HDAC-regulated myomiRs control BAF60 variant exchange and direct the functional phenotype of fibro-adipogenic progenitors in dystrophic muscles

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Fibro-adipogenic progenitors (FAPs) are important components of the skeletal muscle regenerative environment. Whether FAPs support muscle regeneration or promote fibro-adipogenic degeneration is emerging as a key determinant in the pathogenesis of muscular diseases, including Duchenne muscular dystrophy (DMD). However, the molecular mechanism that controls FAP lineage commitment and activity is currently unknown. We show here that an HDAC–myomiR–BAF60 variant network regulates the fate of FAPs in dystrophic muscles of mdx mice. Combinatorial analysis of gene expression microarray, genome-wide chromatin remodeling by nuclease accessibility (NA) combined with next-generation sequencing (NA-seq), small RNA sequencing (RNA-seq), and microRNA (miR) high-throughput screening (HTS) against SWI/SNF BAF60 variants revealed that HDAC inhibitors (HDACis) derepress a “latent” myogenic program in FAPs from dystrophic muscles at early stages of disease. Specifically, HDAC inhibition induces two core components of the myogenic transcriptional machinery, MYOD and BAF60C, and up-regulates the myogenic miRs (myomiRs) (miR-1, miR-133, and miR-206), which target the alternative BAF60 variants BAF60A and BAF60B, ultimately directing promyogenic differentiation while suppressing the fibro-adipogenic phenotype. In contrast, FAPs from late stage dystrophic muscles are resistant to HDACi-induced chromatin remodeling at myogenic loci and fail to activate the promyogenic phenotype. These results reveal a previously unappreciated disease stage-specific bipotency of mesenchimal cells within the regenerative environment of dystrophic muscles. Resolution of such bipotency by epigenetic intervention with HDACis provides a molecular rationale for the in situ reprogramming of target cells to promote therapeutic regeneration of dystrophic muscles.

Keywords: SWI/SNF chromatin remodeling, BAF60, microRNA, HDAC, FAPs, muscular dystrophy

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Duchenne muscular dystrophy (DMD) is an inherited muscular disease that remains fatal despite the increasing knowledge of the biochemical and functional consequences of dystrophin absence in skeletal muscles [Dalkilic and Kunkel 2003]. DMD pathogenesis is characterized by an early stage regenerative response that counterbalances muscle loss but eventually declines as the disease prog-
These data suggest that the intrinsic myogenic potential—mediated formation of myotubes and endogenous muscle—restored the ability of HDACis to promote satellite cell-during disease progression. Importantly, FAPs from regenerative environment that mediates HDACis ability to reprogram the FAP phenotype and supports compensatory regeneration at early stages of DMD progression in mdx mice, thereby providing the rationale for the pharmacological control of the FAP phenotype to counter DMD progression.

**Results**

**HDACis promote a myogenic phenotype in FAPs of young mdx mice**

We began to decipher the molecular events underlying the ability of FAPs to mediate the beneficial effects of HDACis in young mdx mice (Mozzetta et al. 2013) by performing a gene expression microarray analysis. RNA was collected from FAPs immediately after their isolation from muscles of young (1.5-mo-old) mdx mice treated with TSA or untreated. Among the up-regulated transcripts, those implicated in muscle determination and differentiation as well as in muscle contraction and metabolism scored as the top category in FAPs isolated from TSA-treated mdx mice (Fig. 1A,B). Supplementary Fig. S1A,B. The complete list of up-regulated and down-regulated genes is accessible through Gene Expression Omnibus (GEO) series accession number GSE34843 (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE34843).

Interestingly, in TSA-treated FAPs, we noted the induction of the muscle-specific determination factor MyoD and the SWI/SNF chromatin remodeling complex subunit BAFl60C (encoded by Smarcd3) that is required for skeletal myogenesis and MYOD-mediated activation of muscle gene transcription (Simone et al. 2004; Lickert et al. 2004; Forcales et al. 2012; Albini et al. 2013). Although the muscle gene transcripts induced by TSA in FAPs from young mdx mice were expressed at lower levels than in MuSCs (Fig. 1B), MyoD-expressing FAPs eventually differentiated into MYHC-positive myotubes rather than Oil red O-positive adipocytes when cultured in adipogenic conditions [adipogenic differentiation medium] for 6 d (Fig. 1C–F). Gene ontology and network analysis revealed the up-regulation of components of pathways (e.g., Wnt and Notch) implicated in muscle development and adult myogenesis (Brack and Rando...
as well as signals recently involved in FAP activation by eosinophils during muscle regeneration (Supplemental Fig. S1A,B; Heredia et al. 2013). Quantitative RT–PCR (qRT–PCR) analysis confirmed the induction of muscle genes in FAPs from TSA-treated mdx mice over the very low levels detectable in the same cells from control mdx mice (Supplemental Fig. S1C). Western blot analysis confirmed the expression of MYOD and the late contractile protein MYHC in FAPs from young mdx mice exposed to TSA, while the adipogenic protein Perilipin was down-regulated under the same conditions [Fig. 1C].

The up-regulation of muscle genes observed in FAPs from TSA-treated mdx mice could, in theory, be due to contamination with sporadic MuSCs during FACS sorting and their subsequent clonogenic expansion. However, a large proportion of MYOD-expressing FAPs was detected already within 4 d after their isolation from TSA-treated mdx mice [Fig. 1D], timing incompatible with a potential clonogenic expansion of contaminating MuSCs. We further investigated this issue by evaluating the ability
GSE52062). Among the total NASs detected in FAPs accessible through GEO series accession number GSE52062 (Fig. 2A). The complete list of NASs in FAPs from young statistically significant increase in NASs in response to TSA HDACis (Mozzetta et al. 2013), did not show any statistically significant increase in NASs in FAPs from TSA-treated mdx mice, as compared with untreated controls. An increased frequency of NA sites (NASs) was detected in FAPs from young mdx mice exposed to TSA [Fig. 2A], a treatment leading to the functional and histological recovery of dystrophic muscles [Minetti et al. 2006; Mozzetta et al. 2013] and to the up-regulation of muscle genes [see Fig. 1]. In contrast, FAPs from old mdx mice, which are resistant to the beneficial activity of HDACis [Mozzetta et al. 2013], did not show any statistically significant increase in NASs in response to TSA [Fig. 2A]. The complete list of NASs in FAPs from young and old mdx mice with or without TSA treatment is accessible through GEO series accession number GSE52062 (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE52062). Among the total NASs detected in FAPs from TSA-treated young mdx mice, ~70% were induced de novo, while 30% were previously existing NASs that were further increased by TSA [Fig. 2B]. Interestingly, the E-box sequences typically bound by myogenic basic helix–loop–helix factors and detected in previous chromatin immunoprecipitation (ChIP) combined with deep sequencing [ChIP-seq] analysis of MYOD–chromatin binding in C2C12 myoblasts [Cao et al. 2010] were found to be associated with high frequency with both de novo and “increased” NASs detected in FAPs from TSA-treated young mdx mice [Fig. 2C,E]. When we matched the NASs activated by TSA in FAPs from young mdx mice with the changes in gene expression revealed by microarray analysis [Fig. 1B], we identified 3.04% of the total NASs that were found on putative regulatory elements [within 20 kb of the transcription start site (TSS)] of up-regulated and down-regulated genes [Fig. 2D,E]. The remaining 96.5% of the TSA-induced NASs apparently did not coincide with up-regulated or down-regulated genes [Fig. 2D,E]. Global gene ontology analysis of the overlap between NASs and genes up-regulated in FAPs from young mdx mice treated with TSA revealed a strong enrichment in muscle-related genes and components of signaling pathways, such as the WNT signaling, that promote skeletal myogenesis [Table 1; Supplemental Fig. S3A–C]. Among these genes, we again noted MyoD and BAF60C (Table 1). We focused on these two key activators of skeletal myogenesis as “proxy” to validate the ability of TSA to activate the myogenic program in FAPs via histone hyperacetylation and chromatin remodeling at muscle-specific loci. Indeed, qRT–PCR analysis confirmed the ability of TSA to induce the expression of Myod1 and Baf60c in FAPs from young but not old mdx mice [Supplemental Fig. S3D]. We used ChIP with anti-acyethylated H3 [AceH3] antibodies and formaldehyde-assisted isolation of regulatory element (FAIRE) to monitor the histone acetylation and chromatin structure at the NASs detected in Myod1 and Baf60c genes. This analysis showed an increased histone acetylation and chromatin accessibility in coincidence with the NASs detected at MyoD1 and Baf60C loci [Fig. 2F]. In contrast, TSA could not induce the expression of Hoxa7 [Supplemental Fig. S3D], which was not annotated among the genes up-regulated in FAPs from TSA-treated young mdx mice. The lack of Hoxa7 induction correlated with the absence of NASs and lack of histone hyperacetylation and chromatin remodeling on the regulatory sequence of Hoxa7 in FAPs from TSA-treated mdx mice [Fig. 2F]. Collectively, the results of the combined gene expression profile and NA-seq analysis revealed a latent myogenic feed-forward circuitry in FAPs from young mdx mice that is derepressed by HDACis and relies on the activation of the core components of the transcriptional myogenic machinery, MYOD and BAF60C. Conversely, the constitutive repression of MYOD and BAF60C appears to compromise the activation of this network in FAPs from old mdx mice.

Identification of HDACi-induced myogenic microRNAs (miyomiRs) that target the BAF60 variants A and B

We next analyzed the genomic distribution of the largest portion of TSA-induced NASs (>95%) that did not associate with putative regulatory elements within 20 kb of

HDACis promote genome-wide changes of chromatin structure and focal remodeling at muscle loci in FAPs from young mdx mice

The induction of BAF60C and MyoD in FAPs from TSA-treated mdx young mice suggests that these two factors could assemble to form the pioneer complex that recruits the SWI/SNF complex, which promotes chromatin remodeling at muscle loci, as previously shown in skeletal myoblasts [Forcales et al. 2012] and during conversion of human embryonic stem cells into skeletal muscle cells [Albini et al. 2013]. We used a nuclease accessibility (NA) assay coupled to next-generation sequencing (NA-seq) to monitor chromatin remodeling at the genome-wide level in FAPs from TSA-treated mdx mice, as compared with untreated controls. An increased frequency of NA sites (NASs) was detected in FAPs from young mdx mice exposed to TSA [Fig. 2A], a treatment leading to the functional and histological recovery of dystrophic muscles [Minetti et al. 2006; Mozzetta et al. 2013] and to the up-regulation of muscle genes [see Fig. 1]. In contrast, FAPs from old mdx mice, which are resistant to the beneficial activity of HDACis [Mozzetta et al. 2013], did not show any statistically significant increase in NASs in response to TSA [Fig. 2A]. The complete list of NASs in FAPs from young and old mdx mice with or without TSA treatment is accessible through GEO series accession number GSE52062 (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE52062). Among the total NASs detected in FAPs from TSA-treated young mdx mice, ~70% were induced de novo, while 30% were previously existing NASs that were further increased by TSA [Fig. 2B]. Interestingly, the E-box sequences typically bound by myogenic basic helix–loop–helix factors and detected in previous chromatin immunoprecipitation (ChIP) combined with deep sequencing [ChIP-seq] analysis of MYOD–chromatin binding in C2C12 myoblasts [Cao et al. 2010] were found to be associated with high frequency with both de novo and “increased” NASs detected in FAPs from TSA-treated young mdx mice [Fig. 2C,E]. When we matched the NASs activated by TSA in FAPs from young mdx mice with the changes in gene expression revealed by microarray analysis [Fig. 1B], we identified 3.04% of the total NASs that were found on putative regulatory elements [within 20 kb of the transcription start site (TSS)] of up-regulated and down-regulated genes [Fig. 2D,E]. The remaining 96.5% of the TSA-induced NASs apparently did not coincide with up-regulated or down-regulated genes [Fig. 2D,E]. Global gene ontology analysis of the overlap between NASs and genes up-regulated in FAPs from young mdx mice treated with TSA revealed a strong enrichment in muscle-related genes and components of signaling pathways, such as the WNT signaling, that promote skeletal myogenesis [Table 1; Supplemental Fig. S3A–C]. Among these genes, we again noted MyoD and BAF60C (Table 1). We focused on these two key activators of skeletal myogenesis as “proxy” to validate the ability of TSA to activate the myogenic program in FAPs via histone hyperacetylation and chromatin remodeling at muscle-specific loci. Indeed, qRT–PCR analysis confirmed the ability of TSA to induce the expression of Myod1 and Baf60c in FAPs from young but not old mdx mice [Supplemental Fig. S3D]. We used ChIP with anti-acyethylated H3 [AceH3] antibodies and formaldehyde-assisted isolation of regulatory element (FAIRE) to monitor the histone acetylation and chromatin structure at the NASs detected in Myod1 and Baf60c genes. This analysis showed an increased histone acetylation and chromatin accessibility in coincidence with the NASs detected at MyoD1 and Baf60C loci [Fig. 2F]. In contrast, TSA could not induce the expression of Hoxa7 [Supplemental Fig. S3D], which was not annotated among the genes up-regulated in FAPs from TSA-treated young mdx mice. The lack of Hoxa7 induction correlated with the absence of NASs and lack of histone hyperacetylation and chromatin remodeling on the regulatory sequence of Hoxa7 in FAPs from TSA-treated mdx mice [Fig. 2F]. Collectively, the results of the combined gene expression profile and NA-seq analysis revealed a latent myogenic feed-forward circuitry in FAPs from young mdx mice that is derepressed by HDACis and relies on the activation of the core components of the transcriptional myogenic machinery, MYOD and BAF60C. Conversely, the constitutive repression of MYOD and BAF60C appears to compromise the activation of this network in FAPs from old mdx mice.

Identification of HDACi-induced myogenic microRNAs (miyomiRs) that target the BAF60 variants A and B

We next analyzed the genomic distribution of the largest portion of TSA-induced NASs (>95%) that did not associate with putative regulatory elements within 20 kb of
the TSSs of up-regulated and down-regulated genes. Supplemental Figure 4 shows that most of these NASs are associated with noncoding genomic sequences that included miRs, long noncoding RNAs [lincRNAs], and long and short interspersed noncoding elements [LINES and SINES, respectively]. Among these noncoding transcripts, we focused initially on miRs, although they represented only a small percentage (1%), since a previous report showed a HDAC-dependent control of miRs in myoblasts isolated from dystrophic muscles (Cacchiarelli et al. 2010). To establish a functional relationship between HDACis, activation of miR, and BAF60-mediated chromatin modifications that activate the myogenic program in FAPs from young mdx mice, we used high-throughput screening (HTS) of a library of miRs against the untranslated region [UTR] sequences of the Baf60 [Smarcd] variants a, b, and c linked to the luciferase (Fig. 3A). We initially performed this screening in HEK293 cells. Among the miRs identified by this screening, we noted two miRs that have been previously implicated in the activation of the myogenic program—miR-1.2 and miR-206 (Sokol and Ambros 2005; Chen et al. 2006; Kim et al. 2006)—as BAF60B targeting miRs (Fig. 3B; Supplemental Fig. S5A). In a complementary approach, among the miRs induced by TSA in FAPs from young mdx mice, small RNA sequencing (RNA-seq) analysis again identified miR-206 as well as miR-133a, another miR that has been implicated in the regulation of skeletal myogenesis (Chen et al. 2006) (Fig. 3C).
Table 1. TSA-induced NAS at E-box sequences in regulatory regions of genes up-regulated by TSA

<table>
<thead>
<tr>
<th>Name</th>
<th>Function</th>
<th>Position</th>
<th>Description</th>
<th>Transcription factors with binding sites</th>
<th>Fold change</th>
<th>Fold change</th>
<th>Fold change</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atp2a2</td>
<td>Calcium pump located in muscle cells</td>
<td>Promoter</td>
<td>Increased</td>
<td>Myf6/MyoD1/Myog</td>
<td>9.0</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Fam65b</td>
<td>Promotes myogenic cell differentiation</td>
<td>Intron 1</td>
<td>Increased</td>
<td>MyoD1/Myog</td>
<td>14.0</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Fglr4</td>
<td>Required for the development of skeletal muscle lineages</td>
<td>Promoter</td>
<td>Increased</td>
<td>Myf5/Myf6/MyoD1/Myog</td>
<td>2.49</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Lmcd1</td>
<td>Z-disc-associated protein</td>
<td>Exon–intron 4</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>2.88</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Myh11</td>
<td>Structural constituent of muscle</td>
<td>Exon–intron 20</td>
<td>De novo</td>
<td>Myf5/Myf6/MyoD1/Myog</td>
<td>3.85</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Myod1</td>
<td>Transcription factor involved in muscle differentiation</td>
<td>Exon–intron last</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>6.14</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Neb</td>
<td>Maintains the structural integrity of sarcomeres</td>
<td>Intron 71</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>18.5</td>
<td>ns</td>
<td>–37.00</td>
</tr>
<tr>
<td>Smarcd3</td>
<td>SWI/SNF subunit involved muscle gene activation</td>
<td>Promoter</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>4.3</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Speg</td>
<td>Required for myocyte cytoskeletal development</td>
<td>Exon–intron 14</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>3.83</td>
<td>ns</td>
<td>–3.96</td>
</tr>
<tr>
<td>Synpo2</td>
<td>Actin-bundling protein in the Z-disc of skeletal muscle</td>
<td>Intron last</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>4.27</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Tcf15</td>
<td>Regulates musculoskeletal patterning</td>
<td>Promoter</td>
<td>Increased</td>
<td>Myf5/MyoD1/Myog</td>
<td>2.83</td>
<td>ns</td>
<td>ns</td>
</tr>
<tr>
<td>Tnni2</td>
<td>Regulates striated muscle contraction</td>
<td>Promoter</td>
<td>De novo</td>
<td>Myf5/MyoD1/Myog</td>
<td>2.17</td>
<td>ns</td>
<td>ns</td>
</tr>
</tbody>
</table>

(ns) Not significant.
Figure 3. Identification of miRs targeting the alternative BAF60A and BAF60B variants. (A) Schematic representation of the pGK plasmid transfected into HEK293 cells for miR sensor screening. The 3’ UTRs of target genes were cloned downstream from the luciferase coding sequence. (B) Volcano plots representing primary data for each of the three miR screens. The significance versus fold change for each of the 875 miRs tested is plotted. miR activities plotted as percent inhibition relative to siRNA against Smarcd1 (Baf60a), Smarcd2 (Baf60b), or Smarcd3 (Baf60c) (100% inhibition) and scrambled sequence control (0% inhibition) (X-axis) and P-value from t-test (Y-axis). miRs that block Luc-Smarcd1, Luc-Smarcd2, or Luc-Smarcd3 production (>20% cutoff for Smarcd1 and Smarcd2; >40% cutoff for Smarcd3; P < 0.11) are shown in green. From this group of hits, miR-1 and miR-206 are highlighted in yellow and purple, respectively. The primary screen was performed using 10 nM miRs in triplicate. (C) Manually assembled list representing small RNA-seq results of miRs induced in FAPs isolated from TSA-treated versus control (CTR) 1.5-mo-old mdx mice. (D) Representation of overlapping results from miR HTS described in B and small RNA-seq described in C. (E) BAF60A, BAF60B, and BAF60C protein levels in TSA-treated versus control (CTR) FAPS were measured by Western blot analysis. (F) MyomiR (miR1.2as, miR133, and miR206) endogenous expression in MuSCs during differentiation (growth medium [GM] > differentiation medium [DM]) (left panel) and in FAPs isolated from young 1.5-mo-old mdx mice treated with vehicle (CTR) or TSA for 15 d (right panel). (G) BAF60 subunits’ relative expression in MuSCs during differentiation (left panel) and in FAPs isolated as described in B after control (CTR) or TSA treatment (right panel). Data are represented as average ± SEM of three different experiments (n = 3); (*) P < 0.05; (**) P < 0.01; (***) P < 0.001.
complete list of miRs expressed in FAPs from young mdx mice with or without TSA treatment is accessible through GEO series accession number GSE51933 [http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE51933]. miR-1, miR-133, and miR-206, collectively referred to as myomiRs, are components of a miR cluster that regulates a wide array of target transcripts in skeletal muscle cells during development and adult life and have been implicated in the pathogenesis of MDs and other muscular disorders [Eisenberg et al. 2009; Greco et al. 2009; Williams et al. 2009]. Thus, the combination of two independent approaches (HTS and small RNA-seq) indicated a potential relationship between TSA-induced myomiRs and regulation of BAF60 variants in FAPs (Fig. 3D).

By using a bioinformatic search for predictive UTR targets of these myomiRs, we consistently noted the presence of Baf60a and Baf60b [but not Baf60c] UTR sequences [Supplemental Fig. S5B]. Indeed, Munsterberg and colleagues [Goljanek-Whysall et al. 2012] have recently identified BAF60B among the targets of miR-1 and miR-206 in skeletal myoblasts. We directly tested the effect of myomiRs on the Baf60a, Baf60b, or Baf60c 3′ UTR linked to the luciferase gene in HEK293 cells. All myomiRs (miR-1, miR-133, and miR-206), when independent of expression by transfection, down-regulated the luciferase activity of Baf60a-UTR-luc and Baf60b-UTR-luc constructs coexpressed in these cells [Supplemental Fig. 5C]. This effect was specific, as mutations that disrupted the recognition sequence of myomiRs in Baf60a and Baf60b mRNAs were sufficient to abrogate miR-mediated reduction of luciferase activity [Supplemental Fig. S5C]. In contrast, Baf60c-UTR-luc was not reduced by myomiR expression in HEK293 cells [Supplemental Fig. S5C]. We further investigated whether exposure to a HDACi (TSA) could replicate the effect of myomiRs on Baf60-UTR-luc activity in C2C12 myoblasts, which provide a more relevant biological context. Supplemental Figure S5D shows that TSA down-regulated the activity of Baf60a-UTR-luc and Baf60b-UTR-luc but not Baf60c-UTR-luc.

We next established a direct correlation between myomiR induction and the expression levels of BAF60 variants in FAPs isolated from young mdx mice exposed or not to TSA. BAF60A and BAF60B protein levels were down-regulated, while BAF60C was up-regulated, in FAPs isolated from muscles of mdx mice exposed to TSA (Fig. 3E). The up-regulation of myomiRs in FAPs of mdx mice exposed to TSA [Fig. 3F, right panel] coincided with the down-regulation of Baf60a and Baf60b transcripts [Fig. 3G, right panel]. In contrast, Baf60c transcripts were induced in FAPs of mdx mice exposed to TSA [Fig. 3G, right panel], consistent with the lack of a myomiR recognition sequences in Baf60c UTR and with the ability of TSA to promote chromatin remodeling and transcription at the Baf60c locus [see Fig. 2]. Of note, a pattern of myomiR and BAF60 variant modulation similar to that detected in FAPs exposed to TSA was also observed in differentiating MuSCs isolated from young mdx mice [Fig. 3F,G, left panels], once again suggesting interesting analogies between MuSCs and the FAPs from regenerating muscles of mdx mice exposed to TSA.

We then used antagonimirs to evaluate the specificity of myomiR-mediated down-regulation of BAF60A and BAF60B in TSA-treated FAPs. Transfection of specific antagonimirs blunted the TSA-mediated down-regulation of Baf60a and Baf60b, but not Baf60c, and prevented the expression of muscle differentiation markers [such as Myh3] in FAPs isolated from young mdx mice (Fig. 4A). Consistently, blockade of each myomiR by ectopic expression of specific antagonimirs completely reverted TSA-mediated induction of MYHC-positive myotubes [Fig. 4B,C] and the inhibition of adipogenic differentiation in FAPs from young mdx mice [Fig. 4B,D]. These results indicate the essential function of myomiRs in mediating the HDACi-directed conversion of FAPs into the myogenic lineage.

Reprogramming of FAPs into a promyogenic phenotype by modulation of the myomiR–BAF60 network

We next tested whether ectopic expression of individual myomiRs could affect the function and phenotype of FAPs isolated from young mdx mice. Previous studies have used coculture experiments to show that the promyogenic activity of FAPs relies on their ability to implement MuSC-mediated formation of multinucleated myotubes during muscle regeneration [Joe et al. 2010]. Moreover, a transwell coculture assay demonstrated that FAPs from young mdx mice promote the ability of MuSCs to differentiate into myotubes, possibly via soluble factors, with this activity being further augmented by HDACIs [Mozzetta et al. 2013]. We therefore used transwell cocultures to test whether transfection of myomiRs would directly influence the ability of FAPs to functionally interact with MuSCs. The myomiRs were individually expressed in FAPs isolated from young mdx mice [Fig. 5A], prior to transwell coculture with MuSCs [Fig. 5B]. Notably, each myomiR increased the ability to promote the formation of larger, MYHC-expressing multinucleated myotubes from cocultured MuSCs [Fig. 5C,D]. The effect of each myomiR was comparable with that of TSA and resulted in an increased ability of FAPs to stimulate MuSC fusion into myotubes, as compared with mock transfected, control FAPs [Fig. 5C,D]. In a parallel assay, we monitored the effect of myomiR expression on the intrinsic ability of mdx-derived FAPs to adopt a fibro-adipogenic or the alternative myogenic lineage that was observed in response to TSA [Fig. 1]. Lineage and differentiation potential were scored by the formation of MYHC-positive myotubes and/or perilipin-expressing adipocytes under adipogenic culture conditions. A significant proportion (~10%) of FAPs from young mdx mice were typically differentiated into perilipin-expressing adipocytes under adipogenic culture conditions. A significant proportion (~10%) of FAPs from young mdx mice was not detected in FAPs isolated from uninjured muscles of wild-type or old mdx mice (data not shown). Treatment with TSA increased the formation of MYHC-positive myotubes [Fig. 5E,F] and the concomitant inhibition of perilipin-expressing adipocytes [Fig. 5E,G]. Ectopic expression of each myomiR replicated the effect of TSA, with increased formation of MYHC-positive myo-
tubes and the reciprocal reduction of perilipin-expressing adipocytes (Fig. 5E–G). These data support the notion that TSA-mediated up-regulation of myomiRs switches FAP fate toward the myogenic lineage.

Lineage determination is programmed at the transcription level by changes in chromatin structure that are typically promoted by the SWI/SNF chromatin remodeling complex. The heterogeneous composition of SWI/SNF has been recently regarded as a key determinant of localized chromatin remodeling to support lineage-specific transcription and ultimately the decision to adopt alternative phenotypes within mesodermal-derived populations (for review, see Puri and Mercola 2012). In both skeletal and cardiac muscle, the formation of the BAF60C-based SWI/SNF complex is essential for the transcriptional activation of cardiac or skeletal myogenic genes [Lickert et al. 2004; Takeuchi and Bruneau 2009; Forcales et al. 2012; Albini et al. 2013]. Moreover, reconfiguring the SWI/SNF composition by miR-mediated targeting of alternative variants of BAF subunits has previously been shown to regulate neurogenesis [Lessard et al. 2007; Yoo et al. 2011]. Thus, we hypothesized that myomiR-mediated down-regulation of Baf60A and Baf60B could exert an effect equivalent to myomiR expression in FAPs from young mdx mice. Indeed, shRNAi-mediated down-regulation of Baf60a and Baf60b was sufficient to replicate the effect of ectopic expression of myomiRs in FAPs shown in Figure 5. Reducing the levels of Baf60a or Baf60b by shRNAi [Fig. 6A] enhanced FAPs’ ability to promote MuSC-mediated formation of myotubes in transwell coculture experiments [Fig. 6B–D] and promoted FAP differentiation into MYHC-positive myotubes with an efficiency comparable with that of TSA [Fig. 6E,F] or ectopic myomiR expression (cf. Figs. 6E,F and 5E,F). Importantly, shRNAi-mediated down-regulation of

**Figure 4.** AntagomiRs against myomiRs counteract the HDACi activity in FAPs. ([A] qRT–PCR detecting relative amounts of Baf60a, Baf60b, Baf60c, and Myh3 transcripts in FAPs from 1.5-mo-old mdx mice transfected in vitro with antagomir-1 [a-miR1], antagomir-133 [a-miR133], antagomir-206 [a-miR206], or scrambled antagomir [mock] in either the absence [black bars, CTR] or presence of TSA added after transfection [gray bars, TSA]. [B] FAPs’ myogenic and adipogenic potential after TSA treatment and/or antagomiR transfection was assessed by immunofluorescence for MYHC [green] and Oil red O [red]. Nuclei were counterstained with DAPI [blue]. [C] Graph showing the quantification of the myogenic differentiation of FAPs in the experiment represented in B. We measured the percentage of nuclei in MYHC-positive cells per field. [D] Graph showing the quantification of the adipogenic differentiation of FAPs in the experiment represented in B. We measured the percentage of the Oil red O-positive cells per field.
Figure 5. MyomiR expression induced a myogenic phenotype in FAPs and enhanced their ability to stimulate MuSC fusion into myotubes. 

[A] qRT–PCR to check miR1.2, miR133, and miR206 expression in FAPs isolated from young mdx mice transfected with psp65-miR1.2, psp65-miR133a, and psp65-miR206 constructs as compared with FAPs transfected with the psp65-null vector (CTR). 

[B] Transwell coculture scheme. MuSCs FACS-isolated from young mdx mice (MuSCs CTR) were seeded at the bottom of the well. FAPs isolated from young mdx mice (control [CTR] or TSA-treated for 15 d) were seeded on a transwell insert. Control FAPs were transfected with psp65-null vector (CTR), psp65-miR1, psp65-miR133, and psp65-miR206. Six hours after FAP transfection, all transwells were put in coculture with MuSCs. 

[C] The myogenic differentiation of MuSCs—alone or after coculture with FAPs—was assessed by immunostaining for MYHC (green). Nuclei were counterstained with DAPI (blue). 

[D] Graph showing the fusion index of the experiment represented in C. We measured the percentage of nuclei that were in MYHC− or MYHC+ mononucleated myotubes (n < 2), the percentage of nuclei that were inside myotubes containing between two and five nuclei (2 < n < 5), and the percentage of nuclei inside myotubes containing more than five nuclei (n > 5). 

[E] FAPs’ myogenic and adipogenic potential after TSA treatment or myomiR transfection was assessed by immunofluorescence for MYHC (green) and PERILIPIN (red). Nuclei were counterstained with DAPI (blue). 

[F] Graph showing the quantification of the myogenic differentiation of FAPs in the experiment represented in E. We measured the percentage of nuclei in MYHC-positive cells. 

[G] Graph showing the quantification of the adipogenic differentiation of FAPs in the experiment represented in E. We measured the percentage of Perilipin-positive cells per field. Data are represented as average ± SEM of three different experiments (n = 3); (*) P < 0.05; (**) P < 0.01.
Figure 6. Down-regulation of Baf60a or Baf60b exerts an effect equivalent to myomiR expression in FAPs. (A) qRT–PCR to check Baf60a, Baf60b, and Baf60c down-regulation in FAPs isolated from young mdx mice and transfected with pSuper short hairpin Baf60a (shBaf60a), pSuper short hairpin Baf60b (shBaf60b), and pSuper short hairpin Baf60c (shBaf60c), as compared with FAPs transfected with the pSuper-null vector (CTR). (B) Transwell coculture scheme: MuSCs FACS-isolated from young MDX mice (MuSCs CTR) were seeded at the bottom of the well. FAPs isolated from young mdx mice (control [CTR] or TSA-treated for 15 d) were seeded on a transwell insert. FAPs were transfected with control (CTR), shBaf60a, shBaf60b, and shBaf60c pSuper vectors. Six hours after transfection, FAPs were put in coculture with MuSCs. (C) The myogenic differentiation of MuSCs—alone or after coculture with FAPs—was assessed by immunostaining for MYHC (green). Nuclei were counterstained with DAPI (blue). (D) Graph showing the fusion index of the experiment represented in C. We measured the percentage of nuclei that were in MYHC− or MYHC+ mononucleated myotubes [n < 2], the percentage of nuclei that were inside myotubes containing between two and five nuclei [2 < n < 5], and the percentage of nuclei inside myotubes containing more than five nuclei [n > 5]. (E) FAPs’ myogenic and adipogenic potential after TSA treatment or BAF60 variant down-regulation was assessed by immunofluorescence for MYHC (green) and PERILIPIN (red). Nuclei were counterstained with DAPI (blue). (F) Graph showing the quantification of the myogenic differentiation of FAPs in the experiment represented in E. We measured the percentage of nuclei in MYHC-positive cells. (G) Graph showing the quantification of the adipogenic differentiation of FAPs in the experiment represented in E. We measured the percentage of the PERILIPIN-positive cells per field. Data are represented as average ± SEM of three different experiments [n = 3], [*] P < 0.05, [**] P < 0.01.
Likewise, knocking down Baf60a and Baf60b reduced the extent of adipogenic differentiation of FAPs (Fig. 6E–G), as also observed with overexpression of myomiRs [cf. Figs. 6E–G and 5E–G]. In contrast, shRNAi-mediated knockdown of Baf60c did not alter the phenotype of FAPs from young mdx mice (Fig. 6E) but abrogated the promyogenic effects of TSA [Fig. 6C,D], indicating an essential role of BAF60C in the activation of the myogenic program upon HDAC blockade.

These results support the concept that HDAC-regulated expression of myomiRs controls the mutually exclusive incorporation of BAF60 variants into distinct SWI/SNF complexes that can alternatively direct adipogenic or myogenic programs in FAPs from young mdx mice. Interestingly, the resistance to TSA of FAPs from muscles of mdx mice at late stages of disease progression, when the regeneration potential is exhausted, suggests that FAPs’ response to HDACis is influenced by regeneration signals. We therefore tested whether induction of regeneration by cardiotoxin (CTX)-mediated muscle injury could enable FAPs of young wild-type mice to adopt the myogenic lineage in response to TSA. Supplemental Figure 6 shows that TSA does not affect the phenotype of FAPs from uninjured muscles of wild-type mice but does promote myogenesis at the expense of adipogenesis in FAPs from CTX-injured regenerating muscles (Supplemental Fig. S6A–E), an effect reminiscent of that observed in young mdx mice. Consistently, FAPs from injured muscles exposed to TSA showed an increased ability to support the myogenic potential of MuSCs (Supplemental Fig. S6F,G). Moreover, ectopic expression of each of the myomiRs (miR-1.2, miR-133a, and miR-206) individually promoted myogenesis and inhibited adipogenesis only in FAPs from CTX-injured wild-type muscles (Supplemental Fig. S7A–D). Likewise, knocking down Baf60 variants Bafl0a and Bafl0b, but not Bafl0c, promoted myogenesis and inhibited adipogenesis only in FAPs from CTX-injured wild-type muscles (Supplemental Fig. S7E–H).

Collectively, these results indicate that regeneration signals—such as those released by injured wild-type muscles and dystrophic muscles at the early, compensatory stages of disease progression—instruct FAPs to adopt the myogenic lineage in response to HDACis. This evidence explains the reported stage-dependent beneficial effect of HDACis in mdx mice [Mozzetta et al. 2013].

BAF60C is an essential component of the chromatin machinery that directs the promyogenic phenotype of FAPs in response to TSA treatment

The data shown in Figure 6 point to the key role of BAF60C in directing the promyogenic phenotype of FAPs. To conclusively demonstrate the role of BAF60C in directing the phenotype of FAPs from dystrophic muscles, we crossed Baf60c<sup>fl/fl</sup> mice with mdx mice to generate mdx<sub>Baf60c<sup>fl/fl</sup></sub> mice. FAPs were isolated from these mice as previously described [Mozzetta et al. 2013], and Baf60c was heritably deleted by infection with adenovirus expressing CRE (AdCRE) followed by culture in adipogenic conditions (see scheme in Fig. 7A). Mdx<sub>Baf60c<sup>fl/fl</sup></sub> FAPs infected with control adenovirus (AdCTR) displayed the expected phenotype, yielding occasional MYHC-positive mononucleated cells (Fig. 7B,D) within a predominant population of Oil red O-positive adipocytes (Fig. 7E,G). In vitro treatment with TSA increased the number of MYHC-positive myotubes, with formation of multinucleated myotubes (Fig. 7B,D), albeit to a reduced extent as compared with FAPs from in vivo treated young mdx mice (cf. Fig. 1). This difference is possibly due to the negative, nonspecific effect of adenoviral infection on the myogenic potential of FAPs. TSA also inhibited the formation of Oil red O-positive adipocytes (Fig. 7E,G). Adenoviral (AdCRE)-mediated deletion of Baf60c (Fig. 7C) reduced the basal number of MYHC-positive cells and abrogated TSA-mediated induction of myogenesis (Fig. 7B,D). However, while BAF60C deficiency increased the number of Oil red O-positive adipocytes, it did not effectively counter the ability of TSA to prevent the activation of the adipogenic program in FAPs [Fig. 7E,G]. This was also confirmed by qRT–PCR analysis of gene expression, as the TSA-mediated repression of adipogenic genes, such as Ppary (<sub>PPARgamma</sub>) and Cebp (<sub>CEBPalpha</sub>), was observed in either control (AdCTR) or Baf60c-deficient (AdCRE) FAPs derived from mdx<sub>Baf60c<sup>fl/fl</sup></sub> mice (Fig. 7F).

These results support the conclusion that BAF60C is an essential component of the chromatin-modifying machinery that promotes the promyogenic phenotype in the FAPs of young mdx mice exposed to HDACis. However, while the absence of BAF60C is sufficient to enhance the adipogenic potential of FAPs from mdx mice, BAF60C does not appear to be required for HDAC-mediated repression of the adipogenic program in FAPs.

Discussion

Our study has identified an HDAC-regulated network that controls the “functional” phenotype of FAPs from dystrophic muscles. This network consists of myomiRs that target two alternative variants of the SWI/SNF chromatin remodeling complex, BAF60A and BAF60B, thereby favoring the formation of a BAF60C-based SWI/SNF complex that confers on MYOD the ability to activate the myogenic program [Forcales et al. 2012; Albini et al. 2013]. HDAC-mediated repression of myomiRs has been previously reported in dystrophic muscles [Cacchiarelli et al. 2010]. Furthermore, myomiRs have been extensively reported to regulate skeletal myogenesis by targeting individual regulatory components of the myogenic network [Braun and Gautel 2011]. Interestingly, treatment with HDACis [Minetti et al. 2006] or up-regulation of miR-206 [Liu et al. 2012] invariably exerts therapeutic effects in muscles of mdx mice. Our results reveal a previously unrecognized link between HDACs, myomiRs, and chromatin remodeling that underlies the transition of FAPs from the “promyogenic” phenotype that supports compensatory regeneration at early stages of DMD to a “pathological” phenotype typical of the late stages of the disease. We show that reversible HDAC-mediated repression of myomiRs and of the two central components of the transcriptional machinery that activates
muscle gene expression—BAF60C and MYOD—underlies the latent myogenic phenotype in FAPs observed in young mdx mice. This potential appears to be a specific feature of FAPs from regenerating dystrophic muscles, as it was not observed in wild-type mice or old mdx mice. As such, this observation reveals an unanticipated lineage “plasticity” of FAPs that correlates with the highly regenerative environment typical of dystrophic muscles at the early stages of the disease. In contrast, FAPs isolated from old mdx mice, in which the regeneration potential declines, showed no myogenic potential. Thus, we speculate that the latent myogenic phenotype of FAPs from regenerating muscles of mdx young mice might reflect the acquisition of a bipotency that is likely adopted in

Figure 7. Genetic ablation of BAF60C in FAPs prevents the HDACi-mediated myogenic phenotype in FAPs but not the repression of the adipogenic program. [A] Schematic representation of the experimental setting. Cells were isolated from young Baf60c<sup>fl/fl</sup> mdx mice and, after 3 d in growth medium (GM), were infected with VQAd Cre/Cyc/GFP virus [AdCRE] (Viraquest, Inc.) or VQAd CMV eGFP virus [AdCTR] (Viraquest Inc.). Twenty-four hours after infection, cells were treated with TSA. After treatment, cells where switched to differentiation medium (DM). [B] FAPs’ myogenic potential was assessed by immunofluorescence for MYHC (green). Nuclei were counterstained with DAPI (blue). [C] Relative expression of Baf60c transcript in FAPs infected with AdCTR or AdCRE and treated or not with TSA. [D] Graph showing the quantification of the myogenic differentiation of FAPs in the experiment represented in B. We measured the percentage of nuclei in MYHC-positive cells. [E] FAPs’ adipogenic differentiation was assessed by Oil red O staining. [F] qRTPCR of Ppar<sub>γ</sub> and Cebp<sub>a</sub>. [G] Relative quantification of the Oil red O area measured as pixels squared per field. Data are represented as average ± SEM of three different experiments (n=3); (*) P < 0.05; (**) P < 0.01.
response to regenerative cues and is resolved into the myogenic phenotype by HDAC inhibition. In contrast, an impairment of the reciprocal interactions between environmental signals, FAPs, and MuSCs contributes to the progressive impairment of the regeneration niche in dystrophic muscles. In this regard, it should be noted that Guttridge and colleagues (He et al. 2013) have recently shown that muscle interstitial cells expressing the same surface markers that define FAP identity (SCA1^{pos}, PDGFRα^{pos}, and CD34^{pos}) adopt the myogenic lineage upon conditions that trigger cachexia. However, these cells fail to fully differentiate into myotubes because of an adverse environment imposed by cachexia [He et al. 2013].

The constitutive fibro-adipogenic phenotype of FAPs from old mdx mice and the related resistance to HDACis appears to be epigenetically determined, as suggested by the striking changes in chromatin accessibility revealed by NA-seq in FAPs isolated from old mdx mice, as compared with FAPs from young mdx mice. Our results indicate that the relative levels of BAF60 variants is a key determinant of the changes in the epigenetic landscape that ultimately directs FAP lineage commitment and that myomiRs are essential upstream regulators of the individual BAF60 variant levels. Previous reports have linked miRs with alternative lineage determination [Yin et al. 2013b; for review, see Lessard and Crabtree 2010]. In particular, work from Crabtree and collaborators [Lessard et al. 2007, Yoo et al. 2011] has elucidated a miR-mediated regulation of BAF53 and BAF45 variants that controls sequential stages of neurogenic differentiation and can even mediate conversion of fibroblasts into neurons. These data are in analogy with our observation that a myomiR–BAF60 variant axis controls lineage potency of FAPs (Figs. 5, 6) and collectively support the notion that miR-regulated composition of SWI/SNF chromatin remodeling complexes determines the epigenetic landscape that controls the cell identity of tissue progenitors [Puri and Mercola 2012].

Our data clearly indicate that the propensity of FAPs to adopt a myogenic phenotype invariably coincides with the activation of the MYOD–BAF60C axis and myomiRs. Although the role of MYOD and BAF60C in the myogenic conversion of somatic and embryonic stem cells has been reported [Davis et al. 1987; Weintraub et al. 1989; Albini et al. 2013], the myogenic conversion of FAPs by myomiRs [Fig. 5] is unprecedented. Interestingly, myomiR-mediated conversion of FAPs from dystrophic muscles is highly reminiscent of miR-mediated conversion of cardiac fibroblasts [Jayawardena et al. 2012]. In particular, the dependence of myomiR function on signals from regenerating young dystrophic muscle [this study] mirrors the emerging view that miR-based lineage conversion in multiple contexts is influenced by signals released by injured tissues, such as acute ischemic myocardium or regenerating skeletal muscle. This is consistent with an essential role of regeneration cues in generating an epigenetic landscape that is permissive for lineage conversion. Our data suggest that when FAPs are exposed to these instructive signals, HDAC-mediated repression of the Myod1–myomiR–BAF60 network maintains a “lineage plasticity,” which can be experimentally resolved by perturbations such as HDAC blockade, myomiR overexpression, or BAF60A and BAF60B knockdown that bias the network toward the formation of the MYOD/BAF60C-based SWI/SNF complex. Signal-dependent activation of this complex by regeneration cues, such as those that activate the p38 pathway, has been previously reported [Simone et al. 2004, Forcades et al. 2012]. As myomiR expression is induced by the combined activity of MYOD and the SWI/SNF complex [Mallappa et al. 2010] and myomiRs favor the composition of the myogenic BAF60C–SWI/SNF complex, by targeting the alternative BAF60A and BAF60B variants, FAP reprogramming into the myogenic phenotype appears to rely on a HDAC-supervised feed-forward circuitry composed by external signals, MYOD, myomiR, and selection of specific BAF60 variants. This is consistent with the robust fibroblast reprogramming into cardiac cells induced by the combined activity of cardiogenic factors and miRs [Nam et al. 2013].

Our data show that FAPs from mdx mice exposed to HDACis adopt the myogenic lineage and have an enhanced ability to support MuSC activity, an effect collectively referred as to induction of the promyogenic phenotype. While the relationship between these two effects is not conclusively established by the present study, it is important to note that both are dependent on the up-regulation of BAF60C and myomiR-mediated down-regulation of BAF60A and BAF60B. However, given the reported evidence that FAPs from injured muscles of wild-type mice do not adopt the myogenic lineage but retain the ability to support MuSC-mediated regeneration [Joe et al. 2010], it is likely that these two events could be separately activated. Likewise, the ability of HDACis to promote the myogenic phenotype and repress the fibro-adipogenic lineage seems to rely on distinct mechanisms, as the genetic ablation of BAF60C impairs HDACi-mediated activation of the myogenic phenotype but not HDACis’ ability to repress the fibro-adipogenic program in FAPs. Given the heterogeneity of FAPs [Pannèrec et al. 2013], it will be interesting to determine whether different fates adopted by these cells rely on the transition from one subpopulation to another.

Overall, we reported the first evidence of an HDAC–myomiR–BAF60 network that controls, in concert with environmental cues, the identity and activity of a cell type [FAPs] that plays a key role in muscle regeneration and is implicated in the pathogenesis of MDs. Future studies will shed further light on the pharmacological potential to reprogram the phenotype of these cells by epigenetic drugs that can be used in the treatment of DMD.

Materials and methods

Animals and in vivo treatments

Wild-type C57/BL10 and C57Bl10 mdx mice were purchased from Jackson Laboratories. Baf60c^{fl/fl} C57/BL10 mice were crossed with mdx C57/BL10 mice to generate the Baf60c^{fl/fl}
mdx mice. Further details on treatments are provided in the Supplemental Material.

**Cell preparation and FACS isolation**
MuSCs were isolated as TERT19+/CD45-/CD31-/α7INTEGRIN+/SCA-1+ cells; FAP cells were isolated as TERT19-/CD45-/CD31-/α7INTEGRIN-/SCA-1+ cells, as described in Mozzetta et al. (2013). Further details on FACS sorting and culture conditions are provided in the Supplemental Material.

**Luciferase assay**
HEK293 and C2C12 cells were transfected with psiCheck 2 Renilla luciferase reporter vectors containing BAF60A, BAF60B, or BAF60C 3′ UTR fragments. Further details on construct preparation and luciferase analysis are provided in the Supplemental Material.

**Western blot**
Western blot was performed using antibodies against the following proteins: BAF60A (BD), BAF60B (Santa Cruz Biotechnology), BAF60C (Forcales et al. 2012), MYOD (Santa Cruz Biotechnology), MYHC [MF20, Developmental Studies Hybridoma Bank (DSHB)], PERILIPIN [Sigma-Aldrich], and TUBULIN (Cell Signaling). Further experimental details are provided in the Supplemental Material.

**Immunofluorescence**
Primary antibodies used for immunofluorescence were against MYOD [1:20, Santa Cruz Biotechnology], MYHC [1:20, MF-20, DSHB], PERILIPIN A/B [1:100, Sigma-Aldrich], Sca-1-FITC [1:100, ebioScience], and α7-Integrin-PE [1:100; eBioscience]. Oil red O was used to stain lipids, as described in Mozzetta et al. (2013). Further experimental details are provided in the Supplemental Material.

**qRT–PCR**
Total RNA was extracted with Trizol, and 0.5–1 μg was retrotranscribed using the TaqMan reverse transcription kit [Applied Biosystems]. Real-time qPCR was performed using primers and conditions provided in the Supplemental Material.

**FAIRE**
Chromatin for FAIRE analysis was prepared as previously described in Albini et al. (2013). Further experimental details and primers used are provided in the Supplemental Material.

**ChIP**
ChIP assay was performed as previously described in Albini et al. (2013). The following antibodies were used: anti-acetylated Histone 3 [AcH3; Millipore] and normal rabbit IgG [IgG; Santa Cruz Biotechnology]. Further experimental details and primers for RT–PCR are provided in the Supplemental Material.

**miR HTS**
For miR screening, HEK293 cells were cotransfected with 10 nM Ambion pre-miR miR precursor human version 2.0 miR library and 100 ng of SMARCD 3′ UTR target sensor plasmids [Fig. 3A] per well in white 384-well plates (Greiner) in triplicate using Lipofectamine 2000 (Invitrogen) following the manufacturer’s protocol. Further experimental details are provided in the Supplemental Material.

**Gene expression analysis**
RNA from 1 × 10⁶ freshly sorted cells was extracted using the RNaseasy kit (Qiagen) following the manufacturer’s protocol. We analyzed Agilent mouse GE 4x44K version 2 (single-color) arrays using the manufacturer’s reader and collected primary data using the supplied scanner software. Further experimental details on data analysis are provided in the Supplemental Material.

**NA-seq**
The library for NA-seq was prepared as described in Gargiulo et al. (2009). Further details on data analysis are provided in the Supplemental Material.

**Small RNA-seq**
C57Bl10 mdx mice [1.5 mo old] were treated for 15 d with TSA. RNA was harvested from freshly isolated FAPs using Trizol reagent, and small RNA libraries were prepared for sequencing using standard Illumina protocols. Further experimental details are provided in the Supplemental Material.

**Statistical analysis**
Data are presented as mean ± SEM. Comparisons were made using the Student’s t-test assuming a two-tailed distribution, with significance being defined as P < 0.05 (*), P < 0.01 (**), and P < 0.001 (***).

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HDAC-regulated myomiRs control BAF60 variant exchange and direct the functional phenotype of fibro-adipogenic progenitors in dystrophic muscles

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