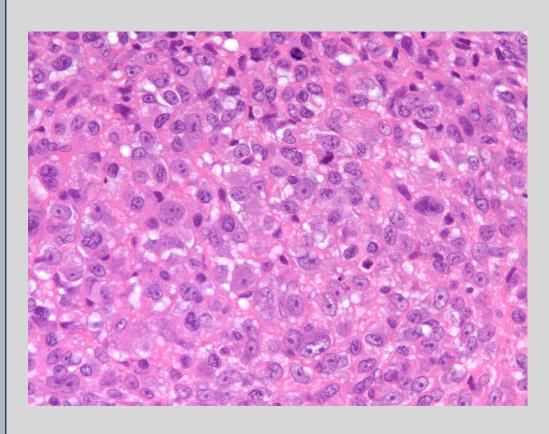
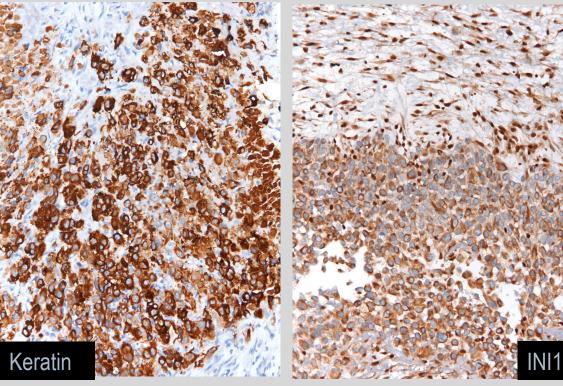


# Indications for Nuclear Imaging in Epithelioid Sarcoma – An Analysis of the **SEER Registry**

# Introduction

- Nuclear imaging, especially PET, has been widely adopted, though its role in staging and surveillance imaging in Epithelioid Sarcoma (ES) cases remains largely undefined
- ES is a mesenchymal tumor that affects patients of all ages<sup>2</sup>
- ES is a high-grade cancer that commonly metastasizes to regional lymph nodes, lung, bone, brain, and the scalp<sup>1, 4</sup>
- ES has an overall risk of metastasis at presentation of 53%<sup>3</sup>
- Metastatic disease is associated with worse outcomes
- Treatment of ES may involve radiation, chemotherapy, and surgery





## Aim

• To better define potential indications for the use of nuclear imaging in the staging of Epithelioid Sarcoma, by evaluating a patient's risk for metastasis based on presenting characteristics.

## Methods

- Using the Surveillance, Epidemiology, and End Results (SEER) database provided by the National Cancer Institute (NCI), we identified a cohort of 565 patients with primary epithelioid sarcoma from 2004 to 2015
- Histologic subtype of ES was determined using the International Classification of Disease for Oncology
- Our primary outcome was presence of detectable metastatic disease at presentation
- We identified patients as having localized, regional, and distant metastatic disease, and defined non-metastatic disease as having localized or regional disease.
- We analyzed our cohort according to the following patient characteristics: age, sex, race, and history of cancer
- We analyzed our cohort according to the following tumor characteristics: size and primary location at presentation
- We analyzed our cohort according to the following socioeconomic factors based on each patients county of residence: median family income, poverty status, and level of education

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## Results

Table 1. Patient and Tumor Characteristics of Patients with Non-metastatic and Metastatic **Epithelioid Sarcoma** 

| Variable                                      | Non-met             | Met                 | <i>p</i> value |
|---|---------------------|---------------------|----------------|
|   | n = 419             | n = 146             |                |
| Age at Diagnosis (%)                          |                     |                     | 0.001          |
| 0-14 years                                    | 24 ( 5.7)           | 6 ( 4.1)            |                |
| 15-29 years                                   | 82 (19.6)           | 22 (15.1)           |                |
| 30-59 years                                   | 206 (49.2)          | 54 (37.0)           |                |
| > 59 years                                    | 107 (25.5)          | 64 (43.8)           |                |
| Sex n (%) male                                | 223 (53.2)          | 84 (57.5)           | 0.421          |
| Race <sup>1</sup> n (%)                       |                     |                     | 0.285          |
| White   | 344 (82.7)          | 113 (77.4)          |                |
| Black   | 43 (10.3)           | 22 (15.1)           |                |
| Other   | 29 (7.0)            | 11 (7.5)            |                |
| History of Cancer n (%) yes                   | 58 (13.8)           | 21 (14.4)           | 0.981          |
| <b>Population</b> $\leq 20,000^{2}$ (%)       | 24 ( 5.7)           | 7 ( 4.8)            | 0.843          |
| Education <sup>3</sup> % (mean (SD))          | 14.56% (5.98%)      | 14.61% (6.30%)      | 0.938          |
| Poverty Status <sup>4</sup> % (mean (SD))     | 15.03% (4.91%)      | 14.88% (6.07%)      | 0.768          |
| Median Family Income <sup>5</sup> (mean (SD)) | \$71,306 (\$16,452) | \$73,295 (\$17,896) | 0.219          |
| Tumor Size <sup>6</sup> n (%)                 |                     |                     | < 0.001        |
| $\leq$ 25 mm                                  | 88 (26.1)           | 8 ( 8.8)            |                |
| 26-50mm                                       | 88 (26.1)           | 16 (17.6)           |                |
| 51-75mm                                       | 67 (19.9)           | 15 (16.5)           |                |
| 76-100mm                                      | 39 (11.6)           | 21 (23.1)           |                |
| > 100 mm                                      | 55 (16.3)           | 31 (34.1)           |                |
| Primary Tumor Site n (%)                      |                     |                     | < 0.001        |
| Extremity                                     | 217 (52.7)          | 41 (31.3)           |                |
| Axial   | 142 (34.5)          | 49 (37.4)           |                |
| Head/Neck                                     | 29 ( 7.0)           | 8 ( 6.1)            |                |
| Viscera                                       | 24 ( 5.8)           | 33 (25.2)           |                |

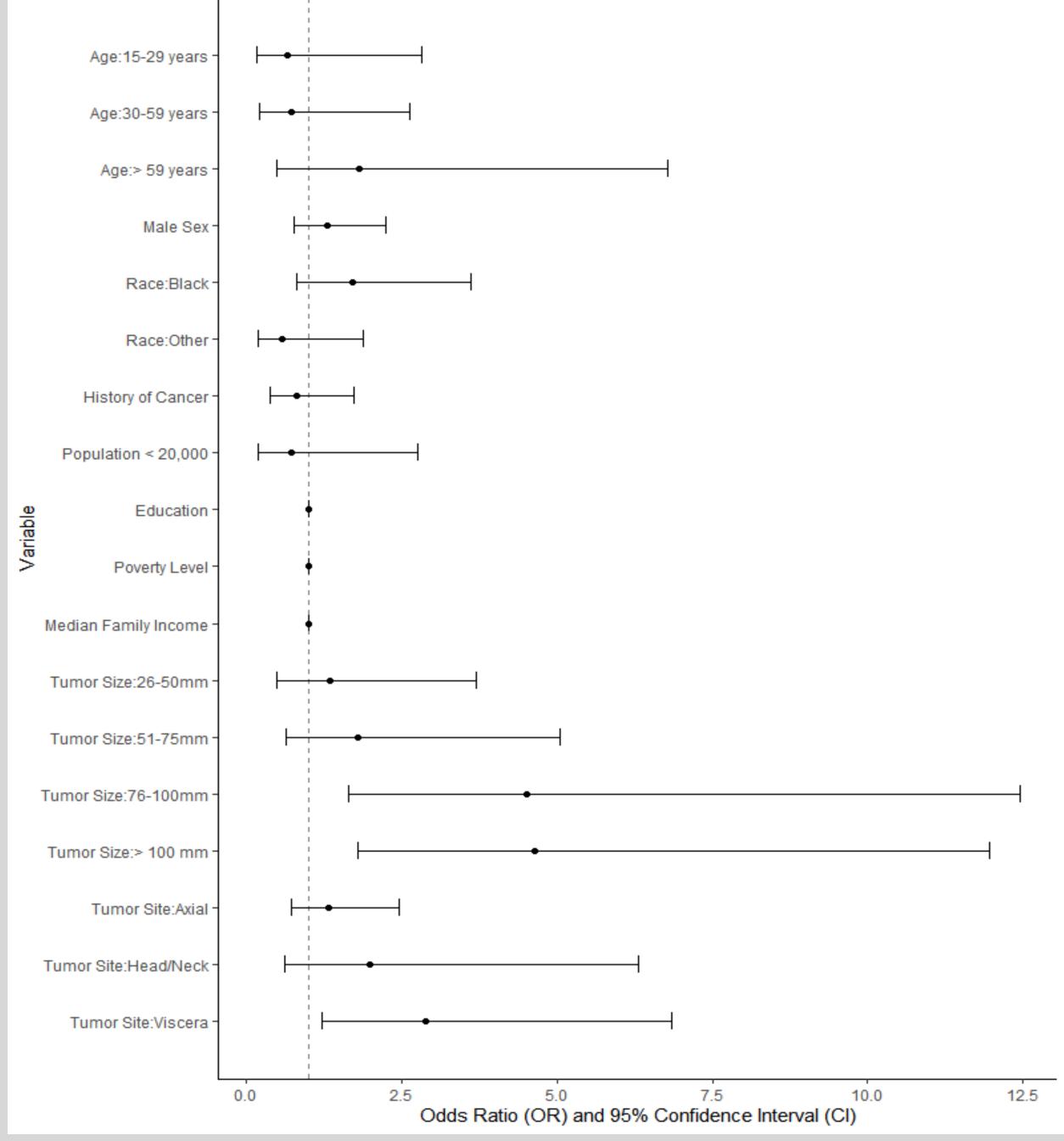
<sup>1</sup>Asian. Pacific Islander, American Indians, Alaska Natives, and Unknown <sup>2</sup>Setting based on Rural-Urban Continuum Code 2013

<sup>3</sup>Percent of persons over 25 years old with less than 12 years of education, United States Census 2013 <sup>4</sup>Percent below Federal Poverty Level, United States Census 2013

<sup>5</sup>Median family income, United States Census 2013

<sup>6</sup>Tumor size missing for 178 (28.6%) cases

#### Figure 1. Forest Plot of Odds Ratios (OR) and 95% Confidence Intervals (CI) From the Multivariate Regression Model



Age Reference Group: 0-14 years Race Reference Group: White Tumor Size Reference Group:  $\leq 25 \text{ mm}$ Tumor Site Reference Group: Extremity

|                | (OR), 95% Confidence I<br>Presentation per Patient | <b>^</b> |
|----------------|--|----------|
|                | Per            |          |
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| Variable                          | OR         | 95% CI       | <i>p</i> value |
|-----------------------------------|------------|--------------|----------------|
|                                   | n = 419    | n = 146      |                |
| Age at Diagnosis                  |            |              |                |
| 0-14 years                        | 1.00 (Ref) | -            | -              |
| 15-29 years                       | 0.66       | 0.15 - 2.81  | 0.580          |
| 30-59 years                       | 0.72       | 0.19 - 2.62  | 0.621          |
| > 59 years                        | 1.82       | 0.48 - 6.7   | 0.370          |
| Male Sex                          | 1.30       | 0.75 - 2.24  | 0.343          |
| Race <sup>1</sup>                 |            |              |                |
| White                             | 1.00 (Ref) | -            | -              |
| Black                             | 1.70       | 0.80 - 3.61  | 0.162          |
| Other                             | 0.57       | 0.17 - 1.87  | 0.359          |
| History of Cancer                 | 0.80       | 0.37 - 1.73  | 0.574          |
| Population $\leq 20,000^2$        | 0.72       | 0.19 - 2.74  | 0.638          |
| Education <sup>3</sup>            | 1.00       | 0.99 - 1.00  | 0.483          |
| Poverty Level <sup>4</sup>        | 1.00       | 0.99 - 1.00  | 0.488          |
| Median Family Income <sup>5</sup> | 1.00       | 0.99 - 1.00  | 0.189          |
| Tumor Size                        |            |              |                |
| $\leq$ 25 mm                      | 1.00 (Ref) | -            | -              |
| 26-50mm                           | 1.34       | 0.49 - 3.69  | 0.563          |
| 51-75mm                           | 1.79       | 0.64 - 5.03  | 0.262          |
| 76-100mm                          | 4.52       | 1.64 - 12.44 | 0.003          |
| > 100 mm                          | 4.62       | 1.79 - 11.96 | 0.001          |
| <b>Primary Tumor Site</b>         |            |              |                |
| Extremity                         | 1.00 (Ref) | -            | -              |
| Axial                             | 1.32       | 0.71 - 2.45  | 0.374          |
| Head/Neck                         | 1.97       | 0.62-6.30    | 0.248          |
| Viscera                           | 2.88       | 1.21 - 6.85  | 0.016          |
|                                   |            |              |                |

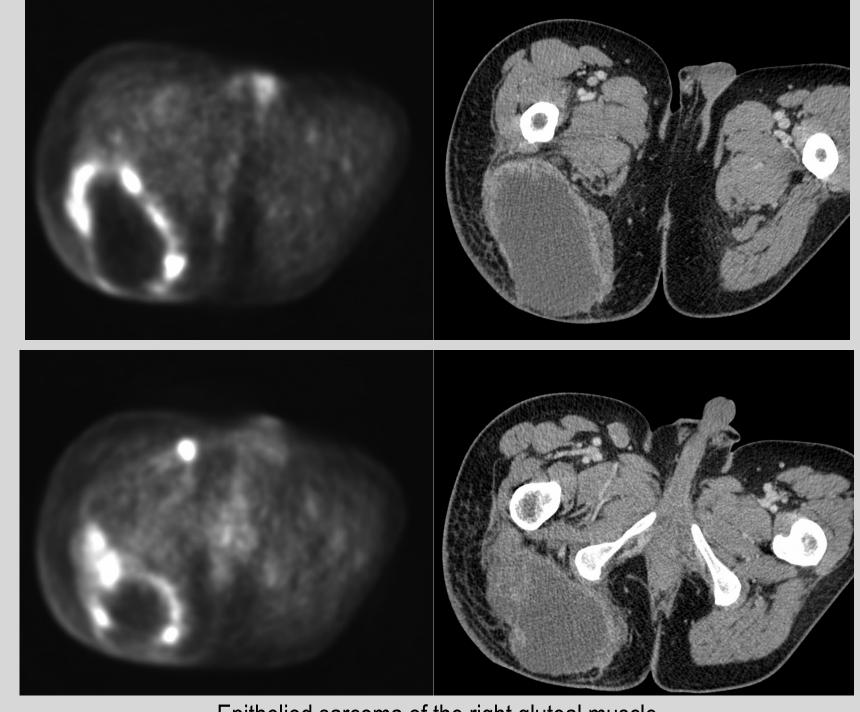
<sup>1</sup>Asian, Pacific Islander, American Indians, Alaska Natives, and Unknown <sup>2</sup>Setting based on Rural-Urban Continuum Code 2013 <sup>3</sup>Percent of persons over 25 years old with less than 12 years of education, United States

2013 <sup>4</sup>Percent below Federal Poverty Level, United States Census 2013 <sup>5</sup>Median family income, United States Census 2013

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nor size and primary tumor site are significantly associated a greater risk of detectable metastatic ES at presentation. • Other factors evaluated did not correlate with metastatic ES at presentation

Our findings suggest that patients who present with tumors greater than 76 mm and/or those with visceral primary tumors, should prompt a more thorough imaging workup to evaluate for metastatic disease.



 A small subset of patients was excluded due to a lack of staging data. • The SEER database categorizes location of tumors as localized, regional, or distant. Regional localization could not be further evaluated, and due to this, we included regional tumors as non-metastatic disease.

Socioeconomic variables were gathered at the county level, which may not accurately represent the lived realities of the patients included.

**Charlie Callif** 



## Conclusion

Epitheliod sarcoma of the right gluteal muscle

#### Limitations

### Work Cited

Chase, D.R. and F.M. Enzinger, *Epithelioid sarcoma. Diagnosis, prognostic* indicators, and treatment. The American journal of surgical pathology, 1985. 9(4): p. 241-263.

Fisher, C., *Epithelioid sarcoma of Enzinger*. Advances in anatomic pathology, 2006. **13**(3): p. 114-121.

Jawad, M.U., J. Extein, E.S. Min, and S.P. Scully, *Prognostic factors for survival in* patients with epithelioid sarcoma: 441 cases from the SEER database. Clin Orthop Relat Res, 2009. **467**(11): p. 2939-48

Spillane, A.J., J.M. Thomas, and C. Fisher, *Epithelioid sarcoma: the* clinicopathological complexities of this rare soft tissue sarcoma. Annals of surgical oncology, 2000. 7(3): p. 218-225.

## **Contact Information**