

The HDAC inhibitor 4b ameliorates the disease phenotype and transcriptional abnormalities in Huntington's disease transgenic mice

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Transcriptional dysregulation has emerged as a core pathologic feature of Huntington's disease (HD), one of several triplet-repeat disorders characterized by movement deficits and cognitive dysfunction. Although the mechanisms contributing to the gene expression deficits remain unknown, therapeutic strategies have aimed to improve transcriptional output via modulation of chromatin structure. Recent studies have demonstrated therapeutic effects of commercially available histone deacetylase (HDAC) inhibitors in several HD models; however, the therapeutic value of these compounds is limited by their toxic effects. Here, beneficial effects of a novel pimelic diphenylamide HDAC inhibitor, HDACi 4b, in an HD mouse model are reported. Chronic oral administration of HDACi 4b, beginning after the onset of motor deficits, significantly improved motor performance, overall appearance, and body weight of symptomatic R6/2^{300Q} transgenic mice. These effects were associated with significant attenuation of gross brain-size decline and striatal atrophy. Microarray studies revealed that HDACi 4b treatment ameliorated, in part, alterations in gene expression caused by the presence of mutant huntingtin protein in the striatum, cortex, and cerebellum of R6/2^{300Q} transgenic mice. For selected genes, HDACi 4b treatment reversed histone H3 hypoacetylation observed in the presence of mutant huntingtin, in association with correction of mRNA expression levels. These findings suggest that HDACi 4b, and possibly related HDAC inhibitors, may offer clinical benefit for HD patients and provide a novel set of potential biomarkers for clinical assessment.

chromatin | neurodegenerative | epigenetic | therapeutic | trinucleotide

Huntington's disease (HD) is an autosomal-dominant neurodegenerative disorder caused by a CAG repeat expansion within the coding region of the HD gene (*Htt*), resulting in a mutant protein (htt) with a lengthened polyglutamine tract (1). Mutant htt protein has been shown to disrupt transcription by multiple mechanisms, but it is unclear which are most important to pathology (2–4). By interacting with specific transcription factors, htt can alter the expression of clusters of genes controlled by those factors. For example, several genes driven by Sp1, which has been shown to interact with htt (5, 6), show decreased mRNA expression in human HD and in mouse models of HD (7). Alternatively, htt may have more global effects on transcription by disrupting core transcriptional machinery (8, 9) or by altering posttranslational modifications of histones, resulting in condensed chromatin structure (10–13). Understanding the basis for transcriptional dysregulation is important for choosing appropriate drug-discovery strategies.

Manifestations of transcriptional dysregulation are evident from several gene-profiling studies, which have revealed alterations in the expression of large numbers of genes in the brains of different HD mouse models and in human subjects with HD (7, 14–16). Many of the expression changes in mouse models are observed in early stages of illness before the onset of symptoms, suggesting that gene expression alterations may be pathogenic.

Because of the extent of gene expression alterations in HD, most of which are decreases in expression, agents that improve transcriptional activity on a broad scale may represent an important therapeutic approach for HD. In addition, the evidence for chromatin-based transcriptional repression in HD suggests that inhibitors of histone deacetylase (HDAC) enzymes, which act in concert with histone acetyltransferase enzymes to modulate gene transcription, may represent useful treatments for HD.

Previous studies have examined the potential therapeutic effects of the HDAC inhibitors suberoylanilide hydroxamic acid (SAHA) (17), sodium butyrate (18), and phenylbutyrate (19) in HD mouse models. Despite showing promise in ameliorating the phenotype in different HD mouse models, the utilities of these compounds, as well as their analogues, are limited by toxicity. Toxicity studies of various HDAC inhibitors, including SAHA, have demonstrated widespread effects in human cancer cells *in vitro*, including activation of proapoptotic and inhibition of antiapoptotic pathways, stimulation of cell differentiation, and induction of growth arrest (20–22). These features have led to the approval of SAHA for use in human cancer clinical trials (22); however, such properties might be expected to exacerbate symptoms in neurodegenerative disorders, such as HD.

We have developed a class of benzamide-type HDAC inhibitors that show promising results in Friedreich's ataxia disease models (23, 24). These compounds are structurally related to the well-known HDAC inhibitor SAHA but are not hydroxamic acids and, unlike SAHA, were found to increase expression of the frataxin gene in lymphocytes from Friedreich's ataxia patients (23). From a panel of these novel HDAC inhibitors, we have further characterized the therapeutic potential in HD mice for one selected compound, HDACi 4b. Our cell culture findings indicate that HDACi 4b shows a low toxicity profile, whereas our *in vivo* studies on R6/2 transgenic mice, which is the most widely used model for preclinical trials (25, 26), demonstrate therapeutic efficacy in preventing motor deficits and neurodegenerative processes. We further report that HDACi 4b treatment ameliorates gene expression abnormalities detected by microarray analysis in these mice.

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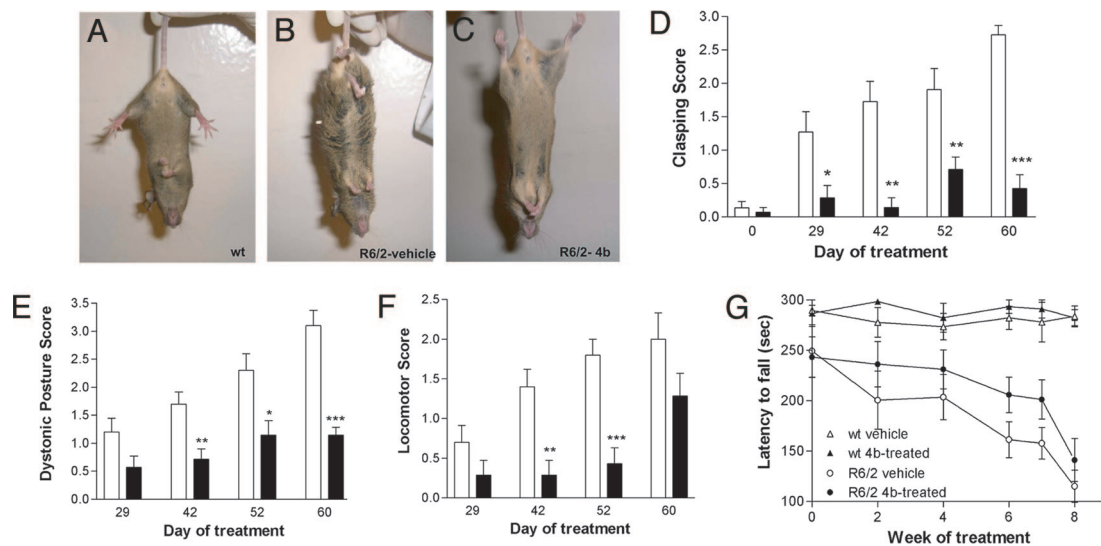


Fig. 1. Behavioral phenotypes of vehicle- and HDACi 4b-treated mice. Hindlimb clasping phenotype of WT mice (A) and R6/2^{300Q} transgenic mice treated with vehicle (B) or HDACi 4b (C). The differences between HDACi 4b-treated (filled bars) and vehicle-treated (open bars) R6/2^{300Q} transgenic mice are shown for the clasping phenotype (D), dystonic posture (E), and locomotor activity (F) over the time of drug treatment. Two-way ANOVAs were used to determine the effect of HDACi 4b treatment and treatment duration on each of these measurements. *, $P < 0.05$; **, $P < 0.001$; ***, $P < 0.0001$. (G) Rotarod performance of HDACi 4b-treated and vehicle-treated WT and R6/2^{300Q} transgenic mice throughout the treatment duration. Time-point 0 reflects baseline performance. Two-way ANOVA revealed significant differences between WT and R6/2^{300Q} transgenic mice with time ($P < 0.0001$), as well as a significant effect of drug treatment in R6/2^{300Q} transgenic mice ($P < 0.05$). Bars represent mean score \pm SEM ($n = 7$ to 8 per group).

Results

***In Vitro* Toxicity Profile of HDACi 4b.** We evaluated the cytotoxic effects of HDACi 4b treatment on cell cycle parameters in human lymphoblast cell cultures. Cells were treated with increasing concentrations of HDACi 4b (1–125 μ M) for 72 h and then assessed by FACS analysis of propidium iodide-stained nuclei. This analysis demonstrated no cell-cidal effects at concentrations < 50 μ M and only cell-static effects at concentrations > 20 μ M [supporting information (SI) Fig. S1]. No apoptotic effects of HDACi 4b were observed, except at concentrations > 50 μ M (Fig. S1), which are 10-fold higher than that previously reported for SAHA using similar cell types and methodologies (27). Importantly, at the highest concentration of 0.125 mM HDACi 4b, only 14% of the total cells gated were observed to be apoptotic (Fig. S1). Given an IC_{50} value of ≈ 1 μ M for HDACi 4b-mediated inhibition of HDAC activity (as measured in HeLa cell nuclear extracts), the concentrations imparting toxic effects are 20–50-fold higher.

HDACi 4b Improves Disease Phenotype in R6/2^{300Q} Transgenic Mice.

We first verified that HDACi 4b can modify the histone acetylation status in the CNS *in vivo*. A single injection of HDACi 4b (100 mg/kg, s.c.) in normal mice resulted in a nearly twofold increase in CNS histone H4 acetylation (Fig. S2). We next developed a formulation for improving water solubility of HDACi 4b by complexing with 2-hydroxypropyl- β -cyclodextrin (HOP- β -CD), similar to that reported previously (17); this allowed for chronic dosing of HDACi 4b in drinking water, corresponding to a dose of 150 mg/kg/day. R6/2^{300Q} transgenic mice and WT littermate controls were treated with HDACi 4b for 67 days beginning at 4 months of age, which is after the onset of motor dysfunction, as evidenced by rotarod deficits at baseline training (Fig. 1G). The effect of HDACi 4b administration on the R6/2^{300Q} phenotype was assessed by several measurements, including motor function, physical appearance, and body weight.

Motor function abnormalities were determined by measuring clasping phenotype, general locomotion, and rotarod perfor-

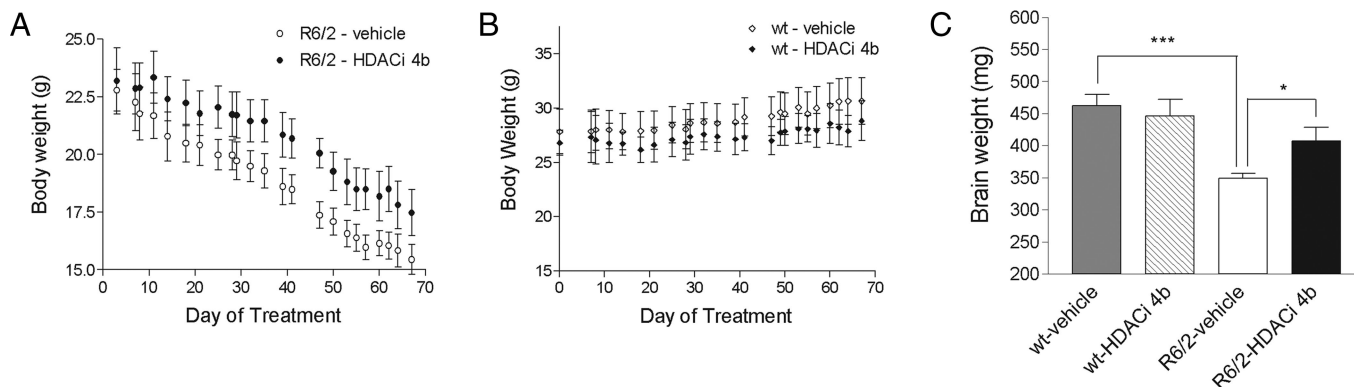


Fig. 2. Body weights over time of R6/2^{300Q} transgenic (A) and WT (B) mice throughout HDACi 4b treatment. Symbols represent mean body weight \pm SEM in vehicle-treated mice (open symbols) and HDACi 4b-treated mice (filled symbols). R6/2^{300Q} transgenic mice treated with HDACi 4b weighed significantly more than vehicle-treated R6/2^{300Q} mice as determined by two-way ANOVA ($P < 0.0001$). (C) Effect of HDACi 4b treatment on overall brain weights of WT and R6/2^{300Q} transgenic mice. Significant differences were determined by one-way ANOVA and Student's *t* test. ***, $P < 0.001$; *, $P < 0.05$.

mance. As expected, R6/2^{300Q} transgenic mice exhibited significant deficits in motor behavior with increasing treatment duration (i.e., age) (Fig. 1). However, significant prevention or amelioration of these deficits were observed with HDACi 4b treatment for the hindlimb clasping test [$F(1,80) = 64.07$; $P < 0.0001$], generalized locomotor behavior [$F(1,60) = 26.81$; $P < 0.0001$], and rotarod performance [$F(1,59) = 4.77$; $P = 0.032$] (Fig. 1). Most notably, as shown in Fig. 1A, HDACi 4b-treated mice show almost no clasping phenotype. We also assessed the effects of HDACi 4b on physical appearance, which can be quantified, in one sense, by monitoring changes in dystonic posture exhibited by R6/2^{300Q} transgenic mice. We found that HDACi 4b treatment significantly prevented the development of kyphosis in R6/2^{300Q} transgenic mice [two-way ANOVA: $F(1,60) = 43.76$; $P < 0.0001$] (Fig. 1E). We further found that HDACi 4b treatment resulted in improved coat appearance, gait stability, and overall motility in R6/2^{300Q} transgenic mice (Movie S1, Movie S2, and Movie S3).

The effects of HDACi 4b treatment on body weight on each genotype were determined by using two-way ANOVA. All R6/2^{300Q} transgenic mice showed a decrease in body weight with duration of treatment (i.e., age) [$F(22,228) = 9.63$; $P < 0.0001$]. However, when comparing the two groups of transgenic mice, we found a significant effect of HDACi 4b treatment to attenuate body-weight decline [$F(1,228) = 48.36$; $P < 0.0001$] (Fig. 2A). HDACi 4b also had a small but significant effect on body weight in WT mice [$F(1,256) = 9.24$; $P < 0.002$] (Fig. 2B).

HDACi 4b Shows Neuroprotective Effects. At the end of drug treatment, which was close to the lifespan of the mice (6 months of age), neuroprotective effects of HDACi 4b treatment were assessed. Overall, there was a 22.9% reduction in brain weight of vehicle-treated R6/2^{300Q} mice compared with WT littermates at age 6 months (356.4 ± 10.19 mg vs. 462 ± 18.32 mg for R6/2^{300Q} and WT mice, respectively, $P = 0.003$) (Fig. 2). However, brains from HDACi 4b-treated mice weighed significantly more when compared with those from vehicle-treated R6/2^{300Q} mice (407.3 ± 21.74 mg vs. 356.4 ± 10.19 mg for drug-treated and vehicle-treated R6/2^{300Q} mice, respectively, $P = 0.045$) (Fig. 2C). Furthermore, HDACi 4b treatment ameliorated the gross striatal atrophy and ventricular enlargement that was observed in the vehicle-treated R6/2^{300Q} mice (Fig. 3). Remarkably, brains from HDACi 4b-treated mice were indistinguishable from those from vehicle- or HDACi 4b-treated WT mice (Fig. 3). No apparent difference in the number of huntingtin-positive striatal nuclear aggregates, as determined by immunohistochemistry using the EM48 anti-huntingtin antibody, was observed between vehicle- and drug-treated R6/2^{300Q} mice (data not shown), consistent with previous studies using sodium butyrate, phenylbutyrate, or SAHA (17–19).

HDACi 4b Treatment Ameliorates Gene Expression Abnormalities in R6/2^{300Q} Transgenic Mice. To evaluate whether HDACi 4b treatment alters gene expression profiles in the CNS of WT and R6/2^{300Q} transgenic mice, we performed microarray analysis on striatum, cortex, and cerebellum of 5-month-old vehicle- and HDACi 4b-treated mice. Hierarchical clustering of the samples based on interarray Pearson correlation showed sample clustering according to brain region and genotype, as expected (Fig. S3). Consistent with results from previous microarray studies (16), R6/2^{300Q} transgenic mice showed a significant number of differentially expressed genes, with cortex and striatum showing an overrepresentation of down-regulated genes. The top expression differences observed in transgenic vs. WT mice at $P < 0.005$ ($n = 988$, 724, and 979 probes for striatum, cortex, and cerebellum, respectively) are listed in Dataset S1.

Because our R6/2 line carries a greater CAG repeat than the parent line, we assessed to what extent these mice recapitulate

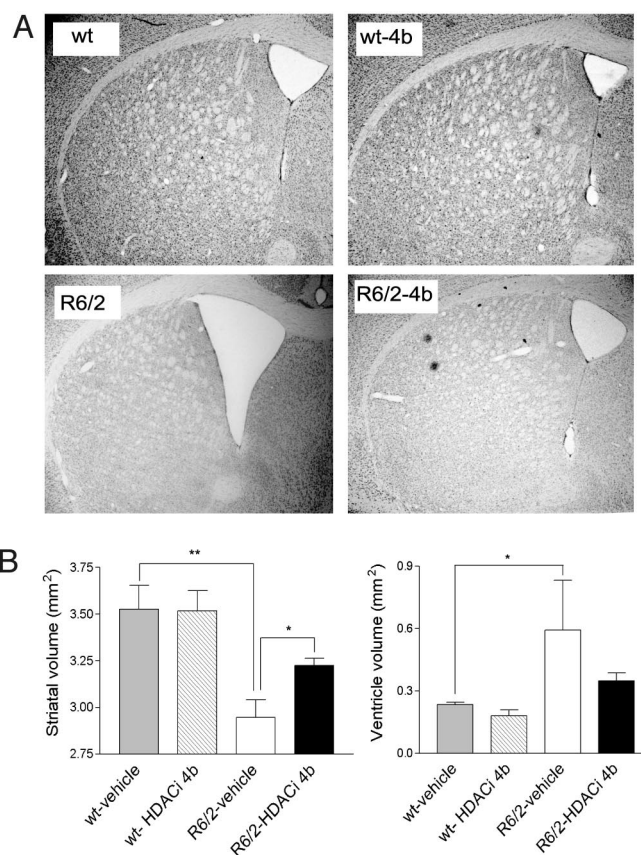


Fig. 3. Effects of HDACi 4b on brain gross morphology. Volumetric analysis of striatum and lateral ventricles was performed on mice at 6 months of age. (A) Representative images of brain at ≈ 0.62 bregma for the indicated conditions. (B) Quantification of striatal area and lateral ventricles. Bar graphs represent mean \pm SEM for WT mice and R6/2^{300Q} mice ($n = 6$ per group). Significant differences were determined by one-way ANOVA and Student's t test. **, $P < 0.001$; *, $P = 0.05$. The effect of drug treatment on ventricular volume did not reach significance.

the gene expression profiles detected in human HD by comparing gene transcriptome profiles from the striatum of R6/2^{300Q} transgenic mice with previously published microarray data from human HD caudate samples (15). Human orthologues of the top 250 differentially expressed mouse genes in the striatum (including both increases and decreases) were mapped and their expression differences screened in the human HD dataset provided by Hodges *et al.* (15). In our dataset, 77 of the top 142 down-regulated genes (54.2%) and 39 of the top 80 up-regulated genes (39%) showed concordance expression changes, defined as those showing a significant difference in the human dataset ($P < 0.05$) and the same direction of change in both mouse and human. A concordance coefficient was calculated, taking into account both concordant and discordant changes, as described previously (16) (SI Methods). The concordance coefficient for our dataset was 0.378 for down-regulated genes and 0.327 for all genes, which is consistent with previous studies reporting statistically significant concordance coefficients for other HD mouse models of 0.190 to 0.490 (with the related R6/1 and R6/2 lines equaling 0.350 and 0.405/0.490, respectively) (16). Hence, our findings indicate that R6/2^{300Q} transgenic mice exhibit expression profiles highly correlated with those observed in human HD.

HDACi 4b treatment caused a less severe effect on overall gene expression, compared with the dramatic effect caused by the presence of the mutant *Htt* gene. Of the top disease-induced gene expression abnormalities (at $P < 0.005$), $\approx 20\%$ showed a

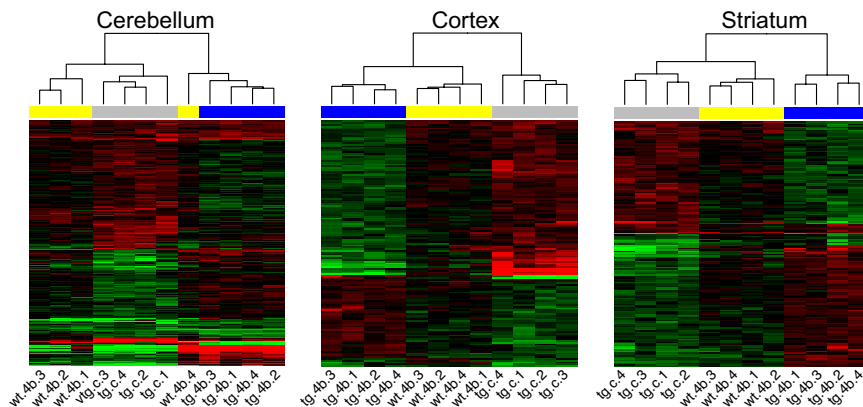


Fig. 4. Heatmaps depicting the probes that are differentially expressed in R6/2^{300Q} mice vs. WT ($P < 0.05$, gray bar) and in HDACi 4b-treated vs. untreated R6/2^{300Q} transgenic mice ($P < 0.05$, blue bar). The HDACi 4b effect on WT mice in these genes is marked by the yellow bars. Log₂-transformed ratios were subjected to two-dimensional hierarchical clustering. Each colored pixel represents an individual ratio from a single subject (four replicates per condition) compared with the average of controls. Relative decreases in gene expression are indicated by green and increases in expression by red. The vast majority of these genes show a trend toward normalization after HDACi 4b treatment. Complete normalization is reached for 100 (31%, cerebellum), 45 (33%, cortex), and 56 (33%, striatum) probes.

change toward the WT level after HDACi 4b administration (Fig. S4). To determine more inclusive lists of potentially therapeutically relevant expression changes, we identified the intersection between disease- and drug-induced expression changes at $P < 0.05$, which resulted in a list of 320, 167, and 137 genes for the cerebellum, striatum, and cortex, respectively. Depending on the region, 85%–94% of these genes showed a trend toward normalization, with one-third being completely normalized by HDACi 4b treatment (Fig. 4). These genes (provided in Dataset S2) could be considered as potential candidate biomarkers for treatment effectiveness. Interestingly, the expression of different sets of genes was altered by HDACi 4b treatment in WT compared with HD transgenic mice in each brain region, with only a <10% overlap of drug-induced differential gene expression between the genotypes (Fig. S5), indicating tissue-specific effects. HDACi 4b treatment did not have any effects on the expression of the endogenous *Htt* gene in R6/2^{300Q} transgenic mice, nor the human *HTT* transgene, as determined by real-time PCR analysis (data not shown).

Expression differences for a diverse group of selected genes

were validated by real-time PCR analysis. From the microarray data, the cerebellum showed the most robust treatment effect, which is interesting in light of studies suggesting that the cerebellum may play a more significant role in the symptomatology of HD than previously thought (28). Expression deficits for aquaporin 1 (*Aqp1*), claudin 1 (*Cldn1*), calmodulin-like 4 (*Calml4*), folate receptor 1 (*Folr1*), chloride intracellular channel 6 (*Clic6*), and ectonucleotide pyrophosphatase/phosphodiesterase 2 (*Enpp2*), detected in the cerebellum of 5-month-old R6/2^{300Q} transgenic mice compared with age-matched WT mice, were almost completely reversed by HDACi 4b treatment (Fig. 5A), validating the array. We next assessed the histone acetylation status of these down-regulated genes in the same sets of mice by using ChIP analysis in combination with real-time PCR. Compared with WT mice, hypoacetylation of histone H3 was detected specifically in association with these six down-regulated genes in R6/2^{300Q} transgenic mice (Fig. 5B). After treatment with HDACi 4b, significant increases in histone H3 acetylation was detected for *Cldn1*, *Calml4*, *Folr1*, and *Clic6* (Fig. 5B). Significant effects of HDACi 4b treatment on the expression of

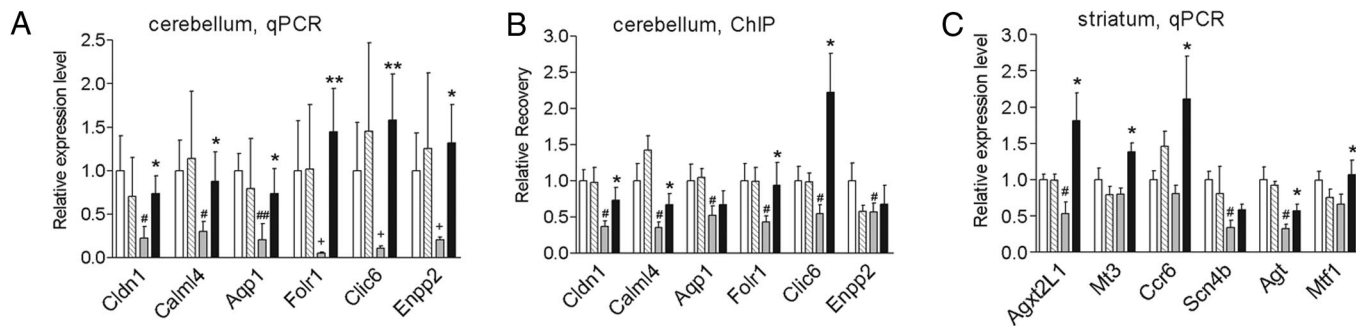


Fig. 5. HDAC inhibitor treatment reverses mRNA abnormalities and increases acetylated H3 in association with genes down-regulated in R6/2^{300Q} transgenic mice. Real-time PCR analysis was performed for the indicated genes on RNA samples from the cerebellum (A) or striatum (C) of vehicle- (white bars) or HDACi 4b-treated (striped bars) WT mice and vehicle- (gray bars) or HDACi 4b-treated (filled bars) R6/2^{300Q} transgenic mice. Data are depicted as fold change of the mean expression level \pm SEM ($n = 4$ mice per group). The relative abundance of each gene expression was normalized by using *Hprt*. (B) Chromatin immunoprecipitation was performed with anti-acetylated histone H3 (ACh3) and anti-histone H3 antibodies on the cerebellum of vehicle- or HDACi 4b-treated WT and R6/2^{300Q} transgenic mice (same key as in A and C). Real-time PCR was performed with primers for the promoter region of *Cldn1* and *Aqp1* and the transcribed region of *Calml4*, *Folr1*, *Clic6*, and *Enpp2*. Recovery was normalized for *Gapdh*, and data are shown as the ratio of the recovery for ACh3 and H3 and normalized for vehicle-treated WT mice \pm SEM. Student's *t* tests were used to determine significant differences in gene expression or recovery levels. The effect of drug treatment on expression of the sodium channel, type IV- β gene (*Scn4b*) in the striatum of HD mice did not reach significance. # denotes significantly different values between R6/2^{300Q} transgenic and WT mice at $P < 0.05$, with + denoting $P < 0.08$; * denotes significantly different values between HDACi 4b- and vehicle-treated R6/2^{300Q} mice at $P < 0.05$ and ** at $P < 0.01$.

alanine-glyoxylate aminotransferase 2-like 1 (*Agxt2l1*), metallothionein 3 (*Mt3*), chemokine receptor 6 (*Ccr6*), metal response element binding transcription factor 1 (*Mtf1*), and angiotensinogen (*Agt*) in the striatum of R6/2^{300Q} mice were also validated by real-time PCR analysis (Fig. 5C).

Discussion

In this study, we report beneficial effects of a novel benzamide-type HDAC inhibitor, HDACi 4b, on the disease phenotype of R6/2^{300Q} transgenic mice at the behavioral, physical, pathologic, and molecular levels. Importantly, we find very low toxic effects of HDACi 4b associated with these outcomes or at therapeutic doses *in vitro*. HDACi 4b-treated mice showed significantly improved movement and coordination and delayed weight loss over vehicle-treated animals, despite initiation of treatment after the onset of motor deficits. This finding is in contrast to previous studies investigating the effects of phenylbutyrate on N171–82Q transgenic mice, in which beneficial effects on these parameters, including rotarod, were not found with treatment beginning after symptom onset (19). HDACi 4b treatment also dramatically improved the physical appearance of, and prevented dystonic posture exhibited by, R6/2^{300Q} transgenic mice (see [Movie S1](#), [Movie S2](#), [Movie S3](#), and Fig. 1). Such effects have not been reported in response to other HDAC inhibitors previously tested in HD mouse models (17–19). Overall, these findings suggest that HDACi 4b treatment may slow the progression of symptoms in humans, who are often diagnosed after symptom presentation.

At the molecular level, we find that HDACi 4b treatment results in both increases and decreases in gene expression of a limited number of genes, rather than having global effects on transcription. This effect is likely because of specificity of HDACi 4b for particular HDAC isoforms, of which at least 11 are expressed in the rodent brain (29). These findings are also consistent with previous studies indicating that commercially available HDAC inhibitors can have both positive and negative effects on gene expression (reviewed in ref. 22). Because HDAC inhibition is expected to result primarily in increased gene transcription, the down-regulation of certain genes may represent a secondary effect.

The therapeutic effects of HDACi 4b likely result from effects on histone acetylation and gene transcription of only a subset of genes that are abnormally expressed in disease. Previous studies have demonstrated hypoacetylation of histones specifically in association with the dopamine D2 receptor and enkephalin, which are down-regulated in R6/2 transgenic mice (30). The authors of these studies further demonstrated reversal of histone hypoacetylation and decreased expression of these genes after treatment with the HDAC inhibitor phenylbutyrate (30). In the present study, we show similar effects for six genes in response to HDACi 4b treatment. In addition, we find complete reversal of the htt-induced transcription aberrations for subsets of genes in all three brain regions. These genes may provide a set of potential biomarkers for clinical assessment of drug treatment effectiveness. It is also possible that beneficial effects of HDACi 4b result from alterations in different genes that compensate for deficits in other systems that have been shown to be disrupted in HD. Support for this hypothesis comes from our pathways analysis results, which indicated that HDACi 4b treatment caused a decrease in the expression of genes associated with cell death, cell cycle, and the immune response ([Table S1](#)).

Interestingly, the expression of different sets of genes was altered by HDACi 4b treatment in WT mice compared with HD transgenic mice within a given brain region. Furthermore, greater numbers of genes were affected in brains of WT mice compared with HD transgenic mice. This may suggest that mutant htt protein binds to and/or disrupts the nucleosome, resulting in a state of chromatin condensation that cannot be overcome by modifying the acetylation status of histones.

In summary, our findings suggest that therapies aimed at modulating transcription may provide clinical benefits to HD patients, with minimal toxic effects. In particular, we suggest that HDACi 4b treatment may prove useful in slowing the progression of symptoms in humans.

Materials and Methods

Animals. An R6/2 line of the CBA × C57BL/6 strain origin (Jackson Laboratories) (31) has been maintained at The Scripps Research Institute by breeding male heterozygous R6/2 mice with F1 hybrids of the same background. At the age of 4 weeks, mice were genotyped according to the Jackson Laboratories protocol to determine hemizyosity for the HD transgene. The CAG repeat lengths in these mice were verified by commercial genotyping (Laragen) and found to be close to 300 (291–296); hence, we have designated these mice as R6/2^{300Q} transgenic mice. Characterization of the disease progression in these mice reveals significant rotarod deficits by 12 weeks of age and survival between 6 and 7 months of age.

Drug Treatment. HDACi 4b [*N*'-(2-aminophenyl)-*N*'-phenylheptanediamide] was synthesized as described previously (23). R6/2^{300Q} transgenic mice were individually housed and maintained on a normal 12-h light/dark cycle with lights on at 6:00 a.m. Food and water were available *ad libitum*. Groups of mice ($n = 8$ per genotype and drug treatment) were administered HDACi 4b (1 mg/ml) for 67 days in solutions that replaced drinking water beginning at 4 months of age. HDACi 4b was dissolved with HOP- β -CD as an excipient (17); control mice received Arrowhead distilled water containing an equal volume of drug vehicle. The volume of consumed liquid was estimated a few days before the start of treatment, and the concentrations of drug solutions were prepared according to the amount consumed. There were significant differences in the amounts of drug/vehicle solutions consumed per mouse per day across genotype and drug groups (3.66 ± 0.08 ml vs. 3.24 ± 0.11 ml per day for WT vs. transgenic mice, respectively, $P = 0.001$; 3.62 ± 0.09 ml vs. 3.14 ± 0.12 ml for HDACi 4b- vs. vehicle-treated mice, respectively, $P = 0.003$). However, given the smaller weights of the transgenic mice, this volume corresponded to approximately the same dose of ≈ 150 mg/kg/day, most of which was consumed over the 12-h dark cycle. Mice were individually housed to ensure consumption of fluid and food for each mouse. Drug/water consumed was recorded every day, and body weights were recorded two to three times per week. New drug was given every 4–5 days. For microarray experiments, groups of mice ($n = 4$ per genotype and drug condition) were injected s.c. with HDACi 4b (150 mg/kg/day) or an equal amount of vehicle (50:50, DMSO:PBS) once per day for 3 days. Mice were killed 6 h after the final injection, and brains were removed for microarray processing. All procedures were in strict accordance with the National Institutes of Health *Guidelines for the Care and Use of Laboratory Animals*.

Motor Behavioral Assessments. Mice were tested on an AccuRotor rotarod (AccuScan Instruments) during the light phase of the 12-h light/dark cycle by using an accelerating rotation paradigm (4–20 rpm). Mice were trained on the rotarod before drug treatment (3 months of age) to establish a behavioral baseline. At baseline, significant differences in rotarod performance are detected in R6/2^{300Q} transgenic mice (288.4 ± 6.3 s vs. 250.0 ± 14.6 s, respectively, $P = 0.02$). Behavioral semiquantitative assessment of motor and postural abnormalities were scored according to a previously described motor scale (32), which includes the widely used assessment of hindlimb clasp (31), evaluation of truncal dystonia (kyphosis), and general locomotion (including rearing and grooming behavior). Assessments were made by two independent observers who were blind to genotype and treatment status. Details of these tests are provided in [SI Methods](#).

Neuropathologic Evaluations of HDACi 4b Treatment. At the end of the drug treatment, mice were killed by brief isoflurane exposure and intracardially perfused with cold 4% paraformaldehyde/PBS. After perfusion, brains were weighed and then frozen at -70°C . Striatal and lateral ventricle volumetric analyses were performed on 25- μm brain sections. Sections 125 μm apart spanning the striatum from approximate bregma 1.18–0.38 mm were obtained by microscopic capture using Axiovision LE software (Carl Zeiss). Areas of striatum and lateral ventricle were manually traced from both sides, and volumes were recorded in square millimeters.

Microarray Analysis. Total RNA was extracted from cortex, striatum, and cerebellum from WT and R6/2^{300Q} transgenic mice treated with vehicle or HDACi 4b. Four replicates were run per sample category, for a total of 48 arrays. RNA quantity was assessed with Nanodrop (Nanodrop Technologies)

and quality with the Agilent Bioanalyzer (Agilent Technologies). Total RNA (200 ng) was amplified, biotinylated, and hybridized on Illumina Mouse Mouseref-8 Expression Beadchips v1, querying the expression of $\approx 22,000$ Refseq transcripts, as per the manufacturer's protocol. Slides were scanned by using Illumina BeadStation, and the signal was extracted by using Illumina BeadStudio software. Raw data were analyzed by using Bioconductor packages (33). Quality assessment was performed looking at the interarray Pearson correlation, and clustering based on top variant genes was used to assess overall data coherence. Contrast analysis of differential expression was performed by using the LIMMA package (34). After linear model fitting, a Bayesian estimate of differential expression was calculated. Approximately half of the genes on the array were called Present, resulting in a false discovery rate of 6%. Data analysis was aimed at (i) identifying transcriptional changes in the R6/2^{300Q} transgenic compared with WT mice and (ii) the effect of drug treatment on gene-expression abnormalities. For the initial analyses, the threshold for statistical significance was set at $P < 0.005$. In a second analysis, we used a less stringent criteria of $P < 0.05$ for both the disease and drug treatment effects, followed by an absolute log₂ ratio of >0.20 . Pathways analysis was performed by using the Functional Analysis Annotation tool in Ingenuity Pathways Analysis software (Ingenuity Systems).

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Real-Time PCR Experiments. Real-time PCR experiments were performed as described previously (14) and in *SI Methods*, using the primer sequences shown in *Table S2*.

Chromatin Immunoprecipitation. ChIP assays on brain homogenates were performed as described in previous studies (23). For each immunoprecipitation experiment, the amount of lysate corresponding to 25–50 μ g of total DNA was incubated with antibodies directed against histones H3 and H4 or acetylated H3 and H4 (Upstate Biotechnology). The recovered DNA was quantified by real-time PCR analysis using primer pairs directed against promoter regulatory or transcribed regions of the indicated genes (*Table S2*).

Statistical Analyses. The data were analyzed with one- or two-factor ANOVA and Student's *t* tests, using GraphPad Prism Software. For PCR analyses, given that the direction of expression change was predicted *a priori* from the microarray findings, one-tailed tests were used.

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