Haploinsufficiency of the schizophrenia and autism risk gene \textit{Cyfip1} causes abnormal postnatal hippocampal neurogenesis through microglial and Arp2/3 mediated actin dependent mechanisms

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\begin{abstract}
Genetic risk factors can significantly increase chances of developing psychiatric disorders, but the underlying biological processes through which this risk is effected remain largely unknown. Here we show that haploinsufficiency of \textit{Cyfip1}, a candidate risk gene present in the pathogenic 15q11.2(BP1–BP2) deletion may impact on psychopathology via abnormalities in cell survival and migration of newborn neurons during postnatal hippocampal neurogenesis. We demonstrate that haploinsufficiency of \textit{Cyfip1} leads to increased numbers of adult-born hippocampal neurons due to reduced apoptosis, without altering proliferation. We show this is due to a cell autonomous failure of microglia to induce apoptosis through the secretion of the appropriate factors, a previously undescribed mechanism. Furthermore, we show an abnormal migration of adult-born neurons due to altered Arp2/3 mediated actin dynamics. Together, our findings throw new light on how the genetic risk candidate \textit{Cyfip1} may influence the hippocampus, a brain region with strong evidence for involvement in psychopathology.
\end{abstract}

\section*{Introduction}

Many psychiatric conditions show high heritability. Recent genetic studies in schizophrenia for example have identified up to 160 loci that increase risk for the disease\cite{1,2}, enriched within synaptic\cite{3}, histone modifying and immune system genes\cite{4}. \textit{CYFIP1} is a genetic risk factor for schizophrenia, autism and developmental delay, by virtue of its presence in the penetrant 15q11.2(BP1–BP2) copy number deletion\cite{5}. Loss of one copy of this interval leads to substantially increased risk for disorder in both Western\cite{6,7} and Han Chinese populations\cite{8}. \textit{CYFIP1} haploinsufficiency is likely a major contributor to the 15q11.2 (BP1–BP2) psychiatric phenotype due to evidence of \textit{CYFIP1}'s involvement in a range of synaptic functions, including key roles in dendritic spine morphology and branching\cite{9-11}. \textit{CYFIP1} protein influences synaptic function in two main ways. \textit{CYFIP1} interacts with fragile X mental retardation 1 (FMRP), the protein gene product of FMR1, to suppress translation of up to 800 target transcripts\cite{12}. Secondly, it interacts with Wiskott–Aldrich syndrome protein family member 1 (WAVE1), a key mediator of cytoskeleton dynamics, to modulate ARP2/3 dependant actin branching\cite{13}.

In the present work we focus on microglia and adult hippocampal neurogenesis (AHN) as important sites of \textit{CYFIP1} action and as potential contributors to psychiatric phenotypes arising from 15q11.2(BP1–BP2) copy number deletion. Adult-born hippocampal neurons, up to 700 of which are estimated to be born per day in humans\cite{14}, develop through a well-defined series of
cellular events, starting from proliferative stem cells through intermediate progenitors, migratory neuroblasts and immature neurons, before eventually forming mature granule cells, which distribute through the dentate gyrus\(^{15}\). In rodent models, adult-born hippocampal neurons have been shown to be involved in a number of behavioural processes, spanning pattern separation\(^{16,17}\), spatial navigation\(^{18,19}\), memory turnover\(^{20}\), and acquisition\(^{21}\) and extinction\(^{22}\) of fear memory. These behaviours are affected in psychiatric disorders including schizophrenia\(^{23–26}\). Antipsychotic drugs, specifically atypical antipsychotics, also impact AHN\(^{27}\) and importantly, a number of functionally diverse schizophrenia risk genes, including \textit{DISC1}\(^{28}\), \textit{SREB2/GPR85}\(^{29}\), \textit{CACNA1C}\(^{30,31}\), \textit{DGCR8}\(^{32}\) and \textit{miR137}\(^{33}\), converge in their ability to modify AHN. Additionally, post-mortem findings from schizophrenia patients reveal changes in cell proliferation and expression of immature neuronal markers in the hippocampus\(^{34–36}\).

Immune changes are another common hallmark of several psychiatric disorders, including schizophrenia, where evidence from epidemiological data\(^{37,38}\) and findings showing increased inflammatory factors in the circulation\(^{39}\), and in post-mortem brain\(^{40}\) of schizophrenia patients, implicate altered immune function in the development of disease symptoms. Mechanistic insights linking immune changes to risk for psychopathology are however largely lacking. Recent interest has focused on microglia, the resident macrophage-like immune cells of the central nervous system, with data suggesting possible alterations in microglial functioning in both those at high risk and those diagnosed with schizophrenia\(^{41–48}\). The causal mechanism(s) by which microglia might contribute to pathogenesis remain unclear, though there is increasing evidence that, in addition to canonical immune functions, microglia also modify neuronal networks through synaptic pruning during development\(^{15}\) and into adulthood\(^{46}\). This microglial-mediated pruning is important for normal brain and behavioural functioning\(^{47}\). Additionally, microglia can modulate neuronal firing through direct cell–cell contacts\(^{48}\), and have been shown to be capable of phagocytosing both developing\(^{49}\) and adult\(^{50}\) neurons.

In this work we used a translationally relevant haploinsufficient mouse model to reveal hitherto undescribed effects of manipulating \textit{Cyifp1} on the survival and migration of newborn neurons in dentate gyrus hippocampus, key cellular processes in the development and function of the adult hippocampus. We also reveal the mechanisms mediating these effects, highlighting CYFIP1 effects on microglia mediated apoptosis and Arp2/3 dependent actin dynamics, respectively. We suggest these findings may have relevance to how \textit{Cyifp1} haploinsufficiency can influence hippocampal function in the context of the markedly increased risk for psychiatric disorder arising from the 15q11.2(BP1–BP2) copy number deletion.

**Materials and methods**

For full Materials and methods, please see Supplementary Information.

**Cyifp1 heterozygous knockout mice**

\textit{Cyifp1}\(^{+/-}\)-mice (MGI:5002986, referred to hereafter as \textit{Cyifp1}\(^{+/-}\)) were maintained heterozygously on a C57/BL6J background, for genotyping examples, see Fig. S1. Animals used were of mixed sex and 8–12 weeks old. \textit{Cyifp1}\(^{+/-}\) and wild-type littermates were kept in conventional cages with 12 h light–dark cycle and ad libitum access to water and food at all time. All experiments were performed in accordance with the Animal (Scientific Procedures) Act 1986, and approved by the local animal welfare committee.

**Tissue harvesting and immunohistochemistry**

Paraformaldehyde fixed brains from 8 to 12-week-old \textit{Cyifp1}\(^{+/-}\) and wild-type littermates were prepared as previously described\(^{51}\). Free floating 40 μm cryostat sections were used for immunohistochemistry. Immunohistochemistry was performed as previously described\(^{51}\), with a 1:12 stereotactic sampling rate. Detection of cleaved-caspase 3 used a modified protocol, where primary antibodies were incubated for 48 h in blocking solution at 4°C. Primary and secondary antibodies are described in Table S1.

**Primary hippocampal progenitor culture**

Hippocampal progenitors were isolated from P7 to P8 animals. Isolated hippocampi were dissociated in papain and progenitors were enriched through OptiPrep density gradient centrifugation. Cells were cultured on poly-L-lysine and laminin in the presence of EGF and FGF2.

**Primary microglial culture**

Microglia were isolated from P7 to P8 whole brain mixed glia through the shake-off method.

**Immunocytochemistry**

Fixed cultures were blocked and permeabilized with 5% donkey serum and 0.1% Triton X100 in PBS (PBST) for 30 min at room temperature. Primary antibodies were applied in the same solution overnight at 4°C. Antibodies are detailed in Table S1. Following three washes in PBST, relevant secondary antibodies were applied in PBST for 2 h at room temperature. For identification of microglia, Alexa 568 conjugated IB4 (ThermoFisher I21412) was added to secondary antibody solutions at 2.5 μg/ml. Cells were washed, counterstained with DAPI, and mounted for microscopy.
Conditioned medium
Microglia conditioned medium was prepared by incubating 5 × 10⁵ microglia per ml in progenitor medium for 24 h. 25% conditioned medium was added to hippocampal cultures.

Time-lapse imaging and cell tracking
Primary hippocampal progenitors were plated on poly-D-lysine and laminin coated 24 well plates, and cultured for 5 days. At 5 DIV, cultures were imaged on an inverted Leica DMI600B microscope, at 37 °C, with 5% CO₂ in air. Cells were imaged for 2 h, with image acquisition every 5 min. Total distance covered was determined through manual cell tracking using the MTrackJ plugin in ImageJ. A minimum of 50 cells were analysed per condition per animal.

Actin analysis
Cells were fixed at 5 DIV with 4% paraformaldehyde in PBS, at 4 °C for 30 min, permeabilised with 0.1% Triton X100 in PBS for 15 min, and incubated with 300 nM Alexa Fluor 647 conjugated phalloidin (ThermoFisher A22287) for 30 min. Fluorescence intensity was measured in ImageJ.

Arp2/3 inhibition
Cells were treated with 50 μM CK-548 or 250 μM CK-666 in DMSO, or vehicle, for 2 h prior to time-lapse imaging. Drugs were present during imaging.

Results
Cyfip1+/− animals have normal proliferation but increased numbers of adult-born immature neurons in the hippocampus
We first investigated cell proliferation in the dentate gyrus of the hippocampus. Animals received a single injection (i.p.) of 100 mg/kg BrdU 6 h prior to sacrifice. No significant differences in cells positive for BrdU (197.3 ± 11.1 vs. 182.9 ± 11.7 cells/mm², t(27) = 0.86, p = 0.41) or the cell cycle marker Ki67 (422.4 ± 24.9 vs. 371.8 ± 30.8 cells/mm², t(27) = 1.27, p = 0.25) were found (Fig. 1A–C), indicating no effects on cell proliferation. However, staining for the early neuronal marker doublecortin (DCX) revealed a significant increase of immunoreactive cells in the Cyfip1+/− animals (367.0 ± 21.9 vs. 550.6 ± 39.4 cells/mm², t(15) = 4.38, p = 6.21 × 10⁻⁴, Fig. 1D, E), consistent with the presence of increased numbers of immature neurons. In a 30 day BrdU pulse-chase protocol, we observed increased numbers of mature postmitotic BrdU+/NeuN+ labelled cells in the dentate gyrus of Cyfip1+/− animals (42.9 ± 7.6 vs. 61.5 ± 3.9 cells/mm², t(10) = 4.54, p = 6.8 × 10⁻⁴), as well as increased proportion of NeuN+ cells in the total BrdU+ population (51.0 ± 2.7% vs. 60.5 ± 0.8%, t(10) = 3.95, p = 0.002) (Fig. 1F–H). These data indicated that the excess numbers of immature neurons seen in the Cyfip1+/− animals were maintained into maturation. We found no significant effects of sex of the animal (see Supplementary Results and Fig. S2).

Increased neuronal numbers in Cyfip1 haploinsufficiency is due to decreased apoptosis
To investigate the cell-intrinsic effects of Cyfip1 haploinsufficiency independent of niche effects we used primary hippocampal progenitor cultures. The in vitro data corroborated the ex vivo finding, showing an increase in the fraction of DCX immunoreactive cells in the cultures prepared from Cyfip1+/− animals (5.5 ± 0.5% vs. 8.8 ± 1.4%, t(16) = 2.78, p = 0.014, Fig. 2A, B) in the absence of any effects on proliferation (see Supplementary Results and Fig. 1F). There were no differences in the proportions of earlier progenitors of the 1/2a GFAP+/nestin+ type (9.4 ± 1.2% vs. 10.0 ± 1.7%, t(35) = 0.76, p = 0.79), or 2b GFAP−/nestin+/DCX− types (22.1 ± 1.6% vs. 20.2 ± 2.5%, t(35) = 0.67, p = 0.51), (Fig. 2C–E), indicating no effects on early stages of neurogenesis. We next assessed general cell viability and survival. We saw a significant increase in the proportion of viable MitoTracker+ cells (82.4 ± 1.2% vs. 89.6 ± 0.7%, t(14) = 3.23, p = 0.002) and fewer propidium iodine positive dead/dying cells (7.7 ± 0.6% vs. 5.4 ± 0.5%, t(16) = 2.44, p = 0.026) (Fig. 2F–H). This suggested an apoptotic mechanism. Indeed, we found significantly fewer cleaved-caspase-3 immunoreactive cells in vivo in the dentate gyrus of Cyfip1+/− animals (2.1 ± 0.2 vs. 1.2 ± 0.1 cells/mm², t(23) = 3.01, p = 0.006, Fig. 2I, J). To confirm this, we investigated apoptosis specifically in the DCX+ populations in our cultures. We found a marked reduction in the fraction of DCX+ cells positive for cleaved-caspase-3 (25.9 ± 5.5% vs. 7.2 ± 2.5%, t(16) = 2.32, p = 0.034, Fig. 2K, L), again demonstrating a high level of concordance between effects in cell culture and intact brain tissue. We found no sex effects in any of the phenotypes investigated in the primary cultures (see Supplementary Results and Fig. S4).

Together, these data suggested Cyfip1 haploinsufficiency impaired a normal homeostatic apoptotic mechanism for modulating newborn neuron number, leading to a greater number surviving to go on to maturity.

Evidence that apoptosis of newly born neurons is mediated by microglia secreted factors
Given the known roles microglia play in earlier stages of neurogenesis, we speculated that microglia may play a role in the survival and death of adult newborn immature neurons. First, we established that there were no genotype differences in density of Iba1+ microglia in hippocampus ex vivo or in vitro (Fig. S5, in either sex). To investigate the
role of microglia in neuronal apoptosis, we depleted microglia from wild-type primary hippocampal progenitor cultures using the specific microglia toxin Mac-1-SAP, targeting CD11b expressing cells. We established the efficacy of the toxin against primary mouse microglia (Fig. 3A).

Depletion of microglia from hippocampal cultures led to an increase in the proportion of DCX+ cells (7.5 ± 1.6% vs. 12.7 ± 0.3%, t(7) = 3.19, p = 0.019, Fig. 3B), and a specific decrease in the fraction of apoptotic DCX+ cells (13.1 ± 3.1% vs. 2.7 ± 1.7%, t(7) = 2.93, p = 0.026, Fig. 3C).

Fig. 1 Cyfp1 haploinsufficiency increases neurogenesis in vivo in the absence of changes in proliferation. A Experimental scheme, proliferation rates were studied with a 6 h pulse of BrdU. B Representative immunohistochemistry showing distribution and location of proliferative (Ki67+) and dividing (Ki67+BrdU+) cells in wild-type and Cyfp1+−/− animals (n = 4 each), respectively. C Quantification showed no difference in numbers of proliferative cells or number of cell divisions. D Representative staining for DCX in wild-type and Cyfp1+−/− animals (n = 8 each), respectively. E Quantification showed a significantly larger number of immature neurons in the Cyfp1+−/− animals. F Experimental scheme, cell maturation was studied with a 30 day BrdU pulse-chase. G Representative images showing BrdU+ and NeuN+ cells. Arrowheads indicate BrdU+ cells H Cyfp1+−/− animals (n = 6) had significantly higher numbers of BrdU+NeuN+ cells than the wild-type animals (n = 5). I The fraction of BrdU+ cells which had matured into NeuN+ cells was significantly increased in the Cyfp1+−/− animals. All data are shown as mean ± SEM.
Fig. 2 Primary hippocampal progenitor cultures prepared from P7 to P8 brain confirm ex vivo findings and indicate that increased neuronal numbers in Cyfp1<sup>+/−</sup> animals is due to reduced apoptosis. A Immunoreactivity for immature neuron marker DCX and hippocampal marker Prox1 confirm cultures contain hippocampus-derived developing neurons. B Cyfp1<sup>+/−</sup> animals showed a significantly higher proportion of DCX<sup>+</sup> cells in the total DAPI<sup>+</sup> population (n = 17). C Representative staining for progenitor markers GFAP and nestin showed cultures contained a mixed population of progenitors at differing stages of development. D There was no significant difference in the proportion of early GFAP+/nestin<sup>+</sup> cells (n = 36). E The proportion of later GFAP<sup>−</sup>/nestin<sup>+</sup> cells (n = 36) was also unaffected. F Live imaging showed cultures are largely comprised of healthy, MitoTracker<sup>+</sup> cells, with some dying PI<sup>+</sup> cells. G A small but significant increase in the proportion of MitoTracker<sup>+</sup> cells was observed in the Cyfp1<sup>+/−</sup> cultures (n = 15). H Conversely, a significant decrease in the proportion of PI<sup>+</sup> cells was seen in the Cyfp1<sup>+/−</sup> cultures (n = 14). I Examples of apoptotic cells in the dentate gyrus, as marked by immunoreactivity for cleaved-caspase 3. J Cyfp1<sup>+/−</sup> animals (n = 13) have significantly fewer apoptotic cells than wild-type animals (n = 12). All data are shown as mean±SEM. K Immunoreactivity for apoptotic marker cleaved-caspase with DCX and nestin showed specific neuronal apoptosis in primary cultures. Arrowheads indicate an apoptotic neuron. L The fraction of immature neurons undergoing apoptosis was significantly lower in the Cyfp1<sup>+/−</sup> cultures (n = 17).
without any associated changes in the nestin+ population (50.7 ± 2.7% vs. 53.8 ± 1.0%, \( t_{17} = 1.07, p = 0.32 \), Fig. 3D), phenocopying the Cyflp1\(^{+/−}\) cultures, and showing a significant role for microglia in regulating DCX numbers. Conversely, addition of exogenous microglia on a semi-permeable membrane, allowing only for soluble factors to influence neuronal apoptosis, had opposite effects. Under these conditions we observed, a decreased proportion of DCX+ cells (5.9 ± 0.8% vs. 1.9 ± 0.5%, \( t_{15} = 4.36, p = 6.5 \times 10^{-4} \), Fig. 3F), increased apoptosis in the DCX+ population (14.8 ± 4.7% vs. 51.7 ± 9.0%, \( t_{15} = 3.27, p = 0.006 \), Fig. 3G) and no effect on the nestin+ population.
Haploinsufficiency of Cyp1 in hippocampal progenitor cultures 2 h after isolation until 3 days in vitro. Dose–response curves of microglial death after Mac-1-SAP administration show near-complete depletion of Iba1+ microglia was achieved within 48 h (n = 6). B Cultures where microglia were depleted (n = 4) showed a higher proportion of DCX+ cells than controls (n = 4). C Correspondingly, the fraction of DCX cells undergoing apoptosis, as marked by cleaved-caspase 3 staining, was significantly decreased in the absence of microglia. D The proportion of nestin+ cells was unaffected by microglia depletion. E Experimental setup, membrane inserts containing wild-type microglia prepared from P7–8 brain were added to cultures 2 h after progenitor isolation and were present throughout the remainder of the experiment. Due to the experimental conditions, no cell–cell contact between microglia and progenitors was possible, but secreted factors were free to diffuse. F The presence of microglia-containing inserts (n = 8) significantly decreased the proportion of DCX expressing cells compared to controls (n = 8). G Conversely, the added microglia significantly increased apoptosis in DCX+ cells. H No effect was seen in nestin expressing cells. I Experimental design, wild-type hippocampal progenitors were exposed to conditioned medium from either wild-type or Cyop1+/- microglia. J In wild-type cultures, conditioned medium from wild-type microglia induced a significant reduction in the proportion of DCX+ cells, whereas medium from Cyop1+/- microglia was unable to do this (n = 4 in all conditions). K An inverse pattern was seen in neuronal apoptosis where wild-type microglia medium increased apoptosis in DCX+ immature neurons but Cyop1+/- medium did not. L Nestin+ cells were not significantly affected by the presence of microglial factors. M Experimental design, Cyop1+/- hippocampal progenitors were exposed to conditioned medium from either wild-type or Cyop1+/- microglia. N As seen in the wild-type cultures, a significant decrease in DCX+ cells was observed only after stimulation with medium from wild-type microglia (n = 4 in all conditions). O Similarly, medium from Cyop1+/- microglia was unable to induce apoptosis in Cyop1+/- progenitors. P Again, the nestin+ population was unaffected. All data are shown as mean ± SEM.

58.4 ± 3.3% vs. 62.7 ± 3.0%, t(15) = 0.98, p = 0.34, Fig. 3H). These experiments provide converging evidence for a previously unknown function of microglia in inducing apoptosis of immature neurons in the hippocampus, which is mediated, at least in part, by soluble factors.

Haploinsufficiency of Cyp1 to support apoptosis of newborn immature hippocampal neurons

Having established a novel microglial dependent mechanism for regulation of neuronal survival, we next tested whether effects on this process could underlie the effects of Cyp1 haploinsufficiency on neuronal survival. To directly address the potential role the induction of apoptosis by microglial secreted factors in an unbiased way, we used a cross-genotype conditioned medium approach. Medium obtained from isolated wild-type and Cyp1+/- microglia preparations was added to both the wild-type and Cyp1+/- hippocampal progenitor enriched cultures (see Fig. 3I, M for schematic of design) and cell proportions and levels of apoptosis were determined. In the wild-type progenitor cultures, adding conditioned medium from wild-type microglia resulted in a significant reduction in the proportion of DCX+ cells, compared to unstimulated cells, whereas Cyp1+/- conditioned medium had no effect (5.4 ± 0.9% vs. 2.9 ± 0.2% vs. 5.4 ± 0.3% for unstimulated, wild-type stimulated and Cyp1+/- stimulated respectively, main effects of genotype F(2,9) = 6.31, p = 0.019, Fig. 3I). Post hoc testing confirmed significant differences between controls and wild-type medium (p = 0.031) and between those stimulated with wild-type and Cyp1+/- medium (p = 0.033). This pattern of effects was mirrored inversely in the fraction of DCX+ cells that were also immunopositive for cleaved-caspase-3, with only conditioned medium from wild-type microglia cultures increasing apoptosis (16.1 ± 3.6% vs. 34.5 ± 3.5% vs. 14.8 ± 2.5%, main effects of genotype F(2,9) = 11.57, p = 0.003, Fig. 3K). Again, post hoc testing showed the wild-type stimulated condition was significantly different from controls and Cyp1+/- stimulated condition (p = 0.007 and p = 0.005, respectively). Consistent with the conclusion that these effects were limited to the DCX+ immature neuron population of cells, the proportion of cultures that was nestin+ was unchanged across all conditions (65.0 ± 5.2% vs. 60.7 ± 4.4% vs. 57.1 ± 4.1%, F(2,9) = 0.75, p = 0.50, Fig. 3L).

Importantly, identical patterns of effects were seen in the Cyp1+/- hippocampal progenitor cultures. Using an identical experimental design (Fig. 3M), we found that conditioned medium from wild-type microglia, but not from Cyp1+/- microglia, decreased the proportion of DCX+ cells (8.1 ± 0.6% vs. 5.1 ± 0.2% vs. 9.0 ± 0.7%, main effects of genotype F(2,9) = 13.82, p = 0.002, Fig. 3N) with post hoc analysis confirming wild-type microglia medium stimulated conditions were significantly different from control and Cyp1+/- stimulated conditions (p = 0.008 and p = 0.002, respectively). Similarly, only conditioned medium from wild-type microglia was associated with a significant increase in the proportion of apoptotic DCX+ cells (12.8 ± 2.3% vs. 28.3 ± 5.0% vs. 13.0 ± 0.9%, main effects of genotype F(2,9) = 6.12, p = 0.021, Fig. 3O) with post hoc analysis confirming the wild-type stimulated condition was significantly different from the unconditioned medium controls and Cyp1+/- stimulated condition (p = 0.007 and p = 0.005, respectively). As before, no effect of any of the medium conditions was seen on the nestin-positive population (64.7 ± 2.6% vs. 64.0 ± 2.3% vs. 57.5 ± 3.3%, F(2,9) = 2.13, p = 0.18, Fig. 3P). These data provide converging evidence for induction of neuronal
apoptosis through factors secreted by microglia a process which is disrupted in Cytfp1^+/− animals.

**Proliferative cells are abnormally positioned in the Cytfp1^+/− hippocampus**

As well as having to survive, newborn cells have to migrate to the correct location to functionally integrate into existing hippocampal circuitry. We noted that the gross positioning of newborn cells appeared different in the Cytfp1^+/− animals. Cells undergoing proliferation are normally largely limited to the subgranular zone (SGZ) of the dentate gyrus, apart from some radially migrating neuroblasts undergoing their final divisions. Indeed, the majority of K67+ cells were found in the SGZ, however a significantly larger fraction was found in the SGZ in the Cytfp1^+/− animals (Fig. 4A, B, 80.4 ± 1.1% vs. 88.2 ± 0.6%, t14 = 6.001, p = 3.2 × 10^-5). As well as an increased fraction remaining in the SGZ, Cytfp1^+/− K67+ cells were also found significantly less far out into the granular zone (Fig. 4C, 17.0 ± 1.4 µm vs. 11.9 ± 1.0 µm, t14 = 2.999, p = 0.009). These data suggested a migration deficit in Cytfp1^+/− animals.

**Cytfp1^+/− adult-born neurons show impaired migration into the granular layer**

In order to gain direct evidence of abnormal migration in Cytfp1^+/− animals we tracked the final positioning of newly born neurons using a 30 day BrdU pulse-chase paradigm followed by staining for BrdU/Neun. Neurons born in the SGZ normally undergo a radial migration outwards into the granular layer. Consistent with a migration deficit in the Cytfp1^+/− animals we found a greater proportion of BrdU+/NeuN+ cells had remained in the SGZ in (Fig. 4D, E, 52.2 ± 3.6% vs. 66.7 ± 2.8%, t12 = 3.197, p = 0.008) and those cells that had migrated had moved a smaller distance into the granular zone (Fig. 4F, 13.6 ± 2.3 µm vs. 6.0 ± 0.7 µm, t12 = 3.549, p = 0.004). When we further analysed the location of the BrdU+/NeuN+ cells that had left the SGZ by dividing the dentate gyrus into four quartiles mapping on to the route taken by migrating cells, we found that the distribution of cells over these quartiles was significantly different (Fig. S6, χ^2 = 25.7, p = 1.1 × 10^-5) and, as would be anticipated with a migration deficit, BrdU+/NeuN+ cells in the Cytfp1^+/− animals were skewed towards the two most proximal quartiles (i.e. those closest to the starting point in the SGZ). Finally, we examined the putative migration phenotype observed in vivo by performing time-lapse imaging in primary hippocampal cultures. Over the 2h imaging period, Cytfp1^+/− cells covered significantly less distance than wild-type cells (Fig. 4G, H, 133.0 ± 4.7 vs. 90.7 ± 12.0 µm, t100 = 3.993, p = 0.003). Both the in vivo and in vitro findings were consistent with a migration deficit due to Cytfp1 haploinsufficiency.

**Actin dynamics are altered in Cytfp1^+/− cells and inhibition of Arp2/3 activity rescues migration deficits in vitro**

As CYFIP1 contributes to the regulation of actin branching and polymerisation through the WAVE1 complex, a function crucial for cytoskeletal reorganisation and cell migration, we speculated that the Cytfp1^+/− related migration phenotype was related to alterations in actin dynamics. A main determinant of actin function in the context of cytoskeleton reorganisation is the ratio of filamentous (F) to globular (G) actin. We first checked this ratio in primary hippocampal cultures, staining with fluorescently labelled phalloidin for F-actin and DNase1 for G-actin, and found a significant increase in F-G-actin ratio in Cytfp1^+/− cells (Fig. 5A, B, 0.67 ± 0.3 vs. 1.48 ± 0.28, t5 = 2.904, p = 0.034). An increase in F-G-actin ratio would be consistent with concurrent increased activity in Arp2/3 brought about by an expected reduction in CYFIP1-WAVE1 complexing under conditions of Cytfp1 haploinsufficiency. Hence, we further speculated that treatment with Arp2/3 inhibitors would influence, and potentially rescue, the abnormal cell migration phenotypes in Cytfp1^+/− cultures. To do this we used two drugs; CK-548, which induces a conformation change inhibiting actin binding and CK-666, which stabilises the inactive form of Arp2/3.

Analysis by time-lapse imaging showed that both drugs were effective in influencing cell migration. However, the effects were essentially opposite in wild-type and Cytfp1^+/− cultures. In wild-type cultures, treatment with Arp2/3 inhibitors had a significant effect on migration (Fig. 5C, vehicle: 107.3 ± 5.6, CK-548:76.3 ± 3.9, CK-666: 65.2 ± 5.4 µm, F(2,24) = 18.91, p = 1.17 × 10^-5), with both CK-548 and CK-666 showing significant decreases in distance moved (p = 5.8 × 10^-4 and 1.1 × 10^-5, respectively). In Cytfp1^+/− cultures, there was also a significant effect on migration by Arp2/3 inhibitors (Vehicle: 68.1 ± 4.4, CK-548:117.2 ± 8.8, CK-666:112.0 ± 9.8 µm, F(2,6) = 11.39, p = 0.009). In contrast to wild-type cultures, both CK-548 and CK-666 caused significant increases in distance moved (p = 0.011 and p = 0.019, respectively), restoring migration to wild-type levels and suggesting excessive Arp2/3 activity was responsible for the migration deficit in Cytfp1^+/− cells.

**Discussion**

We have shown changes in the survival and migration of newly born neurons in hippocampus in a model of Cytfp1 haploinsufficiency, recapitulating the lowered gene dosage found in the pathogenic 15q11.2(BP1–BP2) copy number deletion. We report that Cytfp1^+/− animals have larger numbers of surviving newly born neurons in hippocampus, in the absence of increases in numbers of stem cells or rate of proliferation, and provide evidence that this is the result of a specific failure in microglia-induced
apoptosis (see summary of proposed mechanism in Fig. S7). These newly born neurons also show an Arp2/3 dependent deficit in migration, resulting in a failure to reach their appropriate locations within the dentate gyrus.

Our study is the first to our knowledge to demonstrate a direct effect of microglial soluble factors on the survival of immature neurons as a homoeostatic mechanism. In contrast to earlier findings examining the effect of microglia on adult neurogenesis, we show this occurs in the absence of overt inflammatory disease or microglial activation. Our data indicate this is a mechanism susceptible to pathogenic changes, as evidenced by the effects of Cyfip1 haploinsufficiency on the ability of microglia to regulate apoptosis of immature neurons.
The effects of Cyfip1 haploinsufficiency were highly cell-type specific and appeared limited to the stage of cell maturity where neuronal progenitors are fully committed to a neuronal phenotype, a stage of development characterised by high levels of apoptosis. Whilst we have demonstrated that soluble factors from microglia are involved in controlling numbers of immature neurons during adult neurogenesis, and that haploinsufficiency of Cyfip1 leads to reduced apoptosis via an effect on factors secreted from microglia, the soluble factor(s) responsible remain to be identified. Future proteomic and pharmacological work will serve to identify and validate the identity of the responsible mediator(s). Cyfip1 has pleiotropic roles but the potential links between CYFIP1, actin remodelling, and cytokine secretion are especially interesting. A focus for CYFIP1 action in microglia gains further evidence from a recent preprint by showing the effect of cell specific deletion of Cyfip1 on microglia morphology, movement and function.

In parallel, we have shown that adult-born neurons in Cyfip1−/− animals show a significantly impaired migration phenotype. Here our data point to a key role for CYFIP1 effects on actin dynamics mediated by Arp2/3 activity. We hypothesise that lowered dosage will lead to reduced engagement of CYFIP protein with the CYFIP-WAVE1-Arp2/3 pathway resulting in reduced suppression of Arp2/3 activity and abnormal actin cytoskeleton remodelling required for migration. We show evidence to support this hypothesis though, as predicted, an increase in F-G-actin ratio and rescue of the migration phenotype by pharmacological inhibition of Arp2/3 activity. The activation of the WAVE-Arp2/3 system is complex and, can take place in at least two different Rac1 mediated pathways. It is unlikely that CYFIP1 is the sole

Fig. 5 Disruption of Arp2/3 mediated actin dynamics is responsible for migration deficit. A Fluorescently labelled phalloidin and DNase show the amount of filamentous and globular actin, respectively. B Quantification of fluorescence intensity shows Cyfip1−/− cells have an increased F- to G-actin ratio. C Treatment with Arp2/3 inhibitors CK-548 or CK-666 in wild-type cells significantly reduces the distance moved by cells during live imagine, whereas treatment of Cyfip1−/− cells restores distances to wild-type levels.
regulator of this activation, but our data point towards a significant role in this process.

The combined effects of failure to undergo homoeostatic apoptosis and failure to migrate to the correct location is likely to have profound effects on the functionality of adult-born neurons in hippocampus. We observed lasting effects up to 30 days post division, when adult-born neurons will already have local connections. The prolonged survival and expression of mature markers of these ectopic cells makes it likely these cells will be active, indeed, it is known that synaptic NMADRs signalling is required for neuronal survival prior to this developmental timepoint, making it likely these cells are integrated in some form in the local circuitry. The exact functional consequences on a network level are difficult to predict and will require further electrophysiological study, both on the individual cell and network level. Haploinsufficiency of Cyfip1 has been shown to have disease relevant behavioural consequences in both mice and rat models, but to date, much of the neurobiology and behaviour in these models has remained unexplored. Our data would indicate a focus on hippocampal circuitry would prove of interest. Neuronal migration in the hippocampus has, to our knowledge, not been examined in patient populations. However, as AHN does take place in humans, and the biochemical pathways relevant to this study are well conserved, it is likely similar deficits would be observed in 15q11.2BP1–BP2 carriers.

The 15q11.2(BP1–BP2) deletion is not solely associated with schizophrenia, it also markedly increases risk for autism spectrum disorders, ADHD and developmental delay, and epilepsy. Whether altered microglia functioning due to Cyfip1 haploinsufficiency, impacting on AHN, also plays a part in these disorders is an open question, but all have been associated with alterations in immune system and microglia functioning. Likewise, alterations in migration would have equally profound consequences in other disorders.

As noted previously, a number of other schizophrenia risk genes have also been shown to modify AHN in model systems. Whilst the effects are distinct from those of Cyfip1, each risk gene impacting on neurogenesis in its own way, this convergence of function may be indicative of a common pathogenic route for a number of diverse psychiatric risk genes.

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N.H. and L.S.W. designed the study. N.H. designed and performed experiments, and drafted the paper. J.C. performed animal husbandry/genotyping. L.J.W. optimised and assisted with primary culture. W.P.G. and L.S.W. initiated the study and assisted with data interpretation. M.J.O., W.P.G., J.H. and L.S.W. critically appraised the manuscript. All authors approved the final paper.

Conflict of interest
The authors declare no competing interests.

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