DOPAMINE- AND cAMP-REGULATED PHOSPHOPROTEIN OF 32 kDa (DARPP-32)-
DEPENDENT ACTIVATION OF ERK AND MAMMALIAN TARGET OF RAPAMYCIN
COMPLEX 1 (TORC1) SIGNALING IN EXPERIMENTAL PARKINSONISM
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Running head: DARPP-32-dependent activation of ERK and mTORC1

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**Background:** DARPP-32 is implicated in L-DOPA-induced dyskinesia.
**Results:** PKA-dependent phosphorylation of DARPP-32 in a distinct subset of striatal neurons is
required for L-DOPA-induced activation of ERK and mTORC1.
**Conclusion:** PKA-dependent phosphorylation of DARPP-32 plays a critical role in dyskinesia and
associated signaling alterations.
**Significance:** The PKA/DARPP-32 cascade is a key target for the treatment of dyskinesia.

Dyskinesia, a motor complication caused by prolonged administration of the
antiparkinsonian drug L-DOPA, is
accompanied by activation of cAMP signaling and hyperphosphorylation of the
dopamine- and cAMP-regulated phosphoprotein of 32 kDa (DARPP-32).
Here, we show that the abnormal phosphorylation of DARPP-32 occurs specifically in medium spiny
neurons (MSNs) expressing dopamine D1 receptors (D1R). Using mice in which DARPP-32 is
selectively deleted in D1R-expressing MSNs, we demonstrate that this protein is required for
L-DOPA-induced activation of the extracellular signal regulated protein kinases 1 and 2 (ERK) and the mammalian
target of rapamycin complex 1 (mTORC1) pathways, which are implicated in dyskinesia. We also show that mutation of
the phosphorylation site for cAMP-dependent protein kinase on DARPP-32 attenuates L-DOPA-induced dyskinesia and
reduces the concomitant activations of ERK and mTORC1 signaling. These studies demonstrate that, in D1R-expressing MSNs,
L-DOPA-induced activation of ERK and mTORC1 requires DARPP-32 and indicates the importance of the
cAMP/DARPP-32 signaling cascade in dyskinesia.

Parkinson’s disease (PD) is a frequent
neurodegenerative disorder characterized by
the progressive death of the dopaminergic
neurons of the substantia nigra pars compacta
(1). In the dorsal striatum, the area preferentially innervated by the substantia nigra pars compacta, medium spiny neurons
(MSNs) react to the loss of dopamine by
increasing their sensitivity to dopaminergic agonists (2). This phenomenon is particularly
evident at the level of the MSNs that directly
innervate the output nuclei of the basal
ganglia. These cells are generally referred to as striatonigral MSNs and are selectively
enriched in dopamine D1 receptors (D1Rs)
(3). The development of sensitized D1R-
mediated transmission in striatonigral MSNs
has been proposed to play a critical role in the
generation of the motor side effects, or
dyskinesia, produced by repeated administration of L-DOPA, the most common
antiparkinsonian medication (4,5).

In the dopamine-depleted striatum, but not
in the normal striatum, L-DOPA promotes the
phosphorylation of the dopamine- and cAMP-
regulated phosphoprotein of 32 kDa (DARPP-32)
(6,7), a key component of the canonical
cAMP/cAMP-dependent protein kinase (PKA) signaling cascade activated by D1Rs (8). DARPP-32 has been causally linked to the emergence of L-DOPA-induced dyskinesia (LID) (7). In agreement with this idea, it has been recently shown that LID is decreased by selective inactivation of DARPP-32 in striatonigral MSNs (9).

In addition to the cAMP/PKA/DARPP-32 pathway, L-DOPA-mediated stimulation of D1Rs activates the extracellular signal regulated kinases 1 and 2 (ERK) (7,10-12) and the mammalian target of rapamycin complex 1 (mTORC1) (13), which are critical regulators of transcription and translation (14,15). Abnormal activation of ERK and mTORC1 has been implicated in the development of L-DOPA-induced dyskinesia (LID) (7,13,16,17). It has been proposed that DARPP-32 is involved in the increase of ERK phosphorylation observed in the striatum following stimulation of D1Rs (7,18). Previous work also showed that, in mice lesioned with 6-hydroxydopamine (6-OHDA), a toxin used to model PD in rodents), inhibition of ERK prevents the ability of a D1R agonist to activate mTORC1 (13). Taken together, these findings suggest that L-DOPA may promote mTORC1 signaling via sequential and coordinated activation of DARPP-32 and ERK. However, the existence of a link between DARPP-32 and ERK signaling in the dorsal striatum has been challenged (10), prompting a more detailed analysis of the involvement of DARPP-32 in the regulation of ERK and mTORC1. In this study, we provide evidence demonstrating that L-DOPA increases PKA/DARPP-32 signaling selectively in striatonigral MSNs and that PKA-dependent phosphorylation of DARPP-32 is implicated in LID and in the activation of the ERK and mTORC1 cascades associated with this condition.

**EXPERIMENTAL PROCEDURES**

**Animals**—In this study we used mice expressing Flag-tagged DARPP-32 selectively in D1R-expressing striatonigral neurons and Myc-tagged DARPP-32 selectively in dopamine D2 receptor (D2R)-expressing striatopallidal neurons (DI-DARPP-32-Flag/D2-DARPP-32-Myc transgenic mice) (19), mice in which DARPP-32 is conditionally deleted in D1R-, or D2R-expressing MSNs by means of the loxP/Cre recombinase system (D32^{F/F}D1RCre^{+} and D32^{F/F}D2RCre^{+} conditional knock out mice) (9) and knock-in mice expressing a mutated form of DARPP-32, in which the phosphorylation site for PKA (Thr34) is substituted with an Ala (Ala34Thr mutant mice) (20).

**Drugs**—L-DOPA (Sigma-Aldrich Sweden AB) was injected at a dose of 10 or 20 mg/kg in combination with the peripheral DOPA decarboxylase inhibitor, benserazide hydrochloride (Sigma-Aldrich Sweden AB) (7.5 or 12 mg/kg). Both drugs were dissolved in physiological saline (0.9% NaCl) and injected intraperitoneally in a total volume of 10 ml/kg of body weight. When mice were not treated with L-DOPA, they received an equivalent volume of vehicle.

**6-OHDA lesion**—Mice were anaesthetized with a mixture of fentanyl citrate (0.315 mg/ml), fluanisone (10 mg/ml) (VetaPharma, Leeds, UK), midazolam (5 mg/ml) (Hameln Pharmaceuticals, Gloucester, UK) and water (1:1:2 in a volume of 10 ml/kg) and mounted in a stereotaxic frame (David Kopf Instruments, Tujunga, CA) equipped with a mouse adaptor. 6-OHDA-HCl (Sigma-Aldrich, Sweden AB) was dissolved in 0.02% ascorbic acid in saline at a concentration of 3 µg of freebase 6-OHDA/µl. Each mouse received two unilateral injections of 6-OHDA (2 µl/injection) into the right dorsal striatum as previously described (7), according to the following coordinates (mm) (21): AP +1, ML - 2.1, DV -3.2 and AP +0.3, ML -2.3, DV -3.2. Animals were allowed to recover for 3 weeks before behavioral evaluation and drug treatment. This procedure leads to a decrease in striatal tyrosine hydroxylase (TH) immunoreactivity ≥ 80% and to a marked akinesia affecting the side of the body contralateral to the lesioned striatum (7,13).

**Abnormal involuntary movements (AIMs)**—Mice were treated for 10 days with one injection per day of L-DOPA (20 or 10 mg/kg) plus benserazide (12 or 7.5 mg/kg). AIMs were assessed after the last injection (day 10) using a previously established scale (22). Twenty minutes after L-DOPA administration, mice were placed in separated cages and individual dyskinetic behaviors (i.e., AIMs) were assessed for 1 min every 20 min, over a period of 120 min. AIMs were classified into four subtypes: locomotive AIMs (contralateral

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**Note:**

1. The text is a continuation of a scientific article discussing the role of DARPP-32 in L-DOPA-induced dyskinesia and related signaling pathways.
2. The methods section includes details on the experimental design, including the use of transgenic and knock-in mice to study the effects of L-DOPA and 6-OHDA on striatal function.
3. The results section would typically follow, detailing the observed effects and mechanisms underlying the phenomena discussed in the methods section.
4. The conclusions would summarize the findings and implications for future research.
5. The reference section would list the sources used in the study.
turns), axial AIMS (dystonic posturing of the upper part of the body toward the side contralateral to the lesion), limb AIMS (abnormal movements of the forelimb contralateral to the lesion), and orolingual AIMS (vacuous jaw movements and tongue protrusion). Each subtype was scored on a severity scale from 0 to 4: 0, absent; 1, occasional; 2, frequent; 3, continuous; 4, continuous and not interruptible by outer stimuli.

Flag and Myc-tagged DARPP-32 immunoprecipitations—Tagged DARPP-32 was immunoprecipitated from acutely dissected striata as previously described (18). Briefly, mice were sacrificed using focused microwave irradiation and bilateral striata from each mouse were rapidly dissected and frozen on liquid nitrogen. Striata were then sonicated in lysis buffer with protease and phosphatase inhibitors and homogenates were incubated simultaneously with EZView Red anti-Flag M2 affinity gel (Sigma) and Myc antibody–coupled (Novus) magnetic beads (Invitrogen) overnight at 4°C. Anti-Flag beads were separated from the anti-Myc beads using a magnetic particle concentrator (Invitrogen). Anti-Flag and anti-Myc beads were separately washed and bound proteins were eluted by boiling in sample buffer. The unbound homogenate was retained for the total striatum sample.

Western blotting—For the studies with D32F/F1DRCre+ and D32F/F2DRCre+ conditional knock out mice, the animals were treated with L-DOPA plus benserazide and killed by decapitation 30 min later. The heads of the animals were cooled in liquid nitrogen for 6 seconds and the brains were removed. The dorsal striata were dissected out on an ice-cold surface, sonicated in 750 µl of 1% SDS and boiled for 10 min. Aliquots (5 µl) of the homogenate were used for protein determination using a BCA (bicinchoninic acid) assay kit (Pierce ESRB, Oud Beijerland, the Netherlands). Equal amounts of protein (30 µg) for each sample were loaded onto 10% polyacrylamide gels. Proteins were separated by sodium dodecylsulfate-polyacrylamide gel electrophoresis and transferred overnight to PVDF membranes (Amersham Pharmacia Biotech, Uppsala, Sweden) (23). The membranes were immunoblotted using antibodies against phospho-Thr34-DARPP-32 (1:750) (24), phospho-Ser10-acetyl-Lys14-histone H3 (1:50000) (Millipore AB, Solna, Sweden), phospho-Thr202/Tyr204-ERK (1:2000), phospho-Ser235/236-ribosomal protein S6 (rpS6) (1:1000), phospho-Ser240/244-ribosomal protein S6 (1:1000) (Cell Signaling Technology, Beverly, MA, USA) and phospho-Ser845-GluA1 (1:1000) (PhosphoSolutions, Aurora, CO, USA). Antibodies against histone H3 (1:50000, Abcam), ERK (1:1000), rpS6 (1:1000), GluA1 (1:1000) (Cell Signaling Technology, Beverly, MA, USA) and DARPP-32 (1:750) (25) that are not phosphorylation state specific were used to estimate the total amount of protein. Detection was based on fluorescent secondary antibody binding and quantified using a LiCor Odyssey infrared fluorescent detection system (LiCor, Lincoln, NE). The levels of each phosphoprotein were normalized for the amount of the corresponding total protein detected in the sample.

Statistical analysis—Data were analyzed using two-way ANOVA, in which treatment and cell type or treatment and genotype were the independent variables, followed by Bonferroni-Dunn post hoc test for specific comparisons.

RESULTS

L-DOPA increases DARPP-32 phosphorylation in striatonigral MSNs. Previous work carried out in 6-OHDA-lesioned mice showed that systemic administration of L-DOPA activates ERK and mTORC1 specifically in the D1R-expressing MSNs of the striatonigral pathway (13, 26, 27). Therefore, we started by examining whether the increase in DARPP-32 phosphorylation produced by L-DOPA and implicated in dyskinesia (6, 7) occurred in the same population of striatal neurons. To address this question D1-DARPP-32-Flag/D2-DARPP-32-Myc transgenic mice were lesioned unilaterally with 6-OHDA and injected for 10 days with 20 mg/kg of L-DOPA, a procedure which induces dyskinesia (7). LID was evaluated by scoring four types of AIMS immediately after the last injection of L-DOPA. At each time point, the scores for all types of AIMS were totaled and the average score was 28.9 ± 2.3, with a median value of 31. This value was used to divide the mice into moderately dyskinetic (total AIMS score below the median
value of 31) and severely dyskinetic (total AIMs score ranging above the median value of 31). The following day, the animals were injected with L-DOPA to induce DARPP-32 phosphorylation and killed 30 min later. It should be noted that chronic L-DOPA by itself does not affect protein phosphorylation in 6-OHDA-lesioned mice. Thus, when mice were killed 24 hrs following the last of 10 daily injections of L-DOPA we did not find any alterations in the levels of P-Thr34 DARPP-32 (65 ± 10% vs. 100 ± 47% in control unlesioned mice treated with saline), P-Ser845-GluA1 (61 ± 9% vs. 100 ± 26% in control mice), P-Thr202/Tyr204-ERK2 (106 ± 6% vs. 100 ± 3% in control mice), or P-Ser235/236-rpS6 (104 ± 4% vs. 100 ± 3% in control mice).

Lesion with 6-OHDA per se did not affect phosphorylation of DARPP-32 at Thr34 (Fig. 1A and B). Similarly, 6-OHDA did not produce any significant change in the phosphorylation of DARPP-32 at Thr75, a site implicated in the modulation of PKA activity (Fig. 1C and D). When L-DOPA was administered to 6-OHDA-lesioned D1-DARPP-32-Flag/D2-DARP-32-Myc mice, a large increase in the phosphorylation of DARPP-32 was detected in DARPP-32 immunoprecipitated from the D1R-expressing striatonigral MSNs (p < 0.001 versus unlesioned control mice; two-way ANOVA, followed by Bonferroni-Dunn test) (Fig. 1B). In contrast, no change in Thr34 phosphorylation was observed in DARPP-32 immunoprecipitated from D2R-expressing MSNs (Fig. 1B). We did not find any change in Thr75 phosphorylation in response to L-DOPA (Fig. 1A and B). Notably, the levels of phospho-Thr34 DARPP-32 immunoprecipitated from the D1R-expressing MSNs of mice with severe AIMs were significantly higher than those detected in mice with moderate AIMs (Fig. 1A and B). Thus, cell-type specific analysis of DARPP-32 phosphorylation demonstrates that LID is accompanied by increased phosphorylation of DARPP-32 at Thr34 and that this change occurs specifically in the striatonigral MSNs of the direct pathway.

Lack of DARPP-32 in striatonigral, but not in striatopallidal, MSNs inhibits L-DOPA-induced ERK and mTORC1 signaling. The studies performed in D1-DARPP-32-Flag/D2-DARPP-32-Myc transgenic mice revealed that the increase in DARPP-32 phosphorylation implicated in LID occurred in the same neurons (i.e. the striatonigral MSNs) where activation of ERK had been previously demonstrated (26,27). In order to determine the involvement of DARPP-32 in L-DOPA-induced ERK phosphorylation, we employed D32^{F/F}D1RCre^{+} mice and D32^{F/F}D2RCre^{+} mice, in which DARPP-32 is conditionally deleted in striatonigral and striatopallidal MSNs, respectively (9). In agreement with previous work (9), Western blot analysis revealed a reduction of DARPP-32 by 73.5 ± 0.5% in D32^{F/F}D1RCre^{+} mice versus D32^{F/F}Cre^{+} mice and by 48.7 ± 3.5% in D32^{F/F}D2RCre^{+} mice versus D32^{F/F}D2RCre^{+} mice, reflecting loss of DARPP-32 protein from a sub-population of MSNs in each mouse line. D32^{F/F}D1RCre^{+} mice, D32^{F/F}D2RCre^{+} mice and D32^{F/F}Cre^{+} littermates were lesioned unilaterally with 6-OHDA, treated for 10 days with 10 mg/kg of L-DOPA and killed 30 min after the last injection. In a previously published study, we have shown that the resultant dyskinetic response is strongly attenuated in D32^{F/F}D1RCre^{+} mice, but not in D32^{F/F}D2RCre^{+} mice (9). Administration of L-DOPA did not affect ERK phosphorylation in the intact striata (data not shown). This result is in line with several previous studies showing that L-DOPA, or dopaminergic agonists, do not affect ERK phosphorylation in the normal, unlesioned striatum (7,16,27,28). In contrast, following lesion with 6-OHDA, L-DOPA increased phospho-ERK in control, D32^{F/F}Cre^{+} mice (Fig. 2A). This effect was reduced in D32^{F/F}D1RCre^{+} mice, which lack DARPP-32 specifically in striatonigral MSNs, but preserved in D32^{F/F}D2RCre^{+} mice, which lack DARPP-32 in striatopallidal MSNs (Fig. 2A). We also examined the state of phosphorylation of the Lys14-acetylated form of histone H3, a downstream target of ERK (7,29). L-DOPA produced a large increase in phospho-Ser10-acyetyl-Lys14-histone H3 in the striata of 6-OHDA-lesioned D32^{F/F}D2RCre^{+} mice (Fig. 2B). This effect was attenuated in D32^{F/F}D1RCre^{+} mice, whereas it was maintained in D32^{F/F}D2RCre^{+} mice (Fig. 2B).

Increased ERK signaling has been implicated in the activation of mTORC1 in striatonigral neurons (13). Therefore, we examined whether DARPP-32, by virtue of its ability to regulate ERK, was also implicated in
the L-DOPA-induced phosphorylation of rpS6, a downstream target of mTORC1 (30). We found that L-DOPA induced phosphorylation of rpS6 at two sites, Ser235/236 and Ser240/244, was reduced in the striata of 6-OHDA-lesioned D32^{f/f} D1RCre mice, which lack DARPP-32 specifically in striatonigral MSNs (Fig. 2D and E). In contrast, genetic inactivation of DARPP-32 in the D2R-expressing MSNs of the striatopallidal pathway did not alter the ability of L-DOPA to increase phosphorylation of S6 at Ser235/236 and Ser240/244 (Fig. 2D and E).

In the same experiments we also examined the ability of L-DOPA to induce PKA-mediated phosphorylation of the GluA1 subunit of the glutamate AMPA receptor, which is increased in association with dyskinesia (7). We found that this effect of L-DOPA was also attenuated in D32^{f/f} D1RCre mice, but not in D32^{f/f} D2RCre mice (Fig. 2C).

PKA-dependent phosphorylation of DARPP-32 at Thr34 is required for LID and for the concomitant increase in ERK and mTORC1 signaling. The analysis of the changes of DARPP-32 phosphorylation in D1-DARPP-32-Flag/D2-DARPP-32-Myc transgenic mice indicate that LID is accompanied by increased Thr34 phosphorylation. In order to establish unequivocally the importance of this phosphorylation site in dyskinesia and to determine its involvement in the regulation of ERK and mTORC1 signaling, we made use of Thr34Ala DARPP-32 knock-in mice. Thr34Ala mutant mice and wild-type littermates were lesioned with 6-OHDA and injected for 10 days with L-DOPA (10 mg/kg). LID was determined on day 10 by assessing the severity of AIMS (cf. Materials and Methods). We found that AIMS were significantly lower in the Thr34Ala mutant mice as compared to wild-type mice (Fig. 3A). The following day, the animals were treated with L-DOPA and killed 30 min later. In wild-type mice, administration of L-DOPA in combination with 6-OHDA lesion resulted in a large increase in the state of phosphorylation of ERK, Lys14-acetylated histone H3, rpS6 and GluA1 (Fig. 4). These responses were all reduced in the striata of Thr34Ala mutant mice (Fig. 4).

**DISCUSSION**

In this study, we utilized various transgenic mouse lines to demonstrate the existence of a link between PKA-mediated phosphorylation of DARPP-32 and the concomitant activation of ERK and mTORC1 signaling produced in the dorsal striatum by L-DOPA. We also demonstrate that this regulation occurs exclusively in a well-defined sub-group of MSNs, corresponding to the D1R-expressing striatonigral neurons of the direct pathway.

Previous work suggested that the ability of L-DOPA to activate PKA and DARPP-32 signaling in D1R-expressing MSNs was implicated in LID (7,16). It was also found that, in addition to the activation of this canonical dopaminergic cascade, administration of L-DOPA increased the phosphorylation/activation of ERK (7,10-12) and that this effect represented an additional mechanism contributing to the development of dyskinesia (7).

The present results demonstrate that activation of ERK/mTORC1 requires intact PKA/DARPP-32 signaling and that PKA-mediated phosphorylation of DARPP-32 is required for the full expression of LID. First, we show that the increase in phospho-Thr34-DARPP-32 occurs in the same neuronal population, i.e. the striatonigral MSNs, in which abnormal ERK and mTORC1 regulation has been observed (26,27,13). Second, we show that cell-type specific inactivation of DARPP-32 in striatonigral MSNs strongly reduces the ability of L-DOPA to increase the phosphorylation of ERK and its downstream target, histone H3. Third, we provide evidence that a similar reduction in L-DOPA-mediated ERK and histone H3 phosphorylation occurs following a mutation of DARPP-32 that prevents its activation by PKA. Finally we demonstrate that abolishment of PKA-mediated phosphorylation of DARPP-32 reduces dyskinetic behavior.

The involvement of DARPP-32 in the regulation of ERK exerted by L-DOPA is in line with the observation that the increase in ERK phosphorylation produced by dopaminergic drugs, such as cocaine and amphetamine, is lost in DARPP-32 deficient mice and in Thr34Ala mutant mice (31). PKA-mediated phosphorylation at Thr34 converts DARPP-32 into an inhibitor of protein phosphatase-1 (32). Inhibition of protein phosphatase-1 has been proposed to modify
the state of phosphorylation of protein kinases and protein phosphatases which control the state of phosphorylation of ERK, such as the mitogen activated protein kinase/ERK kinase and the striatal-enriched protein phosphatase (31). The present data, indicating that mutation of Thr34 on DARPP-32 decreases the ability of L-DOPA to promote ERK signaling, are compatible with the existence of a similar mechanism of regulation in the dorsal striatum. Thus, sensitized D1R transmission, developed in response to dopamine depletion, would potentiate the effect of L-DOPA on PKA and DARPP-32, leading to the concomitant phosphorylation/activation of ERK. This idea is supported by evidence indicating that inhibition of PKA prevents the ability of L-DOPA to increase ERK phosphorylation in a rat model of PD (33). However, using a similar experimental model, it has been recently reported that a large reduction in L-DOPA-induced activation of PKA and DARPP-32 does not affect ERK phosphorylation (34). These results suggest that the ability of L-DOPA to promote ERK signaling depends on basal, rather than stimulated, PKA activity and DARPP-32 phosphorylation. Further studies will be necessary to clarify this issue.

The involvement of DARPP-32 in the development of LID has been previously shown using DARPP-32 knock out mice (7), as well as D32f/fD1RCre+ and D32f/fD2RCre+ mice (16). In these studies it was found that dyskinesia was significantly reduced following systemic or cell-specific inactivation of DARPP-32. The present results, showing that LID is reduced in Thr34Ala mutant mice, support this idea and indicate that the involvement of DARPP-32 in dyskinesia depends on PKA-mediated phosphorylation at Thr34.

One important finding of this study is that phosphorylation of DARPP-32 is required for the L-DOPA-induced activation of mTORC1 recently described in the striatonigral MSNs of dopamine-depleted mice (13). Enhanced mTORC1 signaling, which leads to the phosphorylation of rpS6 at multiple sites (30) and to increased 5’-cap-dependent initiation of mRNA translation (14), has been implicated in dyskinesia (13). The involvement of phospho-Thr34-DARPP-32 in the control of striatal mTORC1 may be a direct consequence of the ability of PKA/DARPP-32 to promote ERK signaling. Indeed, activation of ERK is thought to promote the stimulation of the small heterotrimeric GTP-binding protein Rheb, which activates mTORC1 (35,36). In support of this view, it has been shown that pharmacological inhibition of ERK prevents the ability of D1R agonist SKF81297 to increase mTORC1-dependent phosphorylation of rpS6 in the dopamine-depleted striatum (13).

In conclusion, we show that the activation of three distinct signaling pathways, previously implicated in the development of LID, occurs along the same intracellular cascade. This cascade is induced specifically at the level of the striatonigral MSNs, which express D1Rs, and includes the activation of PKA/DARPP-32, ERK and mTORC1. The finding that DARPP-32 is involved not only in the control of ERK, but also in mTORC1, highlights the key role played by this phosphoprotein in striatal signaling and provides information for the development of efficacious and selective interventions to be used in the treatment of the motor side effects produced by antiparkinsonian drugs.

REFERENCES


FOOTNOTES

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Abbreviations: PD, Parkinson’s disease (PD); MSNs, medium spiny neurons; D1R, dopamine D1 receptor; D2R, dopamine D2 receptor; DARPP-32, dopamine- and cAMP-regulated phosphoprotein of 32 kDa; PKA, cAMP-dependent protein kinase; ERK, extracellular signal regulated kinases 1 and 2; mTORC1, mammalian target of rapamycin complex 1; LID, L-DOPA-induced dyskinesia (LID); 6-OHDA, 6-hydroxydopamine; AIMs, abnormal involuntary movements; rpS6, ribosomal protein S6.
**FIGURE LEGENDS**

Fig. 1. LID is accompanied by increased levels of phospho-Thr34-DARPP-32 in D1R-expressing MSNs. *D1-DARPP-32-Flag/D2-DARPP-32-Myc* transgenic mice were lesioned unilaterally with 6-OHDA and treated for 10 days with 20 mg/kg of L-DOPA. AIMs were determined immediately after the last injection of L-DOPA. The following day, mice were treated with vehicle or L-DOPA and killed 30 min later. Levels of phospho-Thr34-DARPP-32 (A) and phospho-Thr75-DARPP-32 (C) were determined in the striata of control mice (unlesioned, treated with vehicle) and mice lesioned with 6-OHDA and treated chronically with vehicle or L-DOPA. This latter group was divided into low dyskinetic (L Dys) and high dyskinetic (H Dys) (cf. Results). Data are expressed as percent of control (n=4-12). **p<0.01 vs. control; one-way ANOVA followed by Bonferroni-Dunn test. DARPP-32 was immunoprecipitated from D1R- and D2R-expressing MSNs using anti-Flag and anti-Myc antibodies, respectively, and the levels of phospho-Thr34-DARPP-32 (B) and phospho-Thr75-DARPP-32 (D) were determined by western blotting. Data are expressed as percent of control (n=4-12). ***p<0.001 vs. control, †††p<0.001 versus L Dys group; one-way ANOVA followed by Bonferroni-Dunn test.

Fig. 2. Selective deletion of DARPP-32 in striatonigral MSNs decreases L-DOPA-induced phosphorylation of ERK, histone H3, GluA1 and rpS6. *DARPP-32<sup>F/F</sup>D1RCre<sup>+</sup> mice, DARPP-32<sup>F/F</sup> mice and DARPP-32<sup>F/F</sup>D2RCre<sup>+</sup> littermates were lesioned unilaterally with 6-OHDA, treated for 10 days with 10 mg/kg of L-DOPA and killed 30 min after the last injection. Top rows show representative autoradiograms obtained using antibodies against total or phosphorylated ERK (A), histone H3 (B), GluA1 (C), rpS6 (D and E). Bottom rows are summary of results showing means ± SEM (n=6). *p<0.05, **p<0.01 and ***p<0.001 vs. unlesioned (UL) *DARPP-32<sup>F/F</sup>Cre<sup>+</sup> and DARPP-32<sup>F/F</sup>D2RCre<sup>+</sup> mice treated with L-DOPA; †p<0.05, ††p<0.01 and †††p<0.001 vs. 6-OHDA lesioned DARPP-32<sup>F/F</sup>Cre<sup>+</sup> mice treated with L-DOPA (L); two-way ANOVA, followed by Bonferroni-Dunn test. A significant interaction between treatment and genotype was found in DARPP-32<sup>F/F</sup>D1RCre<sup>+</sup> mice (F<sub>(1, 20)</sub>= 8.9, p<0.01 for phospho-ERK 1, F<sub>(1, 20)</sub>= 11.47, p<0.01 for phospho-ERK2; F<sub>(1, 22)</sub>= 8.32, p<0.01 for phospho-Ser10-acetyl-Lys14-histone H3; F<sub>(1, 20)</sub>= 7.44, p<0.05 for phospho-Ser845-GluA1; F<sub>(1, 20)</sub>= 7.64, p<0.05 for phospho-Ser235/236-rpS6; F<sub>(1, 20)</sub>= 7.74, p<0.05 for phospho-Ser240/244-rpS6). A significant effect of the treatment was found in DARPP-32<sup>F/F</sup>D2RCre<sup>+</sup> mice [F<sub>(1, 20)</sub>= 26.64, p<0.001 for phospho-ERK 1, F<sub>(1, 20)</sub>= 30.95, p<0.001 for phospho-ERK2; F<sub>(1, 22)</sub>= 36.98, p<0.001 for phospho-Ser10-acetyl-Lys14-histone H3; F<sub>(1, 20)</sub>= 35.44, p<0.001 for phospho-Ser845-GluA1; F<sub>(1, 20)</sub>= 80.3, p<0.001 for phospho-Ser235/236-rpS6; F<sub>(1, 20)</sub>= 44.41, p<0.001 for phospho-Ser240/244-rpS6].

Fig. 3. Mutation of Thr34 on DARPP-32 decreases LID. Wild type (WT) and T34A mutant mice were lesioned unilaterally with 6-OHDA and treated for 10 days with 10 mg/kg of L-DOPA. AIMs were determined immediately after the last injection. (A) Time profile of the sum of locomotive, axial, limb and orolingual AIMs scored every 20 min over a period of 120 min after the last drug administration. (B) Sum of total AIMs scored during all observation periods. *p<0.01 versus WT, two-way ANOVA followed by Bonferroni-Dunn test. A significant effect of the genotype was found (p<0.05, F<sub>(1, 20)</sub>= 10.8).

Fig. 4. Mutation of Thr34 on DARPP-32 decreases L-DOPA-induced phosphorylation of ERK, histone H3, GluA1 and rpS6. Thr34Ala mutant mice and wild-type littermates were lesioned with 6-OHDA, treated for 11 days with 10 mg/kg of L-DOPA and killed 30 min after the last injection. Top rows show representative autoradiograms obtained using antibodies against total or phosphorylated ERK2 (A), GluA1 (B), histone H3 (C), rpS6 (D and E). Data are represented as means ± SEM (n=7-10). *p<0.05, **p<0.01 and ***p<0.001 vs. unlesioned mice treated with L-DOPA (open bars); †p<0.05, ††p<0.01 and †††p<0.001 vs. 6-OHDA-lesioned mice treated with L-DOPA; ‡ p < 0.05 vs. unlesioned mice treated with L-DOPA; two-way ANOVA, followed by Bonferroni-Dunn test. A significant interaction was found between treatment and genotype [F<sub>(1, 26)</sub>= 5.75, p<0.05 for phospho-ERK2; F<sub>(1, 28)</sub>= 6.82, p<0.05 for GluA1; F<sub>(1, 20)</sub>= 20.12, p<0.001 for phospho-Ser10-acetyl-Lys14-histone H3; F<sub>(1, 28)</sub>= 19.74, p<0.001 for phospho-Ser235/236-rpS6; F<sub>(1, 28)</sub>= 4.26, p<0.05 for phospho-Ser240/244-rpS6].
Fig. 3

A

B

ALMs score/observation period

Time (min)

0 20 40 60 80 100 120

24

20

16

12

8

4

0

WT

T34A

Total ALMs score

30

25

20

15

10

5

0

**
Dopamine- and cAMP-regulated phosphoprotein of 32 KDa (DARPP-32)-dependent activation of ERK and mammalian target of rapamycin complex 1 (TORC1) signaling in experimental parkinsonism
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