Rhes influences striatal cAMP/PKA-dependent signaling and synaptic plasticity in a gender-sensitive fashion

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Mechanisms of gender-specific synaptic plasticity in the striatum, a brain region that controls motor, cognitive and psychiatric functions, remain unclear. Here we report that Rhes, a GTPase enriched in medium spiny neurons (MSNs) of striatum, alters the striatal cAMP/PKA signaling cascade in a gender-specific manner. While Rhes knockout (KO) male mice, compared to wild-type (WT) mice, had a significant basal increase of cAMP/PKA signaling pathway, the Rhes KO females exhibited a much stronger response of this pathway, selectively under the conditions of dopamine/adenosine-related drug challenge. Corticostriatal LTP defects are exclusively found in A2AR/D2R-expressing MSNs of KO females, compared to KO males, an effect that is abolished by PKA inhibitors but not by the removal of circulating estrogens. This suggests that the synaptic alterations found in KO females could be triggered by an aberrant A2AR/cAMP/PKA activity, but not due to estrogen-mediated effect. Consistent with increased cAMP signaling, D1R-mediated motor stimulation, haloperidol-induced catalepsy and caffeine-evoked hyper-activity are robustly enhanced in Rhes KO females compared to mutant males. Thus Rhes, a thyroid hormone-target gene, plays a relevant role in gender-specific synaptic and behavioral responses.

The Ras homolog enriched in striatum (Rhes) is a small GTP-binding protein discovered by subtractive hybridization1 and highly expressed throughout the dorsal striatum and nucleus accumbens of rodent brain2–4. Rhes transcription is regulated by thyroid hormones during development4–7 and by dopamine (DA) in the adult rat brain8. Early studies in cell lines3,9 indicated that Rhes, most likely by binding to Gαi, reduces G-protein-coupled receptor (GPCR)-mediated accumulation of cAMP10. Accordingly, Rhes knockout (KO) mice have

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increased striatal cAMP/PKA activity. In addition to its influence on cAMP accumulation, Rhes acts as a selective striatal E3-ligase of mutant huntingtin, the summolation of which promotes neurotoxicity. These in vitro observations together with in vivo findings are consistent with the hypothesis that Rhes-mediated functions are common striatal targets for selective neurodegeneration in basal ganglia disorders.

Consistently, it has been found that Rhes mediates AKT-mediated signaling, participates in intracellular iron homeostasis and, most notably, modulates mammalian target of rapamycin complex 1 (mTORC1). This critical pathway is associated, among other processes, with L-DOPA-induced dyskinesia (LID) in animal models of Parkinson disease. These findings support the hypothesis that putative anti-Rhes drugs may attenuate the neurological manifestations of Huntington disease and LID.

Despite its growing role in striatal pathologies, the function of Rhes GTPase in striatal physiology remains less understood.

Previous studies indicated that lack of Rhes resulted in sex-sensitive behavioral phenotypes in mutants, but the mechanisms are unknown. Here we dissected the mechanisms of the gender-specific differences associated to Rhes deletion, focusing on striatal cAMP/PKA-dependent signaling and tested the differential motor responses elicited by DA and adenosine agonist/antagonists. Overall, our study indicates that Rhes orchestrates gender-sensitive alterations in striatal signaling, synaptic plasticity, and behavioral responses.

**Results**

**Rhes mRNA expression in GABAergic medium spiny neurons of mouse and human striatum.** To identify the striatal neurons that express Rhes, we performed double in situ hybridization (ISH) on coronal sections along the rostro-caudal extent of the striatum of wild-type (WT) mouse brains (Fig. 1A–C). As a marker of GABAergic medium spiny neurons (MSNs) we used protein phosphatase 1 regulatory subunit B (PPP1R1B), also known as Darpp32, a bifunctional signal transduction molecule, whose phosphorylation and function is regulated by dopaminergic and adenosine receptor stimulation.

As expected, Rhes expression was observed in virtually all Darpp32-positive striatal neurons, corresponding to the projecting MSNs (Fig. 1D–G, arrows). Then, we performed double ISH using RASD2 and PPP1R1B riboprobes, respectively Rhes and Darpp32 human orthologs, to analyze mRNA distribution of this GTPase in the MSNs of human striatum (Fig. 1H,I). Consistent with the results in mouse brain, we found that RASD2 mRNA was highly enriched in the putamen nucleus and expressed in virtually all the PPP1R1B-positive projecting MSNs (Fig. 1I, arrows).

**Lack of Rhes unmasks a gender-dependent modulation of striatal cAMP/PKA signaling.** In order to analyze whether the lack of Rhes modulates CAMP/PKA signaling in a gender-specific manner, we evaluated the phosphorylation state of a selective PKA substrate, the glutamate AMPA receptor subunit at Ser845 residue (p-Ser845-GluA1), in both male and female mutants versus their matched controls, under basal and challenged conditions. Administration of SKF 81297, a DA D1R full agonist, increased p-Ser845-GluA1 levels in males of both genotypes (Fig. 2A, left panel; two-way ANOVA, treatment effect: F(1,32) = 7.027, p = 0.0092, Fisher’s post-hoc). Similarly, p-Ser845-GluA1 level in mutants (expressed as fold-induction) was similar to WT animals (Fig. 2A, right panel). This observation suggests that lack of Rhes in males mainly perturbs striatal tonic CAMP/PKA activity, as indicated by the significantly higher basal p-Ser845-GluA1 levels in KO striata (p = 0.0001). Although SKF 81297 induced an apparently higher striatal p-Ser845-GluA1 level in mutants (p = 0.0145, Fisher’s post-hoc), its relative increase compared to baseline (expressed as fold-induction) was similar to WT animals (Fig. 2A, right panel). This observation suggests that lack of Rhes in females mainly perturbs striatal phasic CAMP/PKA activity, as indicated by the significantly higher basal p-Ser845-GluA1 levels in KO striata (p = 0.0001). As seen with D1R agonist, despite the higher A2AR/cAMP/PKA-dependent signaling in both WT and KO males (Fig. 2C, left panel; two-way ANOVA, treatment effect: F(1,29) = 24.466, p < 0.0001), its relative increase compared to baseline (expressed as fold-induction) was similar to WT animals (Fig. 2C, right panel; p = 0.0483, Fisher’s post-hoc). In contrast to what observed in male mutants, lack of Rhes in females did not result in altered basal striatal p-Ser845-GluA1 content (Fig. 2B,D, left panel). However, in KO females both SKF 81297 and haloperidol administration induced higher p-Ser845-GluA1 levels, when compared to WT treated controls (two-way ANOVA, genotype × treatment interaction, SKF 81297: F(1,29) = 7.027, p = 0.0129; haloperidol: F(1,32) = 22.383, p < 0.0001). As a consequence, in KO females, the comparable basal activity along with an exaggerated phasic D1R- and A2AR-dependent cAMP/PKA signaling resulted in a greater fold-induction of this pathway (Fig. 2B,D, right panels; SKF 81297, p = 0.0175; haloperidol, p = 0.0004).

Overall, our findings indicate that lack of Rhes in females, rather than influencing the tonic striatal CAMP/PKA activity as seen in male mutants, selectively increases the phasic activation of this pathway.

**Lack of Rhes does not affect intrinsic membrane properties and corticostriatal long-term depression.** Based on the strict control of glutamatergic inputs on MSNs activity, we first analyzed the striatal AMPAR and NMDAR subunit expression in WT and KO animals. Western blotting analysis indicated that protein levels of both NMDAR (GluN1, GluN2A and GluN2B) and AMPAR (GluA1 and GluA2A/3) subunits are not altered in knockouts of both genders (Fig. 3A,B; p > 0.05, per each protein). Next, we examined the effects of Rhes deletion in modulating synaptic properties and plasticity of striatal...
Figure 1. *Rhes* mRNA expression in both mouse and human striatal MSNs. (A–C) *In situ* hybridization (ISH) of coronal sections showing *Rhes* expression along the rostro-caudal extent of the mouse striatum. (D–G) Double ISH of mouse caudate/putamen (CPu) using specific antisense riboprobes for *Rhes* (D,F) and *Darpp32* (E,G), showing that *Rhes* is expressed in *Darpp32*-positive striatal MSNs. Boxed regions in (D,E) are shown at higher magnification in both brightfield (F) and fluorescence (G), and neurons strongly expressing both *Rhes* and *Darpp32* are indicated by arrows. (H,I), Double ISH analysis of *RASD2* and *PPP1R1B* expression on cryosections from post-mortem human brain. (H) Nissl stained specimen highlighting the neuroanatomical region of the putamen and globus pallidus nuclei. (I) Brightfield photomicrograph image of the boxed region in (H) showing co-localizing *RASD2* (purple) and *PPP1R1B* (silver grains) mRNA expression in MSNs (arrows). Abbreviations: gp, globus pallidus nucleus; pu, putamen nucleus. Scale bar: 2.5 mm (A–C,H); 75 μm (D,E,I); 15 μm (F,G).
MSNs in mutant mice. Based on intracellular and patch-clamp recordings in corticostriatal slices, we failed to find any significant difference between genotypes in the intrinsic membrane properties of MSNs (Fig. 3C,D, upper panels; Student’s t-test, p > 0.05). Moreover, analysis of activity-dependent synaptic plasticity in striatal MSNs revealed that the delivery of a high frequency stimulation (HFS) to corticostriatal fibers in presence of Mg²⁺ (1.2 M) was able to induce comparable long-term depression (LTD) in the MSNs of both genotypes, without any evident effect of gender (Fig. 3C,D, bottom panels; two way ANOVA, gender effect, male: F(1,136) = 0.71, p > 0.05; female: F(1,136) = 0.41, p > 0.05).

Figure 2. Effect of dopaminergic drugs on striatal cAMP/PKA activity in Rhes male and female mutant mice. (A,B) Striatal p-Ser845-GluA1protein levels, in male (A) and female (B) WT and KO mice, following 5 mg/kg SKF 81297 (WT, n = 9 males, n = 7 females; KO, n = 9 per gender) or vehicle (Veh) treatment (WT, n = 9/gender; KO, n = 9 males, n = 8 females) with relative fold-induction. (C,D) Striatal p-Ser845-GluA1 levels, of male (C) and female (D) WT and KO mice after 0.5 mg/kg haloperidol (Hal) treatment (WT, n = 8 males, n = 9 females; KO, n = 9/gender) or Veh (WT, n = 8 males, n = 9 females; KO, n = 9/gender), with relative fold-induction. Top panels show representative blots comparing the different genotypes and treatments. *p < 0.05, **p < 0.01, ***p < 0.0001, compared with Veh-treated group within genotype. #p < 0.05, ##p < 0.01, compared with WT group. All data are expressed as mean ± SEM. Genotypes are as indicated.
Figure 3. Intrinsic membrane properties and long-term depression in Rhes mutant mice. (A,B) Western blotting analysis on the striatal glutamate AMPAR and NMDAR subunit levels in male (A) (n = 6/genotype per GluA1, GluA2/3, GluN1, GluN2B; n = 5 WT, 6 KO per GluN2A), and female (B) (n = 6/genotype per GluA1, GluA2/3, GluN2A; n = 4 WT, 5 KO per GluN1; n = 6 WT, 5 KO per GluN2B) mice. The top panels show representative blots comparing the different genotypes, for each protein detected. All data are expressed as mean ± SEM. Genotypes are as indicated. (C,D) Current-voltage graphs (left) and representative traces (right) obtained after applying hyperpolarizing and depolarizing steps of current to MSNs recorded from WT (n = 6 males, n = 7 females) and KO (n = 10 per gender) mice. Time-courses (top) and example traces (bottom) of the post-synaptic responses measured in MSNs recorded from slices of male KO (n = 6) and WT (n = 5) mice (left panel) and female KO (n = 4) and WT (n = 5) mice (right panel), showing no difference in high-frequency stimulation (HFS)-induced LTD (Student's t-test, pre- vs. 30 min post-HFS, **p < 0.01).
Lack of Rhes selectively affects corticostriatal long-term potentiation in A2AR/D2R-positive MSNs in a gender-specific manner. Then, we investigated the corticostriatal long-term potentiation (LTP) in male and female mutants. Delivery of HFS to corticostriatal fibers in Mg²⁺-free medium, a condition that favors NMDA receptor activation, was able to induce LTP with comparable amplitude and time course in WT and KO male mice (Fig. 4A; two-way ANOVA, genotype effect, $F_{(1,189)} = 0.42, p > 0.05$). Conversely, LTP amplitude was significantly altered in a subpopulation of MSNs recorded from mutant females. Specifically, although HFS-induced LTP was similar to controls in 12 out of 27 neurons, a complete loss of potentiation was found in the remaining 15 MSNs recorded (Fig. 4B; two-way ANOVA, genotype effect, $F_{(2,493)} = 30.29, p < 0.001; \text{time \times genotype interaction, } F_{(1,272)} = 8.71, p < 0.001$).

To explain the synaptic plasticity defects found in KO females, we hypothesized that an exaggerated phasic cAMP/PKA activation associated to HFS might occlude LTP expression in MSNs that did not show any apparent activity-dependent plasticity. To test this idea, low frequency stimulation (LFS) protocol was applied to corticostriatal fibers during baseline in striatal slices of WT and KO females. While in WT this stimulation protocol did not induce any change in EPSP amplitude, as expected, in KO females LFS decreased the EPSP amplitude in 56% of the recorded MSNs. This abnormal synaptic depotentiation seen in female mutants was similar to that observed in control animals after the induction of a HFS-dependent LTD$^{29}$. Further, in KO females, the remaining MSNs (44%) responded with a short-term potentiation. (Fig. 4C; two-way ANOVA, genotype effect, $F_{(2,234)} = 19.51, p < 0.001; \text{time \times genotype interaction, } F_{(1,192)} = 7.33, p < 0.001$).

Finally, given the evidence of a heterogeneous population of striatal MSNs (SP/D1R-expressing direct and the A2AR/D2R-expressing indirect pathways$^{30}$), we performed post hoc immunohistochemical analysis to assess the identity of MSNs subtype in KO females not showing any apparent activity-dependent LTD. We found that the MSNs that did not respond to HFS with a physiological magnitude of LTP in KO females were adenosine A2AR-positive cells that belong to the D2R-expressing striato-pallidal pathway (Fig. 4D–F). Moreover, to determine whether circulating ovarian hormones could explain the gender-specific dichotomous response to HFS, we analyzed LTP in the MSNs of ovariectomized (OVX) KO females. Similar to what observed in intact mutants, HFS of corticostriatal fibers induced LTP only in 6 out of 12 MSNs recorded from OVX KO females (Fig. 4G; two-way ANOVA, time × genotype interaction, $F_{(17,170)} = 4.90, p < 0.001$). In addition, no significant difference was found in the striatal expression of Estrogen Receptor subunits (ERα and ERβ) between genotypes (Fig. 4H-I; $p > 0.05$ per each protein).

Altered striatal LTP in Rhes mutant females in A2AR/D2R-positive neurons is associated to exaggerated cAMP/PKA signaling. In light of our biochemical studies, we postulated that the excessive phasic cAMP/PKA signaling induced by HFS in female mutants might prevent the normal LTP expression in A2AR/D2R-positive MSNs. To verify this idea, we explored in WT and KO female striatal slices the consequences of changes in cAMP/PKA activity and A2AR activation on corticostriatal LTP expression. As expected, in WT slices patch-clamp recordings of MSNs showed that intrapipette application of PKA inhibitor Rp-cAMPS (100 μM) was able to block LTP$^{31}$, in all the recorded cells (Fig. 5A), confirming that an intact cAMP/PKA activation is required for striatal HFS-induced LTP. In contrast, based on the hypothesis that an abnormal high A2AR/cAMP/PKA activity prevented HFS-induced LTP in Rhes KO females, we demonstrated that intrapipette application of a PKA inhibitor, at a dose that blocked LTP in all the MSNs of WT, was able to normalize synaptic plasticity in SP-negative MSNs (data not shown) of mutant females (Fig. 5B; two-way ANOVA, group effect, $F_{(1,170)} = 40.53, p < 0.001$). We next evaluated whether non-physiological A2AR activation in WT could affect the induction of LTP. To this aim, bath application of A2AR agonist (CGS21680, 3 μM) 10 minutes prior to HFS protocol revealed the existence of two different subsets of MSNs, one normally responding to HFS with a LTP amplitude similar to control, and the other being defective in this form of synaptic plasticity (Fig. 5C; two-way ANOVA, group effect, $F_{(2,119)} = 12.83, p < 0.01$). Therefore, the dichotomous MSNs behaviors under A2AR agonist strengthened the idea that, differently from SP/D1R-positive MSNs, non-physiological increase of cAMP/PKA signaling in A2AR/D2R-bearing neurons prevents its activity-dependent plasticity, as seen in Rhes KO females.

To further assess the role of A2AR activation on striatal LTP expression, we recorded the effects of the A2AR antagonist ZM241385 (1 μM) in WT slices. Intracellular and patch-clamp recordings showed that following blockade of A2AR, LTP was induced only in SP/D1R-positive MSNs (Fig. 5D). In contrast, the remaining A2AR/D2R-positive MSNs, which did not respond with LTP, paradoxically showed a LTD (Fig. 5D; two-way ANOVA, group effect, $F_{(1,272)} = 26.78, p < 0.001$). Similar to what observed in WT slices, application of ZM241385 in KO females did not affect a regular HFS-induced LTP in SP/D1R-positive MSNs (Fig. 5F). In contrast, ZM241385 in A2AR/D2R-positive MSNs (Fig. 5G) of KO females failed to induce LTD (Fig. 5E; two-way ANOVA, group effect, $F_{(1,408)} = 28.27, p < 0.001$), unlike reported in controls.

Rhes mutants display altered motor responses to dopamine and adenosine drugs in a gender-specific fashion. Here we investigated the gender effect of Rhes deletion on behavioral responses induced by DA and adenosine drugs. At all doses tested, the DA D1R agonist, SKF 81297, similarly enhanced horizontal motor activity in males of both genotypes (Fig. 6A; three-way ANOVA with repeated measures, treatment effect: $1.25 \text{mg/kg, } F_{(1,413)} = 3.3633, p < 0.0001; 2.5 \text{mg/kg, } F_{(1,413)} = 86.567,$
Figure 4. Long-term potentiation in MSNs of Rhes mutant mice and morphological characterization.
(A) Time-course plots of EPSP amplitude (left) and pairs of traces (right) demonstrate the presence of HFS-induced LTP in MSNs recorded from slices of male WT (n = 5) and KO (n = 7) mice (Student’s t-test, pre- vs. 30 min post-HFS, **p < 0.01). (B) Time course graphs of the EPSC amplitude (left) and pairs of traces (right) show the induction (n = 12) or the complete loss (n = 15) of LTP in MSNs from female KO mice compared with age-matched WT animals (n = 5) (Student’s t-test at 30 min post-HFS, WT vs. KO with LTP, p > 0.05, WT vs. KO with no LTP, ***p < 0.001). (C) In KO females application of a low-frequency stimulation (LFS) induced a decrease of EPSP amplitude in 5 of 9 MSNs recorded and a mild potentiation in the remaining 4 MSNs, whereas the same protocol induced no change of synaptic activity in all the MSNs (n = 7) recorded from WT females (Bonferroni post-hoc: WT vs. KO with LTP, *p < 0.05; **p < 0.01; ***p < 0.001; Student’s t-test at 30 min post-HFS: WT vs. KO with a depression of EPSP, ***p < 0.001).
(D–F) Confocal laser scanning microscopy (CLSM) images of double-labeled immunofluorescence for biocytin and adenosine receptor (A2AR). Biocytin immunolabeling is visualized in red streptavidin-Cy3 immunofluorescence (D) and A2AR is visualized in green Cy2 immunofluorescence (E). The colocalization of A2AR and biocytin is showed in merged panel (F) as yellow fluorescence. (G) HFS protocol applied in the MSNs of ovariectomized (OVX) KO females induced LTP in 6 of 12 MSNs recorded (Student’s t-test at 30 min post-HFS: WT vs. KO OVX with no LTP, *p < 0.05).
(H,I) Striatal ERα and ERβ, determined by Western blotting, in WT and KO male (n = 6 /genotype per ERα; n = 6 WT, 5 KO per EPβ) (H) and female (n = 5 WT, 6 KO per ERα; n = 6 /genotype per ERβ) (I) mice. The top panels show representative blots comparing the different genotypes. All data are expressed as mean ± SEM. Genotypes are as indicated.
Figure 5. Altered striatal LTP in Rhes KO females associated to exaggerated A2AR/cAMP/PKA signalling. (A) The LTP of the MSNs recorded with intrapipette application of PKA inhibitor Rp-cAMPs was blocked in all the recorded cells (n = 4) (Student’s t-test at 30 min post-HFS: WT in Rp-cAMPs, pre vs post HFS, p > 0.05). (B) LTP induction in MSNs of KO females recorded with intrapipette application of Rp-cAMPs (Student’s t-test at 30 min post-HFS: KO with no LTP, n = 7, vs. KO with LTP, n = 5, ***p < 0.001). (C) Bath application of A2AR agonist CGS21680 in slices from WT females revealed the existence of two different subsets of MSNs showing LTP and no LTP (Student’s t-test at 30 min post-HFS: WT with no LTP, n = 5, vs. WT with LTP, n = 4, ***p < 0.001). (D) A2AR antagonist ZM241385 bath applied in slices of WT females induces LTP in n = 12 cells and LTD in the remaining n = 6 neurons (Student’s t-test at 30 min post-HFS: WT with LTD vs. WT with LTP, ***p < 0.001). (E) ZM241385 application in KO neurons induced a regular LTP (n = 16) in SP/D1R-positive MSNs and no change (n = 10) in A2A/D2R-positive MSNs (Student’s t-test at 30 min post-HFS: KO with no LTP vs. KO with LTP, **p < 0.01). (F,G) Confocal laser-scanning microscopy images of triple-labeled immunofluorescence for biocytin, SP and D2 receptor. SP is visualized in green, D2R is visualized in light blue-Cy5 fluorescence and biocytin immunolabeling is visualized in streptavidin-Cy3 fluorescence, the merged image is shown on the bottom right of the panels.
p < 0.0001). Likewise, SKF 81297 treatment in females also induced hyperlocomotion in both genotypes at either dose tested (Fig. 6B; 1.25 mg/kg, F(1,330) = 10.644, p = 0.0018; 2.5 mg/kg, F(1,390) = 14.095, p = 0.0003). However, female mutants showed a significant higher hyperactivity at 2.5 mg/kg drug dose (genotype effect, F(1,390) = 5.456, p = 0.0221), compared to control treated animals.

Next, we analyzed the effect of the DA D2R antagonist haloperidol, on catalepsy. We found no major treatment effect in males following 0.1 mg/kg drug administration in both genotypes (Fig. 6C). On the other hand, both 0.25 and 0.5 mg/kg haloperidol administration induced a higher cataleptic response in male KO mice, compared to WT (Fig. 6C; two-way ANOVA, genotype effect, 0.25 mg/kg, males,
Lack of Rhes does not affect hippocampal long-term potentiation. In addition to striatum, Rhes mRNA expression is also seen in both murine and human hippocampus (Fig. 8A), but its role in hippocampal LTP is unclear. Rhes was expressed in the granule cell layer of the dentate gyrus (DG) and in the pyramidal cell layer of Ammon's horn (Fig. 8A,b–d). Autoradiographic images using radioactive \(^{35}\)S ISH also showed expression of RASD2 in the hippocampal formation of human brain samples (Fig. 8A,f). At high magnification, we observed autoradiographic grains on the granule cells of the DG and in the hilus (Fig. 8A,g,h). Pyramidal neurons of fields CA3, CA2, and CA1 of Ammon's horn and the subiculum were also specifically labeled (Fig. 8A,i–l).

To determine whether alterations of striatal LTP in KO mice were area-specific or a widespread phenomenon associated to lack of Rhes, we measured basal synaptic transmission, short-term plasticity and LTP at CA1 hippocampal synapses in Rhes mutants (Fig. 8B,C). Overall, differences in the input-output relationship (\(p > 0.05\) for all stimulation intensities) and in the paired-pulse ratio (\(p > 0.05\) at all inter-pulse intervals) were not observed either in male (Fig. 8B) or female (Fig. 8B) mutants. In addition, theta-burst stimulation (TBS) (Fig. 8B,C), delivered to Schaffer collateral fibers, produced comparable levels of CA1-LTP in male (fEPSP amplitude at 60 min post-TBS: WT 155 \(\pm\) 10%; KO 146\% \(\pm\) 7%, left panel) and female KO mice when compared to gender-matched controls (fEPSP amplitude at 60 min post-TBS: WT 146\% \(\pm\) 9%; KO 143\% \(\pm\) 12%; right panel). Indeed, two-way ANOVA confirmed no significant effect of gender (\(F_{1,16} = 0.146, p > 0.05\)) or genotype (\(F_{1,16} = 0.125, p > 0.05\)). Thus Rhes predominantly affects synaptic plasticity associated with striatum.

**Discussion**

In the present work, we studied how lack of Rhes influences the striatal cAMP/PKA signaling-related synaptic and behavioral responses under both basal and challenged conditions, in a gender-dependent manner. We analyzed a Rhes KO mouse line\(^{23}\), in which the PGK-neo cassette had been removed in order to avoid any potential interference on gene transcriptional regulation\(^{35–37}\). In accordance with previous data\(^2\), we report that lack of Rhes does not affect basal motor abilities in both male and female mutant mice (data not shown). Similarly, electrophysiological data revealed that Rhes mutant mice display no difference in intrinsic and synaptic basal properties of striatal MSNs, when compared to controls. Moreover we did not find any apparent alteration in striatal LTD between genotypes in both genders.

Conversely, lack of Rhes was found to be critical in the expression of striatal MSNs LTP in a gender-sensitive manner. Consistently, while WT and KO males display similar LTP, in females only in a subset of D1R-positive MSNs we recorded comparable LTP amplitudes between genotypes. Based on the knowledge that LTP is a form of corticostriatal plasticity that requires intact cAMP/PKA signaling\(^{31}\), the inability of certain MSNs to potentiate EPSP amplitude in response to HFS apparently contradicts
the greater striatal cAMP-PKA signaling found in Rhes mutant females. To explain such discrepancy, we hypothesized that the apparent loss of LTP in mutants might be generated by an exaggerated increase in PKA activity, which in turn leads to constitutive saturated LTP. Consistent with this idea, in mutant females, LFS protocol of corticostriatal fibers during baseline transmission causes significant "depotentiation" or mild potentiation in a ratio that reflects the proportion of MSNs responding to HFS with a lack of LTP or with a LTP expression similar to control, respectively.

Several striatal responses in animals and humans are affected by gender through the action of various factors, including circulating estrogens. Therefore, we evaluated the potential involvement of this these sexual hormones in the expression of altered synaptic plasticity seen in female mutants. Overall, electrophysiological experiments excluded a direct implication of estrogens in determining the LTP defects as we confirmed in OVX mutants the existence of two independent striatal subpopulations of MSNs that either express LTP or undergo a saturated state. In addition, we also ruled out an altered estrogen receptor expression level as the cause for defective LTP, as striatal levels of ERα and ERβ are comparable between genotypes.

In order to identify the receptor specificity of the MSNs fraction in which LTP is saturated in KO females, we performed staining experiments after patch-clamp recordings. Remarkably, in Rhes mutants we found that striatal MSNs that did not express LTP are selectively A2AR/D2R-positive. Thus, in line with the notion that non-physiological phasic A2AR/cAMP/PKA responses found in female mutants

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Figure 7. Medial-to-lateral expression gradient of Rhes, A2AR and D2R in the mouse CPu.
(A–H) Representative pseudocolor autoradiographs from coronal section of mouse brain showing medial-to-lateral expression gradient of (A,E) Rhes mRNA, (B,F) A2AR and (C,G) D2R. (D,H) D1R mRNA appears uniformly distributed. (I) Histogram showing quantification of Rhes, A2AR, D2R and D1R mRNA expression levels obtained after densitometric quantification of autoradiograms. Differences are expressed as percentage of variation compared with the signal measured in the dorsomedial part of the CPu. (J–M), A2AR levels were determined by radioactive ISH (n = 3/genotype for both genders; J,L) and in total striatal extracts by Western blot, in naïve male (n = 6/genotype; K) and female (n = 6/genotype; M) mice. The top panels show representative blots comparing the different genotypes. All data are expressed as mean ± SEM. *p < 0.05, **p < 0.01. Scale bar: 750μm
Figure 8. Expression pattern of Rhes and RASD2 mRNA in mouse and human hippocampus.
(A)(a–d), In situ hybridization experiment on a coronal section of mouse brain showing Rhes expression in hippocampus. Boxed regions in (a) are shown at higher magnification in (b; dentate gyrus, DG, and hilus), (c; CA3) and (d; CA1). (e–l), Nissl staining (e) and radioactive (35S) ISH analysis (f–l) showing expression of RASD2 in the hippocampus in a representative section from post-mortem human brain. RASD2 autoradiographic signal is present in the dentate gyrus (g; DG), hilus (h), CA3 (i), CA2 (j), CA1 (k), and subiculum (l). Scale bar: 1.3 mm (a); 60 μm (b–d, g–l); 1.8 mm (e,f). (B,C) Input/output relationship (left panel) and paired pulse (PP) ratios (center panel) obtained from CA1 area in WT and KO male (B) and female (C) mice. Each group represents n = 6 mice. Superimposed pooled data (right panel) representing normalized changes in the field potential amplitude (fEPSP) (± SEM) induced by TBS between male WT (n = 6) and KO (n = 4) mice (B, right panel) and female WT (n = 5) and KO (n = 5) mice (C, right panel).
may occlude activity-dependent LTP, we revealed that PKA inhibitor strikingly restores LTP in A2AR/ D2R-positive KO MSNs. Moreover, we found that A2AR agonist application unmasks in WT striatum the existence of two different subsets of MSNs, one not affected by such manipulation that responded to HFS with a regular LTP and another population of neurons, sensitive to A2AR stimulation, that fails to express a physiological LTP magnitude.

Altogether, this set of pharmacological experiments on slices from WT female striata reproduces the findings obtained in Rhes mutants and, in turn, demonstrates that A2AR activity drives the direction of corticostriatal plasticity in striato-pallidal pathway by regulating the magnitude of cAMP/PKA activity. In accordance to the consistent role of A2ARs in determining a physiological LTP in striato-pallidal neurons\(^1\), pharmacological blockade of these receptors with ZM241385 prevents HFS-induced LTP, triggering an unexpected LTD induction in WT slices. An explanation for this switch in synaptic plasticity likely relies on the ability of A2AR inhibition to unmask a higher D2R-mediated signaling which, in turn, favors LTD expression\(^2\). In contrast to what observed in WT, we report that ZM241385 administration in KO females prevents any form of synaptic plasticity in A2AR/D2R-expressing MSNs. This suggest that lack of LTD response in Rhes KO striata can be explained by previous results of our group demonstrating that D2R activity, as measured by GTP gamma assay, is slightly compromised in Rhes KO\(^2\).

It is well established that A2AR transmission exerts a pivotal role in modulating the magnitude of haloperidol-induced catalepsy and caffeine-triggered hyperactivity\(^3\). Therefore, in line with an exacerbated A2AR-mediated signaling in Rhes KO, we found significant higher haloperidol-induced catalepsy and caffeine-induced motor simulation responses, when compared to gender-matched controls. Although more pronounced in females, enhanced behavioral responses to haloperidol and caffeine also occurs in Rhes KO males when compared to their sex-matched treated controls. These in vivo observations suggest that, differently to striatal LTD and DA D1R-dependent motor stimulation, lack of Rhes affects striato-pallidal-dependent behaviors with a less prominent gender-sensitive influence. Although we reported that Rhes mRNA medial-lateral expression gradient strictly resembles that of A2AR and D2R, we exclude that abnormal striato-pallidal responses seen in mutants are associated to different A2AR expression since striatal mRNA and protein levels were comparable between genotypes. We also show that Rhes mRNA is expressed not only in striatal Darp32-positive MSNs of rodents and humans, but also in various hippocampal regions. In keeping with this evidence, we demonstrated that hippocampal LTP induction in the CA1 region is unaffected in both genders of mutants, thus strengthening the sex-specific role of Rhes in modulating synaptic plasticity selectively in striatal MSNs. Furthermore, cognitive functions analyzed by object recognition and fear conditioning tasks are comparable between KO and WT mice of both genders (data not shown).

Extending our previous work\(^2\), the present study indicates that lack of Rhes differently impacts on striatal cAMP/PKA signaling in a gender specific fashion, causing a significant basal increase in males while generating a higher response in females, selectively under challenge conditions. In light of a gender-sensitive regulation of cAMP/PKA activity and consistent with a crucial involvement of this signaling pathway in modulating D1R-, D2R- and A2AR-mediated behavioral responses, we explain the sex-related differences in response to SKF81297, haloperidol and caffeine found in KO mice, when compared to their matched-controls.

Similarly, we argue that the striking gender-related difference in LTP expression found in A2AR/ D2R-positive MSNs of mutants depends on the different magnitude of cAMP/PKA signaling triggered by HFS protocol in male and females. Therefore, we propose that yet unknown factors, other than estrogens, should be directly or indirectly involved in the regulation of striatal cAMP/PKA activity underlying the gender specific LTP expression found in Rhes KO.

Altogether our findings, indicating that Rhes plays a crucial gender-specific role in orchestrating striatal-dependent behaviors, increase the functional complexity of this GTP binding protein function.

In addition, considering the enhanced haloperidol-induced catalepsy and caffeine-triggered motor hyperactivity in mutants, we propose that Rhes could emerge as a novel thyroid hormone-target gene to investigate the molecular and cellular determinants associated to the use of these substances in humans.

**Methods**

**Animals.** In order to prevent the possibility that the presence of the PGK-neo cassette may interfere with normal transcriptional regulation within the Rhes locu\(s\)\(^5\)\(^-\)\(^7\)\(^,\)\(^3\)\(^4\), we crossed Rhes\(^{loxP-neo}\) offspring\(^2\) with a CMV-Cre deleter strain in a B6D2 genetic background\(^4\). The validation of Cre-mediated loxP-flanked neo cassette excision was assessed by PCR analysis using the following primers: Rhes-EGFP\(_{\text{forward}}\) 5'-CATGTCCTGAGTTCGTGA-3', and Rhes-Ex2\(_{\text{reverse}}\): 5'- ACCACATGCGGATGAGTTCT-3'. The amplification product was ~400 bp. Mice obtained were then genotyped by PCR using the following primers: Rhes\(_{\text{forward}}\): 5'-TTTAGAATTTCACTGTGT-3' and Rhes\(_{\text{Ex2\_reverse}}\): 5'-ACCACATGCGGAGTTCT-3', for Rhes\(^{+/\text{loxP-neo}}\) (WT) allele (420 bp); Rhes\(_{\text{EGFP\_forward}}\): 5'-CATGTCCTGAGTTCGTGA-3' and Rhes\(_{\text{Ex2\_reverse}}\): 5'- ACCACATGCGGAGTTCT-3' for Rhes\(^{+/-}\) (KO) allele (370 bp). KO mice were viable, fertile and did not show alteration in body weight and basal motor activity (data not shown). Ten- to twelve-week-old male and female WT and KO mice, derived from mating of heterozygous animals (Rhes\(^{+/-}\)), back-crossed to F11 generation to C57BL/6 strain, were used in this study. Mice were housed in groups in standard cages (29 × 17.5 × 12.5 cm) at constant temperature (22°C ± 1°C) and maintained on a 12/12 h light/
In long-term depression (LTD), the intensity was increased to suprathreshold levels. To induce depotentiation, low frequency stimulation (LFS) protocol (2 Hz, 10 min of total duration) was used as previously shown.

**Results.**

Whole-cell patch-clamp recordings were performed as previously shown. Quantitative data on EPSP modifications induced by HFS are expressed as a percentage of the controls, the latter representing the mean of responses recorded during a stable period (15–20 min) before the

**Supplementary Materials.**

Human tissue was obtained from the Brain and Tissue Bank for Developmental Disorders at the University of Maryland, Baltimore. Samples for cryosectioning were dissected in the region of the putamen nucleus and hippocampus of brains from one male (16 years old, 16 h post-mortem) and one female (35 years old, 6 h postmortem) who had no history of primary neurological or psychiatric disorders and processed as previously described. Human samples.

**In situ hybridization (ISH).** ISH analysis was performed according to protocols previously described. Digoxigenin (DIG) (Rhes, 0.5 Kb; RASD2, 2.8 Kb), fluorescein- (Darpp32, 0.6 Kb), or 35S-labelled (PPP1R1B, 1.5 Kb; RASD2, 2.8 Kb; Rhes, 0.5 Kb; A2AR, 1.2 Kb; Drd2, 1.0 Kb Drd1, 0.3 Kb) antisense riboprobes were used. Detailed sequences used for each probe are the following: RASD2, nucleotides 178–2971 (NM_014310.3); Darpp32, nucleotides 144–685 (NM_144828.1); PPP1R1B, nucleotides 23–1500 (NM_181505.3); A2AR, nucleotides 521–1736 (NM_009630.3); D1R49; D2R49; Rhes25. For double ISH on mouse tissue, sections were hybridized simultaneously with DIG- and fluorescein-labelled riboprobes. A two-step chromogenic reaction using NBT/BCIP and HNPP/Fast Red Fluorescent Detection Set (Roche) was performed to visualize the DIG- and fluorescein-labelled riboprobes, respectively. Specimens were counterstained with DAPI. For radioactive ISH, hybridized sections were washed with Biowax MR X-ray films (Kodak) for two to four days. For each probe used, sections from different genotypes and gender were processed in parallel in order to minimize the experimental variability. Double-ISH on human tissue was performed combining DIG-labelled and 35S-radiolabelled riboprobes. After chromogenic reaction using NBT/BCIP for the detection of the DIG-labelled riboprobe, slides were dipped in Kodak NTB emulsion for autoradiography as previously described. Sections were examined using brightfield and darkfield light microscopy.

**Image analysis and quantification.** Quantification analyses were performed in blind and sample identity was not revealed until correlations were completed. For densitometric analyses, 15–20 sections per animal along the whole rostro-caudal extent of the striatum were used and images of autoradiography films resulting from radioactive ISH experiments were scanned at a resolution of 1200 dpi. For the analysis of Rhes, A2AR, D2R and D1R mRNA expression in the striatum of WT animals the Caudate-Putamen (CPu) was divided into four quadrants, dorso-medial (DM), dorso-lateral (DL), ventro-medial (VM) and ventro-lateral (VL), accordingly to Harrison and LaHoste, 2006. Optical Density (OD) for each gene analyzed was evaluated in the four areas of the striatum. Background OD value was determined in structures of the same section devoid of specific signal and subtracted for correction to obtain the Relative OD (ROD) value. Results were expressed as percentage increase/decrease of mRNA-expression in the DM for each probe. Data were analyzed by Student’s t-test. For the analysis of A2AR mRNA expression ROD was evaluated in whole striatum of WT and KO adult male and female mice. Results were expressed as percentage increase/decrease of A2AR mRNA-expression. Analysis was performed by using StatView software (version 5.0.1.0; SAS Institute).

**Electrophysiological recordings from medium spiny neurons (MSNs).** The preparation and maintenance of coronal corticostriatal slices have been previously described. Intracellular recordings with sharp electrodes. For the long-term potentiation (LTP) protocol, at the beginning of intracellular recordings, magnesium ions were omitted from the medium to increase the N-methyl-D-aspartate (NMDA)-mediated component of excitatory post-synaptic potential (EPSP). As conditioning high frequency stimulation (HFS) protocol to induce either long-term depression (LTD) or LTP we used three trains (3 seconds duration, 100 Hz, 20 seconds intertrain interval). During tetanic stimulation, the intensity was increased to suprathreshold levels. To induce depotentiation, low frequency stimulation (LFS) protocol (2 Hz, 10 min of total duration) was used as previously shown.

Whole-cell patch-clamp recordings. Patch-clamp recordings were performed as previously shown. Quantitative data on EPSP modifications induced by HFS are expressed as a percentage of the controls, the latter representing the mean of responses recorded during a stable period (15–20 min) before the
HFS. Current–voltage relationships were obtained by applying steps of current of 50 pA in both hyperpolarizing and depolarizing direction (from −600 to 400 pA). Statistical analysis was performed using Prism 4.0 (GraphPad). Student's t-test was used to compare data within group pre vs. 30 min post-HFS. Group comparisons were analyzed with two-way ANOVA. When a significant interaction was observed the data were further analyzed with Bonferroni post-hoc test.

Extracellular recordings in the hippocampus. Hippocampal slices were prepared as previously described. A single slice was placed on a nylon mesh, completely submerged in a small chamber (0.5 ml) and superfused with oxygenated artificial cerebrospinal fluid (ACSF) at a constant flow rate of 3 ml/min. A bipolar tungsten stimulating electrode was positioned in the CA1 stratum radiatum to stimulate the Schaffer collateral fibers, and extracellular field excitatory post-synaptic potentials (fEPSPs) were recorded with a glass microelectrode (2–3 MΩ) in the stratum radiatum. LTP was evoked by theta burst stimulation (TBS) protocol consisting of 4 trains of 5 pulses at a frequency of 100 Hz, with an inter-train interval of 200 ms. All data are presented as mean ± SEM and n indicates the number of slices. Statistical analysis was performed by two-way ANOVA with gender and genotype as between subject factors. Statistical significance was set at p < 0.05.

Tissue processing and double immunofluorescence of striatal MSNs. Tissue processing and double immunofluorescence were performed according to protocols previously described. See Supplementary Materials for detailed information.

Behavior. Drug-induced motor responses. Mice were randomly assigned to 1.25, 2.5 mg/kg SKF 81297 or vehicle (Veh)-treated groups, habituated to the experimental cage (35 × 25 × 30 cm) for 60 min, and then injected with drug or Veh and placed in the same cage, as previously described. The horizontal motor activity was evaluated during 10-min intervals over a 1 h-test session, through a computerized video tracking system (Videotrack, Viewpoint S.A., Champagne au Mont d'Or, France). The effect of SKF 81297 on locomotor activity (expressed in cm) was used as dependent variable and analyzed by three-way (genotype × treatment × time) ANOVA with repeated measures.

Caffeine-induced locomotor hyperactivity was performed according to a previously validated protocol. Briefly, WT and KO animals were randomly assigned to caffeine (7.5 mg/kg), or Veh-treated groups. The horizontal motor activity was evaluated during 10-min intervals over a 2 h-test session. Effect of caffeine on locomotor activity (expressed in cm) was used as dependent variable and analyzed by three-way (genotype × treatment × time) ANOVA with repeated measures.

Catalepsy. Catalepsy in mice was assessed according to a previous protocol. To evaluate the haloperidol-induced cataleptic response, 60 min before drug treatment WT and KO mice of both genders were placed one per cage for adapting to the new environment. Animals were randomly grouped to be treated, respectively, with 0.1, 0.25, 0.5 mg/kg haloperidol (Hal), or vehicle (Veh). Catalepsy was measured 30, 60 and 120 min after drug or vehicle injection, by gently placing the front limbs of the animals over a 4 cm-high horizontal bar. The intensity of catalepsy (expressed in s) was assessed by measuring the time each mouse remained with limbs completely immobile for a maximum of 300 s. Data were analyzed between genotypes, at each haloperidol dose tested, by two-way (time × genotype) ANOVA with repeated measures. Other independent groups of female mice were used to assess the effect of 8-(3-Chlorostyryl)-caffeine (CSC) in haloperidol-induced catalepsy, according to Chen et al. Mice were co-administered with Veh plus 0.5 mg/kg haloperidol or haloperidol plus 2.5 mg/kg CSC (administered 10 min before haloperidol injection). Catalepsy was evaluated 60 min after treatment and analyzed by two-way ANOVA, followed by Fisher’s post-hoc comparison.

Western blotting. Male and female mice of both genotypes were treated with 5 mg/kg SKF 81297, 0.5 mg/kg haloperidol (Hal) or Veh, and killed by decapitation 30 min later, according to a previous protocol. Normalized values of phospo-GluA1 at residue Ser845 levels were averaged and used as dependent variable. Data were analyzed between genotypes by two-way (treatment × genotype) ANOVA, followed by Fisher’s post-hoc comparison, when required. Striatal levels of glutamate NMDAR (GluN1, GluN2A, GluN2B), AMPAR (GluA1, GluA 2/3) subunits, as well as estrogen (α and 3) and adenosine (A2AR) receptors, were analyzed in naïve WT and KO animals of both genders. Normalized values of each target were used as dependent variable and analyzed between genotypes by Student’s t-test. For detailed information see Supplementary Materials.

All behavioral and biochemical values are expressed as mean ± SEM. Statistics analyses were based on parametric tests as the data were normally distributed. Normality was checked with Shapiro-Wilk test (p < 0.05).

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Author Contributions
FN., V.G., B.P. and A.U. designed research; FN., D.V., F.E. and A.D.M. performed behavioral and biochemical experiments; B.Pc., S.M. and G.M. performed ISH experiments under the supervision of M.P.; V.G., VP., M.M., G.S., C.S. performed electrophysiological research under R.N. and B.P. supervisions; C.G. performed the immunostaining experiments; F.N., D.V., V.G., M.P. and B.P. analyzed data and made figures; F.N., V.G., B.P. and A.U. wrote the paper.

Additional Information
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