



# Patient Presentation for Radiology-Pathology Correlate

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# Initial presentation



- Patient TM is a 55 year old post-menopausal female, presenting to UVA ED via transfer from outside hospital. Only relevant history is of tubal ligation.
- Presenting symptoms
  - Nausea, 4 episodes of nonbloody nonbilious emesis
  - Worsened abdominal pain: 10/10, constant sharp lower abdomen
  - 8 – 9 lbs unintentional weight loss over past week
- Recent symptoms
  - LLQ abdominal pain: 4 mo. history, cramping & occasionally sharp, radiates to left back
  - Constipation, Increased belching
  - LE swelling
- Evaluated at MJH
  - Large heterogeneous left abdominal mass appearing to arise from left ovary
  - Leukocytosis 15,000
  - Lactic acidosis 1.6
- Transferred to UVA



- Medical Hx
  - Anxiety & Depression
  - Hypertension
  - OA
- Surgical Hx
  - Tubal ligation
- Family Hx
  - MGM: stomach cancer
- Social Hx
  - Never smoker
  - Occasional EtOH
  - Married, 3 children, 11 grandchildren
  - Runs a pre-school out of her home



Heterogeneous and enlarged left ovary, continuous with a superiorly extending heterogeneously attenuating mass

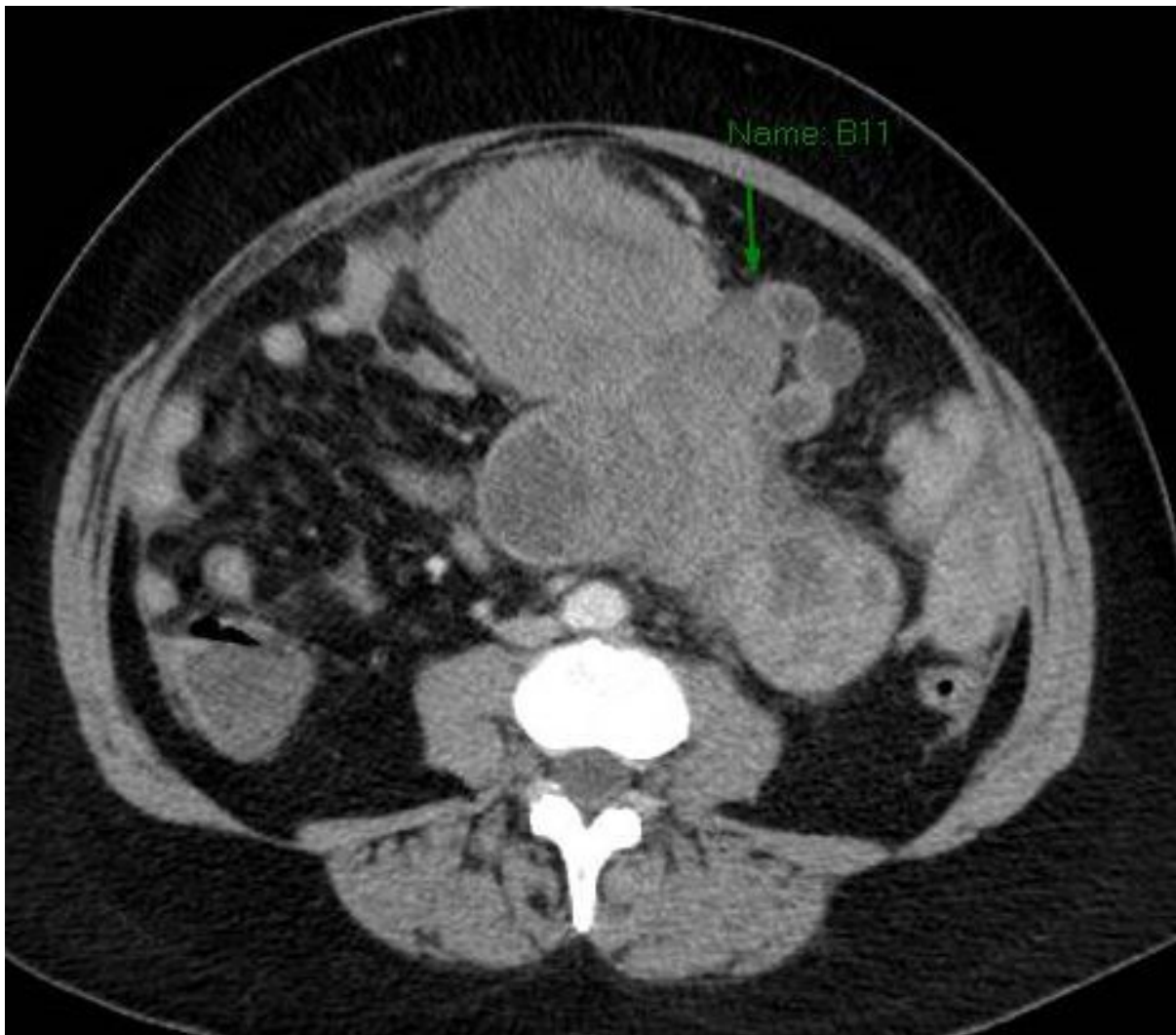
Likely arising from the left ovary





Large multilobulated heterogeneously attenuating mesenteric mass noted, measuring approximately 11.2 x 12.8 x 15 cm

No clear fat plane separates mass from left psoas muscle



No clear fat plane  
separates mass from  
loops of small bowel

Peritoneal  
carcinomatosis

Highly suspicious for  
metastatic ovarian  
cancer



- Informed consent obtained
- Labs
  - PT: 15.1
  - INR: 1.3
  - Plts: 260
  - CA-125: slightly elevated
  - CEA: normal



- US guided FNA
- Anterior approach
- Patient supine

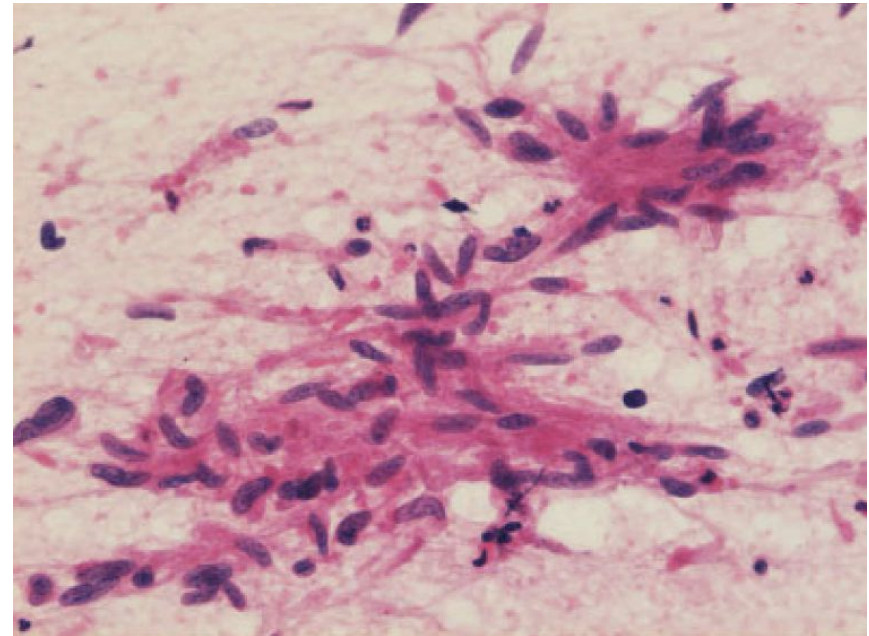
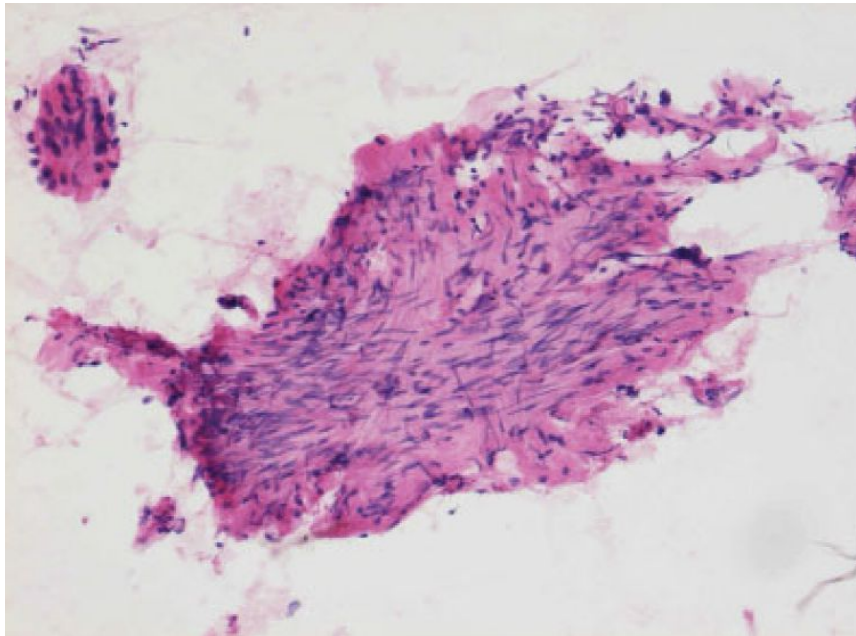




# FNA Findings



- Spindle cells?!?
- 2 core biopsies taken





- Core findings:
  - Cellular spindle cell proliferation
  - Frequent mitotic figures (up to 40 per 10 hpf)
  - Nuclear pleomorphism
  - Tumor necrosis
- Differential: leiomyosarcoma vs GIST
- Immunohistochemistry:
  - Positive staining with SMA and desmin
  - Negative for c-kit (CD117), Dog-1 and CD34
- Diagnosis of leiomyosarcoma supported

# Leiomyosarcoma



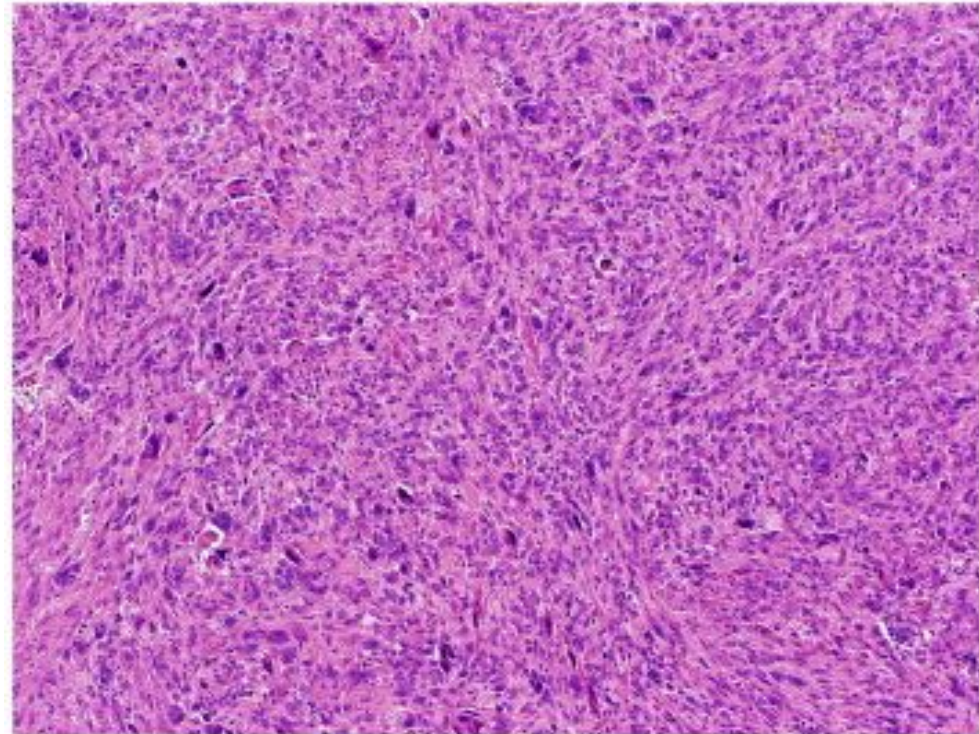
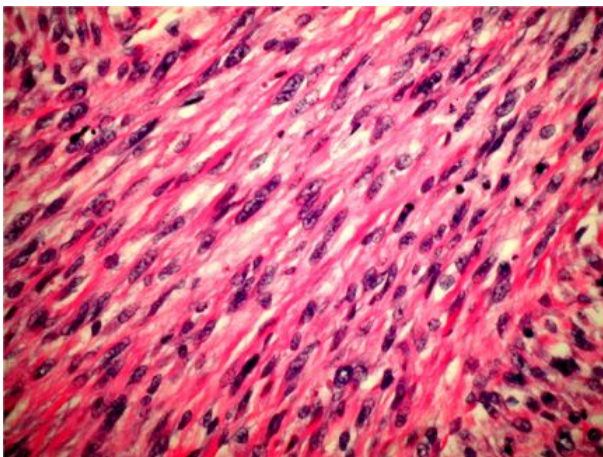
- Most common uterine sarcoma; accounts for 1 – 2% of uterine malignancies
- Aggressive, associated with poor prognosis and high risk of recurrence (53 - 71%)
- Vast majority are sporadic; leiomyomas are not precursors
- Clinical-pathologic features
  - Peri- or postmenopausal age
  - Extrauterine extension
  - Large size (over 10 cm)
  - Infiltrating border
  - Necrosis





- Pathologic features
  - Hypercellularity
  - Severe nuclear atypia
  - High mitotic rate

Well-differentiated

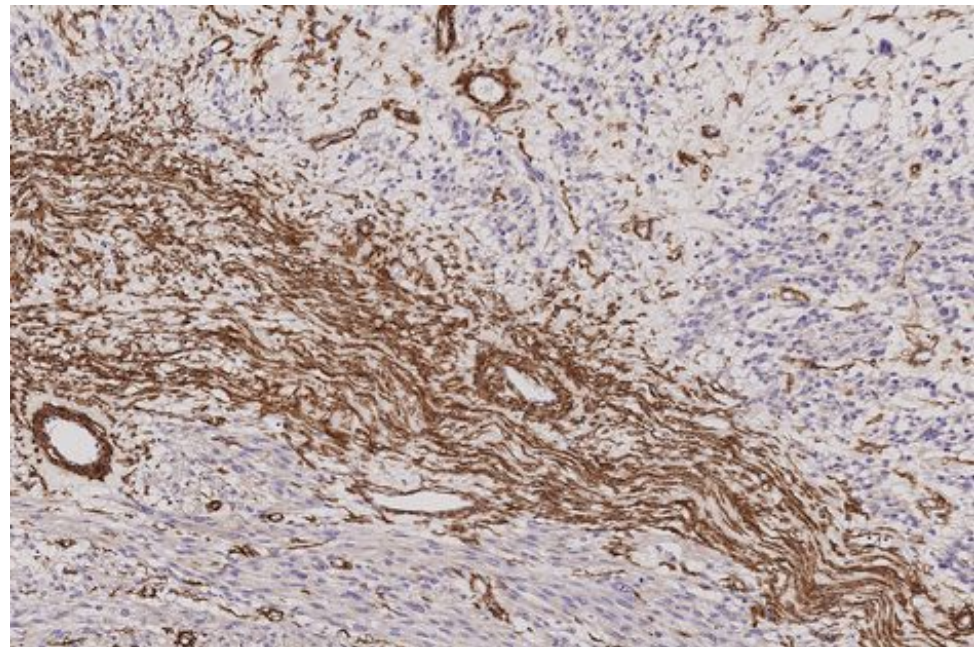


Spindle cell variant





- Usually express smooth muscle markers
  - Histone deacetylase 8 (HDCA8)
  - Smooth muscle actin (SMA)
  - Desmin
  - H-caldesmon



# Malignant?



- Malignancy judged based on Stanford criteria: (1) cellular atypia, (2) mitosis, (3) coagulative necrosis
- Staged using the 2009 International Federation of Gynecology and Obstetrics (FIGO) staging system

Group	Mitotic Index (MI) (per 10 HPF)	Atypia	Coagulative tumor cell necrosis	Designation	Metastatic or recurrent disease
I	≥5 to <20	None or mild	None	Leiomyoma with increased MI	1/89
II A	<10	Diffuse, moderate or severe	None	Atypical leiomyoma with low risk percent or recurrence	2/46
II B	≥10	Diffuse, moderate or severe	None	Leiomyosarcoma	4/10
III	≤20	Diffuse, moderate to severe	Present	Leiomyosarcoma	19/33
IV A	<10	None to mild	Present	Smooth muscle tumors of low malignant potential, limited experience	1/4
IV B	≥10	None to mild	Present	Leiomyosarcoma	3/4
V	≥1 to ≤20	Multifocal, moderate to severe	None	Atypical leiomyoma, limited experience	0/5



- Overall survival rate 15 - 25%, with median survival of 10 months (in one study)
- Cohort of 2017 National Cancer Database study of N = 7455 showed 10 - 14% 5-year survival in metastatic leiomyosarcoma
- Size, grade, and stage help in prognostication
  - Survival decreases dramatically with increasing tumor size
- Early and complete surgical resection remains the best-evidenced effective treatment
  - Once disease has spread beyond uterus, likelihood of achieving long-term survival is low with current therapeutics
- Influence of adjuvant therapy on survival is uncertain
  - Radiation may be useful for local recurrences
  - Chemotherapy increases survival of women with metastatic leiomyosarcoma, although the survival benefit may not persist and long-term survival, or chance of cure, remains low



- Case follow-up

- Results of biopsy conveyed by GynOnc to patient TM via telephone encounter
- Planning to do chemotherapy due to size of mass and involvement of great vessels

- References

- D'Angelo E, Prat J. Uterine sarcomas: a review. *Gynecol Oncol.* 2010 Jan;116(1):131-9.
- Seagle BL, Sobecki-Rausch J, Strohl AE, Shilpi A, Grace A, Shahabi S. prognosis and treatment of uterine leiomyosarcoma: A national Cancer Database study. *Gynecol Oncol.* 2017 Apr;145(1):61-70.
- Domanski H.A., Akerman M., Rissler P., Gustafson Pelle. Fine-Needle Aspiration of Soft Tissue Leiomyosarcoma: An Analysis of the Most Common Cytologic Findings and the Value of Ancillary Techniques. *Diagnostic cytopathology.* 2006 Aug;34(9).
- UpToDate. Treatment and prognosis of uterine leiomyosarcoma. Accessed August 7, 2017.
- Smooth Muscle Actin, Alpha (SMA). Leica Biosystems. Accessed August 7, 2017. <<http://www.leicabiosystems.com/ihc-ish-fish/immunohistochemistry-ihc-antibodies-novocastra-reagents/primary-antibodies/products/sma-alpha-smooth-muscle-actin/>>

- Questions?