Traf2 and NcK interacting kinase (TNiK) contains serine-threonine kinase and scaffold domains and has been implicated in cell proliferation and glutamate receptor regulation in vitro. Here we report its role in vivo using mice carrying a knock-out mutation. TNiK binds protein complexes in the synapse linking it to the NMDA receptor (NMDAR) via AKAP9. NMDAR and metabotropic receptors bidirectionally regulate TNiK phosphorylation and TNiK is required for AMPA expression and synaptic function. TNiK also organizes nuclear complexes and in the absence of TNiK, there was a marked elevation in GSK3β and phosphorylation levels of its cognate phosphorylation sites on NeuroD1 with alterations in Wnt pathway signaling. We observed impairments in dentate gyrus neurogenesis in TNiK knock-out mice and cognitive testing using the touchscreen apparatus revealed impairments in pattern separation on a test of spatial discrimination. Object-location paired associate learning, which is dependent on glutamatergic signaling, was also impaired. Additionally, TNiK knock-out mice displayed hyperlocomotor behavior that could be rapidly reversed by GSK3β inhibitors, indicating the potential for pharmacological rescue of a behavioral phenotype.

These data establish TNiK as a critical regulator of cognitive functions and suggest it may play a regulatory role in diseases impacting on its interacting proteins and complexes.

Introduction

Central to understanding the molecular basis of cognitive functions are the signaling mechanisms connecting neurotransmitter receptors to intracellular pathways regulating transcription, translation, and changes in electrical properties of neurons. It has become apparent that many of the proteins that participate in these pathways are physically organized within the cytosol into multiprotein complexes that act as molecular machines exploiting their different protein components to perform regulatory functions (Husi et al., 2000; Scott and Pawson, 2009). Within many signaling complexes are protein kinases that phosphorylate the nearby proteins and thereby orchestrate a variety of cellular functions (Scott and Pawson, 2009). How neuronal signaling complexes function is poorly understood and there are very few examples of studies where the dysfunction of signaling complexes has been studied following a mutation in the intact animal.

Toward these issues, we were intrigued by Traf2 and NcK interacting kinase (TNiK), a protein with both scaffolding and kinase domains that had been implicated in postsynaptic signaling as well as in regulation of cell proliferation (Mahmoudi et al., 2009; Shitashige et al., 2010). TNiK is expressed in the nervous system but its role in vivo is currently unknown. A recent study showed that activation of NMDA receptors (NMDARs) regulates phosphorylation of TNiK (Coba et al., 2009). Moreover, knockdown of TNiK in primary cultured neurons decreases surface GluA1 levels (Hussain et al., 2010) and alters the synchrony of network activity (MacLaren et al., 2011), suggestive of a postsynaptic signaling function at excitatory synapses. TNiK has also been implicated in controlling dendritic outgrowth mediated by a ternary complex involving the E3 ubiquitin ligase Nedd4-1, Rap2A, and TNiK (Kawabe et al., 2010). In non-neuronal cells TNiK modulates cell proliferation by regulating activation of Wnt signaling cascade through its ability to interact with β-catenin and phosphorylate the transcription factor Tcf7l2 (Mahmoudi et al.,...
2009; Shitashige et al., 2010). It is unknown if TNIK plays any role in neurogenesis or brain development. Finally, a link between TNIK and schizophrenia has also been suggested based on the observation that TNIK binds Disrupted in Schizophrenia 1 (DISC1) in vitro resulting in decreased TNIK levels and kinase activity (Wang et al., 2010). Human genetic studies have not identified mutations in TNIK, although several association studies have suggested TNIK to be involved in schizophrenia, attention deficit hyperactivity disorder, and general cognitive function (Potkin et al., 2009; Shi et al., 2009; Ayalew et al., 2012; Elia et al., 2012).

Here we address the in vivo role of TNIK by examining mice carrying a knock-out mutation in TNIK and show the mutation leads to dysregulation of key synaptic and nuclear signaling mechanisms. We identify complexes of proteins associated with TNIK in the posttranslational density and the nucleus and show that the TNIK mutation has a dramatic impact on the regulation of GSK3β and phosphorylation of proteins within the complexes. We assessed the requirement of TNIK for synaptic plasticity, neuronal development and specific aspects of higher order cognitive processing using a computerized touchscreen apparatus (Bussey et al., 2012) and find evidence that TNIK plays a role in multiple cognitive functions through both synaptic and nuclear signaling pathways.

Materials and Methods

Generation of TNIK mutant mice. The targeting vector was constructed using the AB2.2 genomic DNA BAC clone. The vector containing 6.9 and 2.9 kb of 5' and 3' homology arms, respectively, replaced 2.6 kb of TNIK genomic DNA (X8438374 to X8440972; Ensembl Build 55) containing part of exon 6 and 7 that encoded the kinase domain with IRES-lacZ-neo reporter cassette. The targeting construct was electroporated into E14TG2a embryonic stem (ES) cells. G418 (neo-resistant) clones were screened for homologous recombination by long-range PCR using the Expand Long Template PCR system (Roche Cat No 1186184001) with PCR primer \( (5'\text{-GAGCTATTCCAGAGTATGGA-3'}\) ) and primer Y \( (5'\text{-CAGAGGCTTCTGATCTCC-3'}\) ) that correspond to sequence in the IRES-lacZ-neo cassette and sequence outside the 2.9 kb flanking region, respectively. The correctly targeted ES cells were injected into C57BL/6 blastocysts to create chimeric mice, which were bred with 129S5 C57BL/6J blastocysts to create chimeric mice, which were bred with 129S5/SvEvTac mice to generate heterozygous (+/−) TNIK mutant mice. Those F1 heterozygous mice had been backcrossed with 129S5 mice 1–2 times before being used for intercrossing. Genotyping PCR consisted of a 540 bp product amplified from the wild-type (wt) allele using a forward primer A \( (CAAGTCTCTCTCATTAGTGG) \) and primer B \( (CAAGGCTTTGCTCATTCCTC) \) downstream of the cassette. A 700 bp product was amplified from the targeted allele using primer B with forward primer X, within the selection cassette. After enzymatic amplification for 35 cycles (45 s at 94°C, 45 s at 55°C, and 1 min at 72°C), the PCR products were size fractionated on a 2% agarose gel in 1× Tris-borate-EDTA buffer. Animals were treated in accordance with the UK Animal Scientific Procedures Act 1986.

Biochemical assays. Posttranslational density (PSD) and nuclear fractions from mouse hippocampus were prepared using a three-step protocol as previously described (Coba et al., 2009). In brief, tissue was homogenized in ice-cold buffer (10 mM Tris-HCl, pH 7.4/320 mM sucrose, 20 mM β-glycerol phosphate, 1 mM sodium orthovanadate, and 50 mM sodium fluoride) and protease inhibitors (Roche-Complete protease inhibitor mixture). The homogenates were centrifuged at 800 × g for 10 min at 4°C and pellet resuspended in 50 mM HEPES, pH 7.4, 40 mM β-glycerol phosphate, 1 mM sodium orthovanadate, and 50 mM sodium fluoride and protease inhibitors (Roche-Complete protease inhibitor mixture). The homogenates were centrifuged at 16,000 × g for 15 min at 4°C and supernatants used for nuclear extract assays. Data presented as mean ± SEM and statistical significance determined using unpaired Student’s t tests.

Hippocampal slice electrophysiology. Synaptic physiology was assessed using a multi-electrode array system and whole-cell voltage-clamp techniques as described previously (Kopanitsa et al., 2006; Carlisle et al., 2008). Eight MEA1060-BC set-ups were run in parallel and monopolar stimulation of Schaffer collateral/commissural fibers through array electrodes was performed by STG2008 stimulus generator (Multi Channel Systems). For long-term potentiation (LTP) experiments, a single principal recording electrode was picked in the proximal part of CA1 stratum radiatum. To stimulate control and test pathways, two stimulation electrodes were assigned on the subicular side and on the CA3 side of stratum radiatum, respectively. The distance from the recording electrode to the test stimulation electrode was 400–510 μm and to the control stimulation electrode 280–510 μm. To evoke orthodromic field EPSPs (fEPSPs), test and control pathways were activated in succession at a frequency of 0.02 Hz. Baseline stimulation strength was adjusted to evoke a response that corresponded to 40% of the maximal attainable fEPSP at the recording electrode located in proximal stratum radiatum. Amplitude of the negative part of fEPSPs was used as a measure of the synaptic strength. To induce LTP, 10 bursts of baseline strength stimuli were administered at 5 Hz to test pathway with four pulses given at 100 Hz per burst (total 40 stimuli). LTP plots were scaled to the average of the first five baseline points. Normalization of LTP values was performed by dividing the fEPSP amplitude in the tetanized pathway by the amplitude of the control fEPSP at corresponding time points. Normalized LTP values averaged across the period of 65–65 min after theta-burst stimulation was used for statistical comparison.

Paired stimulation with an interpulse interval of 50 ms was used to observe paired-pulse facilitation (PPF) in baseline conditions in the test pathway before LTP induction. PPF was calculated by dividing the negative amplitude of fEPSP obtained in response to the second pulse by the amplitude of fEPSP amplitude evoked by the preceding pulse. To assess changes in basalt synaptic transmission input–output relationships were initially compared by mixed model repeated-measures ANOVA and Bonferroni post hoc test implemented in Prism 5 (GraphPad Software) using individual slice data as independent observations. Since several slices were routinely recorded from every mouse, fEPSPmax, PPF, and LTP values between wt and mutant mice were then compared using two-way nested ANOVA design with genotype (group) and mice (subgroup) as fixed effects. Fisher’s F statistic was calculated as mean of squaresGenotype/mean of squaresResidual and genotype effect was considered significant if corresponding probability for the group F statistic was <0.05.

Conventional extracellular recording techniques were used to examine the induction of long-term depression (LTD) by 900 pulse trains of 1 Hz presynaptic fiber stimulation (Carlisle et al., 2008). For whole-cell voltage-clamp techniques, slices were bathed in an oxygenated (95% O2:5% CO2) artificial CSF containing the following (in mS): 124 NaCl, 2.4 KCl, 25 NaHCO3, 1 NaH2PO4, 4.0 CaCl2, 4 MgSO4, 0.1 picromonox, and 10 glucose. EPSCs were recorded using electrodes and were filled with a solution containing 102 mM cesium gluconate, 17.5 mM CsCl, 10 mM TEA, 50 mM HEPES, pH 7.2. MTTX in the extracellular solution was added to the recording chamber. MTTX was added to the extracellular solution in experiments recording mIPSCs. Unpaired Student’s t tests were used to determine statistical significance of differences between whole-cell voltage-clamp measurements in mutant and wt neurons.

Microarray analysis. Hippocampi (TNIK+/− n = 11; wt n = 33) from male and female mice aged 98–174 d were dissected and snap frozen in liquid nitrogen. RNA was extracted using a Qiagen miRNAeasy kit. RNA integrity was verified using the Agilent 2100 Bioanalyzer and all samples had RIN values >7. Unpoled samples were prepared for array analysis using the Illumina TotalPrep-96 RNA Amplification Kit and hybridized to Illumina Mouse WG-6 v2.0 Beadchips. The data were read into the Bioconductor using Lumi (Du et al., 2008) and re-annotated using a previously described dataset (Barbosa-Morais et al., 2010). A Variance Stabilizing Transformation was applied followed by Quantile Normalization. Differential (t) expression of wt and TNIK+/− mice was determined using the empirical Bayes linear model approach implemented in the Limma package (Smyth et al., 2005). Multiple hypothesis testing
Histology. For 5-bromo-4-chloro-3-indolyl-β-galactoside (X-gal) staining, free-floating sections were stained overnight in X-gal (Malford), mounted on pretreated slides, dehydrated, and mounted with DPX mountant. For immunohistochemistry, free-floating sections were incubated in either anti-doublecortin (1:250; Cell Signaling Technology 4604), anti-NeuroD1 (1:500; Abcam ab60704) or anti-Ki67 (1:300; Abcam ab15580). Sections were incubated with either biotinylated anti-mouse or anti-rabbit secondary antibody (1:200; DAKO) for peroxidase staining or with anti-mouse Alexa Fluor 633 or anti-rabbit Alexa Fluor 568 (1:1000) for immunofluorescence. For peroxidase staining, signal was amplified using an avidin–biotin system (Vector Laboratories) before staining in 3,3-diaminobenzidine (Sigma). For quantitation of cell numbers, a minimum of two sagittal brain sections per animal was analyzed from four pairs of TNik⁻/⁻ mice and wt littermates. Total numbers of immunopositive cells in the hippocampus were counted by an experimenter blinded to genotype and statistical significance determined using unpaired Student’s t tests.

Cognitive testing in the rodent touchscreen operant system. Testing was conducted in a touchscreen-based automated operant system that consisted of an operant chamber (21.6 × 17.8 × 12.7 cm) made of clear Perspex walls and a stainless steel grid floor, housed within a sound- and light-attenuating box (40 × 34 × 42 cm) (Med Associates). A pellet dispenser delivering 14 mg dustless pellets (BioServ) into a magazine, a house light, and a tone generator was located at one end of the chamber. At the opposite end of the chamber was a flat-screen monitor equipped with an infrared touchscreen (16 × 21.2 cm) (Craft Data Limited). The touchscreen was covered by a black Perspex “mask” that either had six windows (for two-choice spatial discrimination) or three windows (for object-location paired associates learning) positioned in front of the touchscreen allowing the presentation of stimuli to be spatially localized and prevented the mouse from accidentally triggering the touchscreen. Stimuli presented on the screen were controlled by custom software (“MouseCat,” L.M. Saksida), and responses made via nose pokes at the stimuli were detected by the touchscreen and recorded by the software. Before testing, the body weight of mice was slowly reduced and then maintained at or >85% of free-feeding body weight. Animals were pretrained through several stages to learn to touch stimuli on the screen to obtain a reward as described previously (Brigman et al., 2008).

Two-choice spatial discrimination pattern separation. Adult mice (wt n = 8, TNik⁻/⁻ n = 12) were initially trained to touch one of two illuminated stimuli (squares). Responses at one location (correct) resulted in a reward; responses at the other location (incorrect) resulted in a 5 s timeout (signaled by extinction of the house light). During training, the location of the two illuminated stimuli was separated by an intermediate degree of separation (two empty/dark locations; Creer et al., 2010). Animals were required to complete seven of eight consecutive correct touches (acquisition) following which the reward contingences were reversed so that the previous incorrect location now became correct, and animals had to complete seven of eight consecutive correct touches in the other location (rever-
The correct starting location was counterbalanced between animals in each genotype and in addition, the reversal component of the task decreased the likelihood of animals developing a side bias or the use of nonspatial strategies. Mice were given a maximum of 60 trials/session/d until they reached a training criterion of completing a minimum of one acquisition and one reversal on 3 of 4 consecutive days.

Following training, pattern separation was assessed by varying the distance between the choice locations, either situated by a high degree of separation (i.e., four empty/dark locations between the two illuminated locations) or a low degree of separation (i.e., no empty/dark location between the two illuminated locations). Mice were given unlimited trials/session/d in a 60 min period and were allowed a maximum of one reversal. Spatial separations were held constant during each testing session per day but were varied across testing days. Mice were tested for four sessions on each separation, and the order of separation tested was counterbalanced between animals in each genotype across days.

Average number of trials taken to reach the criterion of seven of eight consecutive correct touches for acquisition and reversal for each separation was calculated. Group differences were analyzed using either an independent samples t test or a mixed between- and within-subjects ANOVA with genotype as the between-subjects factor and separation as the within-subjects factor. A paired samples t test was used for post hoc analysis to assess significant between × within-subjects interaction effect. All values reported represent mean ± SEM.

Object-location paired associates learning. Mice (wt n = 8, TNiK−/− n = 12) were tested for the ability to associate between objects (shapes) and locations on the touchscreen (Talpos et al., 2009). There were three objects (flower, plane, and spider) and three correct spatial locations (left, middle, and right, respectively). For each trial, only two objects were presented; one object in its correct location (S+) and the other object in one of two incorrect locations (S−). There were six possible trial types (Talpos et al., 2009), so that the flower was rewarded only when presented in the left location, the plane was rewarded only when presented in the middle location, and the spider was rewarded only when presented in the right location. A nose poke to the correct S+ resulted in delivery of a reward and incorrect responses resulted in a 5 s timeout period, followed by a correction procedure whereby the trial was repeated until the mouse made a correct choice. Nose pokes to response windows in which no stimulus was presented were ignored. Mice were given 36 trials/session/d for 30 sessions. Group differences were analyzed using a mixed between-within subjects ANOVA with genotype as the between-subjects factor and session block as the within-subjects factor. All values reported represent mean ± SEM.

Results

TNiK in glutamate receptor function in vivo

The TNiK gene was replaced with a LacZ reporter using homologous recombination in mouse ES cells (Fig. 1A,B), producing viable homozygous (TNiK−/−) mice that showed no detectable TNiK protein on immunoblots of brain extracts (Fig. 1C). We observed normal Mendelian transmission in TNiK−/− intercrosses (χ2 test, p = 0.23) and no detectable abnormalities in body size or gross brain morphology (Fig. 1E). In wt mice, TNiK is widely expressed in brain, including the cortex and hippocampus, where strongest staining was observed in dentate gyrus granule cells (Fig. 1D). As previously reported, TNiK protein was
Figure 3. Synaptic physiology in TNiK−/− mice. A, NMDAR-mediated EPSCs are normal in TNiK−/− mice. Traces (top) show EPSCs recorded at −80 and +40 mV in pyramidal cells from a wt (left) and TNiK−/− mouse (right). Calibration bars are 100 pA and 25 ms. The ratios of NMDAR-mediated EPSCs to AMPAR-mediated EPSCs measured at −80 and +40 mV (bottom) are not significantly different in TNiK−/− mice (n = 17; N = 3) compared with wt littermates (n = 17; N = 3). B, Baseline synaptic transmission was slightly but significantly enhanced in TNiK−/− mice. Input–output relationships illustrate averaged fEPSP amplitudes in slices from TNiK−/− (n = 28; N = 10) and wt mice (n = 32; N = 10) in response to stimulation of Schäffer collaterals by biphasic voltage pulses of 0.1–4.2 V (*p < 0.05, #p < 0.01). Amplitudes of fEPSPs evoked by maximum 4.2 V stimulation were significantly increased in TNiK mice (F(1,40) = 9.18; p = 0.004, two-way nested ANOVA, main genotype effect). Representative families of fEPSP traces are shown at right. Calibration bars are 1 mV and 2 ms.

C, Spontaneously occurring IPSCs were recorded in cells voltage-clamped at −70 mV and all-points histograms were generated from 10 s long segments of data. Points corresponding to baseline noise were estimated from the Gaussian curve centered on 0 pA. The difference between this area and the remaining area under the histogram corresponds to the currents generated by the IPSCs (shaded area) and was used to calculate total charge transfer. Traces show mIPSCs recorded before and after bath application of picrotoxin (100 μM). D, Comparison of sIPSC and mIPSC charge transfer in wt (open bars) and TNiK−/− cells (filled bars). Spontaneous IPSCs (recorded in the absence of TTX) in cells from TNiK−/− mice (n = 10 cells from 4 mice) are not significantly different from that seen in wt (Figure legend continues.).
expressed in the PSD (Kawabe et al., 2010; Wang et al., 2010) and the nucleus (Fig. 2A) (Mahmoudi et al., 2009).

As a starting point for understanding the biochemical role of TNiK and the impact of a knock-out mutation, we characterized the protein interactions and complexes formed by TNiK. Since TNiK binds DISC1 (Camargo et al., 2007; Wang et al., 2010), which binds GSK3β (Mao et al., 2009), expression of all three proteins was identified in the nucleus and in the PSD (Fig. 2A).

Using immunoprecipitations (IP) from PSD extracts, these three proteins were co-isolated in NMDAR complexes (Fig. 2B) together with CRMP2 (Fig. 2C), a GSK3β substrate (Cole et al., 2006). These findings are consistent with previous reports showing that CRMP2 binds DISC1 (Camargo et al., 2007) and NMDAR GluN2B subunits (Al-Hallaq et al., 2007). How TNiK is recruited into complexes with the NMDAR is unknown: a direct interaction between TNiK and NMDAR subunits has not been reported in numerous yeast two-hybrid screens using NMDAR GluN1 and GluN2 subunits as bait, thus it seems likely that TNiK interacts with NMDARs via an intermediate protein. A-kinase anchoring protein 9 (AKAP9) (Lin et al., 1998) is a candidate since it binds TNiK (Camargo et al., 2007) and the cytoplasmic domain of GluN1. To test this, AKAP9 mutant mice were generated and characterized, and as shown in Figure 2D, TNiK’s association with GluN1 was reduced to 52% (p < 0.05) and the reverse IP confirmed this result with a 63% decrease (p < 0.05; Fig. 2D).

Since the total levels of GluN1 and TNiK remained unchanged in extracts from AKAP9 mutants (Fig. 2D) and the interaction between TNiK and PSD-95 or GluN2B was not altered in AKAP9 mutants (Fig. 2E), these results support the view that AKAP9 links TNiK to GluN1 in vivo.

To understand the functional coupling of NMDAR and other glutamate receptors to TNiK, we examined the phosphorylation of S735, a proposed autophosphorylation site indicative of TNiK activity (Shitashige et al., 2010; Wang et al., 2010). In PSD extracts from hippocampal slices treated with NMDA, S735 was robustly dephosphorylated (42%, p < 0.01). In contrast, mGluR1 agonist increased S735 phosphorylation (241%, p < 0.01) and AMPAR activation had no effect (p > 0.05) (Fig. 2F–I). In PSD extracts of TNiK<sup>−/−</sup> mice, a decrease in AMPAR GluA1 levels (35%, p < 0.05; Fig. 2J) was observed, consistent with a recent report in primary cultures with TNiK knockdown (Wang et al., 2010). These data show biochemical interactions between TNiK and glutamate receptors, we examined excitatory and inhibitory synaptic transmission and synaptic plasticity in the hippocampus of TNiK<sup>−/−</sup> mice. NMDAR- and AMPAR-mediated synaptic currents, long-term plasticity (LTP and LTD), and sIPSCs and mIPSCs were normal in the hippocampal CA1 region of TNiK<sup>−/−</sup> mice (Fig. 3A–F). We noted, however, that evoked mEPSCs were slightly but significantly enhanced at stronger stimulation intensities compared with wt littermates (Fig. 3B; F<sub>1(9,522)</sub> = 6.48, p < 0.0001). We also observed a significant increase in PPF (188 ± 3% in wt compared with 196 ± 3% in TNiK<sup>−/−</sup> mice, F<sub>1(4,140)</sub> = 5.36, p = 0.026) and a significant decrease in the frequency of AMPAR-mediated mEPSCs in TNiK<sup>−/−</sup> cells (Fig. 3G).

Postsynaptic signaling complexes not only regulate glutamate receptor function and synaptic plasticity, but also transcriptional and other signaling pathways. Thus in the absence of a major
changes were observed in the levels of GluN1, PSD-95, CRMP2, Tau, or NeuroD1 in TNiK⁻/⁻ mice (p > 0.05; Fig. 4A) and the hyperphosphorylation of the GSK3β sites on these proteins was reversed within minutes of applying a pharmacological inhibitor of GSK3β to hippocampal slices (data not shown).

**GSK3β antagonist reversal of locomotor hyperactivity**

Spontaneous locomotor hyperactivity has been reported in mice overexpressing GSK3β (Mohn et al., 1999; Prickaerts et al., 2006; Shen et al., 2008); therefore we tested locomotor activity in four test environments (open field, novel object exploration, elevated plus maze, and locomotor activity chamber). A robust increase in locomotor activity was observed in TNiK⁻/⁻ mice in all tests (Fig. 5A). Next, the ability of this effect to be rapidly reversed with a GSK3β inhibitor was tested. TNiK⁻/⁻ and wt mice were placed in an activity chamber for an initial period of 30 min to assess baseline spontaneous locomotor activity where as expected, TNiK⁻/⁻ mice showed significantly elevated locomotor behavior compared with wt (Fig. 5B; p < 0.001). Within 30 min following administration of the GSK3β inhibitor SB216763, the locomotor activity of TNiK⁻/⁻ mice was significantly reduced (p < 0.05) to the same levels as wt mice. Furthermore, there was no significant difference between wt mice treated with vehicle or SB216763, indicating a unique sensitivity to GSK3β inhibition (p < 0.05; Fig. 5C) was conferred by the TNiK mutation. These results suggest the mechanism underlying the locomotor phenotype in TNiK⁻/⁻ mice is the result of aberrant GSK3β signaling.

**TNiK is required for dentate gyrus neurogenesis**

Because the S274 phosphorylation on NeuroD1 is known to inhibit its transcriptional activity (Chae et al., 2004) and NeuroD1 is required for dentate gyrus neurogenesis through Wnt signaling (Miyata et al., 1999; Kuwabara et al., 2009), and both Wnt/β-catenin defects result in loss of the dentate gyrus (Miyata et al., 1999; Liu et al., 2000), this raises the possibility that TNiK⁻/⁻ mice could have dentate gyrus abnormalities. Moreover, inhibition of Wnt (Wexler et al., 2006; Shen et al., 2008); therefore we tested locomotor activity in four test environments (open field, novel object exploration, elevated plus maze, and locomotor activity chamber). A robust increase in locomotor activity was observed in TNiK⁻/⁻ mice in all tests (Fig. 5A). Next, the ability of this effect to be rapidly reversed with a GSK3β inhibitor was tested. TNiK⁻/⁻ and wt mice were placed in an activity chamber for an initial period of 30 min to assess baseline spontaneous locomotor activity where as expected, TNiK⁻/⁻ mice showed significantly elevated locomotor behavior compared with wt (Fig. 5B; p < 0.001). Within 30 min following administration of the GSK3β inhibitor SB216763, the locomotor activity of TNiK⁻/⁻ mice was significantly reduced (p < 0.05) to the same levels as wt mice. Furthermore, there was no significant difference between wt mice treated with vehicle or SB216763, indicating a unique sensitivity to GSK3β inhibition (p < 0.05; Fig. 5C) was conferred by the TNiK mutation. These results suggest the mechanism underlying the locomotor phenotype in TNiK⁻/⁻ mice is the result of aberrant GSK3β signaling.

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Figure 6. Impaired dentate gyrus neurogenesis and cognition in TNiK−/− mice. A, B, Histograms comparing dentate gyrus granule cell number and cells stained with markers relevant to neurogenesis (see text) in wt and TNiK−/− mice brain sections. Representative images of dentate gyrus cells in wt and TNiK−/− brain sections stained with markers for DCX (A), NeuroD1, and Ki67 (B). Scale bars: (in A) 200 and (in B) 60 μm. C–E, Hippocampus development in TNiK−/− mice. Number of NeuN-positive neurons at P7 (upper, left (Figure legend continues.)
precursor marker expressed in differentiating neurons) (von Bohlen Und Halbach, 2007), we observed significant reductions of both (DCX, 55%, p < 0.01; NeuroD1, 41%, p < 0.001; Fig. 6A, B). The cell proliferation marker Ki67 (Kee et al., 2002) was also reduced (70%, p < 0.01; Fig. 6B). The decrease in dentate neurons in TNiK−/− mice was also observed by postnatal day 7 (43%, p < 0.001) (Neunstaining; Fig. 6C) as was the reduction in cells expressing NeuroD1 (37%, p < 0.01; Fig. 6D) and DCX (77%, p < 0.001; Fig. 6E). Commensurate with a requirement for TNiK during these developmental stages, we examined the postnatal expression profile of TNiK and found strong expression in principal excitatory neurons in all hippocampal subfields from P7 (Fig. 7).

TNiK mutants show cognitive deficits in touchscreen tests

The role of TNiK in behavior is unknown and the abnormalities in the dentate gyrus pointed to the possibility that there could be a role for TNiK in pattern separation. Pattern separation is a specific cognitive function where the ability to form distinct representations from similar inputs closely presented in space and/or time is necessary for decreasing the interference between memories. Impairments in dentate gyrus neurogenesis induced either by X-irradiation or Wnt inhibition lead to impairments in pattern separation (Clelland et al., 2009) and conversely, enhanced pattern separation performance correlates with increased neurogenesis (Creer et al., 2010; Sahay et al., 2011). We therefore asked whether TNiK was required for pattern separation using the same two-choice spatial discrimination task in the touchscreen apparatus that was used to show the role of Wnt (Clelland et al., 2009). Mice were initially trained to discriminate the correct spatial location between two illuminated squares (located in two of six possible locations) and then pattern separation was tested by varying the distance between the choice locations, either situated far apart (high degree of separation) or close together (low degree of separation; Fig. 6F). TNiK−/− mice showed no significant difference to wt mice during initial training (trials to training criterion: wt = 458.1 ± 45.8, TNiK−/− = 576.3 ± 74.2, p = 0.248) and when tested with stimuli separated by a high degree of separation (p > 0.05; Fig. 6F). However, TNiK−/− mice showed a significant impairment in discrimination when the locations were in close proximity (genotype × separation interaction, p < 0.01; low separation, p < 0.05). Previous studies have shown a reduction in dentate gyrus neurogenesis impairs pattern separation without impacting object-location paired associates learning, which can also be measured in the touchscreen apparatus (Clelland et al., 2009). Moreover, intrahippocampal infusions with NMDA or AMPA antagonists impair performance in this task (Talpos et al., 2009). Since TNiK is involved in NMDAR and AMPA function, it was of interest to assess object-location paired associates learning in TNiK mutants. The task measures the ability to associate three objects (flower, plane, and spider) with their correct spatial locations on the screen [left (L1), middle (L2), and right (L3)], respectively (Fig. 6G). We found that TNiK−/− mice were significantly impaired in learning the association of an object and its location (effect of genotype, p = 0.016; Fig. 6G).

Discussion

Our data provide the first direct evidence showing TNiK in the nervous system plays key roles at the synapse and nucleus, and is essential for cognition. We found that TNiK was required for two different forms of learning—pattern separation and object-location paired associates learning—that both rely on processing visual and spatial information. Pattern separation is important for keeping similar memories separate and is known to require the function of the dentate gyrus (Gilbert et al., 2001; Leutgeb et al., 2007; Bakker et al., 2008; Hunsaker and Kesner, 2008). Deletion of the essential NR1 subunit of the NMDAR specifically in dentate granule cells (McHugh et al., 2007) or disruption of neurogenesis through interference with Wnt signaling or irradiation (Clelland et al., 2009) all result in impaired pattern separation. Conversely, increased neurogenesis has been reported to correlate with enhanced pattern separation ability (Creer et al., 2010; Sahay et al., 2011). Although we observe that TNiK−/− mice show a marked impairment in neurogenesis of dentate gyrus granule cells and exhibit deficits in pattern separation, we cannot exclude the possibility that some other signaling deficit or neuroanatomical change within the granule cells might also contribute to this specific cognitive impairment. In addition to pattern separation, TNiK was required for learning object-location paired associates, which is not dependent on dentate gyrus neurogenesis (Clelland et al., 2009), but performance of this task is dependent on NMDAR and AMPAR function (Talpos et al., 2009). These findings suggest that loss of TNiK leads to additional dysfunction outside of the dentate gyrus and thus has wider cognitive impacts.

The molecular functions of TNiK are summarized in Figure 8, highlighting a role for TNiK in both nuclear and postsynaptic complexes. Several lines of evidence indicate that TNiK mediates its role in neurogenesis through nuclear signaling complexes involving Wnt and GSK3β substrates. In the dentate gyrus subgranular zone, Wnt proteins (Grove et al., 1998; Lee et al., 2000) and β-catenin (Gao et al., 2007) are essential for neurogenesis. One of the major effectors of Wnt/β-catenin pathways is the activation of the basic helix-loop helix transcription factor NeuroD1 (Kuwabara et al., 2009), and similar to Wnt/β-catenin defects, the absence of NeuroD1 during hippocampus development results in loss of cells in the dentate gyrus (Miyata et al., 1999; Liu et al., 2000). Furthermore, in adult stages, the conditional deletion of NeuroD1 shows it is indispensable for survival and maturation of adult-born neurons (Gao et al., 2009) and defines NeuroD1 as a critical regulator of adult neurogenesis. The transcription factor activity of NeuroD1 has also been shown to be modulated by heterodimerization with the schizophrenia susceptibility gene TCF4 (Brzózka et al., 2010) and by phosphorylation of S274 by GSK3β (Chae et al., 2004), which inhibits NeuroD1 transcription activity. We found that deletion of TNiK resulted in enhanced phosphorylation of NeuroD1 S274, which would be expected to contrib-
ute to the observed reduction in neurogenesis and number of dentate gyrus neurons. Moreover, we found that NeuroD1-positive cells were decreased in the dentate gyrus of \( TNiK^{-/-} \) mice and this decrease in precursor/progenitor cells was paralleled with a decrease in the number of newly born neurons and total number of dentate gyrus granule cells.

In non-neuronal cells, \( TNiK \) has been described as a critical component of transcriptional regulatory mechanisms, mediated by Wnt signaling, through binding and phosphorylation of the HMG box transcription factor Tcf7l2. \( TNiK \) forms a ternary complex with Tcf7l2 and \( \beta \)-catenin, modulating the transcriptional activation of Wnt target genes (Mahmoudi et al., 2009). In agreement with this, our results show that the deletion of \( TNiK \) results in a dysregulation of mRNAs involved in Wnt pathways. For example, \( CCND1 \) is a characteristic Wnt/\( \beta \)-catenin-modulated gene (Mahmoudi et al., 2009) and was downregulated in the \( TNiK^{-/-} \) mice. \( CCND1 \) was also reported to decrease as a result of \( DISC1 \) knock-down in vivo (Mao et al., 2009) and \( DISC1 \) and \( \beta \)-catenin, both \( TNiK \) interacting proteins, regulate Wnt signaling and neuronal progenitor proliferation (Kuwabara et al., 2009; Mao et al., 2009). Moreover, \( DISC1 \) induces a premature cell cycle exit and neuronal differentiation through GSK3\( \beta \) activity (Mao et al., 2009) and deletion of \( TNiK \) leads to an impairment in cell proliferation as demonstrated by the decrease in the total numbers of Ki67+, cells. These data indicate that \( TNiK \) is a key regulator of neurogenesis and cell proliferation through a combination of mechanisms, including modulation of transcription factor phosphorylation, activation of Wnt pathway and nuclear protein complexes.

The finding that \( TNiK \) was critical for constraining GSK3\( \beta \) was unexpected. In the absence of this constraint, there was increased phosphorylation of both nuclear substrates controlling cellular proliferation and synaptic substrates in the signal transduction complexes formed with scaffold proteins and glutamate receptors. GSK3\( \beta \) controls both neurogenesis and rapid synaptic signaling events (Jope and Roh, 2006; Beaulieu et al., 2009; Luo et al., 2010; Sirerol-Piquer et al., 2011). Aberrant GSK3\( \beta \) signaling has also been implicated in and been a prominent therapeutic target (Jope and Roh, 2006; Beaulieu et al., 2009) for a variety of cognitive disorders including neuropsychiatric, neurodegenerative, and more recently, neurodevelopmental disorders such as Fragile X. Interestingly, \( Fmr1 \) knock-out mice show elevated GSK3\( \beta \) levels, aberrant neurogenesis, and impairments in spatial discrimination similar to that observed in \( TNiK^{-/-} \) mice (Min et al., 2009; Luo et al., 2010; Guo et al., 2012). This is noteworthy as \( Fmr1 \) is expressed widely in hippocampal subfields and moreover, loss of \( Fmr1 \) results in a phenotype where the dentate gyrus is particularly impacted, similar to that of \( TNiK^{-/-} \) mice. Our findings support the view that \( TNiK \) acts in concert with GSK3\( \beta \) and \( DISC1 \) to regulate formation of neural circuits. \( DISC1 \) has been linked to Wnt signaling and neuronal progenitor proliferation (Kuwabara et al., 2009; Luo et al., 2010; Ishizuka et al., 2011) and in agreement with our biochemical data, \( TNiK \) directly binds \( DISC1 \) and \( \beta \)-catenin (Mahmoudi et al., 2009; Wang et al., 2010) forming complexes with GSK3\( \beta \). \( DISC1 \) knockdown induces a premature cell cycle exit and neuronal differentiation through GSK3\( \beta \) activity (Mao et al., 2009) and \( TNiK \) knock-out leads to an impairment in cell proliferation as demonstrated by the decrease in the total numbers of Ki67+ cells. It will be important in future work, using conditional and domain specific mutations in both \( TNiK \) and GSK3\( \beta \), to dissect the multifunctional roles of these interacting kinases. These genetic approaches combined with pharmacological tools will allow the developmental and cell type-specific functions to be elucidated.

We observed several relevant electrophysiological deficits in \( CA3-CA1 \) synapses of \( TNiK^{-/-} \) mice showing a requirement for \( TNiK \) in normal excitatory synaptic function. The frequency of mEPSCs recorded in \( CA1 \) pyramidal cells of \( TNiK^{-/-} \) mice was significantly lower than that seen in wt cells. Decreases in mEPSC frequency can arise from a reduction in the number of excitatory synapses and/or a decrease in the probability of transmitter release from the presynaptic terminal. We found that PPF in the hippocampal \( CA1 \) region was significantly enhanced in slices from \( TNiK^{-/-} \) mice suggesting that \( TNiK \) may influence presynaptic processes involved in glutamatergic synaptic transmission. At the biochemical level, we found that \( TNiK \) was required for GluA1 expression, which is consistent with recent studies using siRNA-mediated \( TNiK \)
knockdown in primary neuronal cultures (Hussain et al., 2010; Wang et al., 2010). The change in GluA1 did not appear to have a major impact on synaptic transmission as we did not observe reductions in mEPSC amplitudes or evoked AMPAR-mediated synaptic responses in hippocampal slices from TNiK/H11002/H11002 mice. It is not surprising that the 35% decrease in GluA1 levels in TNiK/H11002/H11002 mice is not associated with changes in basal synaptic transmission in the hippocampal CA1 region. For example, previous studies have found that both fEPSPs and mEPSCs are normal in GluA1 knock-out mice (Zaminillo et al., 1999), although this was not seen in all studies (Andrásfalvy et al., 2003). This is most likely due to the fact that functional AMPARs can be made from hetero and homomeric combinations of other subunits (GluA2–4). In addition, 95% of the extrasynaptic pool of AMPAR subunits normally contain GluA1 subunits (Lu et al., 2009). Thus there is likely a considerable reserve pool of other AMPAR subunits that could compensate for the modest decrease in total GluA1 levels in TNiK mutants.

TNiK was physically linked to the NMDAR complex and its autophosphorylation site was regulated by both NMDAR and mGluR (in opposing directions). Moreover, in the absence of TNiK there was an elevation of GSK3β phosphorylation on postsynaptic substrates. Plasticities induced by transcription and translation-independent LTP and LTD protocols were normal in TNiK/H11002/H11002 mice. We cannot rule out a potentially important role for TNiK in other forms of LTP such as transcription- or translation-dependent forms of late-phase LTP, which may be relevant to the potential link with fragile X mental retardation protein noted above.

A number of circumstantial lines of evidence suggest TNiK is involved in neuropsychiatric diseases. In addition to its interaction with DISC1, there are links between TNiK and other genes implicated in schizophrenia. NeuroD1 heterodimerizes with the schizophrenia susceptibility protein TCF4 (Brzózka et al., 2010) and TNiK forms a ternary complex with Tcf7l2 (another schizophrenia susceptibility protein; Hansen et al., 2011) and β-catenin, modulating the transcriptional activation of Wnt target genes. We find that TNiK binds and regulates glutamate receptor complexes, which are involved in a range of different cognitive disorders including schizophrenia, autism, and forms of mental retardation (Fernández et al., 2009; Bayés et al., 2011). A recent large-scale study of de novo copy number variation in schizophrenia found a highly

Figure 8. Molecular functions of TNiK. The molecular functions of TNiK in the PSD and nucleus are shown. Proteins are shown as colored shapes and grouped into their synaptic and nuclear locations. Black lines connecting proteins show known protein–protein interactions. Arrows show regulatory interactions: red, phosphorylation (phosphorylated amino acid residue indicated with number and letter); blue, dephosphorylation; green, regulates protein level.
significant enrichment in mutations affecting the MAGUK-associated signaling complexes that include PSD-95, TNiK, and proteins dependent on TNiK function (Kirov et al., 2012). We observed that loss of TNiK impairs object-location paired associates learning, which has been evaluated in humans using computer-automated tests such as the object-place paired associates learning (PAL) test from the Cambridge Neuropsychological Test Automated Battery that is widely used in clinical settings (Barnett et al., 2010). Moreover, schizophrenic patients show robust impairments on PAL and performance on this test correlates with symptom severity (Barnett et al., 2005, 2010). Aberrant neurogenesis is also emerging as a neuropathological change in schizophrenia (Reif et al., 2006, 2007; Toro and Deakin, 2007; DeCarolis and Eisch, 2010) and while no study has yet explicitly examined pattern separation in schizophrenic patients, its importance as a cognitive function of potential relevance to schizophrenia has been argued (Tamminga et al., 2010). It is now emerging that human cognitive disorders arise from mutation or disruption in many postsynaptic signaling proteins, including those interacting with TNiK, and future studies may determine whether TNiK plays a role in modulating these disease phenotypes.

Notes
Supplemental Figure 1, Tables 1 and 2, and details of antibodies used in WB, IP, IHC, and IF assays are available at http://www.genes2cognition.org/publications/tnik. This material has not been peer reviewed.

References


