdU (1:100 in blocking solution, OBT0030, ABD Serotec, UK) for 48 hours at 4°C and Alexa Fluor 488 (5µg/ml in TTBS, donkey anti-rat, Life Technologies, UK) for 2 hours in the dark.
Sections were then incubated with goat anti-double cortin (DCX, 1:100 in blocking solution, Santa 
Cruz Biotechnology, UK, SC8066) for 24 hours at 4°C to identify newly differentiated immature 
neurons, AlexaFluor 647 (10µg/ml in TTBS, donkey anti-goat, Life Technologies, UK) for 2 hours in 
the dark. Sections were then exposed to DAPI (1:3000 in TBS D9542, Sigma UK) for 5 minutes. 
Washed sections were then mounted on to glass microscope slides and coverslipped with 
fluorescence mounting medium (S3023, Dako, UK). The number of cells double labelled with BrdU 
and DCX were counted in the supra- and infrapyramidal blades of each dentate gyrus in the dorsal 
(Bregma -1.72mm to -5.28mm) and ventral (Bregma -5.28mm to -6.72mm) portions of the 
hippocampus according to the atlas of Paxinos and Watson (2009). The total number of double 
labelled cells was estimated by multiplying the total number counted for each area (dorsal or 
ventral, supra- or infrapyramidal blade) by 12. Counts and total volume of the hippocampus was 
analysed using Zen Blue (Carl Zeiss Microscopy, Germany).

Parvalbumin & Doublecortin: All steps were carried out at room temperature unless otherwise 
specified. One in every 12 sections was blocked in blocking solution (1% Triton-X in 0.01M PBS, 10% 
donkey serum) for 2 hours, goat-anti DCX (1:100, SC8066 Santa Cruz) and mouse anti-parvalbumin 
(1:1000, Sigma-Aldrich, P3088) in 0.1% Triton-X, 0.2% donkey serum in 0.01M PBS overnight at 4 °C. 
Sections were washed in 0.1% Triton-X, 0.2% donkey serum in 0.01M PBS at least three times before 
icubation with donkey anti-goat Alexa 647 (10µg/ml in 0.01M PBS, Life Technologies UK) and 
donkey anti-mouse Alexa 555 (1:1000 in 0.01M PBS, Life Technologies UK) for 2 hours in the dark. 
Sections were washed with 0.01M PBS and incubated for ten minutes in the dark with DAPI stain 
(1:3000 in 0.01M PBS, D9542, Sigma UK) for nuclei staining. Sections were washed twice more in 
0.01M PBS before being mounted on standard microscopy slides using Mowiol aqueous mounting 
medium (Sigma, UK) and standard cover slips. Sections were imaged using an epifluorescent 
microscope (Leica DM6000B Upright Timelapse System with Leica Application Suite Advanced 
Fluorescence 3.0.0 build 8134 software, Leica Microsystems). The number of cells were quantified by 
manual counting in the supra and infrapyramidal blades of the dorsal (-2.28mm to – 5.04 mm) and 
ventral (-5.16 to -6.48mm) dentate gyrus. The area was calculated for each region and the number 
of immunopositive cells quantified through visual counting, giving a cell count per mm². The analysis 
was performed using Image J, with the plugin ‘Cell Counter’.

GAD67: All steps were carried out at room temperature unless otherwise specified. Sections were 
washed between each step for 3 x 30 minutes in blocking solution (0.2% Triton-X in 0.01M PBS 
(TTBS) containing 5% donkey serum, 2.5% bovine serum albumin). One in every 12 sections was 
incubated in blocking solution for 160 minutes, GAD67 antibody (1:500 in blocking solution,
MAB5406, Millipore, UK) for 48 hours at 4°C and Alexa 647 (1:1000 in blocking solution, donkey ant-
mouse, Life Technologies UK) for 150 minutes in the dark. Sections were then placed into DAPI
(1:200 in 0.01M PBS, D9542, Sigma UK) for 10 minutes in the dark before being mounted onto glass
slides as above as above. Number of cells labelled with GAD67 were counted through the entirety of
the dentate gyrus (infra- and supra-pyramidal blades), CA1, CA2 and CA3 and hilus in the ventral
(Bregma 5.28mm to -6.72mm) and dorsal (Bregma -1.72mm to -5.28mm) portions of the
hippocampus, according to the atlas of Paxinos and Watson (2009). The number of cells per
measured area was obtained using Zen Blue (Zeiss).

All slides were imaged at x20 using Axio Scan Z1 (Zeiss). Spearmans Rho was used to investigate
correlations between PV+, DCX+ and BrdU+ cells.

Behavior

Pattern separation

Animals

Pattern separation ability was assesses using a modified spontaneous location recognition paradigm
20, 21. Forty two rats (male: 11 control, 10 PPS; female: 12 control, 9 PPS) were food restricted to 85-
90% of their free feeding weight one week before testing (PND 60-100). Animals were handled daily
for 5 minutes during this time.

Apparatus

Testing took place in a black plastic circular arena (45cm high x 75cm diameter) with wood shavings
covering the floor. The arena was placed in the centre of a dimly lit room, and three proximal spatial
cues were placed equidistant around the arena to aid spatial orientation. Objects used were tin cans
(7.5cm x 7.5cm x 11cm) and small beer bottles (5.5cm x 5.5cm x 15cm) with the labels removed. Blu-
tack was used to fix them to the floor of the arena, and they and the arena were cleaned with 70%
ethanol solution between trials. A video camera was suspended directly above the arena and trials
were recorded for later analysis.

Behavior

Animals freely explored the arena for 10 minutes a day for 5 days. On day 6, objects were placed
into the arena. Objects were tin cans for half of the animals from each group and sex, and bottles for
the remaining animals. Animals were tested in two conditions – easy (large spatial separation, LS) and difficult (small spatial separation, SS). Three identical objects, A1, A2 and A3 were placed 12.5cm away from the walls of the arena, 25cm from the centre with either A2 and A3 at 120° (LS) or 50° (SS) apart (Figure 1). Animals explored the arena for 10 minutes (sample phase). 24 hours later, animals were returned for 5 minutes (choice phase). The arena now contained two new objects, A4 and A5, identical in appearance to those used in the sample phase. A4 was in the same (familiar) location as A1, and A5 was placed equidistant to the previous locations of A2 and A3 (novel spatial location, Figure 1). Rodents display an innate desire to explore novel objects and objects in novel locations over familiar ones, thus animals will explore A5 in preference to A1. It will be harder to recognise A5 in the novel location in the SS condition than in the LS condition because SS places greater demand on pattern separation abilities than LS. The amount of time animals spent exploring objects in the sample and choice phases was assessed from video recordings by an observer blind to group. A discrimination ratio was then calculated:

\[
\frac{\text{time exploring novel location object}}{\text{time exploring familiar location object}} \div \text{total exploration time.}
\]

Half of the animals from each group and sex were tested in the LS first then the SS, the other half experienced SS then LS. Exploration was counted as the rat having its head directed at the object (2cm or less), or touching the object with its nose. Sitting on or climbing over the object was not counted.

**Fear conditioning**

**Animals**

Animals were trained in trace, delay or control protocols at PND 60-100. 130 rats (male: 33 control, 36 PPS; female: 30 control, 31 PPS) were used, with sample sizes of 9-13 per protocol per sex. Three days before testing, animals were handled daily and taken to the testing room for habituation to transport and handling.

**Apparatus**

Testing took place in two identical standard modular test chambers (32cmx25.5cmx27cm, Sandown Scientific, UK) with grid floors (19 stainless steel rods, 1cm apart) underneath which was a stainless steel pan. The ceiling, front and back walls of the boxes were clear plexiglass, the sides walls
stainless steel, and the chamber was enclosed within a sound attenuating chamber with a ventilation fan providing a background noise of 63dB and a video camera attached to the inside of the door. The grid floor was connected to a shock generator, and there was a speaker attached to the inside of the sound attenuating chamber. The boxes were connected to a computer and shock, light, sound and video generation were controlled by a computer. One box (context one, C1) was scented with a drop of lavender oil (Botanics Aromatherapy Pure Essential Oil, Boots, UK) placed onto a tissue in the pan, and contained an IR light bar to allow filming as this box was always dark. In the second box (context two, C2), the pan was filled with wood shavings and the walls and ceiling decorated with black stars on a white background, and this box was always light (via houselight in the test chamber). Boxes were cleaned with ethanol wipes between animals, and sawdust/lavender scent replaced. Half of the animals from each group and sex were trained in C1 and the other half in C2, and all trials were video recorded for later analysis. The CS was 15s, 75dB white noise and the US was 0.5s, 0.5mA scrambled footshock in all protocols.

**Behavior**

**Training:** Animals were placed individually into chambers for 120s. Animals were then trained in one of three protocols: trace protocol (TP) - rats received 10 CS-US pairings, with a 30s stimulus free trace interval between the offset of the CS and the onset of the US, delay protocol (DP) - 10 CS-US pairings were presented, here the CS was followed immediately by the US and control protocol (CP) - animals received 10 presentations of the CS and 10 presentations of the US, which were explicitly unpaired. Intertrial intervals (ITI) were 312 s (+/-62s) for trace and delay protocols, 156 s (+/-31s) for the control protocol. ITI length is positively correlated with freezing to later presentations of the CS in trace conditioning, and negatively associated with freezing to a context. This length of ITI was selected as optimal for assessing freezing to cue and context in both trace and delay protocols.

**Context recall 1:** Animals were returned to their training chamber 24 hours later for 10 minutes to assess contextual fear responses.

**Cue recall:** Forty-eight hours after training, animals trained in C1 were placed into C2 and vice versa to reduce contextual associations and maximise responding to the CS. A plastic insert was placed over the bars of the floor to aid the discrimination. Animals received a 120 s acclimation period, followed by 360 s of CS the 240 s post-CS.

**Context recall 2:** Seventy-two hours after training, animals were returned to their training chamber for 10 minutes to assess contextual fear responses.
Freezing during training (baseline, post-shock and trace interval) and recall was analysed from video recordings by an observer blind to group. Freezing was defined as immobility with the exception of movement required for respiration, and was sampled every 10 s.

**Statistical analysis**

All data were checked for homogeneity of variance and normality, and transformed when these conditions were not met. Data were analysed in JMP (statistical software, SAS Institute, Cary, NC, USA) using generalised linear models, with experimental treatment (control or PPS), sex, separation distance (pattern separation only), protocol (trace, delay or control, fear conditioning only) fitted as factors. Litter was nested within group and added as a random factor to account for litter of origin and animal was nested within litter and added as a random factor to account for multiple measurements on the same animal.

**Results**

**Neurogenesis**

PPS resulted in a 46% decrease in cell proliferation in the suprapyramidal blade of the ventral dentate gyrus in males only (F1,8.81=4.9, p=0.05, Figure 2a), as measured by the number of co-labelled BrdU and DCX cells. There were no changes in the infrapyramidal blade or in females (F1,7.541=0.28, p=0.62). Cell survival (as measured by DCX labelled neurons) was increased by 65% in suprapyramidal and infrapyramidal blades of the ventral dentate gyrus after PPS in males (F1,9.29=14.21, p=0.004, Figure 2b) but not females (F1,8.38=1.25, p=0.29). Volume of the dentate gyrus did not vary as a result of PPS (males: F1,8.82=0.01, p=0.91; females: F1,8.31=0.09, p=0.77).

**Behavior**

**Pattern separation:** Animals from both groups spent an equal amount of time exploring all three objects in the sample phase (F1,8.5=0.01, p=0.91). During the choice phase for LS, all animals spent more time exploring the novel object (A5) than the familiar one (A4) (one sample t-test: control males: t=3.62, p=0.005; PPS males: t=3.05, p=0.01; control females: t=3.12, p=0.01; PPS females: t=3.18, p=0.01), and both sexes were able to discriminate between objects equally well during LS (F1,4.46=0.07, p=0.81, Figure 3a), demonstrating that PPS did not affect pattern separation in the easy
condition. During the choice phase for SS, only control males spent more time exploring the novel object than the familiar one (one sample –test: control males: t=4.38, p=0.001), and control males were the only group able to discriminate between novel and familiar objects in SS (F1,37.55=9.3, p=0.006, Figure 3b). Thus, control and stressed females were equally unable to distinguish between the novel and familiar objects in the SS condition, and stressed males showed an impairment in spatial pattern separation in the more difficult SS condition when compared to control males.

**Fear conditioning - context:** Results are shown from the first 90s of each period only. Twenty-four hours after conditioning, male and female control animals experiencing CP and TP demonstrated robust levels of contextual freezing (Figure 4a, data shown for males only). As expected, contextual freezing was markedly decreased in control DP animals where stronger associations to the CS are formed, resulting in decreased contextual freezing 39. However, PPS resulted in increased contextual freezing in DP males (F2,198.3=3.24, p=0.04, Figure 4a) but not females (F2,24.49=0.67, p=0.52). There were no differences between protocols or stressed and control animals in contextual freezing responses 72 hours post conditioning (males: F2,24.76=0.75, p=0.48; females: F2,30.65=2.12, p=0.14).

**Fear conditioning – cue recall:** Forty-eight hours after conditioning cue recall was performed. In males, PPS reduced conditioned freezing during CS presentation in the TP (F2,198.9=3.6, p<0.0001, Figure 4b). In females, PPS increased freezing during CS presentation in the DP (F2,185=3.37, p=0.04, Figure 4c). Overall, DP animals showed a greater freezing response to the CS than CP and TP animals, and TP froze more than CP animals (males: F2,198.9=3.6, p<0.0001; females: F2,24.24=25, p<0.0001). This is expected as DP animals make stronger CS-US associations than TP animals, who make stronger associations than controls. There was no effect of PPS on post CS freezing, but overall, TP froze more than DP animals (males: F2,197.1=3.48, p=0.03; females: F2,27.08=4.2, p=0.03).

To investigate whether differences between the groups were present during encoding, we analysed freezing responses for 60 s after presentation of the 10 US (shocks) during conditioning, and during the 30 second ‘trace’ gap for TP animals (after presentation of CS, sound). All animals, males and females, froze more after US presentation than at baseline (F10,631.1=28.2, p<0.0001), and TP animals froze more after CS presentation from CS 2 (F10,222=17.44, p<0.0001). PPS resulted in lower levels of freezing in male TP animals both after US (F1,11.14=7.06, p=0.02) and CS presentation (F1,10.93=10.74, p=0.007, Figure 4d).

**GABA and GABAergic interneurons**
PPS increased the expression of GABAergic (PV+) interneurons in the suprapyramidal but not infrapyramidal blade of the ventral hippocampus of males only ($F_{1,8.27}=6.83$, $p=0.03$, Figure 5a). There were no differences in females ($F_{1,7.948}=3.2$, $p=0.1$). GAD67 was increased in the ventral but not dorsal hilus in stressed males ($F_{1,311}=18.3$, $p<0.0001$, Figure 5b). Furthermore, following PPS, there was a significant positive correlation between the number of DCX+ and PV+ cells in the suprapyramidal blade of the dentate gyrus only ($r=0.64$, $p=0.04$).

**Discussion**

**Summary**

The present study demonstrates that PPS has a profound effect on the structure and function of the adult male hippocampus. Specifically, we found that PPS decreased production but enhanced survival of new born neurons in the ventral dentate gyrus, and impaired pattern separation and trace fear conditioning - behaviours which depend on neurogenesis. We also found evidence of altered GABA signalling following PPS - expression of PV+ interneurons was increased in the ventral dentate gyrus and GAD67 increased in the ventral hilus. Due to GABA’s role in maintaining quiescence of neural progenitor cells and promoting survival of new born neurons, this provides a potential molecular mechanism underlying alterations in AHN after PPS.

**Neurogenesis**

We predicted that PPS would reduce proliferation (BrdU/DCX co expressing cells) and survival (DCX only) of new born neurons, as prenatal and early postnatal stressors (first 1-3 weeks of life) robustly decrease AHN in adult males. In the present study, PPS decreased proliferation, but resulted in increased survival of new born neurons, and these changes were specific to the ventral dentate gyrus in males. Thus, similar to prenatal and early postnatal stressors, PPS results in decreased cell proliferation in the male dentate gyrus. However, the importance of timing is highlighted, as in contrast to other early-life stress paradigms, PPS increases cell survival. Delineating the contrasting mechanisms of dysfunction produced by stress at different early life time points is crucial to producing effective interventions.

Altered neurogenesis is robustly associated with impairments in hippocampal-dependent behaviours, especially those with a temporal or spatial component. Therefore, the second aim of
the present study was to determine the consequences of PPS on two hippocampal dependent
behaviours, trace conditioning and pattern separation.

Trace and delay fear conditioning

We previously demonstrated that PPS results in impaired hippocampal-dependent
contextual fear conditioning responses in male animals. Although an intact hippocampus is
necessary for this task (at least in one trial studies), other brain regions, such as the amygdala, are
also required. Therefore, we used a trace/delay fear protocol to more precisely dissect the
effects of PPS on the manipulation which is thought to rely on AHN.

Males subjected to PPS displayed lower conditioned fear responses to the CS but only after
training with the trace protocol. This is likely due to deficits on encoding, as lower levels of freezing
were observed in PPS trace protocol males after the second presentation of CS and US during
training. Thus, PPS selectively impacts hippocampal function in males but not females. Hippocampal
lesions impair responses to the CS in trace but not delay protocols, whereas other regions are
equally important in both. Furthermore, the acquisition of trace fear is impaired following
ablation of AHN using optogenetic or infusion methods, yet delay fear responses remain intact. The relative contributions of dorsal and ventral hippocampus to trace conditioning are not
entirely clear. One study found that bilateral infusions of the AMPA/kainate glutamate receptor
antagonist CNQX into the dorsal dentate gyrus disrupts expression of trace fear conditioning (ventral
inactivation was not assessed), whereas others reveal inactivation of the ventral, but not dorsal,
hippocampus impairs acquisition and expression of trace fear.

The effects of increased new born neuronal survival on trace memories are unknown, but
increased neurogenesis disrupts contextual fear memories. Together our data suggests that
PPS-induced alterations in neurogenesis could underlie the impairments in trace fear conditioning
observed in males in the present study.

Males who had experienced PPS froze more to contextual cues 24 hours after training in the
delay condition. The impact of PPS on contextual fear conditioning selectively in males is consistent
with our previous findings, although here the direction of effect is different. There are
methodological differences between the studies which likely account for the disparity in results – a
single presentation of the US in our previous work compared to the present study which used 10 CS-
US pairings. Ablation of AHN results in impaired contextual fear responses in single but not multiple
trial conditioning. The enhanced contextual freezing in delay conditioning we measured may be
modulated by other neural substrates including the amygdala, which is known to be overactive after multiple CS-US pairings in this model. Females froze more to representation of the cue during the delay protocol only after PPS, another behaviour which relies more heavily on the amygdala.

**Pattern separation**

AHN is necessary for successful spatial pattern separation, particularly when stimuli are of increasing similarity. We showed that animals were able to discriminate between stimuli when they were presented at large separation distances, but only control males successfully completed the task at the smaller, more difficult, separation distances. Deficits in pattern separation are particularly evident in psychiatric disorders with a strong hippocampal phenotype, such as post-traumatic stress disorder, schizophrenia and major depressive disorder. To date, no studies have considered the impact of early life stress on pattern separation. This study provides novel evidence for the role of PPS in disrupted pattern separation ability in adult life, and again points to a role between disrupted AHN and impaired hippocampal-dependent behaviour.

Typically, the dorsal hippocampus has been associated with impairments in cognitive and spatial behaviour (such as pattern separation) and the ventral hippocampus with emotional responses (such as those associated with fear conditioning). However, the situation is likely more complex, and the dorsal hippocampus has been shown to play a role in trace fear responses, especially with long trace intervals. Presently, the relative contributions of dorsal and ventral hippocampus to the pattern separation task used in the present study are unknown. However, studies using pattern separation for reward value suggest a crucial role for the ventral dentate gyrus, and one study found expression of the immediate early gene zif268 in both ventral and dorsal dentate gyrus when rats were required to change between different navigational strategies in a plus maze. Further work is required to delineate the relative contributions of dorsal and ventral hippocampus to the pattern separation task used in the current study. Here, we see impairments in both emotional (trace fear) and spatial (pattern separation) tasks correlated with altered neurogenesis in the ventral hippocampus.

**GABAergic Interneurons**

Neural stem cells, the substate for neurogenesis, are largely quiescent, a state maintained through tonic GABA current from parvalbumin (PV+) interneurons. We found that PPS increased PV+
interneurons in the ventral dentate gyrus, and increased GAD67, the rate-limiting enzyme in GABA production, in the ventral hilus in males specifically. This suggests a mechanism whereby GABA could be responsible for suppressing generation of adult-born neurons in the ventral hippocampus in males following PPS.

GABA promotes survival of new born neurons. This may explain why although generation of new neurons was decreased, survival of DCX-expressing neurons, a population aged 24 hours-2 weeks, was markedly increased after PPS in the male ventral dentate gyrus, especially as we find a positive correlation between numbers of PV+ and DCX+ cells in the ventral dentate gyrus. Taken together, altered GABA signalling through increased numbers of PV+ interneurons may underlie aberrant neurogenesis and behaviour observed after PPS. Altered GABA signalling has been previously found in models of PPS, with levels of α1-3 subunits of the ionotrophic GABA A receptor altered in a region specific manner.

Sex differences

Sex-based differences in the development of neuropsychiatric disorders abound, and have been found previously in our PPS model. In the present study, PPS produces fewer changes in females than males. Specifically, PPS enhanced freezing to representation of the CS after a delay protocol only in females, with no other changes evident. This complements research by Toledo-Rodriguez & Sandi (2007), who found that PPS on PND 28-30 impaired extinction to a tone CS in adult males but not females. Interestingly, this study found increased fear responses to a tone in adolescent males, and reduced contextual freezing in adolescent females after PPS, highlighting the importance of timing when assessing the long-term outcomes of early life stress.

PPS decreased the production yet increased the survival of new-born neurons in the infrapyramidal blade of the ventral dentate gyrus of males, with no effect in females. This mirrors the adult situation, where males demonstrate greater sensitivity to the effects of stress hormones, showing more extensive suppression of neural stem cell proliferation in the dentate gyrus. However, adults also show decreased survival of newborn neurons in response to stress, which is in contrast to the increased survival observed in the present study. Prenatal stressors almost universally result in decreased generation and survival of newborn neurons in both males and females, whereas postnatal stress in the first 1-2 weeks of life produces more variable results. Age of assessment is also vital: males experiencing early postnatal stress tend to have enhanced neurogenesis at PND 21, which switches to suppression as the animals age (as measured by DCX-
positive cells). In contrast, females display decreased neurogenesis at PND 21, and this effect disappears in adulthood (DCX-positive cells). Less is known about the age- and sex-specific effects of this early prenatal stress on cell proliferation. Our study demonstrates that PPS produces a decrease in proliferation (measured by BrdU) with an increase in survival (DCX-positive cells) in adult males, with no effects apparent in females, marking the pre-pubertal phase as a time-point with unique consequences for adult hippocampal neurogenesis.

Sex differences were also seen in the pattern separation task, male and female rats performed equally well when the task was easy, but only control males were able to complete the difficult phase. We previously found male animals were superior in a spatial navigation task (water maze) and a number of other studies reveal a male advantage in spatial reference, however others find no difference. The fact that control females could not complete the more difficult pattern separation means assessing the effects of PPS on females in this task was not possible, and alternative tasks are needed to probe further female responses to PPS.

Expression of PV+, GABAergic interneurons and GAD67 were altered in males experiencing PPS, but remained unchanged in females. Prenatal stress decreases PV+ interneurons and GAD67 expression in frontal cortex and hippocampus in early and adult life in males, and alters expression of GABA receptor subunits in a region and sex dependent manner. PPS has previously been shown to decrease expression of GABA_A receptor α1 subunit in the hippocampus of adult males, and elevation is associated with stress resilience, but little is known of the effects in females. Exposure to a more extended pre-pubertal stress paradigm (spanning PND 28-42) resulted in reduced mRNA levels of GAD67 in the central nucleus of the amygdala in male rats, whereas a shorter protocol (PND 28-30) was insufficient to elicit such changes. This suggests that the exact timing and duration of early life stress are likely crucial in determining later outcomes.

Our results add to the literature demonstrating sex differences in response to PPS. In the present study, males display greater responses to PPS, but previous work demonstrates that females are sensitive in different ways. For example, females display learning impairments, exhibit greater compulsivity in a delay discounting task and have reduced preference for sweet solutions (anhedonia). Furthermore, males and females react similarly regarding certain behaviours: both sexes display heightened anxiety and altered cognitive bias after PPS. In humans, sex differences are also observed: associations between childhood maltreatment and incidents of anxiety and depression are higher in women than men, whereas risk for substance use disorder show the opposite pattern. This suggests that males and females are distinct in their vulnerabilities to the experience of stress during development. Understanding these differences is
crucial in the advancement of appropriate interventions and treatments for psychiatric illness associated with experience of childhood maltreatment.

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Conflict of interest

The authors declare that they have no conflict of interest.

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Figure Legends

Figure 1. Apparatus used for pattern separation. Animals explored three objects, A1 A2 and A3 in a circular 75cm arena, for 10 minutes on day 1. The objects were placed 12.5cm from the perimeter. A2 and A3 were either separated by a a) large (52cm) or b) small distance (20.5cm). C) 24 hours later animals were re-exposed to the arena, A1 was replaced with an identical object, A4 and objects A2 and A3 were replaced with A5, placed equidistance to the position of A2 and A3.

Figure 2. Effects of PPS on neurogenesis. A) In males, PPS decreased production of new neurons in the ventral hippocampus, and B) increased survival. *=p<0.05, ***=p<0.001. Error bars represent 1 SE.

Figure 3. Pattern separation in control and PPS animals. A) All animals successfully discriminated the object in the novel from the familiar spatial location at large separation distances, but B) only control males were successful at small separation distances. Bars with different letters are significantly different to one another. Error bars represent 1 SE.

Figure 4. Fear responses in PPS and control animals exposed to three different conditioning protocols. A) PPS males experiencing the delay protocol demonstrated enhanced levels of contextual freezing when compared to their control counterparts in a 24 hour recall test. B) PPS trace protocol males froze significantly less than controls to representation of the CS 48 hours after conditioning. C) PPS delay protocol females froze more than controls to representation of the CS 48 hours after conditioning. D) PPS males experiencing the trace protocol froze less post CS and post US on the conditioning day when compared to controls. Con = control animals, PPS = pre-pubertal stress animals, CP = control protocol, TP = trace protocol, DP = delay protocol. *=p<0.05, **=p<0.01. Error bars represent 1 SE.

Figure 5. Effects of PPS on parvalbumin and GAD67 expression in the hippocampus. A) Parvalbumin positive cells (PV+) were increased in the ventral dentate gyrus after PPS in males, and B) GAD67 (GAD+) was increased in the ventral hilus of PPS males. *=p<0.05, ***=p<0.001. Error bars represent 1 SE.