ALDH Activity Correlates with Metastatic Potential in Primary Sarcomas of Bone

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Abstract

Osteosarcoma (OS), chondrosarcoma (CSA), and Ewings sarcoma (ES) are the three most common primary malignancies of bone, and are rare diseases. As with all sarcomas, the prognosis of these diseases ultimately depends on the presence of metastatic disease. Survival is therefore closely linked with the biology and metastatic potential of a particular bone tumor’s cells. Here we describe a significant correlation of aldehyde dehydrogenase (ALDH) activity and the presence/absence of distant metastases in ten consecutive cases of human bone sarcomas. Additionally, cultured human CSA cells, which are historically chemo- and radio-resistant, may be sensitive to the ALDH inhibitor, disulfiram. While it is premature to draw broad conclusions from such a small series, the importance of ALDH activity and inhibition in the metastatic potential of primary bone sarcomas should be investigated further.

Keywords

Osteosarcoma, Chondrosarcoma, Ewings Sarcoma, Aldehyde Dehydrogenase, Metastases

1. Introduction

Osteosarcoma (OS), chondrosarcoma (CSA) and Ewings sarcoma (ES) are the three most common primary malignancies of bone [1]-[6]. Most examples of “bone cancer” treated by musculoskeletal oncologists are cases wherein cancer has metastasized to the bony skeleton. Examples of primary tumors that metastasize to bone include epithelial neoplasia such as breast, prostate, renal, thyroid, and lung carcinomas [7] [8]. In contrast, primary bone sarcomas are rare diseases of mesenchymal origin that arise within the skeleton itself. As with all
sarcomas, the prognosis of primary bone sarcoma is ultimately determined by the presence or absence of systemic metastatic disease [9]-[12]. The most common site for sarcoma metastasis is to the lung, but other sites such as bone and lymph node also occur [13]-[18]. Modern treatment strategies for OS and ES include neoadjuvant chemotherapy, wide resection of the tumor, and post-operative chemotherapy. Unfortunately, CSA is resistant to both cytotoxic chemotherapy and external beam radiotherapy, and thus is primarily a surgical disease [19] [20]. Regardless of the histologic subtype, the metastatic potential of bone sarcoma cells dramatically impacts patient survival, as cure is much less likely in the setting of systemic disease. While advances in chemotherapy and surgery have improved local control and patient functionality, there have not been significant advances in the survival of patients with bone sarcomas in over two decades [13] [21] [22]. Clearly, a more complete understanding of the biology of bone sarcoma metastases is required if prognoses are to improve.

Aldehyde dehydrogenase (ALDH) is a tetrameric enzyme that oxidizes aldehydes to carboxylic acids and enables cells to resist oxidative stress. ALDH has been implicated as a cancer stem cell marker. Cells with high ALDH levels have demonstrated enhanced tumorigenicity in multiple cancer cell types. In clinical studies, high ALDH activity predicted poor survival in breast and ovarian cancer [23]-[29]. We observed that highly metastatic murine OS cells displayed greater resistance to oxidative stress than less metastatic murine OS cells when challenged with \( \text{H}_2\text{O}_2 \), and hypothesized that diminished ALDH activity in the less metastatic cells might explain this difference. Indeed, we demonstrated and published that ALDH expression and activity are significantly greater in murine OS cells with high metastatic potential than in less metastatic murine OS cells. ALDH-high cells were also more invasive than ALDH-low cells through a semisolid matrigel matrix. Finally, we treated OS cells with the ALDH inhibitor, disulfiram, and observed that disulfiram effectively reduced ALDH activity and altered metastatic OS cell morphology, resulting in fewer filopodia and greater uniformity of shape [15] [16]. These data suggested that ALDH is important to OS metastatic biology, and its inhibition in vitro caused diminished metastatic potential.

We wished to understand the importance of ALDH in the metastatic potential of human bone sarcomas. In order to accomplish this, the ALDH activity of cultured human bone sarcomas cells was assayed and compared with the metastatic histories of the patients from whom they were derived. We also treated primary bone sarcoma cells with disulfiram and doxorubicin (a cytotoxic agent commonly used in the treatment of bone sarcomas), and evaluated their effects on human bone sarcoma cells in vitro.

2. Materials and Methods

2.1. UPMC Musculoskeletal Oncology Research Registry and Tissue Bank

The University of Pittsburgh Medical Center (UPMC) Musculoskeletal Oncology Research Registry and Tissue Bank is an Institutional Review Board-approved data collection system wherein signed written informed consent from each participant allows us to collect the clinical data of patients with benign and malignant bone tumors prospectively from the time of diagnosis throughout the entire course of care. Additionally, patients’ tumor tissue may be harvested as a reagent for laboratory study at the time of biopsy or surgery. We accrued ten consecutive patients with primary bone tumors whose cells had been harvested at the time of biopsy or surgery between October, 2011 and April, 2013. Their demographic data, histologic diagnosis, and metastatic history were evaluated.

2.2. Establishing Bone Sarcoma Cell Lines

Tumor tissue from patients was washed with Dulbecco’s Phosphate-Buffered Saline (DPBS), finely minced, and enzymatically digested at 37°C in 0.2% collagenase-type XI (Sigma-Aldrich) for one hour. The cells were then incubated at 37°C in dispase (2.4 U/ml in HBSS, Invitrogen) for 45 minutes. Cells were then washed with DPBS and centrifuged (2500 rpm, 5 minutes) to obtain a cell pellet, which was then resuspended in proliferation medium (PM-DMEM supplemented with 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin (all from Invitrogen)). The dissociated cells in PM were passed through a 100 µm filter, and inoculated into plastic tissue culture flasks. Cells were maintained at 37°C, 5% \( \text{CO}_2 \), in a humidified incubator and PM was changed every 3 days.

2.3. ALDH Assays via Fluorescence-Activated Cell Sorting (FACS) Analysis of ALDH Activity

The Aldelfluor Kit (STEMCELL Technologies) was used to determine the enzymatic activity of ALDH in cul-
tured bone sarcoma cells. Human bone sarcoma cells were trypsinized, washed with DPBS, and counted using a hemocytometer. Cells were then resuspended in Aldefluor buffer at a concentration of $1 \times 10^6$ cells/mL. Aldefluor buffer contains an ABC transport inhibitor that prevents efflux of the Aldefluor dye. Cells were then incubated at 37°C for one hour, washed in Aldefluor buffer, and maintained in 4°C throughout the process of ALDH assay per the manufacturer’s instructions. High ALDH activity was assessed using the FL1 channel of a BD FACSARia Cell Sorting System and FACSDiva software (version 6.1.2; Becton, Dickinson and Company, San Jose, CA). Collected cells were analyzed for high ALDH activity with fluorescence-activated cell sorting (FACS), according to their fluorescence intensity, which corresponds to their ALDH activity levels, as well as low side scatter (SCC\textsuperscript{h}).

2.4. Disulfiram Treatment

Cultured human tumor cells were trypsinized, washed in DPBS, and counted using a hemocytometer. Cells were then plated in a 12-well dish (10,000 cells/well, in 1 mL of PM). Cells were allowed to adhere to the flask overnight and treated with disulfiram at concentrations of 0, 500 nM, 1 uM, and 1.5 uM. 48 hours later, the cells were fixed with 4% paraformaldehyde for 10 mins. The brightfield images of the cells were taken using a Nikon Eclipse E800 microscope (Melville, NY) equipped with a Retiga Exi digital camera (QImaging). All images were acquired and analyzed using Northern Eclipse software (version 6.0; Empix Imaging).

2.5. Doxorubicin Treatment

Human bone sarcoma cells were evaluated for response to doxorubicin, a cytotoxic chemotherapy often used in the treatment of bone sarcomas. Cells were plated 50,000 cells/well in 12 well plates in proliferation medium. Cells were allowed to proliferate at 37°C for 24 hours. Doxorubicin was then added in varying concentrations (0, 25 uM, 50 uM, 100 uM, 250 uM, and 500 uM) and the cells were incubated for 24 hours. At that point, cells were fixed in 10% formaldehyde for 5 minutes. The brightfield images of the cells were taken using a Nikon Eclipse E800 microscope (Melville, NY) equipped with a Retiga Exi digital camera (QImaging). All images were acquired and analyzed using Northern Eclipse software (version 6.0; Empix Imaging).

2.6. Statistical Analysis

A student’s t test was used to calculate a p value between metastatic and non-metastatic groups. A p value of less than 0.05 was considered the threshold for statistical significance. Statistical analyses were performed by a biostatistician not directly involved with the project.

3. Results

3.1. Bone Sarcoma Patients and ALDH Activity

Table 1 illustrates the age, gender, histologic diagnosis, site of metastasis (where applicable), ALDH activity of each patient’s cultured tumor cells, and standard deviations of the ALDH values when more than one measurement was performed. There were seven males and three females and the average age of the entire cohort was 57.4 years at the time of diagnosis. There were three cases of OS, five cases of CSA, and two cases of ES. The average age of the five patients with OS and EWS was 45 years, and the average age of the five CSA patients was 69.8 years. Of the patients who experienced metastases, 87.5% (seven of eight) experienced pulmonary metastases. Two patients experienced metastases to additional locations besides the lung (brain, bone) and one patient developed lymph node metastases. The average percentage of ALDH-high cells among the patients who had metastases was 18.1, and 2.1 in non-metastatic patients, with standard deviations of 12.6 and 1.3, respectively. According to our statistician, the difference in the percentage of ALDH-high cells between the metastatic and non-metastatic patients did reach statistical significance with a p value of 0.0091. If ALDH-high percentage of 3.1 was used as a threshold value, high ALDH activity correlated 100% with the incidence of metastases.

Figure 1 illustrates a sample ALDH activity assay from patient #4, a 63-year-old female with CSA metastatic to her lungs. 20.3% of her cells displayed high ALDH activity. Figure 2 displays the ALDH activity of patient #3, an 86 years old male with non-metastatic CSA. Note the difference in the percentage of ALDH-high cells, as depicted by the green curves.
Table 1. Summarizes our series of ten consecutive patients with bone sarcomas.

<table>
<thead>
<tr>
<th>Patient age</th>
<th>Gender (M/F)</th>
<th>Histology (OS, CSA, ES)</th>
<th>Site of mets</th>
<th>ALDH</th>
<th>ALDH std dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>70</td>
<td>M</td>
<td>CSA</td>
<td>Lung</td>
<td>23.2</td>
<td>2.7</td>
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<tr>
<td>80</td>
<td>M</td>
<td>CSA</td>
<td>-</td>
<td>3.05</td>
<td>2.85</td>
</tr>
<tr>
<td>86</td>
<td>M</td>
<td>CSA</td>
<td>-</td>
<td>1.2</td>
<td>-</td>
</tr>
<tr>
<td>63</td>
<td>F</td>
<td>CSA</td>
<td>Lung</td>
<td>20.3</td>
<td>-</td>
</tr>
<tr>
<td>19</td>
<td>M</td>
<td>OS</td>
<td>Lymph node</td>
<td>39.4</td>
<td>-</td>
</tr>
<tr>
<td>35</td>
<td>M</td>
<td>ES</td>
<td>Lung, brains</td>
<td>4.5</td>
<td>2.6</td>
</tr>
<tr>
<td>27</td>
<td>M</td>
<td>ES</td>
<td>Lung, bone</td>
<td>26.6</td>
<td>-</td>
</tr>
<tr>
<td>50</td>
<td>M</td>
<td>CSA</td>
<td>Lung</td>
<td>21.2</td>
<td>-</td>
</tr>
<tr>
<td>60</td>
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<td>OS</td>
<td>Lung</td>
<td>3.9</td>
<td>-</td>
</tr>
<tr>
<td>84</td>
<td>F</td>
<td>OS</td>
<td>Lung</td>
<td>5.5</td>
<td>-</td>
</tr>
</tbody>
</table>

Figure 1. Demonstrates the ALDH activity of patient #4, a 63 years old female with CSA metastatic to her lungs. Her percentage of ALDH-high cells was 20.3. The average percentage of ALDH-high cells in patients with metastatic disease was 18.1 (standard deviation = 12.6).

Figure 2. Depicts the ALDH activity of patient #2, an 86 years old male with non-metastatic CSA. His percentage of ALDH-high cells was 1.2. The average percentage of ALDH-high cells in non-metastatic patients was 2.1 (standard deviation = 1.3).

### 3.2. Disulfiram and Doxorubicin Experiments

Both disulfiram and doxorubicin decreased bone sarcoma cell proliferation in a dose-dependent fashion (Figures 3-5).
Figure 3. Demonstrates the response to disulfiram and doxorubicin (Dox) in patient #5, a 19 years old male with OS metastatic to his lymph nodes. Both disulfiram and doxorubicin demonstrated qualitative decreases in cell proliferation.

Figure 4. Shows the responses to disulfiram and doxorubicin (Dox) of patient #2, an 80 years old male with non-metastatic CSA. Both disulfiram and doxorubicin demonstrated qualitative decreases in cell proliferation.
Patient #4, 63F with CSA metastatic to lung

<table>
<thead>
<tr>
<th>Control</th>
<th>Disulfiram 500nM</th>
<th>Disulfiram 1uM</th>
<th>Disulfiram 1.5uM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>Dox 25 uM</td>
<td>Dox 50 uM</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Dox 100 uM</td>
<td>Dox 250 uM</td>
<td>Dox 500 uM</td>
</tr>
</tbody>
</table>

Figure 5. Displays the histology of patient #4, a 63 years old female with CSA metastatic to her lungs. Both disulfiram and doxorubicin (Dox) demonstrated qualitative decreases in cell proliferation.

4. Discussion

Bone sarcomas are rare, primary skeletal neoplasia. The most common forms are OS, CSA, and ES. Like all sarcomas, the prognosis of these diseases is ultimately determined by the presence of metastatic disease [9]-[12]. Others have demonstrated that ALDH expression and activity correlates with prognosis in other neoplasia, but this relationship has not been thoroughly evaluated in sarcomas of bone. We have previously demonstrated in murine OS cells that high ALDH activity correlates with murine OS cell biology [15] [16]. Here we report on the level of ALDH activity in a consecutive series of bone sarcoma patients.

This study has several limitations, not the least of which is the small series (n = 10) of bone sarcoma patients. While we did demonstrate a statistically significant difference (p = 0.0091) between the percentage of ALDH-high cells in metastatic versus non-metastatic patients, we are reluctant to draw any broad conclusions at this time. These data, while novel and encouraging, are very limited. Greater numbers of both metastatic and non-metastatic bone sarcoma patients should be studied in order to confirm or refute the hypotheses that ALDH activity correlates with bone sarcoma metastatic potential, and that bone sarcoma cell metastatic potential can be affected via ALDH inhibition.

As with our murine OS cell data, we appreciated a dose-dependent effect with disulfiram treatment. While this was not completely surprising in our human OS patient, it was interesting to observe this in the CSA patients as well. CSA is remarkably chemo- and radio-resistant. CSA is therefore treated with surgery alone, as other adjuvants do not impact upon survival. We hypothesize that the ability to withstand oxidative stress via high ALDH activity is an essential attribute that sarcoma cells must possess in order to metastasize throughout the body. The feasibility of disulfiram or other ALDH inhibitors as much-needed adjuvants in the treatment of CSA and other bone sarcomas must be investigated, and these experiments are under way. Additionally, the importance of ALDH activity vis-a-vis the metastatic potential of soft tissue sarcoma should also be evaluated with a similar strategy outlined above.

5. Conclusion

In conclusion, we have demonstrated a significant correlation between ALDH activity and metastatic potential.
in a consecutive series of ten patients with OS, CSA, and ES. We have also demonstrated that human OS and CSA cells are sensitive to ALDH inhibition with disulfiram, which raises the possibility of using ALDH inhibition as a therapeutic strategy in bone sarcomas. These data are fascinating, but additional investigations are required to understand more completely the relationship between ALDH and metastatic potential in bone sarcomas.

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