Memory and synaptic deficits in *Hip14/DHHC17* knockout mice

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Palmitoylation of neurotransmitter receptors and associated scaffold proteins regulates their membrane association in a rapid, reversible, and activity-dependent fashion. This makes palmitoylation an attractive candidate as a key regulator of the fast, reversible, and activity-dependent insertion of synaptic proteins required during the induction and expression of long-term plasticity. Here we describe that the constitutive loss of huntingtin interacting protein 14 (*Hip14*, also known as *DHHC17*), a single member of the broad palmitoyl acyltransferase (PAT) family, produces marked alterations in synaptic function in varied brain regions and significantly impairs hippocampal memory and synaptic plasticity. The data presented suggest that, even though the substrate pool is overlapping for the 23 known PAT family members, the function of a single PAT has marked effects upon physiology and cognition. Moreover, an improved understanding of the role of PATs in synaptic modification and maintenance highlights a potential strategy for intervention against early cognitive impairments in neurodegenerative disease.

striatum | hippocampus | posttranslational modification | long-term potentiation | Huntington disease

Palmitoylation is a posttranslational protein modification increasingly recognized as an important regulator of neuronal development, synaptic function, and plasticity (1, 2). Palmitoyl acyltransferase (PAT) enzymes regulate membrane association of proteins by catalyzing the addition of the fatty acid palmitate to cysteines via thioester bonds. Palmitoylation is readily reversible, making it an attractive candidate for a regulator of the rapid synaptic protein trafficking required for synaptic transmission and plasticity. The growing list of palmitoylated neuronal substrates includes scaffolds, ion channels, and vesicle-associated proteins, and their palmitoylation status can have dramatic effects on function and/or localization within membranes. For example, synaptic activity dynamically regulates palmitoylation of postsynaptic density protein-95 (PSD-95), influencing its clustering at postsynaptic sites (3–5), whereas palmitoylation of NMDA- and AMPA-type glutamate receptor subunits regulates their insertion, removal, and stabilization in the postsynaptic membrane (6–9). The importance of this modification within the brain is highlighted by implication of palmitoylation deficits in a number of neurological diseases including Alzheimer's disease, schizophrenia, and Huntington disease (HD) (10).

To date, 23 DHHC proteins have been identified in humans. The PAT HIP14 (DHHC17) is enriched in the brain and has a number of known synaptic substrates including PSD-95, GluA1/2, GAD-65, SNAP-25, and synaptotagmin I (11–13), suggesting roles in pre- and postsynaptic function. Indeed, *Hip14* siRNA reduces PSD-95 clustering in hippocampal cultures (12), and *Hip14* loss-of-function impairs neurotransmitter release in *Drosophila* (14). Interestingly, *Hip14* function is impaired in HD (15, 16), suggesting that some synaptic deficits observed in this disease (17–23) may arise from hypopalmitoylation of *Hip14* substrates. In support of this, we recently reported the generation of

Hip14 knockout mice, which share some interesting similarities to late-stage HD mouse models in behavioral, biochemical, and neuropathological measures (11). However, it remains unknown whether the loss of this single PAT can result in synaptic and plastic deficits reminiscent of disease states.

Here we investigate the consequences of a single PAT knockout on neuronal physiology, synaptic signaling, and plasticity. We report that *Hip14* knockout results in marked cellular and synaptic alterations in striatal spiny projection neurons (SPNs) and impaired hippocampal long-term potentiation (LTP). Furthermore, performance in a hippocampal-dependent learning paradigm was also significantly impaired. In conclusion, the constitutive absence of *Hip14/DHHC17* results in major neurophysiological deficits and associated cognitive dysfunction.

Results

Altered SPN Membrane Properties and Excitability. *Hip14* dysfunction has been implicated in the pathogenesis of HD (11). As striatal physiology is well characterized in various HD models (17, 18), we first used whole-cell patch clamp recordings to examine the consequences of *Hip14* knockout on cellular and synaptic properties of SPNs, the most vulnerable cell type in HD. Due to the ~20% loss of striatal cells in these mice (11), in addition to targeting SPNs based on their characteristic somatic diameter (8–16 μ m) and ellipsoid morphology (Fig. 1A) (24), selection criteria were validated by current-clamp recordings in K^+ -based internal solution. Of 17 cells recorded under these conditions (animals aged postnatal day: WT, 25 ± 1 ; *Hip14*^{−/−},

Significance

Palmitoylation can influence the subcellular localization of various synaptic proteins; this process is therefore increasingly recognized as an important regulator of basal synaptic communication as well as activity-dependent synaptic plasticity. Despite the fact that in vitro experiments indicate that many synaptic proteins can be palmitoylated by more than one palmitoyl acyltransferase (PAT), the enzymes responsible for palmitoylation, we describe in this paper various synaptic, plastic, and cognitive consequences that result from constitutive loss of a single PAT, namely *Hip14/DHHC17*. This reveals an important functional role for this PAT that cannot be compensated for by other existing PATs.

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Fig. 2. Pre- and postsynaptic alterations to SPN glutamatergic synapses in *Hip14*^{-/-} slices. (A) Voltage-clamp recordings of sEPSCs at V_h -70 mV in SPNs. (B) There was a significant interaction between genotype and event amplitude cumulative probabilities, due to more large events in *Hip14*^{-/-} SPNs (Inset). (C) There was a significant genotype effect with increased sEPSC IEIs and reduced mean event frequency in *Hip14*^{-/-} SPNs (Inset). (D) Stimulus evoked EPSCs (eEPSCs) carried by AMPARs (-70 mV, paired pulses, Upper) and NMDARs (+60 mV, single pulse, Lower) at 50 (gray) and 150 μ A (black). *Hip14*^{-/-} eEPSCs were significantly smaller for both AMPA-mediated (E) and NMDAR-mediated (F) currents. Paired-pulse facilitation of AMPAR currents was increased (G), whereas NMDAR eEPSC decay kinetics (weighted τ ; H) were decreased in *Hip14*^{-/-} SPNs. No differences in NMDA:AMPA ratio were observed (I). Repeated measures ANOVA ## interaction, $P < 0.01$; ψ genotype, $P < 0.05$; $\psi\psi$ genotype, $P < 0.01$. * t test, $P < 0.05$ (Bonferroni post-hoc in E). IPI, interpulse interval.

Fig. 4C), consistent with the reduced sEPSC frequency and again suggestive of a loss of functional excitatory synapses. Despite reductions in hippocampal volume (11), there were no significant differences in CA1 or CA3 cell counts between genotypes ($P = 0.11$, two-way ANOVA; Fig. 4D), suggesting volume reductions unrelated to pyramidal cell number. To confirm a loss of synapses in *Hip14*^{-/-} CA1, we analyzed dendritic spine density by Golgi impregnation. In agreement with our conclusions from the electrophysiological data, we observed significantly reduced spine densities in CA1 pyramidal neurons of the stratum radiatum from *Hip14*^{-/-} mice ($P < 0.001$, unpaired t test; Fig. 4E).

Impaired Hippocampal Long-Term Potentiation. Next, we investigated long-term potentiation at hippocampal CA3 Schaffer

collateral to CA1 synapses. In WT animals, high-frequency stimulation (HFS; 100 Hz, $1 \text{ s} \times 3$, 10-s intertrain interval) resulted in LTP whereas LTP was entirely absent in *Hip14*^{-/-} mice ($P = 0.02$, unpaired t test of % potentiation 50–60 min post-HFS, herein; Fig. 4F), despite the clear posttetanic potentiation indicative of intact presynaptic short-term potentiation. To determine whether *Hip14* is essential for LTP or whether the threshold for LTP induction is altered as a result of *Hip14* knockout, we facilitated the NMDAR dependence of this form of plasticity (30) by addition of the NMDAR coagonist glycine ($10 + 2 \mu\text{M}$ strychnine). Under these conditions, LTP was partially restored to a level not significantly different from that in WT ($P = 0.12$; Fig. 4G). Finally, we combined this with a more robust induction protocol ($3 \times$ HFS separated by 5 min each)

Fig. 3. Reduced sEPSC frequency in hippocampal CA1 pyramidal cells. (A) No differences were observed in CA1 current-clamp membrane potential responses (i, Upper) to 50-pA current injection steps (i, Lower and ii). (B) In CA1 voltage-clamp recordings of sEPSCs (-70 mV, i) there were no differences in event amplitudes (ii, Left), but there was a significant genotype effect on IEIs (ii, Right) and reduced mean event frequency in *Hip14*^{-/-} (ii, Right Inset). (C) PPRs were similar, suggesting that decreased event frequency is due to reduced synapse number. (D) AMPAR-mediated rectification was unaltered. (E–G) NMDA:AMPA ratios (E), isolated NMDAR decay kinetics (F), and the ifenprodil (ifen) sensitivity of isolated NMDAR currents (G) were also not significantly different. ψ , genotype. * t test; see text for details.

Fig. 4. Impaired LTP due to increased induction threshold and spatial memory deficits in *Hip14*^{-/-} hippocampus. (A and B) fEPSP recordings in the stratum radiatum demonstrated no significant genotype difference in PPR (A) or fiber volley amplitudes (B). (C) fEPSP slope values were significantly reduced in *Hip14*^{-/-} slices. (D) Total hippocampal cell counts showed no difference between WT and *Hip14*^{-/-} mice. (E) Dendritic spine density was decreased in the stratum radiatum of *Hip14*^{-/-} mice (cells: *n* = 28 for WT, 21 for *Hip14*^{-/-}). (F) Hippocampal slices from *Hip14*^{-/-} mice displayed severe deficits in LTP using a stimulation protocol that induced reliable LTP in slices from WT mice. (G) The addition of glycine (10 μ M) to the bath partially alleviated the *Hip14*^{-/-} LTP deficit to levels not significantly different from WT. (H) Combining glycine (10 μ M) with an enhanced stimulation protocol was sufficient to restore the LTP deficit to WT levels. (I and J) Learning a novel-object location (I) is impaired in *Hip14*^{-/-} mice whereas the ability to recognize a novel object remains intact (J). For F–H, superimposed traces on the *Left* are from WT mice before (black) and 50–60 min after (light gray) high-frequency stimulation (HFS). Superimposed traces on the *Right* are from *Hip14*^{-/-} mice before (black) and 50–60 min after (dark gray) HFS. #interaction, *P* < 0.01; **t* test, *P* < 0.05; *****t* test, *P* < 0.001.

and found that this completely restored LTP in *Hip14^{-/-}* mice to WT levels ($P = 0.65$; Fig. 4H). Thus, the loss of a solitary palmitoyltransferase (*Hip14^{-/-}*) produces significant plasticity deficits, resulting in a markedly increased threshold for LTP induction. However, these deficits can be overcome by strong induction paradigms, demonstrating that the machinery required for LTP is intact, even if severely impaired.

Impaired Hippocampal-Dependent Memory. Serious hippocampal plasticity deficits should be reflected by impaired performance in hippocampal-dependent spatial learning tests such as novel-object location memory (31). As anticipated, WT animals spent more time investigating the relocated object in trial 2 ($P = 0.0002$, unpaired t test; Fig. 4J), demonstrating spatial memory

reflected by clear preference for spatial novelty. In contrast, *Hip14^{-/-}* mice failed to show any preference for the object in trial 2 ($P = 0.11$, unpaired t test; Fig. 4*I*), thus exhibiting a marked impairment in spatial memory. When tested in a perirhinal cortex-dependent novel object recognition task (31), in which one of a pair of familiar objects from trial 1 is replaced in trial 2 (same location), both WT ($P = 0.015$, paired t test) and *Hip14^{-/-}* mice ($P = 0.004$, paired t test) spent significantly more time investigating the novel object (Fig. 4*J*). Thus, although object recognition memory is intact in *Hip14^{-/-}* mice, they exhibit marked deficits in both hippocampal plasticity and spatial memory.

Discussion

Numerous PATs regulate palmitoylation in the mammalian brain. Despite considerable substrate overlap between these PATs, we found a number of electrophysiological and behavioral consequences induced by the loss of Hip14/DHHC17 alone. In the striatum we observed alterations in membrane properties and a reduction in spontaneous excitatory neurotransmission, likely due to both reduced synapse numbers (11) and decreased Pr. Perhaps most striking, however, was the profound impairment of hippocampal LTP and spatial memory in *Hip14*^{-/-} mice. Our data confirm a clear role for palmitoylation in synaptic function and suggest that HIP14 dysfunction in HD may account for a number of synaptic and plastic deficits. Furthermore, our results highlight a major modulatory requirement of Hip14 in the gating of hippocampal synaptic plasticity and the formation of spatial memory.

We previously reported a decrease in excitatory synapses in the striatum of *Hip14*^{-/-} mice (11), suggesting deficits in excitatory transmission to the basal ganglia. Consistently, we found here that SPNs from *Hip14*^{-/-} mice had reduced spontaneous excitatory event frequencies and evoked current amplitude. We, and others, have shown that spine formation and maintenance in SPNs is dependent upon an intact glutamatergic network (32, 33); it is tempting to suggest that synapse reductions in *Hip14*^{-/-} mice result from a breakdown in structural connectivity, driven by impaired transmission. A critical role for Hip14 in transmitter release has been described in *Drosophila* (14), and our conclusions are further supported by the identification of SNAP-25 and synaptotagmin as HIP14 substrates (12, 14). However, no Pr alterations were observed in CA1 neurons, suggesting that release deficits are not global.

Hip14 deletion was sufficient to impair hippocampal LTP and performance on a spatial memory task. We found that fEPSP responses were diminished in *Hip14*^{-/-} stratum radiatum, which, together with the observed decrease in sEPSC frequency and reduction in spine density, is highly suggestive of fewer excitatory synapses. It is unlikely that this alone can account for the LTP deficit, as we saw no correlation between the initial response size and the amount of potentiation 50–60 min post-HFS ($r^2 = 0.11$, $P = 0.42$). Furthermore, the conditions that restored LTP to WT levels (glycine and enhanced stimulation) would be expected to influence existing synapses rather than recruit additional synapses.

The idea that protein palmitoylation can influence synaptic plasticity has been previously suggested (2). As the growing list of palmitoylated neural proteins includes a variety of surface receptors, scaffolding proteins, and vesicle fusion proteins among others, the precise mechanism by which *Hip14* knockout impairs LTP is difficult to pinpoint and remains to be determined. NMDAR subunits are palmitoylated at two cysteine clusters; palmitoylation of cluster 1 enhances synaptic stability whereas palmitoylation of cluster 2 increases retention within the Golgi and reduces surface expression (9). In the present study, NMDA:AMPA ratios were unaffected in both the striatum and hippocampus, suggesting a lack of up- or down-regulation of synaptic NMDARs following *Hip14* knockout and arguing against significant depalmitoylation at cluster 1 (34). Moreover,

we also rule out the possibility that the observed LTP deficit is due to a change in the composition of synaptic NMDARs as NMDAR decay kinetics and ifenprodil sensitivity were unaffected in *Hip14* knockouts.

AMPA subunits, which are HIP14 substrates (13), are palmitoylated at two different cysteines; palmitoylation of the TMD-2 cysteine promotes accumulation in the Golgi and prevents forward trafficking to the surface, whereas depalmitoylation of the C-terminal cysteine enhances PKC phosphorylation of nearby serines, association with 4.1N, and insertion into the membrane (7, 35). We did observe a significant increase in a subset of large AMPAR-mediated EPSCs in *Hip14*^{-/-} SPNs, which may be a consequence of depalmitoylation-induced release from the Golgi. Interestingly, although depalmitoylation of the C-terminal cysteine enhances extrasynaptic insertion events of AMPA receptors, these receptors are not stable as the authors found no evidence of increases in steady-state AMPAR surface expression (7). This suggests that some degree of AMPAR palmitoylation is required for surface retention. Such stability within the membrane would be absolutely essential for LTP, which is significantly impaired at *Hip14*^{-/-} CA1 synapses.

It has also been shown that palmitoylation of scaffolding proteins PSD-95 (28) and A-kinase anchoring protein 79/150 (29) is required for LTP in cultured hippocampal neurons, and recent work has highlighted the importance of certain SNARE family vesicle fusion proteins in LTP expression (36), some of which are recognized or candidate palmitoylated proteins (37). Thus, our observed LTP deficit may result from depalmitoylation of a number of synaptic proteins. We have previously reported reduced PSD-95 and SNAP-25 palmitoylation in brain tissue from *Hip14*^{-/-} mice (11). Although our results do not provide a definitive conclusion regarding the mechanism of impaired LTP, our findings are clearly behaviorally relevant as highlighted by impairments in hippocampal-dependent memory. That said, perirhinal cortex-dependent novel-object recognition (31) was not altered, suggesting that *Hip14* knockout does not result in global memory deficits and that certain neuronal networks are more vulnerable than others.

In vitro expression studies have revealed considerable overlap between PAT substrates. For example, multiple PATs in addition to HIP14, including DHHC2, -3, -5, and -7, are able to palmitoylate PSD-95 (2, 38). Due to this apparent lack of substrate specificity, one might expect a similar overlap of the cellular and behavioral phenotypes of various DHHC-deficient animal lines. However, the PAT-substrate relationship in vivo will depend on the expression levels and localization of both the substrates and the PATs themselves within individual cells. Perhaps it is not surprising, then, that considerable phenotypic differences exist among the DHHC-deficient mice. DHHC8 knockout mice have sexually dimorphic alterations in locomotor activity and prepulse inhibition (39), whereas DHHC-5-deficient mice exhibit impaired fear conditioning and are born at half the expected rate (40). Furthermore, polymorphisms in the human DHHC8 gene are implicated in schizophrenia (39), whereas human DHHC9 and 15 mutations are associated with X-linked mental retardation (41, 42). Thus, despite partial in vitro substrate overlap, relatively specific substrate effects occur in single PAT knockouts. Moreover, in the present study, we found deficits in striatal neurons that were not present in CA1 cells, demonstrating that the loss of a single PAT does not affect all networks equally.

Although wild-type huntingtin interacts with HIP14 and enhances its PAT activity, this interaction is reduced in the presence of mutant huntingtin (16). *Hip14*^{-/-} mice recapitulate some of the behavioral, biochemical, and neuropathological deficits observed in HD, and *Hip14* function is reduced in HD mice (11). Interestingly, nearly all of the striatal cellular and synaptic changes observed here are reminiscent of those in

various HD models, albeit at late stages of phenotype progression (17, 18). In the present study, we were able to observe electrophysiological changes within the striatum of 3- to 6-wk-old *Hip14*^{-/-} mice that do not occur in HD models until months later. Also, SPN loss and striatal volume reductions are evident at birth in *Hip14*^{-/-} mice (11), whereas the striatal degeneration in HD, and in HD mice, progresses over time (17).

It is interesting to note that several HD mouse models demonstrate impaired, but not absolute, loss of LTP at the earliest stages of their phenotype progression (19, 21–23, 43). In addition to our findings here, those findings in HD mice suggest that palmitoylation deficits may underlie early synaptic dysfunction in HD mice, which may contribute to early cognitive deficits reported in prodromal HD carriers (44, 45). Thus, manipulations aimed at enhancing protein palmitoylation may be a valuable strategy to combat early cognitive symptoms in HD.

Materials and Methods

Animals were housed and maintained according to the Canadian Council on Animal Care and all procedures were approved by the University of British Columbia Committee on Animal Care. Experiments were conducted on the same *Hip14*^{-/-} mouse line that was previously characterized (11). All data are expressed as mean $\pm$ SEM, and the statistical analyses used included Student *t* test and two-way ANOVA as detailed in the text.

Electrophysiology. Standard whole-cell patch-clamp striatal recordings were obtained from 400- μ m coronal slices from ~30-d-old male mice as previously described (24), with recording conditions adapted from ref. 46 to contain 10 mM glucose and as detailed in the text. Horizontal hippocampal slices (400 μ m) were prepared as previously described (47), and plasticity experiments were produced as in ref. 30 and as described in the text. Both whole-cell and field responses were evoked by 40- μ s stimulation pulses delivered through a glass micropipette placed within 200–400 μ m of the recording site.

Cell Counts. Mice were terminally anesthetized by i.p. injection of 2.5% (vol/vol) avertin and perfused with 3% (vol/vol) paraformaldehyde/0.15% glutaraldehyde in PBS. Mouse brains were fixed in the same solution for 24 h at 4 °C followed by cryoprotection in 30% (vol/vol) sucrose. Brains were frozen on dry ice and cut into coronal sections (25 μ m) using a cryostat microtome (HM 500 M, Microm International). Every eighth section throughout the hippocampus spanning from Bregma -0.94 to -2.8 mm was stained with an antibody reactive to NeuN (Chemicon), a marker of neuronal nuclei, as described previously (48). CA1 and CA3 hippocampal subregions were delineated in every eighth NeuN-stained section using a 2.5 $\times$ objective, and then cells were counted using a 40 $\times$ objective on a Zeiss Axio Imager M2 microscope with a motorized computer-controlled stage (Ludl Electronics) and a CX9000 Digital CCD video camera (MicroBrightfield). Total estimated number of CA1 and CA3 neurons was obtained using the Fractionator Probe in StereoInvestigator 9, a morphometry and stereology software package (MicroBrightfield) with a 25- $\times$ 25- μ m counting frame and 150- $\times$ 150- μ m grid size. Overall, the coefficient of error was <0.1. All quantification was conducted with the investigator blind to genotype.

Spine Analysis. Animals were transcardially perfused with PBS and Golgi impregnated using the FD Rapid GolgiStain kit (FD Neurotechnologies) following the manufacturer's instructions. Final steps were performed on 150- μ m-thick coronal sections (cut by cryostat at -22 °C) before coverslip and permount application.

Dendrites were imaged with an oil-immersion lens at 63 $\times$ magnification using a Zeiss Axiovert 200M microscope. Analysis was restricted to the posterior stratum radiatum, and dendrite collaterals off the apical dendrite of CA1 pyramidal neurons were chosen as regions of interest (ROI). Images were collected throughout the depth of the tissue at 2- μ m steps in the z-plane and converted to TIFF files, and spines were counted manually in ImageJ with the experimenter blinded to genotype. Long dendritic protrusions lacking a clear head were excluded from the analysis. Two to three ROIs were averaged for each cell, and seven cells were analyzed per animal (animal *n* = 4 for WT; *n* = 3 for *Hip14*^{-/-}).

Novel-Object Location and Recognition Testing. Novel-object testing took place in open 50- $\times$ 50-cm black plastic boxes with 16-cm sides. Testing occurred over a period of 2 d, and the same cohort of animals was used for both the novel-object location task (conducted on day 1) and the novel-object

recognition task (conducted on day 2). Recordings were collected using the automated behavioral analysis software Ethovision (Noldus) via live video collected from a ceiling-mounted video camera. On day 1, mice were exposed to the box for 10 min before being removed for 5 min. Two different novel objects were then placed in the upper corners of the box, far enough from the edges to allow movement around the perimeter. Mice were then reintroduced to the box in the lower left corner, and their movements were recorded for 5 min. The number of investigations of both the novel objects was scored (trial 1). Mice were then removed from the box, and the object in the top right corner was moved to the lower right corner. Mice were reintroduced to the box again for another 5 min, and then the number of investigations of the objects was scored (trial 2). On day 2, the 10-min exposure and trial 1 were identical to that on day 1. However, in trial 2 of day 2, the object in the top right corner was replaced with a different and

unfamiliar object positioned in the top right corner. The number of investigations was recorded for a 5-min period.

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