Lineage of origin in rhabdomyosarcoma informs pharmacological response

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Lineage or cell of origin of cancers is often unknown and thus is not a consideration in therapeutic approaches. Alveolar rhabdomyosarcoma (aRMS) is an aggressive childhood cancer for which the cell of origin remains debated. We used conditional genetic mouse models of aRMS to activate the pathognomonic Pax3:Foxo1 oncogene and inactivate p53 in several stages of prenatal and postnatal muscle development. We reveal that lineage of origin significantly influences tumor histomorphology and sensitivity to targeted therapeutics. Furthermore, we uncovered differential transcriptional regulation of the Pax3:Foxo1 locus by tumor lineage of origin, which led us to identify the histone deacetylase inhibitor entinostat as a pharmacological agent for the potential conversion of Pax3:Foxo1-positive aRMS to a state akin to fusion-negative RMS through direct transcriptional suppression of Pax3:Foxo1.

[Keywords: alveolar rhabdomyosarcoma, Pax3:Foxo1, sarcoma, satellite cell, myoblast, histone]

Supplemental material is available for this article.

Received January 22, 2014; revised version accepted June 3, 2014.
intrinsic are possible nonmyogenic cells of origin of embryonal rhabdomyosarcoma (eRMS) from the adipogenic lineage [Hatley et al. 2012; Kikuchi and Keller 2012]. For the alveolar subtype that is often incurable when metastatic [Malempati and Hawkins 2012; Hawkins et al. 2013], we previously suggested Myf6-expressing (differentiating) cells of the myogenic lineage as the cell of origin [Keller et al. 2004a,b]. These studies had used conditional mouse models triggering in the Myf6Cre lineage the pathognomonic Pax3:Foxo1 chimeric oncogene, which is commonly found in the human disease as a result of a t(2;13) translocation [Arndt and Crist 1999]. Nevertheless, we and others have reconsidered whether mesenchymal stem cells or satellite cells could be an alternate cell of origin [Ren et al. 2008; Charytonowicz et al. 2009; Hettmer and Wagers 2010], and thus we performed the studies described here.

An interesting aspect of the search for cell of origin is that we uncovered a differential susceptibility of the Pax3:Foxo1 locus to be transcribed based on the cell lineage originally transformed. This finding may have translational significance in that the related Pax7 locus in myogenic progenitors is a classic example of a bivalent epigenetic locus [Mozzetta et al. 2011], with different transcriptional activation states depending on degree of myodifferentiation, and thus is potentially amenable to pharmacological intervention. Transcription factors have not typically been considered approachable therapeutic targets, but if transcription of Pax3:Foxo1 itself could be inhibited, then the implicit conversion of fusion-positive aRMS to fusion-negative RMS would have great appeal, given the substantial difference in outcomes between these two clinical groups in retrospective studies [Sorensen et al. 2002; Missiaglia et al. 2012].

Results

The p53 pathway is frequently aberrant in aRMS

We and others have commonly used p53 inactivation in mouse models of aRMS [Keller et al. 2004a], yet the clinical relevance of p53 deregulation at the genetic and/or functional pathway levels is debated [Takahashi et al. 2004; Ognjanovic et al. 2012]. To address this issue, we used metagene analysis to test whether p53 functional inactivation was a clinically relevant cooperative initiating mutation in aRMS [Supplemental Fig. S1A]. Using our previously described metagene analysis and S-score method [Rubin et al. 2011], we analyzed a global gene expression data set [Supplemental Tables 1, 2] of 62 Pax3:Foxo1+ and 24 Pax7:Foxo1+ human aRMS tumors for aberrant signaling of the rhabdomyosarcoma-associated p53 pathway. We found that 85% of Pax3:Foxo1+ tumors exhibited a gene expression signature consistent with the “p53 off” state. Similarly, 75% of Pax7:Foxo1+ tumors also had a p53 off state. Thus, the p53 off state was a common signature in human samples.

Pax3:Foxo1 leads to forms of sarcoma for satellite cells different from any other myogenic lineage

Having established a prominent role for p53 pathway inactivation and Pax3:Foxo1 activation as driver events in the genesis of aRMS, we generated mouse models [Keller et al. 2004a] to mimic these initiating driver events through targeting to specific muscle cell types in fetal and postnatal development (Fig. 1A). Tumors occurred in embryonic muscle lineages [McCre for hypaxial Pax3] and embryonic and fetal muscle lineages [Myf6Cre for Myf6] as well as the postnatal satellite cell lineage [Pax7CreER] (Fig. 1B,C). For the latter, tamoxifen was administered at 30 d of age [adolescence]. The Myf5Cre lineage was embryonic-lethal [usually resulting in exencephaly] (data not shown) in allitters except for one animal [U24014] that later went on to develop a tumor of the lower extremity at 74 d of age. The most susceptible lineages were fetal myogenic progenitors and postnatal satellite cells but not Myf5- or Myf6-expressing postnatal committed myogenic progenitors: Of 18 Myf5CreER, Pax3:Foxo1, p53 mice given tamoxifen at 30 d of life [P30 [postnatal day 30]] and observed 43–510 d [median, 395], only one developed at age 401 d a tumor that was diagnosed as aRMS solid variant of the cranial muscle (Supplemental Fig. S1B–D). Similarly, from eight Myf6CreER, Pax3:Foxo1, p53 mice given tamoxifen at P30 and observed 232–505 d [median, 396], no mice developed tumors. Postnatal lineage tracing of the postnatal Myf6CreER lineage showed marking of myofibers as well as Pax7+ subliminar satellite cells [Supplemental Fig. S2], a postnatal result complementing a report that embryonic Myf6Cre might prime Pax7+ satellite cells [Sambasivan et al. 2013]. However, despite a recent report showing that Myf6Cre is active in early embryonic muscle progenitors that can become satellite cells [Sambasivan et al. 2013], other evidence for the Myf6Cre fetal myoblast population being a more common cell of origin for Pax3:Foxo1 aRMS than a postnatal satellite cell includes [1] the absence of tumors from Myf6CreER, Pax3:Foxo1, p53 mice in which Myf6+ satellite cells can activate Pax3:Foxo1, [2] the absence of aRMS tumors from Pax7CreER, Pax3:Foxo1, p53 mice in which the Pax7+ satellite cell population at large can activate Pax3:Foxo1, and [3] the absence of aRMS tumors from Myf5CreER, Pax3:Foxo1, p53 mice whose nonquiescent satellite cells can activate Pax3: Foxo1 [Biessi et al. 2013].

To address the reported possibility that nonmyogenic, marrow-derived cells might give rise to aRMS—at least when Pax3:Foxo1 is driven by a nonnative promoter [Ren et al. 2008]—we first examined cells of the bone marrow compartment for the presence of cells derived from a Myf6Cre or Pax3CreKI [Pax3] [Engleka et al. 2005] ontogeny using eYFP [Keller et al. 2004a] and RFP [Vintersten et al. 2004] reporters, respectively. This lineage tracing was done without introducing Pax3:Foxo1 or p53 mutations. Neither the Pax3 nor the Myf6 lineage was represented in the bone marrow compartment in more than 0.4% of mesenchymal or hematopoietic cells [Supplemental Fig. S1E,F], and thus this line of experimentation was discontinued.

At the histological level, the embryonic McCre [hypaxial Pax3Cre] and fetal Myf6Cre lineages led to aRMS histology (Fig. 1D). The single animal that survived past birth for the Myf5Cre lineage [U24014] also developed...
Figure 1. Characteristics of tumors from Pax7CreER, Myf5Cre, Myf6Cre, and MCCre mice. (A) Representation of prenatal and postnatal myogenesis in the context of Cre drivers used to trigger tumors in this study. For Pax7CreER, Myf5CreER, and Myf6CreER mice, tamoxifen was administered at P30 (see the Materials and Methods). [multinuc.] Multinucleated. (B) Pax7CreER mice and MCcre mice showed prolonged survival and lower tumor incidence compared with Myf6Cre mice (tumor incidences were MCcre, 40%; Pax7CreER, 38%; and Myf6Cre, 100% at 1 yr after birth). Myf5Cre mice were embryonic-lethal [see the text]. All mice also harbor homozygous Pax3P3Fm/P3Fm and p53F2-10/F2-10 alleles. (C) Location and disease stages in each strain. Pax7CreER tumors were more frequently observed in head/neck and genital areas, whereas MCcre mice developed tumors exclusively in the extremities. The highest frequency of stage IV metastasis (40%) was seen for Myf6Cre mice. The IRSG staging system was scaled to mice such that a 4-mm-diameter mouse tumor was equated with a 5-cm-diameter human tumor [Nishijo et al. 2009a]. (D, top row) Histological analysis of tumors. Myf5Cre, Myf6Cre, and MCcre tumors were small round cells [aRMS], whereas most Pax7CreER tumors showed spindle cell morphology [pleomorphic RMS/pleomorphic spindle cell sarcoma]. (Bottom row) Immunohistochemistry for Myogenin. (E) eYFP protein expression representing the transcriptional activation status of Pax3:Foxo1 fusion gene was at the lower limit of detection in Pax7-CreER tumors. Bars, 10 μM.
aRMS with numerous rhabdoid-type rhabdomyoblasts and diffuse anaplasia (Fig. 1D). In contrast, the Pax7CreER postnatal satellite cell lineage led to tumors diagnosed as pleomorphic RMS/pleomorphic spindle cell sarcoma by histological and immunohistochemical criteria (Fig. 1D). Surprisingly, the eYFP marker of Pax3:Foxo1 was expressed only weakly in Pax7CreER-derived tumors compared with the other tumors (Fig. 1E). Correspondingly, Pax3:Foxo1 expression by RT–PCR was variable and generally low in Pax7CreER tumors compared with mice bearing the Pax3:Foxo1 conditional alleles and other Cre drivers (Fig. 2A).

To begin exploring the potential causes of the low Pax3:Foxo1 expression from Pax7CreER,Pax3,p53 tumors, we evaluated the Cre/LoxP recombination efficiency at the Pax3 locus. By genotyping tumors, Pax3 locus recombination frequency was found to be lower in tumors of Pax7CreER-expressing satellite cell origin than Myf6Cre-expressing prenatal (presumably myoblast) origin. Generally Pax7CreER tumors had only one Pax3 conditional knock-in allele, P3Fm, converted to the active Pax3:Foxo1 form, P3Fa [Fig. 2B; see the Discussion]. Recombination for the p53 locus by Pax7CreER in tumors was uniformly complete for both p53 alleles. Furthermore, the lower Pax3 recombination frequency alone would not account for the difference in histology because Pax3 [P3Fa/WT] tumors from the Myf6Cre lineage presented with an aRMS histology, not a pleomorphic histology (Fig. 2C; Supplemental Table 3). By quantitative analysis, the mean florescence intensity of eYFP in primary tumor cell cultures derived from Pax7CreER,Pax3[P3Fm/P3Fa] tumors was rarely above the level of C2C12-negative control cells except in rare instances (Fig. 2D; see the Discussion). On the other hand, mean florescence intensity of eYFP in primary cell cultures derived from a Myf6Cre,Pax3[P3Fa/P3Fa] tumor was twofold to threefold higher than control C2C12 cells (Fig. 2D). RT–PCR studies also showed higher levels of Pax3:Foxo1 mRNA levels in murine primary tumor cell cultures derived from Myf6Cre,Pax3[P3Fa/P3Fa] tumors [Fig. 2E, green] compared with primary tumor cell cultures from Pax7CreER,Pax3[P3Fa/P3Fa] tumors [U31425 and U28285] (Fig. 2E). Altogether, these results suggest that inherently less transcription occurs from the Pax3 locus in Pax7CreER-derived tumors and decrease the probability that aRMS arises from Pax7+ postnatal satellite cells.

Tumors arising from non-Pax7CreER cell of origin authentically recapitulate human aRMS

To further evaluate the lineages that might give rise to aRMS, we compared expression of validated markers of Pax:Foxo1+ aRMS and eRMS [Wachtel et al. 2006]. Myf6Cre myoblast-derived mouse tumors expressed significantly higher levels of the Pax3:Foxo1-specific aRMS marker TCFap2b, whereas Pax7CreER satellite cell-derived tumors expressed reciprocally higher levels of the eRMS-specific marker Fbn2 [Fig. 3A]. We then turned to principal component analysis [PCA] to compare Pax3:Foxo1-bearing tumors with those in the eRMS–UPS spectrum [Rubin et al. 2011]. Results showed that Pax7CreER satellite cell-derived tumors were highly similar to eRMS/UPS tumor models that we previously reported [Fig. 3B; Rubin et al. 2011]. In contrast, all Pax3:Foxo1-bearing tumors histologically consistent with aRMS from the MCre [Pax3], Myf5Cre, and Myf6Cre lineages formed an independent outgroup [Fig. 3B]. Thus, the Pax7CreER postnatal satellite cell lineage did not appear to be poised for giving rise to aRMS when the Pax3:Foxo1 oncogene at the Pax3 locus is triggered; instead, cells of Pax7CreER satellite cell lineage had a tendency to give a spindle cell-like tumor phenotype. We next hypothesized that lineage-specific drug sensitivity could be best interrogated by an ontogeny series that would include Pax7CreER tumors for which Cre-mediated recombination or transformation occurred after the satellite cell state. From Figure 3B, we expected an intermediate state to be rare, but from the Pax3:Foxo1 data in Figure 2E (i.e., samples U28285, U31423, and U29416), we expected at least some instances to exist. To identify a potential ontogeny series, we first sought to find model primary cell cultures best representing Pax3:Foxo1-harborig satellite cell [Pax7CreER lineage]-derived tumors versus Pax3:Foxo1-harborig myoblast [Myf6Cre lineage]-derived tumors. Comparison of gene expression for these two tumor types is given in Supplemental Table 4. The Pax7CreER lineage was associated with expression of Cav1, Dcn, and Cxcl12, whereas the Myf6Cre lineage was associated with expression of Sct, Stc1, and Ryr3 (Supplemental Table 4). We choose five primary cell cultures—i.e., U29415, U31425, U28285, U31423, and U29416—of Pax7CreER lineage and U20325, U23674, and U21459 of Myf6Cre lineage to test the expression of markers associated with Pax7CreER lineage was highest in U29415 cells followed by U31425 or U28285 and lower expression in Myf6Cre lineage primary cell cultures U20325, U23674, and U21459. Conversely, the expression of markers associated with Myf6Cre lineage showed a reverse gradient pattern where the highest expression was observed in U21459/U23674, followed by U29416 or U20325, U31423, U28285, U31425, and U29415 cells (Supplemental Fig. S3A). These results suggested that inherently less transcription occurs from the Pax3 locus in Pax7CreER-derived tumors and decrease the probability that aRMS arises from Pax7+ postnatal satellite cells.

Alveolar rhabdomyosarcoma cell of origin
satellite cell lineage), suggesting that the cell of origin of the sarcoma might play a role in deciding the treatment strategy for patients. Similar results were observed for the FGFR1–4 inhibitor, LY287445, and Wee1 inhibitor, MK-1775, whereby \textit{Myf6Cre} lineage-derived U23674 was found to be the most sensitive to these two drugs compared with intermediate ontogeny cultures U29416 and U31423 and \textit{Pax7CreER} satellite cell lineage-derived U29415 (Supplemental Fig. S3C,D). Interestingly, when \textit{Pax3:Foxo1} was stably knocked down in U23674 cells

\[ \text{Figure 2. Recombination status of \textit{Pax3} allele differs by cell of origin.} \]

\[ \text{A] Relative expression of \textit{Pax3:Foxo1} in mouse tumor tissue.} \]

\[ \text{B] Genotyping for \textit{Pax3:Foxo1} and p53 recombination by cell of origin (Cre driver). Satellite cell-derived (\textit{Pax7CreER}) primary tumor cell cultures showed reduced efficiency in the recombination of \textit{Pax3:Foxo1} knock-in allele locus compared with myoblast-derived primary tumor cell cultures (\textit{Myf6Cre}). However, p53 locus conditional alleles were uniformly inactivated in all tumors irrespective of Cre driver.} \]

\[ \text{C] Representative H&E staining of tumors with Cre driver and \textit{Pax3} locus genotypes as noted. All tumors had homozygous conditional p53 deletion. \textit{Myf6Cre} lineage tumors with heterozygous or homozygous activation of \textit{Pax3:Foxo1} showed small round cell histology consistent with aRMS. In contrast, \textit{Pax7CreER} tumors with heterozygous or homozygous activation of \textit{Pax3:Foxo1} exhibited pleomorphic spindle cell appearances, similar to tumors with only p53 inactivation—neither of which appearance is consistent with aRMS.} \]

\[ \text{D] Flow cytometry of eYFP for \textit{Pax7CreER}- and \textit{Myf6Cre}-derived primary tumor cell cultures. The mean florescence intensity was lower in \textit{Pax7CreER} rhabdomyosarcoma cells, although select tumors of the \textit{Pax7CreER} lineage (e.g., U31423 and U29416) more resembled \textit{Myf6Cre} in eYFP expression (see the Results).} \]

\[ \text{E] Quantitative RT–PCR (qRT–PCR) studies showing higher levels of \textit{Pax3:Foxo1} mRNA levels in \textit{Myf6Cre}-derived primary tumor cell cultures (U23674 and U57844) compared with \textit{Pax7CreER}-derived primary tumor cell cultures (U29415, U31425, and U31423). Statistically significant difference in \textit{Pax3:Foxo1} expression was observed between \textit{Myf6Cre}- and \textit{Pax7CreER}-derived primary tumor cell cultures with one activated \textit{Pax3:Foxo1} (P3Fa) allele (U57844 and U29415) versus primary tumor cultures with two activated \textit{Pax3:Foxo1} alleles [U23674 and U31425].} \]
(U23674 ShY08), there was no difference in their sensitivity to either SNS-032 (Fig. 3E) or PD0332991 (Fig. 3F) compared with the control cells (U23674 ShC01), further supporting that differential drug sensitivity was due to lineage of origin (i.e., epigenetic memory) and not Pax3:Foxo1 expression level.

The Pax7 and Pax3:Foxo1 loci are epigenetically regulated [differential effects of epigenetic modifiers on Pax3:Foxo1 and Pax7 expression depending on the tumor cell of origin]

We next reasoned that if tumor initiation and tumor phenotype is epigenetically determined, then the tumor phenotype can be epigenetically modified by pharmacological agents. The Pax7 locus in activated satellite cells has a bivalent chromatin domain with the presence of both positive (H3K4me3) and negative (H3K27me3) histone marks (Mozzetta et al. 2011), and we hypothesized that the Pax3 locus may be similar. To gain insight into the epigenetic regulation of the Pax3:Foxo1 locus, we examined active (H3K4me3 and H3K9Ac) and repressive (H3K27me3 and H3K9me3) histone modification marks by chromatin immunoprecipitation (ChIP) in tumor cells representing both Pax7CreER (U29415 and U31423) and Myf6Cre (U23674 and U21459) lineages of origin. For this study, we analyzed two genomic regions upstream (−193:−285) of and downstream (+593:+462) from the transcriptional start site of the targeted loci.

Figure 3. Pax3:Foxo1 and aRMS-specific marker expression in Pax3:Foxo1 tumors. (A) For tumors bearing conditional Pax3:Foxo1 alleles, expression of aRMS-specific classifier Tcfap2b was significantly increased in Myf6Cre-derived tumors (M6-P3F-p53) and comparatively lower in Pax7CreER-derived tumors (P7er-P3F-p53); conversely, expression of cRMS-specific marker Fibrillin2 (Fbn2) was increased in Pax7CreER-derived, Pax3:Foxo1-bearers tumors and comparable with the level of Fbn2 from cRMS derived from satellite cells (P7er-Ptch1-p53) or myoblasts (M6-p53). (B) PCA showing the non-satellite cell-derived, Pax3:Foxo1-bearing tumors as an outgroup and Pax7CreER-derived, Pax3:Foxo1-bearing tumors grouping with spindle cell sarcoma/cRMS tumors bearing Ptch1 and p53 mutations. (C,D) Cell viability assay showing differential drug response of primary tumor cell cultures along the continuum of satellite cell-derived [Pax7CreER] origin versus myoblast-derived [Myf6Cre]. (C) For the Cdk-2, Cdk-7, and Cdk-9 inhibitor SNS-032, the IC50 values were 0.24 μM for U23674, 0.83 μM for U29415, 1.79 μM for U31423, and 2.61 μM for U29415. (D) For the Cdk-4,-6 inhibitor PD0332991, the IC50 values were 2.69 μM for U23674, 3.61 μM for U29415, 4.77 μM for U31423, and 8.15 μM for U29415. (E,F) Cell viability assay showing no difference in the response of U23674 ShY08 cells [Pax3:Foxo1 stable knockdown cells] to SNS-032 [E] and PD0332991 [F] compared with the control knockdown cells, U23674 ShC01. For SNS-032, the IC50 values were 0.1305 μM for ShC01 cells and 0.1730 μM for ShY08 cells. In the case of PD0332991, the IC50 values were 6.90 μM for ShC01 cells and 6.88 μM for ShY08 cells. (s.c.) Satellite cell origin, (m.b.) myoblast origin.
tional start site (TSS) of the Pax3 locus. At the level of histone marks, we observed a different chromatin state of the Pax3:Foxo1 locus in primary tumor cell cultures with different lineages of origin. As shown in Supplemental Figure S4, higher levels of both H3K4me3 and H3K9Ac active chromatin marks are observed in U23674 and U21459 cells compared with U29415 and U31425 tumor cells, correlating with high versus low Pax3:Foxo1 expression, respectively. Also high levels of repressive histone marks H3K27me3 and H3K9me3 were observed in U29415 and U31425 cells compared with U23674 and U21459 cells. Altogether, these data suggest that the observed differences in Pax3:Foxo1 mRNA levels could be explained by a different chromatin state in these loci depending on the tumor lineage of origin. From the RT–PCR studies described above, U29415 was selected as a representative Pax7CreER lineage tumor culture, and U23674 was selected as a representative Myf6Cre lineage tumor culture.

To determine whether the Pax locus could be pharmacologically modified, we first treated murine primary tumor cell cultures U23674 and U29415 with varying doses of 5-Aza 2′-deoxycytidine (DNA methyltransferase inhibitor) or SAHA and entinostat [histone deacetylase (HDAC) inhibitors] to determine a sublethal dose at which cells could be treated to study the effects of these drugs on expression of Pax3:Foxo1 and Pax7 (Supplemental Fig. S5). Cells were treated with drugs for 72 h, after which an MTS assay was performed to assess viability. Based on these studies, U23674 cells were treated with 3 μM 5-Aza-2′-deoxycytidine, 2 μM entinostat, or 0.75 μM SAHA for 24 h, after which RNA was extracted, and the expression of Pax3:Foxo1 and Pax7 was examined by RT–PCR. For U23674 tumor cells of the Myf6Cre myoblast lineage, a decrease in both Pax7 and Pax3:Foxo1 mRNA levels upon treatment with all of the drugs was observed, but the effect was more pronounced for HDAC inhibitors [Fig. 4A]. Interestingly, U29415 tumor cells of Pax7CreER satellite cell lineage showed a paradoxical increase in mRNA levels of Pax7 and Pax3:Foxo1 when treated with 20 μM 5-Aza-2′-deoxycytidine, 15 μM entinostat, and 5 μM SAHA [Fig. 4A]. As expected, the rare tumor cultures downstream from U29415 behaved similarly to the Myf6Cre tumor cultures [Supplemental Fig. S6]. We next determined epigenetic marks of Pax3:Foxo1 by performing ChIP and examining the active histone marks H3K4me3 and H3K9Ac along with the repressive marks H3K27me3 and H3K9me3 previously examined in Supplemental Figure S4. However, examination of these specific chromatin marks did not explain the change in Pax3:Foxo1 transcript levels, implying that other regions may be the targets of chromatin remodeling for entinostat.

Previous studies in mdx mice treated with a TNF-α antibody reported increased levels of Pax7 mRNA in satellite cells derived from myofibers [Palacios et al. 2010]. TNF-α has been shown to activate p38α kinase, which leads to interaction between YY1 and PRC2 (polycomb-repressive complex 2) via threonine phosphorylation of EZH2. These proteins together form a repressive complex at the Pax7 promoter and thus regulate the expression of Pax7 [Palacios et al. 2010]. To investigate whether TNF-α can regulate the expression of Pax7 in U23674 and U29415 tumor cells, cultures were treated with 10 ng/ml recombinant TNF-α for 24 h. RT–PCR data showed a reduction in mRNA levels of Pax7 in both U23674 and U29415 cells. TNF-α also reduced Pax3:Foxo1 levels in both cell cultures (Fig. 4A). We next examined the levels of Pax3:Foxo1 at the protein level. By immunoblotting, concomitant reduction in Pax3:Foxo1 levels in U23674 cells treated with 2 μM entinostat for 24 h was observed [Fig. 4B]. This effect was seen across species for human aRMS cell lines Rh30 (24 h of treatment) and Rh41 [24 h of treatment] (Fig. 4C,D). When murine and human tumor cultures were treated with 2 μM entinostat for 72 h, a pronounced reduction in Pax3:Foxo1 protein levels was observed [Fig. 4E,F]. These results showed in principle that Pax3:Foxo1 and Pax7 transcription and protein levels can be modified pharmacologically. To investigate whether the reduction in Pax3:Foxo1 protein levels upon entinostat treatment is mediated by changes in post-translational stability attributable to direct HDAC–Pax3:Foxo1 interactions, we treated human and murine aRMS cultures [Rh41 for 24 h and U23674 for 72 h] with 2 μM entinostat alone or in combination with MG132 [proteasome inhibitor]. No increase in Pax3:Foxo1 protein level was observed when cells were treated with a combination of entinostat and MG132 compared with cells treated with entinostat alone [Fig. 4G,H]. This result suggests that the reduction in Pax3:Foxo1 levels observed upon entinostat treatment is mediated at the transcriptional [or post-transcriptional] level and not at the post-translational level. Other studies from our laboratory have shown that expression of Pax3:Foxo1 is G2 cell cycle phase-dependent [Kikuchi et al. 2014]. To investigate whether the reduction in Pax3:Foxo1 expression caused by entinostat treatment is cell cycle-dependent, we performed a series of studies suggesting that this was not necessarily the case (Supplemental Fig. S7).

Entinostat slows tumor growth in vivo

We next tested the anti-tumor efficacy of entinostat in an orthotopic allograft model of rhabdomyosarcoma generated by injecting U23674 cells into the cardiotoxin-preinjured gastrocnemius muscle in the right leg of SCID-hairless [SHO-Ptkd<sup>actae</sup> H<sup>actae</sup>] mice. Once the tumors reached 0.25 cm<sup>3</sup>, the mice were treated with a daily dose of 10 mg/kg entinostat by intraperitoneal injection, with 0.25 mg/kg actinomycin D by intraperitoneal injection on day 1 of treatment, a combination of entinostat and actinomycin D, or vehicle [DMSO] (Fig. 5A,B). Actinomycin D is one of the commonly used chemotherapeutic agents for treating rhabdomyosarcoma. The tumors in the entinostat cohort grew significantly slower than either vehicle- or actinomycin D-treated mice (P = 0.03). Mice treated with a combination of entinostat and actinomycin D had a significantly slower rate of tumor growth, and their tumors took more time to reach 1 cm<sup>3</sup>
Figure 4. Pharmacologically modified epigenetic regulation of Pax3:Foxo1. (A) RT–PCR assays showing relative expression of Pax3:Foxo1 and Pax7 in representative satellite cell-derived U29415 and myoblast-derived U23674 murine primary tumor cell cultures upon treatment with DMSO, 5-Aza-2′-deoxycytidine [Aza], SAHA, entinostat [Ent], serum-free medium (SFM), and the cytokine TNF-α (10 ng/mL in serum-free medium) for 24 h. The U29415 primary tumor cell culture (satellite cell origin) showed an increase in Pax7 and Pax3:Foxo1 expression upon treatment with 5-Aza-2′-deoxycytidine, SAHA, and entinostat for 24 h, whereas the U23674 primary tumor cell culture (myoblast origin) showed a decrease in the mRNA levels of Pax7 and Pax3:Foxo1 upon treatment. However, treatment with TNF-α showed a reduction in the expression of Pax7 and Pax3:Foxo1 in both U29415 and U23674 cells. (B) Immunoblot showing reduction in Pax3:Foxo1 levels upon treatment with 2 μM entinostat [Ent] and also with 3 μM 5-Aza-2′-deoxycytidine [Aza] for 24 h in U23674 cells. (C) Reduction in Pax3:Foxo1 level upon treatment with 2 μM entinostat [Ent] for 24 h in Rh30 cells. (D) Western blot showing reduction in Pax3:Foxo1 levels upon treatment with 2 μM entinostat [Ent] for 24 h in Rh41 cells. (E) Immunoblot showing significant reduction in Pax3:Foxo1 levels upon treatment with 2 μM entinostat [Ent] for 72 h in U23674 cells [left panel] and the graphical representation [right panel]. (F) Immunoblot showing a significant reduction in Pax3:Foxo1 level upon treatment with 2 μM entinostat [Ent] for 72 h in Rh30 cells [left panel] and the graphical representation for this blot [right panel]. (G) Western blots showing no increase in Pax3:Foxo1 levels in Rh41 cells when they were treated with a combination of the proteasome inhibitor MG132 (0.1 μM, 0.5 μM, and 1 μM) and 2 μM entinostat for 24 h compared with cells treated with entinostat alone. Expression of Noxa was observed when cells were treated with a combination of MG132 and entinostat in Rh41 cells. (H) Western blot showing no increase in Pax3:Foxo1 levels in U23674 cells even when these cells were treated with a combination of 200 nM MG132 and 2 μM entinostat for 72 h compared with cells treated with 2 μM entinostat alone. Noxa could not be detected even after treating the cells with a combination of MG132 and entinostat for 72 h.
concluded that the four curves are significantly different \((P < 0.001)\) [Fig. 5 A,B]. However, after 5 d of treatment with entinostat, the mice showed symptoms of drug toxicity (>10% loss of body weight), and hence no treatment was administered on day 6, and the subsequent daily dose of entinostat was reduced to 5 mg/kg.

**Discussion**

In our studies, we sought to refine experimentally the range of possibilities for the cell of origin of Pax3:Foxo1+ rhabdomyosarcoma. Parenthetically, however, the cell of origin for Pax7:Foxo1+ aRMS is in no way addressed in this study. Surprisingly, in our experiments, the fetal myoblast [Myf6 lineage] appears most susceptible to formation of metastatic aRMS. In contrast, muscle stem cells [satellite cells] appear least poised to give rise to rhabdomyosarcoma—at least from adolescence onward. With regard to tumor initiation, our data suggest that the Pax3 chromatin structure may be closed [and relatively inaccessible to events like Cre/LoxP DNA recombination] and show that transcription from the Pax3 locus is inactive in tumor cells of this satellite cell lineage. However, this low transcriptional activity could be reversed by pharmacological epigenetic modifiers. Even more importantly, Pax3:Foxo1 can be reduced at the mRNA and protein level in myoblast-derived tumors by the same pharmacological epigenetic modifiers. The implication that cell of origin can influence pharmacological response is interesting, if not provocative. We concede, however, that common drug sensitivities are likely also found between tumors of different origins and that while some cell biological properties that may be transmitted from the cell of origin, as many or more may be the result of convergent, facultative phenotypes for tumor cell growth—even if rhabdomyosarcomas arise from different cell lineages.

What is remarkable is that a prenatal [Myf6Cre] lineage of origin should, in our comparison, be most capable of forming aRMS [at least in the context of p53 loss of function]. Rhabdomyosarcoma is a disease that has a steady incidence throughout childhood [aRMS contrasts dramatically with eRMS, the latter of which peaks with toddler and adolescent growth spurts] [Ognjanovic et al. 2009]. Suggestion of Myf6+ prenatal or postnatal myogenic cells, possibly a terminally differentiating subpopulation, as an aRMS cell of origin was first reported by our group [Keller et al. 2004a,b] and later complemented by in vitro studies of human postnatal myoblasts transformed by Pax3:FoxO1 and other cooperative initiating mutations [Naini et al. 2008]. Congenital fusion-positive aRMS is rarely reported, and congenital aRMS is generally fusion-negative [Godambe and Rawal 2000; Grundy et al. 2001; Sueters et al. 2005], although the Pax3:FoxO1 translocation has been detected in rare neonatal instances [Rodriguez-Galindo et al. 2001]. What might account for the delay? One explanation might be an accumulation of other genetic changes [mutations], but epigenetic changes during childhood and adolescent growth and development might also have a role to play. In fact, epigenetic abnormalities are very common in human cancers—and, in contrast to genetic mutations, these epigenetic changes are potentially reversible.

Can cells other than those of skeletal muscle lineages give rise to aRMS? We are increasingly intrigued by this possibility, although the question needs to be addressed with prudence. A mesenchymal stem cell origin for aRMS has been suggested [Charytonowicz et al. 2009; Hettmer and Wagers 2010]. In support of this possibility,
clinical cases of metastatic tumors without primary tumors have been reported for t(1;13) PAX3:FOXO1 aRMS [Lisboa et al. 2008] and t(2;13) PAX3:FOXO1 aRMS (Sandberg et al. 2001; Lisboa et al. 2008). An experimental study by Ren et al. (2008) tested the possibility that constitutively expressed (ectopic) PAX3:Foxo1 could create aRMS, but this study did not place PAX3:Foxo1 under the authentic control of the PAX3 promoter, which, in light of our studies of a very small (0.4%) bone marrow compartment cell population that had ever expressed Pax3, makes the probability of aRMS arising from mesenchymal stem cells very low. However, a non marrow mesenchymal stem cell of origin remains possible.

An interesting alternative cell of origin for aRMS is the pericyte or mesangioblast. Vascular mimicry in rhabdomyosarcoma suggests a plasticity that may (or may not) have carried over from an angioblast origin [Sun et al. 2004]. An interesting case of primary rhabdomyosarcoma of the pulmonary artery was also recently reported [Suman et al. 2013]. Experimentally, Lagha et al. [2009] reported that in the mouse embryonic dermomyotome, the ratio of Pax3 to Foxc2 balances fate between myogenic and vascular cell fates but that PAX3:Foxo1 suppresses Foxc2. This latter result suggests that, in the susceptible cell, an angiogenic precursor might be converted to a myogenic state, a result that has been shown experimentally for embryonic aortic mesangioblasts [Messina et al. 2009]. The link between vascular precursors and the Pax3 lineage does not come as a surprise given the known role of Pax3 in the migration of cardiovascular neural crest during embryogenesis [Conway et al. 1997]. In a provocative study, Goupille et al. [2011] have shown that smooth muscle cells of arteries express Pax3 and are able to contribute to skeletal muscle fiber formation via in vitro coculture conditions and following cell–cell fusion. This plasticity lends itself to the possibility that mesangioblasts or even smooth muscle cells can transform into myogenic phenotype cancers (rhabdomyosarcomas) under select conditions.

Our results raise a potentially important clinical implication by suggesting that PAX3:FOXO1 levels vary by cell of origin, even adjusting for recombination frequency; thus, PAX3:FOXO1 RT–PCR alone may be less sensitive than cytogenetics or FISH to detect the PAX3:FOXO1 rearrangements. Could some MFH or UPS cases be in fact more than cytogenetics or FISH to detect the PAX3:FOXO1 rearrangements, but that smooth muscle cells of arteries express Pax3 and are able to contribute to skeletal muscle fiber formation via in vitro coculture conditions and following cell–cell fusion. This plasticity lends itself to the possibility that mesangioblasts or even smooth muscle cells can transform into myogenic phenotype cancers (rhabdomyosarcomas) under select conditions.

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Materials and methods

Microarray analysis for S scores and PCA

Human microarray data sets were previously published [Wachtel et al. 2004; Lac et al. 2007; Davicioni et al. 2009]. Patient demographics are presented in Supplemental Table 1, A and B. Metagene and S-score analyses were conducted as previously described [Rubin et al. 2011]. For mouse tumors, gene expression analysis was performed using Illumina’s Mouse Ref 6 Beadchip version 1 [Illumina, Inc.]. These data sets have been deposited in Gene Expression Omnibus as accession entry GSE22520 and are also described in Supplemental Table 1C. Rank invariant normalization was performed on the log2 transformed expression value. Afterward, we applied PCA to all mouse tissue samples with the 19,070 probes (selected with the criterion of average log2 intensity >5.5 and standard deviation >0.1) and then plotted in three-dimensional space for visualization (Fig. 4B). Similar microarray data analysis and the PCA methods were previously described [Rubin et al. 2011]. All bioinformatics tasks were performed with MATLAB/Bioinformatics Toolbox (MathWorks, Inc.) unless otherwise noted.

We downloaded the public domain data sets for fusion-negative RMS reported by Davicioni et al. [2006] from https://array.nci.nih.gov/caarray/project/details.action?project.experiment.publicIdentifier=trich-00099 as well as aRMS rhabdomyosarcoma data sets from previously published reports [Wachtel et al. 2004; Lac et al. 2007]. These fusion-negative RMS and aRMS data sets were designated as the test samples, whereas normal skeletal muscle samples reported by Bakay et al. (2002) were used as the control group. We also downloaded signature-specific data sets as described below. All of the studies were performed on an Affymetrix U133A array platform [Affymetrix]. Sample IDs used are given in Supplemental Tables 1 and 2. Control samples of each examination are the eRMS samples derived from Rubin et al. [2011]. We selected 10 high positive, 10 low positive, and 10 negative control samples for each examination, whereas gene-wise t-tests were performed between the 20 positive controls and normal skeletal muscle in order to apply to S score. S score is a subtype scoring method to quantify each sample’s consistency. By using the S score, we can unambiguously identify sample status and the amplitude of pathways or biological processes that gene signatures have depicted. The detailed S-score method has been previously described [Rubin et al. 2011].

We used previously established gene lists [Rubin et al. 2011] to examine whether human fusion-negative RMS and aRMS tumors had evidence of p53 loss of function, Shh gain of function, pRb loss of function, or Ras activation. Gene signatures of p53 loss of function were derived from gene expression data sets in breast cancer [Miller et al. 2005]. We also downloaded data sets for medulloblastoma samples known to exhibit a Shh gain-of-function signature [Thompson et al. 2006]. We took homolog genes from Rb1 wild-type and homozygous Rb1-deleted fusion-negative mouse sarcomas to be pRb loss-of-function gene signature [Rubin et al. 2011]. For Ras activation, we used gene lists for the activated Ras signature of zebrafish CRMs [Langelau et al. 2007]. The details of obtaining gene signature of each case have been previously described [Rubin et al. 2011].
Mice

All animal procedures were conducted in accordance with the Guidelines for the Care and Use of Laboratory Animals and were approved by the Institutional Animal Care and Use Committee (IACUC) at the University of Texas Health Science Center at San Antonio or Oregon Health and Science University. The Myf6Cre, Myf5Cre, Pax7Cre, MCre, Pax7CreER, Pax3Fox, LUSEAP, LacZ reporter (Rosa26mitosin), Pax3-Cre (Pax3Cre), and conditional p53 mouse lines and corresponding genotyping protocols have been described previously (Sostano 1999; Marino et al. 2000; Jonkers et al. 2001; Keller et al. 2004a; Brown et al. 2005; Nishijo et al. 2009a,b). Myf5CreER have been reported (Biressi et al. 2013), and Myf6CreER mice will be reported in detail elsewhere but are also described in Supplemental Figure S2. For Pax7CreER, Myf5CreER, and Myf6CreER mice, tamoxifen was administered at P30 as previously reported [Keller et al. 2004a]. Tumor-prone mice were visually inspected every 2 d for tumors because of the fulminant onset in these models. Genotyping protocols for Pax3Fox and p53 were performed as described [Keller et al. 2004a].

Mouse primary tumor cell cultures

Mouse primary tumor cell cultures were generated from mouse tumors as described previously [Abraham et al. 2011].

Survival analysis and statistical methods

Kaplan-Meier survival analysis of the mice was performed with the endpoint being the development of tumors (i.e., disease-free survival). Survival plots were created using the survival package in R. Relative expression of Pax3Fox and p53 were compared using the Mann-Whitney test with the Hochberg correction for multiple testing (Fig. 4A). Statistical analyses were performed by Kruskal-Wallis test with the Hochberg correction for multiple testing [Fig. 4A]. Statistical analyses were performed using R 3.0.1 [The R Foundation for Statistical Computing, http://www.r-project.org].

RNA isolation and quantitative RT–PCR (qRT–PCR)

RNA isolation and RT–PCR from mouse tumors were performed as previously described [Nishijo et al. 2009a; Rubin et al. 2011]. Probe sets for mouse samples were Gapdh-Mm99999915_g1, Fbn2_Mm00515742_m1, and Tctap2b_Mm00493468_m1. Statistical analysis was performed by Kruskal-Wallis test with Tukey's multiple testing correction. For drug treatment studies, the myoblast-derived primary tumor cell culture U23674 was treated with either 750 nM SAHA, 3 μM 5-Aza-2'-deoxycytidine, or 2 μM entinostat, and the satellite cell-derived primary tumor cell culture U29415 was treated with either 5 μM SAHA, 20 μM 5-Aza-2'-deoxycytidine, or 15 μM entinostat for 24 h. After drug treatment, total RNA was isolated, and cDNA was synthesized as described above. The expression of Pax3Fox and Pax7 was determined by qRT–PCR using TaqMan primers and probes (mouse Pax7-Mm00834072_m1 on a StepOnePlus real-time PCR system from Applied Biosystems).

Histology, immunohistochemistry, and immunocytochemistry

For MyoD, Myogenin, and Desmin immunohistochemistry, staining was performed as previously described [Rubin et al. 2011]. Immunofluorescence for GFP has also been described previously [Nishijo et al. 2009a]. For immunocytochemistry, murine rhabdomyosarcoma primary tumor cell cultures (U23674) and human rhabdomyosarcoma cell lines (Rh41) were cultured in an eight-well CultureSlide (BD Falcon). The cells were washed with PBS, fixed with 4% paraformaldehyde, permeabilized with 0.25% Triton X-100, and incubated with the antibody overnight at 4°C. The primary antibodies used were mouse anti-Pax3 (R&D Systems), rabbit anti-phospho CDC2-Y15, and rabbit anti-phospho Histone H3 (Cell signaling Technology). After incubation, the cells were washed with PBS and incubated with fluorescein isothiocyanate-conjugated anti-mouse and anti-rabbit IgG (Invitrogen) for 1 h. The cells were then examined by using a Zeiss LSM700 confocal microscope and Zen 2010 imaging software (Carl Zeiss).

**Flow cytometry and cell cycle analysis**

cYFP expression in mouse rhabdomyosarcoma cells and bone marrow aspirates was assessed by flow cytometry. Cells were stained with propidium iodide at 20 μg/mL for dead cell elimination, followed by analysis on a FACSscan cytometer (Becton Dickinson). For cell cycle analysis, primary mouse tumor cell cultures were harvested, washed with PBS, and fixed in cold 70% ethanol for 20 min. After fixation, the cells were washed with PBS and then incubated with RNase A and 20 μg/mL propidium iodide. The cells were then analyzed by using an Influx flow cytometer (Becton Dickinson) and FlowJo software (Tree star, Inc.).

**Drug sensitivity assays**

Murine rhabdomyosarcoma primary tumor cell cultures were plated in a 96-well plate at 2500 per well. After 24 h, the cells were incubated with varying concentrations of entinostat [Selleckchem], SAHA (LC Laboratories), and 5-Aza-2'-deoxycytidine [Sigma-Aldrich] for 3 d. The cytotoxic effect of the drugs was then assessed by using CellTitre 96 AQueous One cell proliferation assay [MTS] and a BioTek Synergy 2 plate reader [BioTek].

**Stable knockdown of Pax3Fox**

Murine primary tumor cell cultures with stable knockdown of Pax3Fox were generated as described earlier [Aslam et al. 2014].

**In vivo studies with entinostat**

Orthotopic allograft mouse models of rhabdomyosarcoma were generated as described earlier [Aslam et al. 2014]. Once the tumors reached 0.25 cm³, the mice were treated with either entinostat (LC Laboratories) at a daily dose of 10 mg/kg by intraperitoneal injection, actinomycin D (Sigma-Aldrich) at 0.25 mg/kg on day 1 by intraperitoneal injection, a combination of entinostat and actinomycin D, or vehicle (DMSO). Once the tumors reached 1.5 cm³ or the mice became sick (10%–15% loss of body weight) due to drug toxicity, the mice were euthanized. After 5 d of treatment with entinostat, the mice showed signs of drug toxicity (loss of body weight), so the treatment was halted.
for a day and then continued with 5 mg/kg daily dose [half the initial dose]. All of the drug studies in mice were performed after receiving approval from the IACUC at Oregon Health and Science University.

Statistical analysis for in vivo studies with entinostat
A repeated measures linear mixed effects model with a compound symmetric autocorrelation assumption to contrast treatments pairwise on day 7 with regard to tumor size was used. Day 7 was chosen for analysis because this was the last day with nearly complete data (actinomycin D, \( n = 7 \), actinomycin D + entinostat, \( n = 8 \), DMSO, \( n = 7 \), entinostat, \( n = 9 \)). The tumor size data were log transformed prior to analysis and corrected the \( F \)-values for multiple comparisons using the Tukey method.

Acknowledgments
We thank Rishi Rikhi and Noah Berlow for assistance with data analysis. This work was sponsored by training awards to K.N. and J.A. from the Scott Carter Foundation, support from Ministerio de Ciencia e Innovación [MICINN] SAF2009-08128, SAF2012-37427 to M.S., and National Cancer Institute awards 5R01CA133229 to C.K., 1R01CA154923 to D.M.L., and R01 AR062185 to T.A.R. We thank Rishi Rikhi and Noah Berlow for assistance with data collection. This work was sponsored by training awards to K.N. and J.A. from the Scott Carter Foundation, support from Ministerio de Ciencia e Innovación [MICINN] SAF2009-08128, SAF2012-37427 to M.S., and National Cancer Institute awards 5R01CA133229 to C.K., 1R01CA154923 to D.M.L., and R01 AR062185 to T.A.R. We thank Rishi Rikhi and Noah Berlow for assistance with data collection. This work was sponsored by training awards to K.N. and J.A. from the Scott Carter Foundation, support from Ministerio de Ciencia e Innovación [MICINN] SAF2009-08128, SAF2012-37427 to M.S., and National Cancer Institute awards 5R01CA133229 to C.K., 1R01CA154923 to D.M.L., and R01 AR062185 to T.A.R. We thank Rishi Rikhi and Noah Berlow for assistance with data collection.

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Lineage of origin in rhabdomyosarcoma informs pharmacological response

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Genes Dev. 2014 28: 1578-1591
Access the most recent version at doi:10.1101/gad.238733.114

Supplemental Material
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