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JNK1 controls dendritic field size in L2/3 and L5 of the motor cortex, constrains soma size, and influences fine motor coordination

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INTRODUCTION

Dendrites are highly specialized, excitable compartments of neurons that are molecularly and functionally distinct from the axon. They receive and compute synaptic input and fine-tune the character of the action potential output (Williams and Stuart, 2003; London and Häusser, 2005). Notably, the computational properties of dendrites are directly influenced by the shape and extent of dendritic trees (Rall, 1977; Jaffe and Carnevale, 1999; London and Häusser, 2005; Spruston, 2008). Genetic disorders that involve well characterized anomalies in dendrite architecture include mental retardation syndromes Down’s, RETT, and fragile-X, where decreased arborization is common (Kaufmann and Moser, 2000; Zoghbi, 2003; Pardo and Eberhart, 2007; Jan and Jun, 2010). Conversely, autism and schizophrenia are associated with overgrowth of dendrites during development. Although these are highly heritable disorders, the genetic basis remains largely unknown, though large-scale genomic studies are starting to reveal candidates (Hyman, 2008; Fromer et al., 2014; Purcell et al., 2014).

c-Jun N-terminal kinase-1 (JNK1) has been implicated in the regulation of dendrite arborization in isolated neurons. For example, in cerebellar granule neurons JNK inhibition increases dendrite complexity (Björkblom et al., 2005), while in cortical and hippocampal neurons, JNK inhibitors reduce dendrite growth (Rosso et al., 2005; Podkowa et al., 2010). Yet, mechanistic details are lacking, and detailed quantitation of dendrite architecture using Sholl analysis in Jnk1 knockout mice has not been reported. Interestingly however, recent human genetics studies report deregulation of JNK pathway genes in several dendrite disorders. For example, JNK1 activity in the cortex is dependent...
on a kinase located on chromosome 16p11.2, a gene susceptibility
locus for autism and schizophrenia (Weiss et al., 2008; McCarthy
et al., 2009). Genetic risk for schizophrenia is associated with
the JNK pathway (Winchester et al., 2012), and the interleukin-
1 receptor accessory protein like-1 gene, implicated in monogenic
forms of mental retardation and autism, signals through JNK
(Pavlowsky et al., 2010). Furthermore, chromosomal transloca-
tions leading to loss of function truncations of JNK3 are associated
with intellectual disability (Shoichet et al., 2006; Baptista et al.,
2008; Kunde et al., 2013). These findings suggest that disruption
of normal JNK function may be central to neuropsychiatric dis-
orders that share irregularities in dendrite shape as a common
hallmark.

To gain molecular understanding of the structural changes
that occur in the brain upon disruption of JNK signaling, we set
out to precisely characterize the dendrite architecture in
Jnk1-/-
mice using three-dimensional Sholl analysis. We generated phos-
phorylation site mutants of HMW-MAP2, a major substrate
of JNK1 in dendrites (Kyriakis et al., 1995; Chang et al., 2003;
Björkblom et al., 2005), and examined whether site-specific phos-
phorylation of HMW-MAP2 by JNK1 altered dendrite shape and
microtubule integrity. Our findings show that JNK1 phos-
phorylates HMW-MAP2 on specific residues in the C-terminal
domain to create a microtubule binding motif, leading to increased
microtubule stabilization. Moreover, our investigation revealed
significant structural alterations in L2/3 and L5 dendrites in the
primary motor cortex of Jnk1-/-
mice. Consistent with these find-
ings, ectopic expression of GFP-HMW-MAP2 by JNK1 altered
dendrite length and complexity in the motor cortex suggesting
that phosphorylation of HMW-MAP2 on these residues has a
major impact on dendritic field. Finally we show that the behav-
ioral consequence of disrupted JNK1 signaling is impaired motor
function.

MATERIALS AND METHODS

ANTIBODIES

Anti-MAP2 (AP20) recognizing HMW-MAP2 was from Leinco
Technologies (St. Louis, MO, USA), Phospho-MAP2 (cat. no.
4544; RRID: AB_2144157) recognizing HMW-MAP2, and PJNK-
Thr183/Tyr185 (cat. no. 9255S; RRID: AB_2235013) were from
Cell Signaling Technology Inc. (Danvers, MA, USA) or from
Biosource (cat. no. 44-682G; RRID: AB_1502039). JNK1 (clone
gno. G151-333; RRID: AB_399158) was from PharMingen (San
Diego, CA, USA). Mouse anti-GFP (cat. no. IL-8; RRID:
AB_10013427) was from Clontech (Mountain View, CA, USA).
Anti-mouse b-tubulin (cat. no. KMX-1; RRID: AB_94650) was from
Chemicon (Temecula, CA, USA) and anti-ankyrin-G was from
NeuroMab (clone N106/36; cat. no. 75-146; RRID:
AB_10673030).

PLASMIDS

pEGFP-HMW-MAP2 and pEGFP-NES-Jun were previously
described (Björkblom et al., 2005; Tararuk et al., 2006). Phospho-
phorylation site mutants of HMW-MAP2, EGFP-HMW-
MAP2T1619A/T1622A/T1625A (abbreviated to GFP-MAP2-AAA)
and EGFP-MAP2T1619D/T1622D/T1625D (abbreviated to GFP-MAP2-
DDD), were prepared by insertional overlapping PCR using
mutagenic primers as previously described (Hongisto et al., 2008).
The phosphorylation site numbering is based on the rat HMW-
MAP2 Uniprot entry P15146. For in uto electroporation, the
CMV promoter in EGF-HMW-MAP2WT and EGF-PHMW-
MAP2T1619D/T1622D/T1625D was changed to a CAG promoter for
optimal expression in brain. MAP2C was isolated by PCR from rat
brain cDNA. It was inserted downstream of the gGEX
6P3 vector (GE Healthcare) using the pGEMTe cloning vector
(Promega). pCDNA3-MKK7-JNK1 was a gift from Roger J. Davis
(HHMI, Worcester, MA, USA).

PHOSPHORYLATION

GST-MAP2 0.1–0.4 μM was phosphorylated with active GST-
JNK1α1 or GST-JNK3α1, of comparable specific activities, in 30 μl
kinase buffer (10 mM PBS pH 7.4, 2 mM EGTA, 1 mM DTT,
10 mM MgCl2, 0.1% v/v Triton X100) supplemented with 5 μCi
of [γ32P]-ATP and 25 μM non-isotopically labeled ATP. The
reaction was carried out for 1 h at 30°C and stopped by the addi-
tion of 4× Laemmli sample buffer. GST-cJun(5–89) at 2.22 μM
concentration was used as a positive control to monitor the cat-
alytic activities of JNK1α1 and JNK3α1. Samples were resolved
by SDS–PAGE, stained with Coomassie brilliant blue, destained
and analyzed by autoradiography and phosphorimaging. Velocity
was calculated using Michaelis–Menten kinetics and was plotted
against substrate concentration. For phosphosite analysis, the
phosphorylated bands corresponding to full length GST-MAP2
were excised from the gel and subjected to in gel digestion and
mass spectrometry analysis as described below. Metabolic labeling
of COS-7 cells using [γ32P]-ATP was carried out as previously
(Björkblom et al., 2012).

TISSUE EXTRACT PREPARATION

Forebrains from mice at postnatal day 0 or 2 were rapidly extracted
after decapitation and post-translational modifications were pre-
served using a heat stabilizer (Denator, Sweden). Frozen tissues
were homogenized using an Ultra Turrax homogenizer in ice-cold
lysis buffer (20 mm HEPES, pH 7.4, 2 mm EGTA, 50 mm β-
glycerophosphate, 1 mm dithiothreitol, 1 mm Na3VO4, 1% Triton
X-100, 10% glycerol, 1 mm Na3VO4, 1% Triton
X-100, 10% glycerol, 1 mm benzamidine, 50 mm NaF, 1 μg/ml
leupeptin, 1 μg/ml pepstatin, 1 μg/ml aprotinin, and 100 μg/ml
PMSF). Lysates were normalized for protein using the Bradford
method before SDS-PAGE.

IN-VEL DIGESTION AND PHOSPHOPEPTIDE ENRICHMENT

In vitro phosphorylated proteins were separated on 12% Crite-
riorn gels (Bio-Rad Laboratories, Hercules, CA, USA), gels were
washed in Milli-Q water, stained 1 h with GelCode (Thermo
Scientific, Rockford, IL, USA), destained overnight in Milli-Q
water. Each lane was manually sliced into five fractions and
slices were destained then reduced and alkylated before diges-
tion with 12.5 μg/ml sequencing grade modified porcine trypsin
(Promega, Madison, WI, USA) overnight at 37°C as previ-
ously described (Björkblom et al., 2012). Peptides were eluted
in 75% ACN, 1% FA. 60 μl of peptides were dried and imme-
diately subjected to phospho-peptide enrichment. The peptides
(ca. 50 μg/sample) were re-suspended in 150 μl binding buffer.
(1 M glycolic acid, 80% ACN, 5% TFA) and mixed with 50 μl homogenous suspension of TiO2 magnetic sepharose beads (GE Healthcare Bio-Science AB, Uppsala, Sweden) that had previously been washed five times in the same buffer. Peptides were equilibrated with the beads, binding for 60 min at RT with gentle rocking. The beads were washed three times with 200 μl washing Buffer (80% ACN, 1% TFA) and peptides were eluted twice adding in total 100 μl 5% NH3 pH 12. The pH of the solution was lowered to <3 adding 5 μl 88% FA prior to sample clean up using C18 UltraMicroSpin columns (The Nest Group Inc., Southboro, MA, USA). Eluted peptides were then dried in a Speedvac, resuspended in 0.1% FA and then immediately analyzed by LC–MS.

IMMUNOBLOT ANALYSIS AND QUANTIFICATION

Cells were stimulated as indicated, washed in PBS, and lysed with Laemmli sample buffer. Samples were resolved on 5% (MAP2) or 10% SDS-PAGE and transferred by semi-dry transfer to nitrocellulose. Blots were developed using the enhanced chemiluminescence detection method. Films were pre-flashed, and non-saturated exposures were digitized by flatbed scanning and quantified by densitometry.

CELL CULTURE AND TRANSFECTION

Mouse embryonic fibroblasts (MEFs) were cultured in minimal essential medium (MEM) supplemented with 10% (v/v) bovine calf serum, 2 mM glutamine, 50 μg/ml penicillin, 50 μg/ml streptomycin, and non-essential amino acids (Sigma) were added. COS-7 cells were cultured in MEM supplemented with 10% (v/v) bovine calf serum, 2 mM glutamine, 50 μg/ml penicillin, and 50 μg/ml streptomycin. All cells were grown with 5% CO2 at 37°C.

IMUNOSTAINING

Hippocampal neurons at 12 days in vitro were fixed with 4% paraformaldehyde for 20 min and permeabilized for 3 min with 1% TX100 in phosphate-buffered saline (PBS). Following block in 10% fetal calf serum, cells were stained using anti-JNK1 (Pharmingen) at 1:100, ankyrin-G (NeuroMab) at 1:800, or P-JNK (Biosource) at 1:200 and detected with Alexa-488/568 conjugated secondary antibodies at 1:500.

MEASUREMENT OF TUBULIN AND HMW-MAP2 POLYMERIZATION

MEF2 cells were transfected with 50% Venus-tubulin and 50% EGFP-HMW-MAP2 variants. After 24 h the cells were washed with pre-warmed PEM buffer (80 mM pipes, 1 mM MgCl2, 2 mM EGTA, pH 6.8, PMSE, 1 μg/ml aprotonin, 1 μg/ml pepstatin, 1 μg/ml leupeptin). Cells were then lysed in +37°C PEM buffer containing 0.15% TX-100. After 1 min lysis, the liquid phase was pipetted from the dish and collected as the soluble fraction. The TX-100 insoluble fraction was collected in laemmli buffer. Fractions were analyzed by immunoblotting and quantified using densitometry.

LUCIFER YELLOW LOADING

Mice at 6–8 months were anesthetized with 0.3 mg/g Pentobarbital (Mebunat 60 mg/ml) mixed 50:50 with 0.9% NaCl. Mice were perfused transcardially on ice bedding using 10–20 ml 0.9% NaCl followed by 25–50 ml of buffer comprising 4% paraformaldehyde and 0.125% glutaraldehyde in 0.1 M Sorensen's phosphate buffer (NaH2PO4–Na2HPO4, pH 7.2). The brain was post-fixed in 50 ml 4.0% PFA in phosphate buffer for 4–12 h at +4°C. Coronal sections of 180–200 μm were sectioned using a vibratome. The nuclei were visualized using DAPI (4',6-diamino-2-phenylindole, dihydrochloride (Invitrogen). Pyramidal neurons in the motor cortex and the hippocampus were located according to the Atlas of C57BL/6 mouse brains (Hof et al., 2000). Selected neurons were injected by iontophoresis with lucifer yellow dye (Invitrogen) using pulled borosilicate glass tubes (World Precision Instruments). The DC current source was 2–6 nA from a dual micro-iontophoresis current generator, model 260 (World Precision Instruments). After dye loading, brain slices were transferred to a slide and mounted using Shandon PermaFluor mounting medium (ThermoFisher).

IN UTERO ELECTROPORATION

For dendrite analysis timed-pregnant mice C57/B6 (WT and Jnk1–/–) carrying E15 embryos were anesthetized with isoflurane (induction, 4%; surgery, 2.0%). Following anaesthetization pre-emptive analgesia was administered subcutaneously with a dose of buprenorphine 0.05–1 mg/kg (Temgesic® from Schering-Plow), and the uterine horns were exposed by laparotomy. GFP-CAG-HMW-MAP2WT and GFP-CAG-HMW-MAP2T1619D,T1622D,T1623D (2 μl) containing 0.8% Fast-Green (wt/vol) was injected into the lateral ventricles of embryos. After soaking the uterine horn with a PBS solution, the embryo’s head was carefully held between tweezer electrodes and DNA was electroporated using a CUY21E square wave electroporator (NEPA Gen). Electrical pulses (45 V, 50 ms) were passed five times at 1 s intervals. Uteri were returned to the abdominal cavity and embryos allowed to develop normally until delivery. Animals were sacrificed at P21. The procedures were approved by the National Animal Experimental Board. Collected tissues were fixed in 4% paraformaldehyde for 24 h, immersed in 30% sucrose (wt/vol) and frozen using isopentane. Cryosections (50 μm) were cut coronally and z-stack images were collected using a Zeiss LSM 780 confocal microscope with 20× air objective. LSM images for dendrite analysis collected from M1 were acquired following “The Mouse Brain in Stereotactic Coordinates, Third edition” by Franklin K.B.J and Paxinos G. Coronal sections for analysis were from the region between 5.78 and 4.42 mm (interaural) with respect to bregma 1.98–0.62 mm. The anterior forceps of the corpus callosum, anterior commissure, and the lateral ventricle was used as a landmark to recognize region of interest. Further dendrite analysis was done with Neurolucida software.

THREE-DIMENSIONAL IMAGING AND SHOLL ANALYSIS

For lucifer yellow loaded cells, 1 μm z sections spanning 40–80 μm were acquired using a Leica TCS SP2 microscope with 20× objective. Spiny neurons with large Hoescht-33342-stained nuclei and
pyramidal morphology were selected for analysis. Basal dendrites were traced manually and subjected to Sholl analysis using Neuroulucida software (MBF, Bioscience). Fine apical dendrites were excluded from the analysis as complete loading including tufts was achieved in too few cells. Scoring was performed blind. One-way ANOVA was used to determine significant differences among data sets. For analysis of in utero electroporated mice, confocal sections 0.7-μm thick were acquired using the Zeiss LSM-780 microscope and a 20× objective. Sections were reconstructed manually using Neuroulucida software (MBF, Bioscience). Neurons with sufficient GFP signal to detect higher order branching were selected for analysis. Branched structure analysis was performed for each tree. Sholl analysis was performed for intersections using 10 μm concentric circles surrounding the cell soma. For soma size analysis, cells with a clear beginning and end point in the z-plane were traced for each stack and cross sectional area and perimeter were evaluated using Neuroulucida. One-way ANOVA was used to determine significant differences for the Sholl intersections while t-test was used for branch order and soma size measures. Statistical analysis was performed in Graphpad Prism (version 5). For all the dendrite analysis, pyramidal cells were pooled into one group, as different pyramidal cells could not be accurately distinguished without full labeling of apical dendrites. Despite this, the difference between genotypes greatly exceeded the differences seen in dendrite complexity that exists for L5 neuronal subtypes (Oswald et al., 2013). The same applies to the changes in soma size.

MOUSE BEHAVIOR

For behavior assessment, male mice were group-housed in standard cages with bedding and nesting material under a 12 h light–dark cycle. Food and water were provided ad libitum. Behavior assessment was started when animals were 8–9 weeks old (beam, rotarod, and suspended wire) and 3 or 7 months old (footprint test). Experiments were performed with 1–2 day intervals between tests. Experiments were carried out between 10:00 and 15:00 h. For all tests (with the exception of the footprint test), video-tracking was used. Scoring was carried out using video and sound material by an experimenter that was blinded to the genotype.

BEAM

Coordination and motor ability was assessed. A mouse was placed in the middle of a horizontal beam (2 cm diameter, 120 cm length) raised 40 cm. Color painted marks divided the beam into 10 cm intervals. Mice were allowed to explore beam for 2 min or until they fell from the beam. Time spent on the beam and the number of crossed marks were calculated. After 1 h interval a second trial was repeated. Eighteen male mice (8 Jnk1-/− and 10 wild-type littermates) were used.

ROTA-ROD

Evaluation of motor ability and motor learning was performed on the rotarod (Ugo Basile, Italy) on 2 days. Every experimental day included three trials with 1 h interval. The mouse was placed on the rotating drum with acceleration (from 4 to 40 r.p.m. over 5 min). The latency to fall (6 min cut-off time) was measured over all sessions. Eighteen male mice (8 Jnk1-/− and 10 wild-type littermates) were used.

SUSPENDED BAR TEST

Mice were placed hanging from their forepaws on the center of a wire coat hanger (diameter 3 mm, length 39 cm) at a height of 41 cm above a container with soft bedding. After one training period, mice were video tracked for 45 s. Afterwards separate latencies were scored: (1) time to finish was the time taken to assume an upright position and reach the corner of the coat-hanger with the forepaws, (2) time to get “up” was the time taken to climb further onto the diagonal bar and touch 5 cm along with the forepaws, and (3) time to fall from the wire. A penalty score of 45 s was added to the “finish” score for mice that fell from the bar or failed to complete the task. A penalty of 45 s was added to the “up” score of mice that failed to reach 5 cm along the diagonal bar.

INVERTED GRID TEST

Mice were placed on a metal grid (spacing 1 cm²) and allowed to grip the grid with four paws. The grid was inverted at an angle of 180° 10 cm above the ground and the time for the mouse to fall onto soft bedding was measured in seconds. The monitoring time was 2 min.

FOOTPRINT TEST

The footprint analysis was performed with 12 wild-type and 12 Jnk1-/− mice at 3 months or with five wild-type and seven Jnk1-/− mice at 7 months. The hind paws of mice were coated with ink and mice were encouraged to walk along a runway with side walls of 10 cm. The runway was covered with paper that was 7.5 cm wide, and 80 cm long. Footprints were used to measure a series of 6–10 sequential steps recorded in three trials per mouse. Stride length was measured between the central pads of two consecutive prints on each side. Stride width was measured between the central pads of two footprints, one on either side. Three trials per mouse were meaned. SEMs are shown.

ANATOMICAL MEASUREMENTS

Body weight of mice at 8- to 9-weeks old was measured before and after the battery of behavioral tests [beam, rotarod, acoustic startle, water-maze (not shown)]. The body weight of 3- and 7-month-old animals was measured before the footprint test and the suspended bar tests were performed. To measure hindlimb and forelimb length, mice after behavioral tests were sacrificed and pinned to a polystyrene platform. The joint between the forelimb and the clavicle was determined by examination. The distance from this joint to the end of the paw was measured. For measurement of hindlimb, the joint between the hindlimb and the hip was felt by examination with fingers. The distance from this join to the tip of the paw was measured. Distance between the forelimbs and hindlimbs was measured in a similar way according to the scheme.

STATISTICAL ANALYSIS

Testing of sample variances was done by Student’s t-test or by one-way ANOVA followed by Bonferroni post hoc test using QI Macros SPC software (KnowWare International, Inc.).
RESULTS
JNK1 PHOSPHORYLATES RAT HMW-MAP2 ON T1619, T1622, AND T1625 IN THE PROLINE RICH DOMAIN (PRD)

Earlier work from our group and others showed that HMW-MAP2 is phosphorylated by JNK1 in vitro (Chang et al., 2003; Björkblom et al., 2005). To determine whether this phosphorylation was specific for JNK1, and to identify which amino acids are phosphorylated, we purified recombinant GST-MAP2 from E. coli and phosphorylated it in vitro. JNK1 and JNK3 both phosphorylated GST-MAP2, while JNK1 was slightly more efficient (Figure 1A), suggesting that either JNK isoform can phosphorylate MAP2 in vivo. To identify the phosphorylation sites on GST-MAP2, we carried out MS/MS analysis of TiO2-enriched phosphopeptides from the JNK1 phosphorylated GST-MAP2. This revealed three JNK1-phosphorylated sites in the C-terminal domain; T1619, T1622, and T1625 of rat HMW-MAP2 (Figure 1B).

Since JNK1 is constitutively active in developing brain (Coffey et al., 2000; Tararuk et al., 2006), we investigated whether any of the JNK1 sites on HMW-MAP2 were constitutively phosphorylated in mouse brain under physiological conditions. MS/MS analysis of...
phospho-peptides from postnatal day 2 brain identified 10 sites with basal phosphate occupancy (Figure 1C). Among these, the JNK1 sites T1620 and T1623 of mouse HMW-MAP2 were clearly phosphorylated. The MS/MS spectrum for the dual phosphorylated peptide is shown (Figure 1C). Mouse HMW-MAP2-T1617, -T1620, and -T1623 (Uniprot P20357) correspond to rat HMW-MAP2 sites -T1619, -T1622, and -T1625 (Uniprot P15146). Nine additional phosphopeptides were identified, among these four were previously described (Collins et al., 2005; Trinidad et al., 2006; Munton et al., 2007), while the remaining phosphopeptides were novel.

HMW-MAP2 encodes 43 (Pro)-X-Ser/Thr-Pro motifs (Sánchez et al., 1995) – putative motifs for JNK phosphorylation. As our analysis of JNK sites on MAP2 were carried out using GST-MAP2 (Figure 1A), which lacks the central domain (≈1360 amino acids) of high-molecular-weight (HMW) MAP2, we wanted to determine whether additional JNK sites could exist in the central domain. To test this, we performed metabolic labeling using GFP-HMW-MAP2 phospho-site mutants (Figure 1D). GFP-HMW-MAP2WT or GFP-HMW-MAP2T1619D,T1622D,T1625D, where the JNK1 phosphorylation sites were mutated from threonine to aspartate, were expressed in COS-7 cells. Metabolic labeling of ATP γ-32P]-ATP allowed us to measure phosphate incorporation. GFP-MAP2WT phosphorylation was increased upon co-expression of the constitutively active JNK1 chimera, MKK7-p38 and the JNK-specific inhibitor SP600125 prevented JNK1-dependent phosphorylation (Figures 1E,F). However, expression of GFP-HMW-MAP2T1619D,T1622D,T1625D led to a greater number of cells with protrusions (Figures 2A,B). Similarly, activation of JNK, increased the number of protrusions in GFP-HMW-MAP2WT-expressing cells (Figure 2C). Strikingly, cells that expressed GFP-HMW-MAP2T1619A,T1622A,T1625A failed to generate protrusions altogether, even in the presence of active JNK1 (MKK7-JNK1; Figures 2A–C). Furthermore, GFP-HMW-MAP2T1619A,T1622A,T1625A remained visibly soluble. These data indicate that JNK1-dependent phosphorylation of T1619, T1622, and T1625 of HMW-MAP2 is necessary and sufficient for protrusion growth.

JNK1 PHOSPHORYLATES HMW-MAP2 ON EQUIVALENT SITES IN MOUSE BRAIN

To test whether JNK1 phosphorylated HMW-MAP2 on these sites in vivo, we utilized a commercial antibody that detects the phosphorylated PRD of HMW-MAP2. We examined the site specificity of this antibody, and found that it recognized GFP-HMW-MAP2WT but not GFP-HMW-MAP2T1619A,T1622A,T1625A (Figure 1E) indicating that it was specific for the JNK1-targeted threonine residues of HMW-MAP2 in a phosphorylated state. We used this antibody to compare the phosphorylation of endogenous HMW-MAP2 from wild-type and Jnk1-/- mice. Phospho-MAP2 levels were decreased in the cortex and hippocampus in Jnk1-/- mice, indicating that JNK1 phosphorylates these sites in immature brain (Figures 1F,G).

GFP-MAP2T1619D,T1622D,T1625D EXPRESSING CELLS EXTEND A GREATER NUMBER OF PROTRUSIONS, WHILE MAP2T1619A,T1622A,T1625A EXPRESSING CELLS COMPLETELY FAIL TO GENERATE PROTRUSIONS

MAP2 binds laterally along microtubule polymers and confers rigidity to dendritic arbors (Felgner et al., 1997). To determine whether the JNK1 phosphorylated residues of HMW-MAP2 influence process formation, we prepared a phospho-mimicry mutant of HMW-MAP2 (GFP-HMW-MAP2T1619D,T1622D,T1625Dγ), and expressed either this, GFP-HMW-MAP2T1619A,T1622A,T1625A or GFP-HMW-MAP2WT in COS-7 cells, which lack endogenous MAP2. Exogenous expression of GFP-HMW-MAP2 in COS-7 cells is sufficient to induce occasional protrusions from the soma (Björkblom et al., 2005; Figures 2A,B). However, expression of GFP-HMW-MAP2T1619D,T1622D,T1625Dγ led to a greater number of cells with protrusions (Figures 2A,B). Similarly, activation of JNK, increased the number of protrusions in GFP-HMW-MAP2WT-expressing cells (Figure 2C). Strikingly, cells that expressed GFP-HMW-MAP2T1619A,T1622A,T1625A failed to generate protrusions altogether, even in the presence of active JNK1 (MKK7-JNK1; Figures 2A–C). Furthermore, GFP-HMW-MAP2T1619A,T1622A,T1625A remained visibly soluble. These data indicate that JNK1-dependent phosphorylation of T1619, T1622, and T1625 of HMW-MAP2 is necessary and sufficient for protrusion growth.

JNK1 regulates dendritic field size in vivo

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Lucifer yellow dye. Many of the pyramidal neurons we traced in L5 had large somas consistent with their being corticospinal projecting neurons of L5B (Oswald et al., 2013). Three-dimensional confocal sections underwent Sholl analysis. Concentric rings were spaced 20 μm apart centered around the soma (as illustrated Figure 3A). Intersections, nodes and length of dendritic arbors were mapped within each concentric ring. Only basal dendrites were measured, as dye loading was frequently incomplete in the apical dendrites and tufts, which were excluded. We started with layer 2/3 (L2/3) neurons in the rostral and caudal forelimb areas (RFA, CFA), medial agranular cortex AGm, and lateral agranular cortex (AGl) of the mouse primary motor cortex (M1; Tennant et al., 2011). Small and medium sized pyramidal cells were included. There was reduced complexity in the upper layer neurons of Jnk1−/− mice, defined as fewer intersections, a smaller number of nodes, and reduced dendritic length (Figures 3B–G). In contrast, the layer 5 (L5) neurons of Jnk1−/− mice showed increased dendrite complexity (Figures 3H–L, N). Moreover, the extent of architecture alterations in the large pyramidal neurons of L5 was greater than that observed in the upper L2/3. Representative
tracings of primary motor cortex neurons from the deeper layers are shown (Figure 3N). The increase in dendrite complexity in L5 may have developed at least in part, to offset the reduced input from L2/3 neurons which display reduced dendrite elaboration. The overall elaboration of dendrites observed in the projection neurons of the primary motor cortex in these mice are consistent with our previous findings in isolated cerebellar granule neurons from Jnk1−/− mice (Björklom et al., 2005), suggesting that a common mechanism exists.

**EXPRESSION OF CAG-EGFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> IN VIVO IN THE MOTOR CORTEX ALTERS THE ARCHITECTURE OF L2/3 DENDRITES**

Pyramidal neuron dendrites of the motor cortex respond to external cues and genetic programs to undergo arborization phases from P7 to P21 (Aizawa et al., 2004; Urbanska et al., 2008; Romand et al., 2011). We showed that GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> facilitates protrusion growth in COS7 cells. To determine if phosphorylation of MAP2 on these sites altered dendrite architecture in vivo, we prepared HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> phosphorylation site mutants under the control of the CAG promoter and introduced them to the developing motor cortex using targeted electroporation of E15 embryos. This resulted in efficient labeling of cells in L2/3 of the motor cortex (Figure 4A), however as this technique labels cells destined for the superficial layers, L5 cells did not express ectopic GFP-HMW-MAP2. Dendrite architecture in L2/3 was analyzed from the mice when they reached postnatal day 21, by which time the basal dendrites have developed most of their adult features (Romand et al., 2011). Basal dendrites were analyzed in entirety for each cell as outlined (Figures 4A–C). Neurons expressing GFP-HMW-MAP2<sub>WT</sub> displayed a similar architecture to L2/3 motor cortex neurons labeled with Lucifer yellow, with no alteration in total dendrite length (Figure 3E compared to Figure 4H), although the first branching occurred closer to the soma in these cells (Figures 4D and 3B). Interestingly however, GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> expressing L2/3 cells produced significantly more elaborate dendrites, with increased intersections (Figure 4D). This large increase in the number of intersections resembles the arborization pattern observed in more mature neurons (Figure 3D; Romand et al., 2011).

Higher order dendrite length (third and fourth order) was considerably increased in motor cortex cells expressing GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> (Figure 4E), although the primary dendrite length remained unchanged (Figures 4E,G). Extensive arbor branching, increasing significantly from the primary to the fourth order dendrites (Figures 4J), was visible in GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub>-expressing cells, and there was a trend towards increased total surface area (Figure 4K). Representative images of L2/3 pyramidal neurons labeled with GFP-HMW-MAP2<sub>WT</sub> or with GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> are shown (Figure 4L). In summary, the overall effect of GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> expression compared to GFP-HMW-MAP2<sub>WT</sub> was to increase the length of higher order dendrites and to increase branching of dendrites at all orders. The overall increase in total dendrite length was 50%, partially explained by the extensive increase in branching (Figure 4H). It is notable that neurons expressing GFP-HMW-MAP2<sub>T1619D,T1622D,T1625D</sub> resided a greater distance from the pial surface than neurons expressing GFP-HMW-MAP2<sub>WT</sub> (Figure 4M). As previously shown, cytosolic JNK1 acts as a negative regulator that retards radial migration (Westelund et al., 2011). HMW-MAP2 phosphorylation by JNK1 may thus prematurely halt the migration.

**CELL SOMA AREA IS INCREASED IN L5 M1 PYRAMIDAL NEURONS IN Jnk1−/− MICE**

Examination of cell soma size in M1 L5 neurons indicated that somas were on average 35% larger in Jnk1−/− mice (Figures 5A,B). This was done by stereological measurement of cell soma cross sectional area and perimeter from three-dimensional images of lucifer yellow loaded pyramidal neurons. The increased soma size observed in Jnk1−/− pyramidal neurons is consistent with increased dendritic load in L5 (Figure 3), as soma size has been shown to correlate with dendritic material load (van Pelt et al., 1996).

**Jnk1−/− MICE SHOW DEFECTIVE MOTOR SKILLS**

Since we observed dendrite complexity alterations in L2/3 and L5 of the primary motor cortex, which controls complex muscle activation patterns (Levine et al., 2012), we tested the performance of Jnk1−/− mice using a battery of behavioral tests assessing motor coordination and strength. These tests require a high degree of motor function and relatively subtle deficits can be revealed. We first tested motor activity and balance of mice on the raised beam. Jnk1−/− mice performed poorly while traversing the beam, showing a significantly reduced latency to fall compared to wild-type mice (Figure 6A). Notably, knockout mice frequently had difficulty with placement of hind paws and forward movement of hind limbs (Figure 6B), indicating impaired coordination and balance. We next monitored motor coordination and grip strength by measuring the latency to fall from a suspended wire (Lalonde et al., 1992; Baldo et al., 2012). Mice were placed hanging by their two forelimbs from a wire coat-hanger. Mice lacking Jnk1 showed a reduced latency to fall from the wire suggesting weakened grip strength (Figures 6C,D). When they did complete the task, Jnk1−/− mice took more time to do so (Figure 6C). During the 45 s time allocated, 57% of Jnk1−/− mice failed to complete the task compared to 29% of wild-type mice. We also measured grip strength using the inverted grid test where both forelimbs and hindlimbs are used to grip to an inverted grid. Mice lacking Jnk1 showed significantly reduced latency to fall compared to wild-type (Figure 6E). Together these data suggest impaired motor coordination and strength in mice lacking Jnk1.

A more detailed examination of motor coordination and synchrony is provided by measuring gait using the footprint test (Brooks and Dunnett, 2009). Manual tracing of footprints revealed a reduction in stride width in 3 month old jnk1−/− mice compared to wild-type (Figure 6F). This difference between genotypes was rectified by 7 months. Interestingly though, in 7 month old knockout mice, the stride length was significantly reduced. While we detected a minor (2%) increase in hindlimb length in older knockout mice, this seems unlikely to account for the more significant (15%) decrease in stride length (Figures 6E,G). The small difference in forelimb and hindlimb lengths may be explained by osteoclast deficits as JNK1 regulates osteoclast differentiation.
FIGURE 3 | Dendrite morphology is disrupted in the primary motor cortex of Jnk1-/− mice. (A) Wild-type (WT) and Jnk1-/− mice at 6–8 months of age were loaded with Lucifer yellow dye and a Sholl analysis was carried out on single dendrites, measuring only basal dendrites from reconstructed three-dimensional images. (B–D) L2/3 neurons from the primary motor cortex (M1) were analyzed for dendrite complexity. The number of intersections, nodes, and length were calculated from single complete dendritic trees. There was a significant decrease in dendrite complexity in Jnk1-/− mice. (E–G) There was no significant change in the total dendrite length or number of nodes, though the total surface area was significantly decreased significantly in Jnk1-/− mice in L2/3 neurons. (H–J) In L5 neurons from M1, there was a significant increase in dendrite complexity exemplified as increased number of intersections, nodes and increased dendrite length in Jnk1-/− brains compared to wild-type. (K–M) The total dendritic tree length, number of nodes per tree and total surface area is shown. (N) Neurolucida tracings from M1 L5 neurons in WT and Jnk1-/− mice are shown. Means ± SEMs are indicated. For L2/3 neurons 81 WT and 61 Jnk1-/− dendritic trees were analyzed from a total of 67 cells. For L5 neurons, 49 WT and 63 Jnk1-/− trees were analyzed from a total of 47 cells. For (B–D,H–J), significance was determined using one-way ANOVA. For (E–G,K,L), significance was determined using Student’s t-test. Significance levels are **p < 0.01, ***p < 0.001.
FIGURE 4 | MAP2T1619D,T1622D,T1625D regulates basal dendrite architecture in vivo. (A) Mice electroporated with DNA at E15 in utero were sacrificed at postnatal day 21. The fluorescent micrograph of a cerebral hemisphere shows expression of GFP in L2/3 of Motor cortex 1 (M1). The midline (ML) is indicated. (B,C) Higher magnification confocal micrographs of L2/3 depicting the neurolcida tracing (B) and the dense network of GFP-expressing cells (C). (D,E) The number of intersections (D) and dendrite length (E) from Sholl analysis of whole cell basal dendrites from mice electroporated in utero with pEGFP-CAG-MAP2T1619D,T1622D,T1625D revealed increased dendrite complexity compared to mice expressing pEGFP-CAG-MAP2WT. (F) There was a significant increase in dendrite length in third and fourth order branches. (G) Primary dendrite length however remained unchanged. (H) There was a significant increase in total dendrite length in MAP2T1619D,T1622D,T1625D expressing mice. (I) The average number of nodes per dendritic tree was increased in Jnk1-/- mice. (J) The number of nodes was significantly increased at every dendrite order. (K) The surface area measurements from neurons in GFP-MAP2WT or GFP-MAP2T1619D,T1622D,T1625D expressing mice are shown. (L) Representative Sholl traces from mice expressing GFP-MAP2WT or GFP-MAP2T1619D,T1622D,T1625D are shown. Statistical analysis in (D–F,J) were two-way ANOVA with post hoc Bonferroni test of significance for individual points. *p < 0.05; **p < 0.01; ***p < 0.005. (G–K,M) were analyzed by Student’s t-test. *p < 0.05; ***p < 0.001; ****p < 0.0001. Twenty-four dendritic trees were analyzed from four separate mice. (Ten from WT and 14 from GFP-T1619D,T1622D,T1625D–expressing mice. (M) The distance from the pia of neurons expressing GFP-MAP2WT or GFP-MAP2T1619D,T1622D,T1625D is indicated. Neurons expressing GFP-MAP2T1619D,T1622D,T1625D remained further from the pial surface in 21d mice.
Jnk1-/− mice. As the Jnk1-/− mice exhibited significantly impaired behavior. Finally, we measured ventral length, but found no differences (not shown).

We also subjected mice to the rotarod test (Figure 6I). There was no phenotypic difference between wild-type and knockout mice when they were placed on an accelerating rod, suggesting that gross motor skills were intact. Similarly, Jnk1-/− mice showed no locomotive defect in the open field test (not shown). The rotarod test is considered a less sensitive test of coordination (Brooks and Dunnett, 2009), with the drawback that heavy mice perform poorly compared to lighter mice. As the Jnk1-/− mice are 14.3 ± 1.7% lighter than wild-type mice (Figure 6I), it is possible that an offset due to reduced body weight masks an underlying deficit in coordination and balance control. Of significance, the footprint test of gait is one of the few tests that translates directly from animal to human studies (Brooks and Dunnett, 2009). It crosslinks microtubule protofilaments, facilitated by its C-terminal PRD (Hirokawa et al., 1996; Sanchez et al., 2000; Al-Bassam et al., 2002). While HMW-MAP2 is a highly phosphorylated protein, the outcome of site-specific phosphorylation has not been previously tested either in vitro or in vivo. Our data indicates that JNK1 phosphorylates T1617, T1620, and T1623 (equivalent to T1619, T1622, and T1625 in rat HMW-MAP2) in the cortex and hippocampus of early postnatal mice, as phosphorylation on these sites is reduced in Jnk1-/− mice. Site-directed mutagenesis indicates that the functional consequence of this phosphorylation is to switch HMW-MAP2 from a form that interacts poorly with microtubules and fails to support process growth, to a form that binds avidly and promotes outgrowth (Figures 1 and 2).

In vivo, we show that ectopic expression of GFP-HMW-MAP2T1619D,T1622D,T1625D (phospho-mimicry form) in L2/3 of M1, substantially increases dendrite arborization (Figures 4D, I, J). This is likely to be physiologically meaningful because (i) these sites are phosphorylated in vivo in brain (Figures 1C–G) and (ii) phosphorylation of this domain increases during development and correlates with increased branching of cultured neurons (Diez-Guerra and Avila, 1993). Moreover, ectopic expression of pseudo-phosphorylated HMW-MAP2 augments arbor length, consistent with earlier findings in MAP2 knockout mice where arbor length was reduced (Harada et al., 2002). The extensive increase in higher order dendrite branching upon ectopic expression of phospho-mimicry HMW-MAP2 is somewhat unexpected. However, new arbor formation requires coordinated interplay between microtubules and actin (Farah and Leclerc, 2008), and MAP2 is capable of binding and crosslinking actin, and co-localizes with actin at sites of protrusion sprouting (Dehmelt et al., 2003; Roger et al., 2004). Thus, this phosphorylation may provide a switch whereby arborization of dendrites can be regulated. For example, the pseudo-phosphorylated state of GFP-HMW-MAP2 may protect those dendrites from pruning that otherwise occurs between P7 and P21 (Aizawa et al., 2004; Urbanska et al., 2008; Romand et al., 2011). Together these results indicate that MAP2 mediates JNK1 regulation of dendrite morphology in L2/3 of the motor cortex.

Receptive fields of dendritic arbors determine where, and to what extent the neuron receives synaptic input. In contrast to L2/3, JNK1 substantially decreases the total length and branching of basal dendrites in L5 (by 20–30%, Figures 3H–L), while only moderately increasing the dendritic field in L2/3 (Figures 3B–G). The increased dendritic field in L5 of Jnk1-/− mice may result
FIGURE 6 | *Jnk1*−/− mice show impaired motor skills. (A) Wild-type (WT) and *Jnk1*−/− mice were subjected to the beam walking test. Two trials were undertaken and time to fall measured. In the second trial *Jnk1*−/− mice showed a shorter latency to fall than WT mice. Eight male mice of each genotype were tested. (B) *Jnk1*−/− mouse paws often slipped to the side in the beam test. WT mice did not show this behavior. Representative images from four individual mice are shown. (C) To determine muscle and grip strength, mice were subjected to a grip test using a suspended wire. The results show the latency to “fall,” time to “finish,” and time to “up” (reach 5 cm on the diagonal bar with the forepaws) as illustrated. *Jnk1*−/− mice showed a significantly shorter latency to fall than WT. Furthermore, 57% of *Jnk1*−/− mice failed to complete the course compared to only 29% of WT mice. Seventeen male WT and 21 male *Jnk1*−/− mice were tested. (D) Representative images of mice in the “start,” “finish,” and “up” positions. (E) Fore limb and hind limb strength was measured using the latency to fall of mice hanging on an inverted grid. Five WT and 5 *Jnk1*−/− mice were measured. (F) Gait was assessed by measuring distances between footprints from 3 month (3 m) and 7 month (7 m) mice. *Jnk1*−/− mice at 3 m displayed a significantly narrower stride width than WT mice. By 7 m the difference in stride width was no longer significant however stride length was significantly reduced in *Jnk1*−/− mice. Twelve mice from each genotype were analyzed. (G) A table displaying the anatomical measurements of WT and *Jnk1*−/− mice at 3 m and 7 m is shown. Measured data ± SEM from 11 WT and 12 *Jnk1*−/− mice at 3 m and from five WT and seven *Jnk1*−/− mice at 7 m, are shown. Where significant difference to WT exists, *p*-values are indicated. N.s. indicates that differences compared to controls were not significant. (H) The scheme illustrates the measurement strategy used to determine length of forelimbs and hindlimbs. (I) Mice were subjected to the accelerating rotarod test. There was no phenotypic difference between genotypes. Data shown are meaned from eight WT and eight *Jnk1*−/− male mice. (J) The average weight of 8-week old WT and *Jnk1*−/− mice before and after the battery of behavioral tests is shown. *Jnk1*−/− mice were significantly lighter than WT. For all graphs, means ± SEMs are shown. Levels of significance determined by ANOVA or Student’s *t*-test, are as follows: ** *p* < 0.01; *** *p* < 0.001; **** *p* < 0.0005.
Aside from irregularities in M1, these behaviors could in principle (Gilman et al., 1981), is reduced in young adult mice lacking altered stride width, a classic hallmark of cerebellar dysfunction. There is a pretext for a cerebellar component, as targeted deletion of Jnk1-/- mice exhibited impaired hindlimb placement, gripping the beam from the sides instead of placing their paws on the upper surface. Appropriate limb placement requires a reflex to tactile stimulation that is mediated by the motor cortex (Metz and Whishaw, 2002), as well as motor coordination (Young and Kaas, 2012). Analysis of gait revealed that in older mice lacking Jnk1-/-, stride length was significantly decreased (Figure 6). On the beam, Jnk1-/- mice exhibited impaired hindlimb placement, gripping the beam from the sides instead of placing their paws on the upper surface. Appropriate limb placement requires a reflex to tactile stimulation that is mediated by the motor cortex (Metz and Whishaw, 2002), as well as motor coordination (Young and Kaas, 2012). Analysis of gait revealed that in older mice lacking Jnk1-/-, stride length was significantly decreased (Figure 6).

Irregular growth or maintenance of dendrites, contributes to the development of psychiatric disease (Jan and Jan, 2010). Given that increased risk for these disorders is associated with genetic abnormalities in JNK signaling (Coffey, 2014), our findings in Jnk1-/- mice are important as they demonstrate the molecular and behavioral consequences of interfering with JNK pathway signaling and provide grounds for improved understanding of the molecular underpinnings of psychiatric disorders.

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