

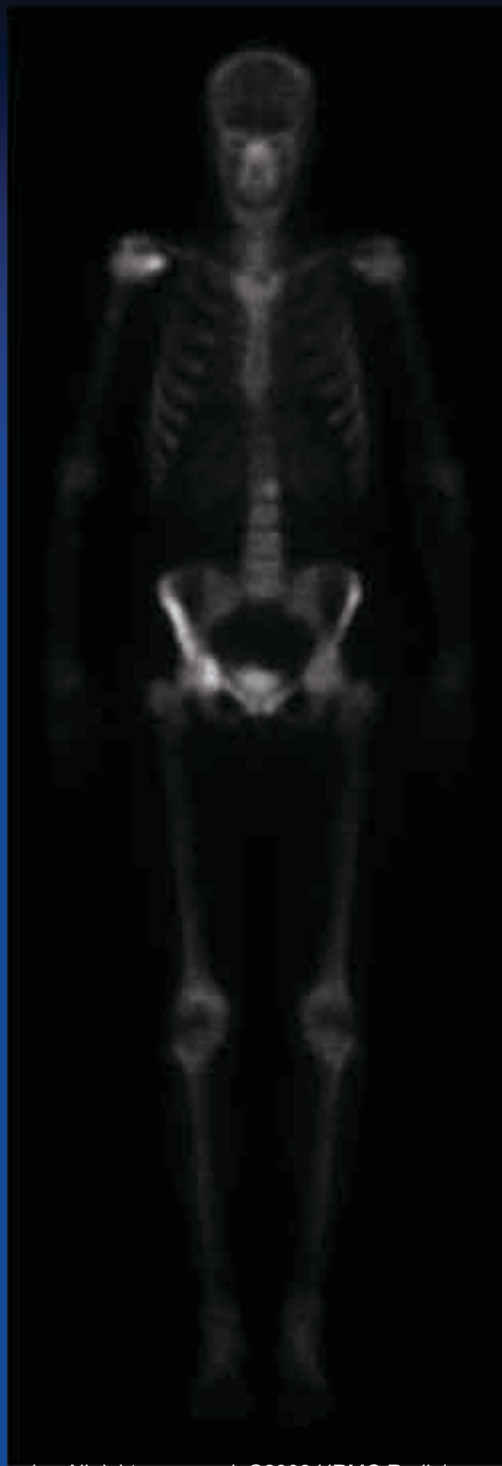
# Rad-Path Conference

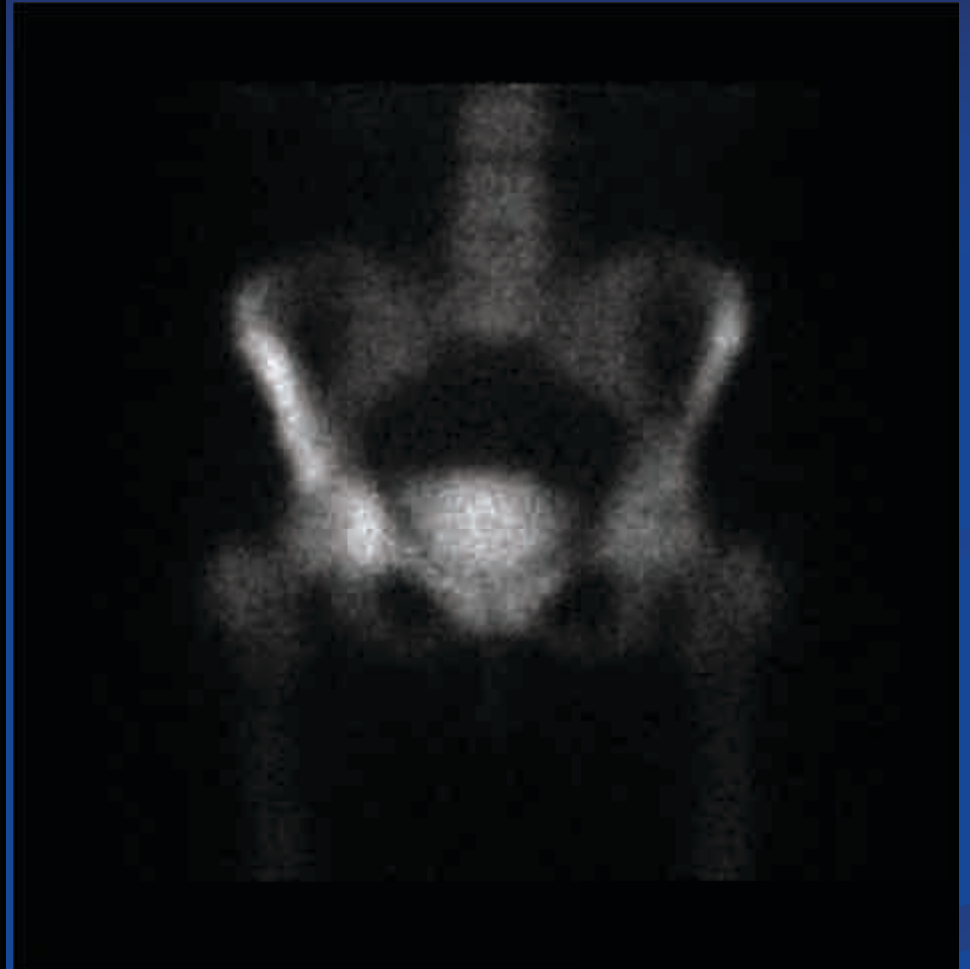
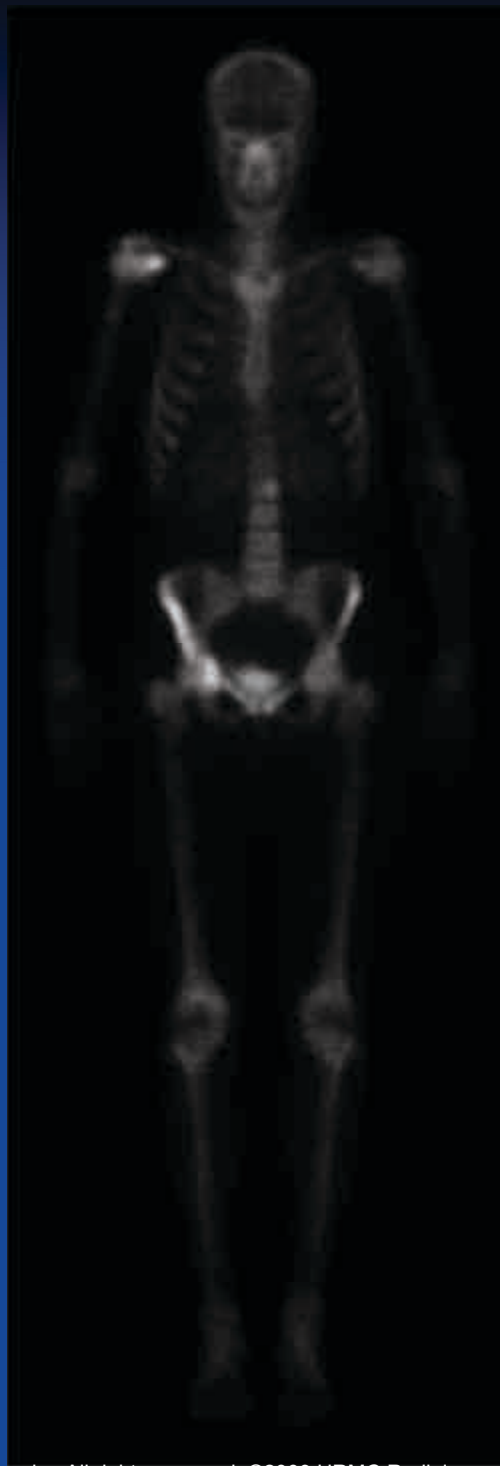
June 26, 2009

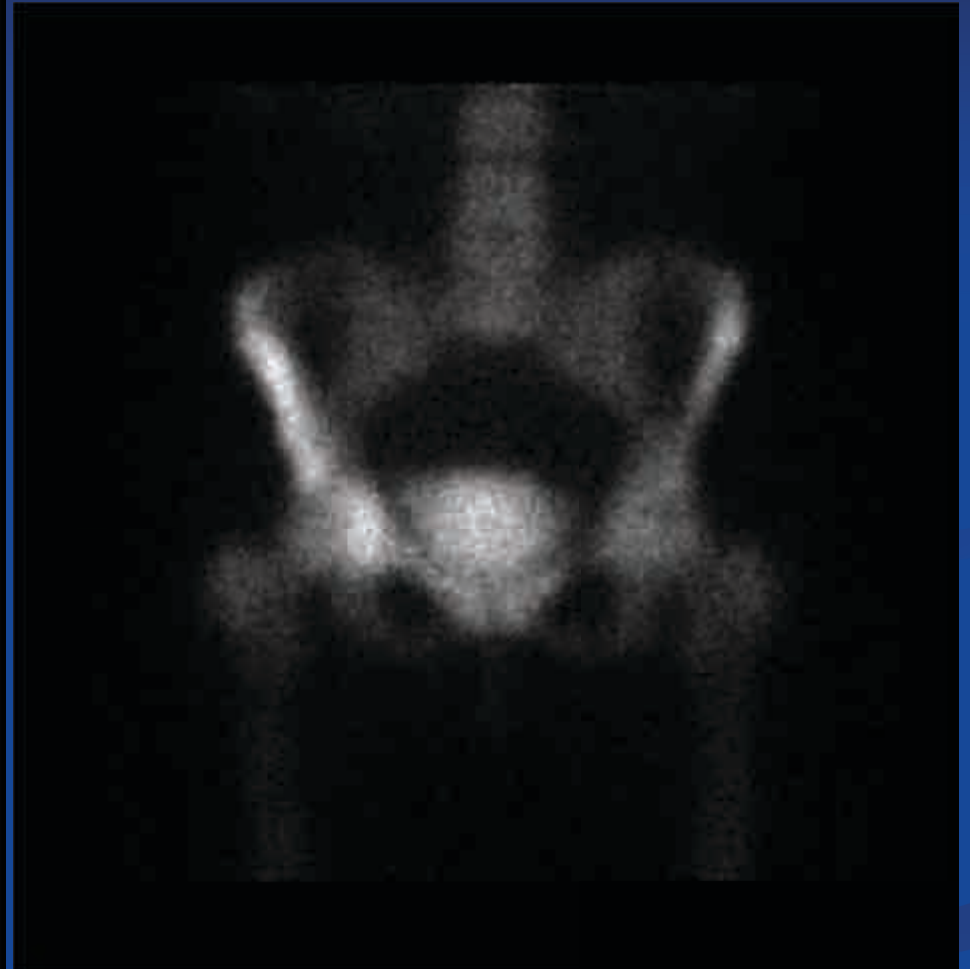
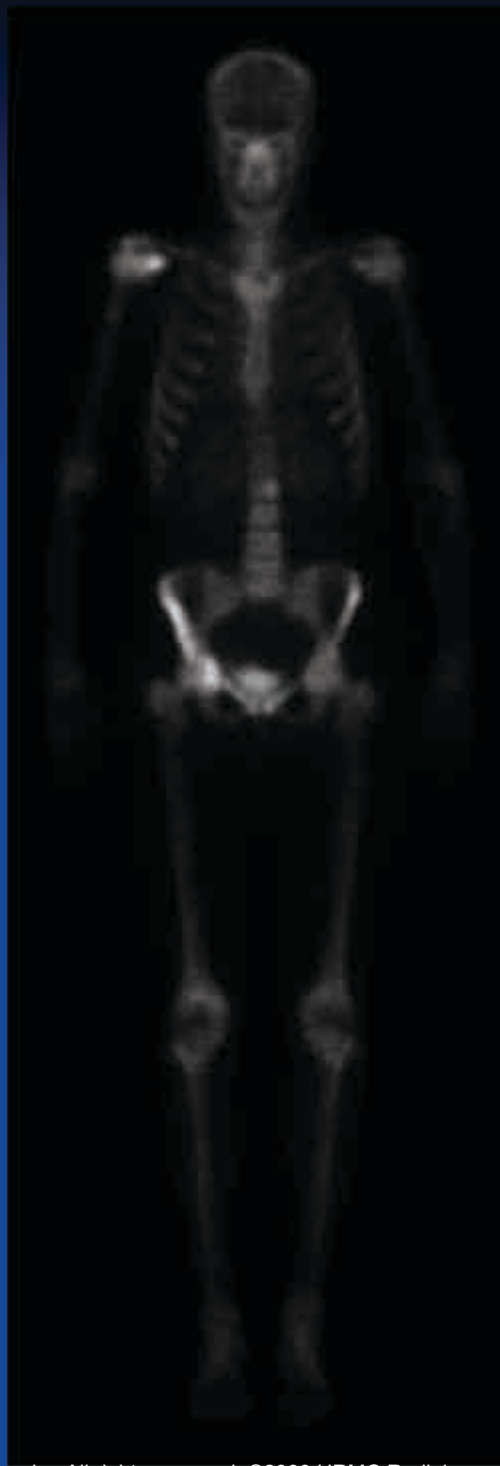
S. Cassar, N. Johnson, and S. Canacci

# Case 1

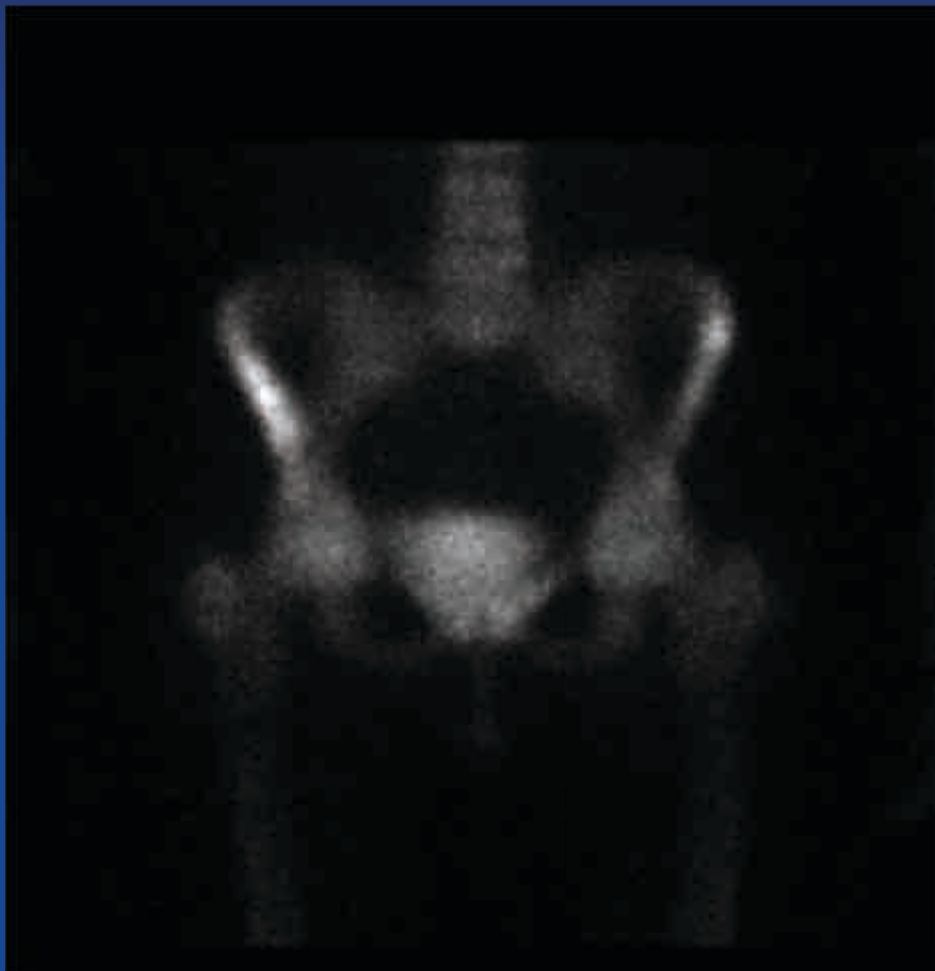
- 44 yo female with a history of breast cancer s/p bilateral mastectomies, chemotherapy, and radiation therapy, now with right groin and upper arm pain for approximately 6 to 8 weeks



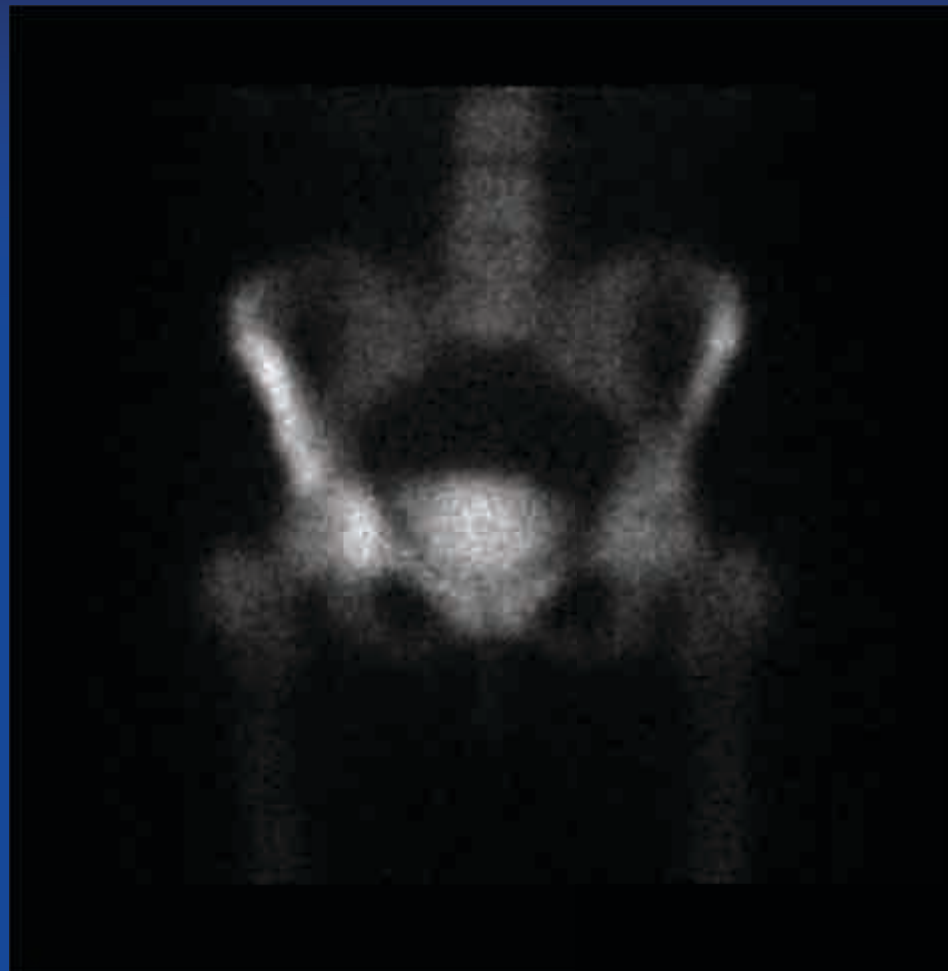




4/09/2009

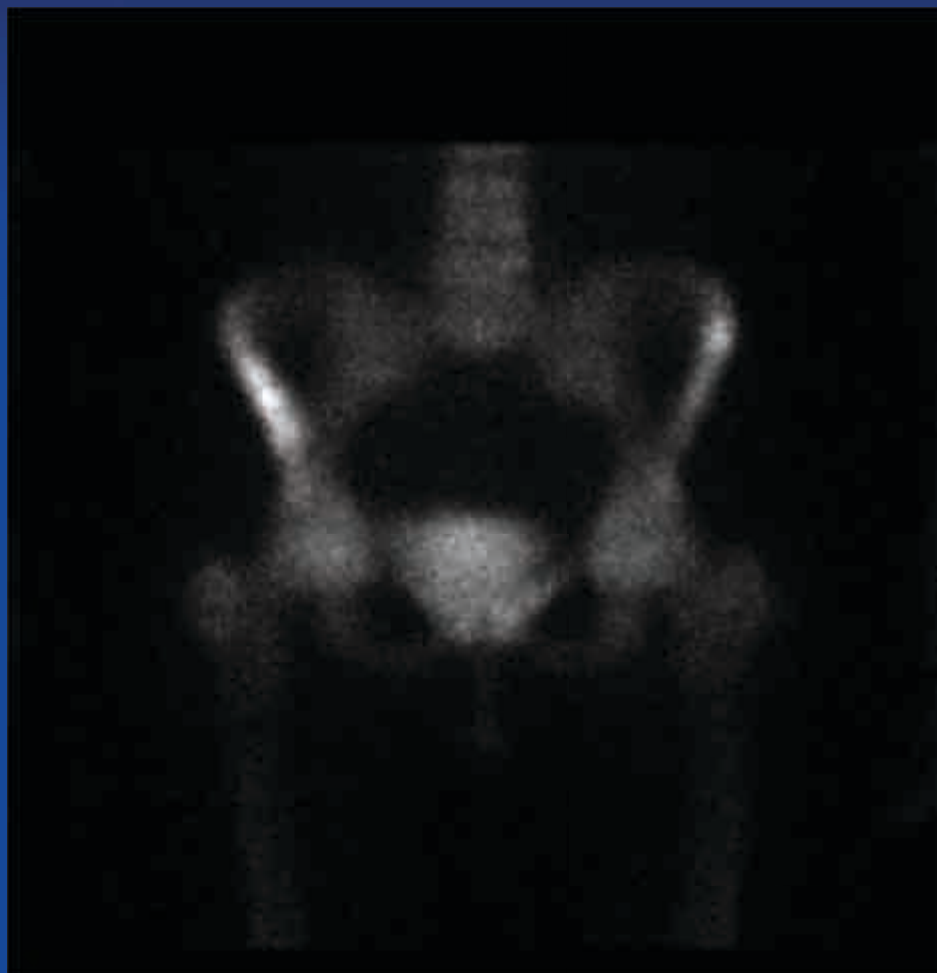


7/21/2008

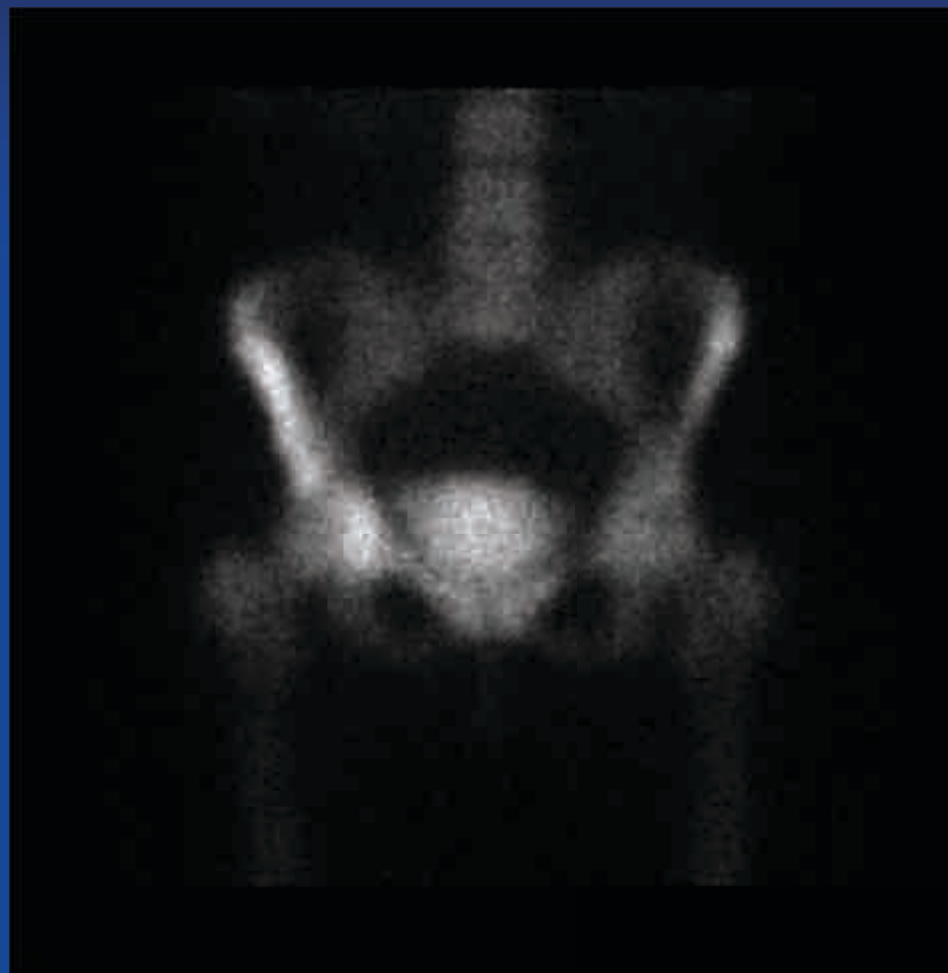


4/09/2009

Interval increase in intensity in right anterior iliac focus suggestive of worsening of metastatic disease.



7/21/2008

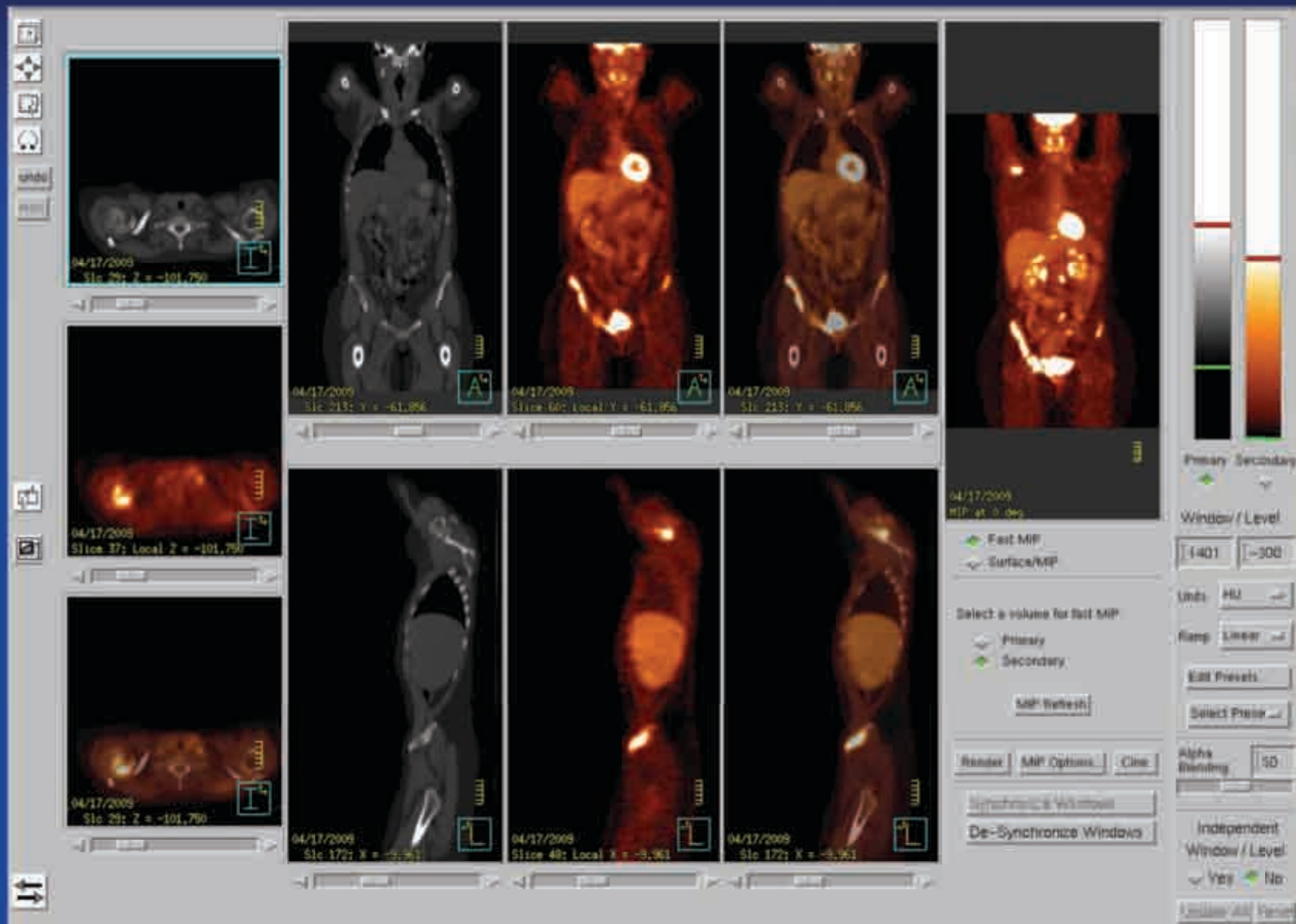


4/09/2009

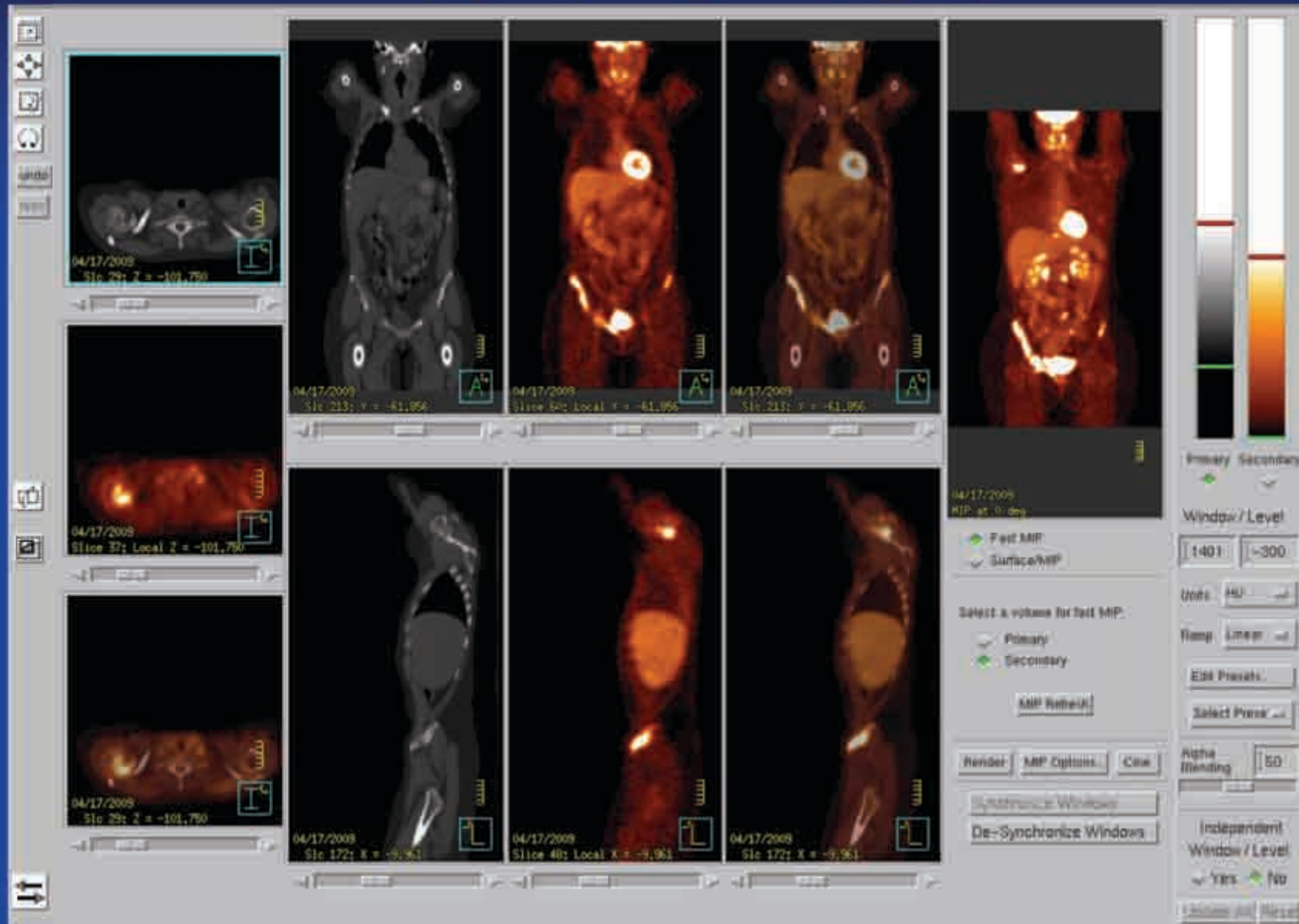


- She had an MRI of her hips performed at Borg and Ide Imaging, which revealed multiple abnormal signal intensities in the pelvic bones and hips, right greater than left, which was highly concerning for metastatic disease.
- Next step? (Hint: restaging)





Hypermetabolic foci of metastatic disease in the bilateral shoulders, right humeral head, spine, and pelvis, right greater than left.



4/17/2009

# Differential Diagnosis

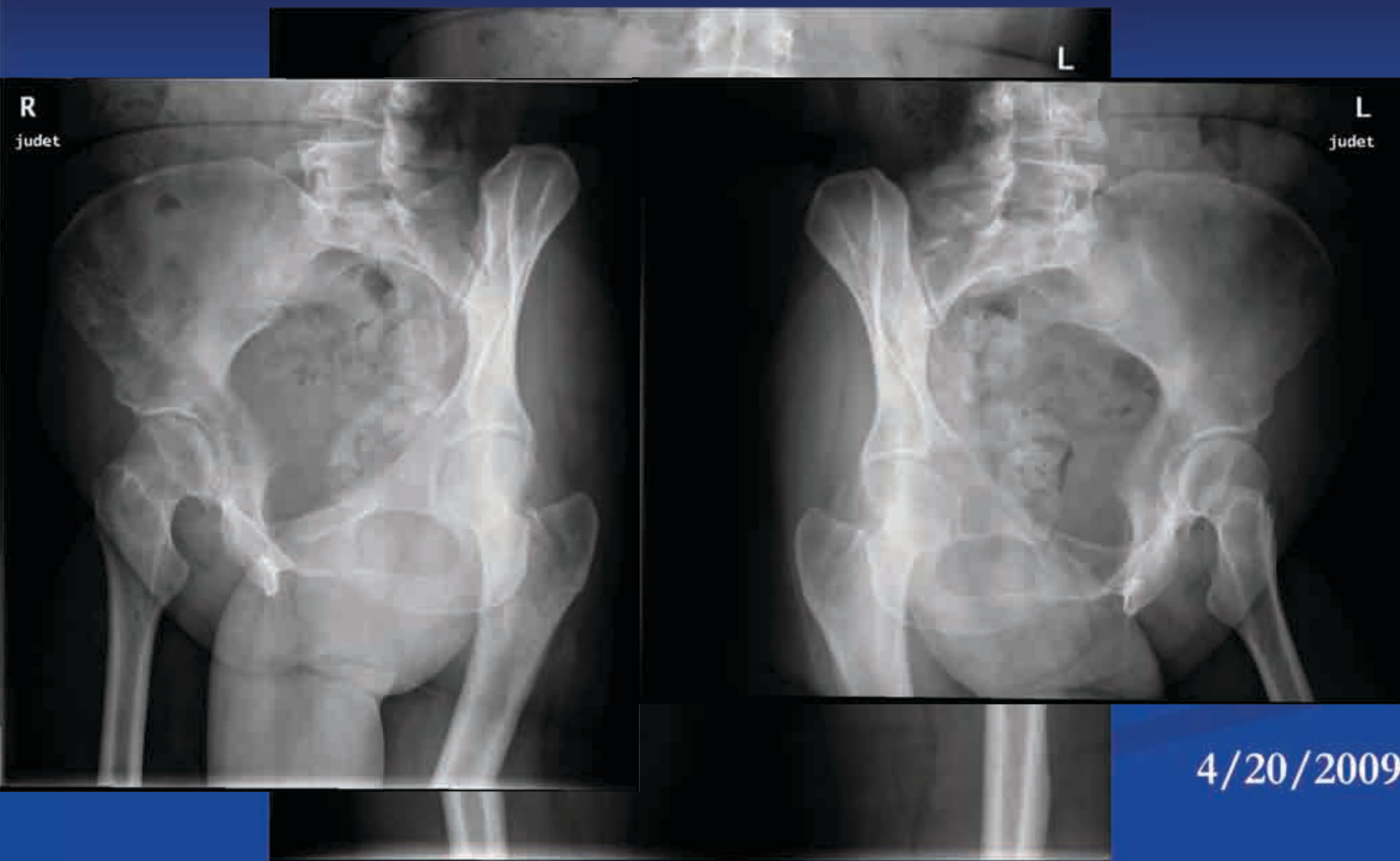
- Metastatic breast cancer
- Metastatic breast cancer
- Metastatic breast cancer
- ...
- Infection/Inflammation (sarcoid, degenerative)
- Trauma
- Second primary neoplasm (thyroid, lung, colon)



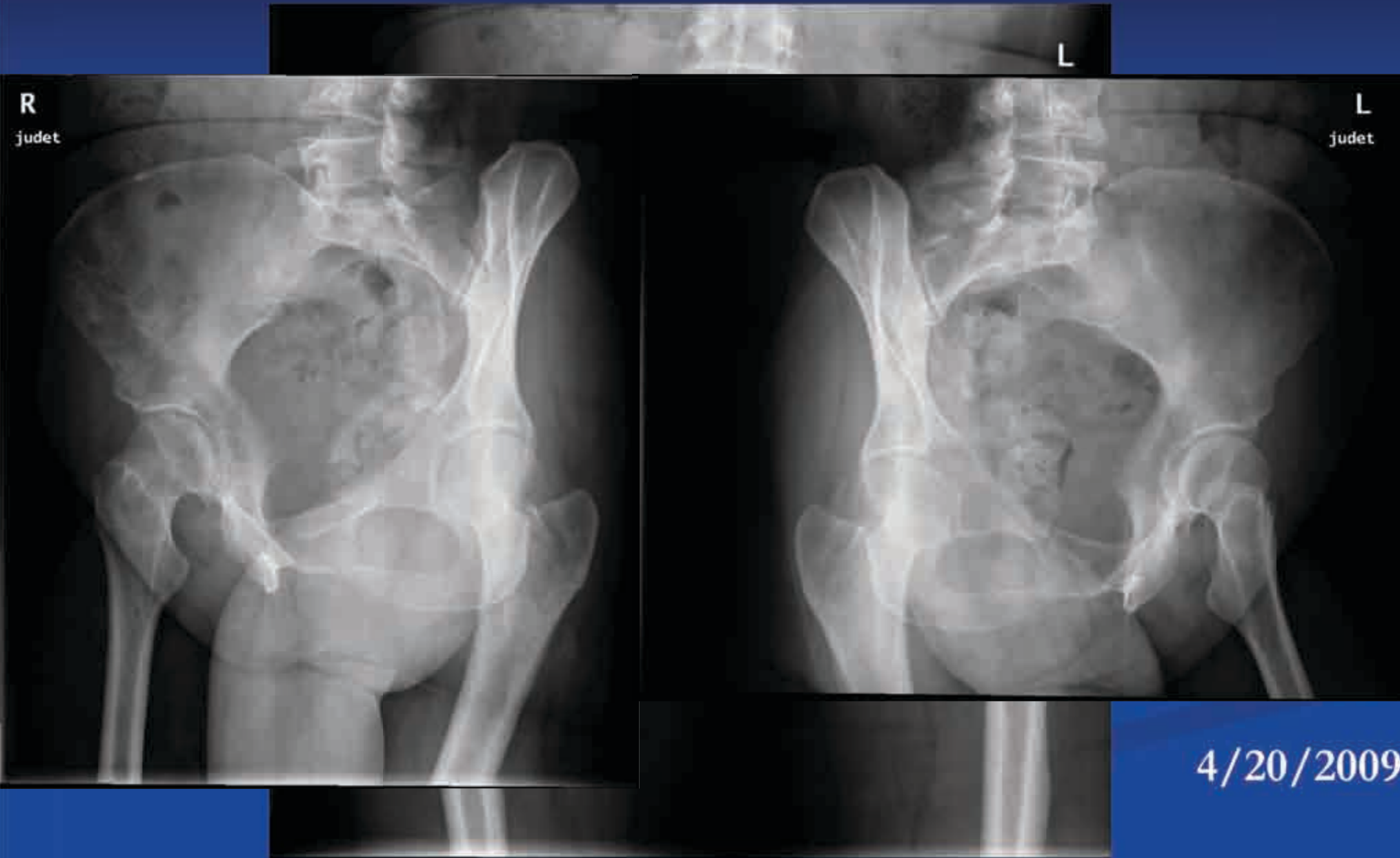
- Orthopedic surgeon consulted for consideration of surgery for stabilization of her pelvis
- He requested plain films and a CT scan of the pelvis (with 3 mm cuts) – note the recent PET-CT was done with 5 mm slices – to look for fracture



4/20/2009

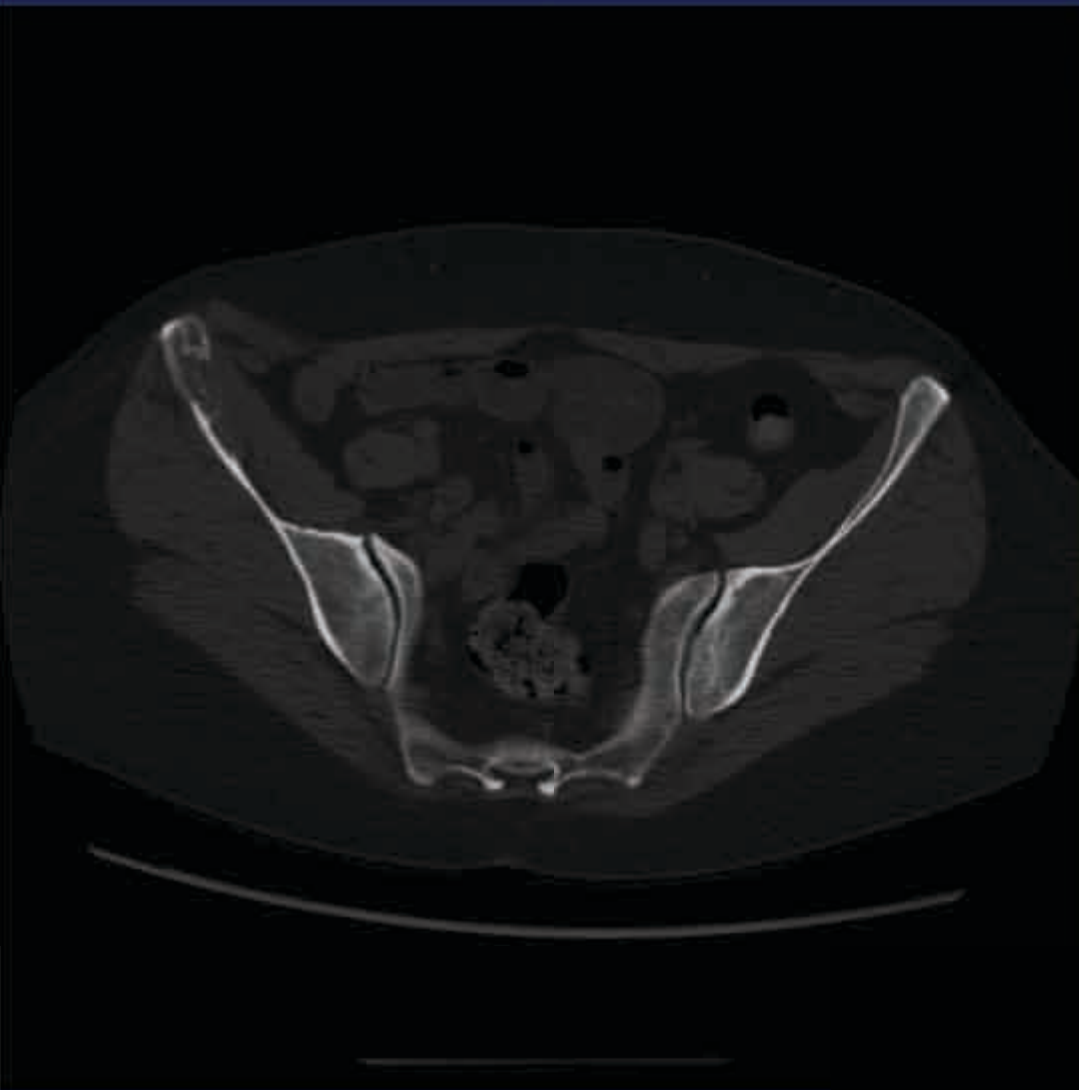


Increased sclerosis of the right acetabulum and lateral ileum and two sclerotic foci in the proximal right femur consistent with metastases.

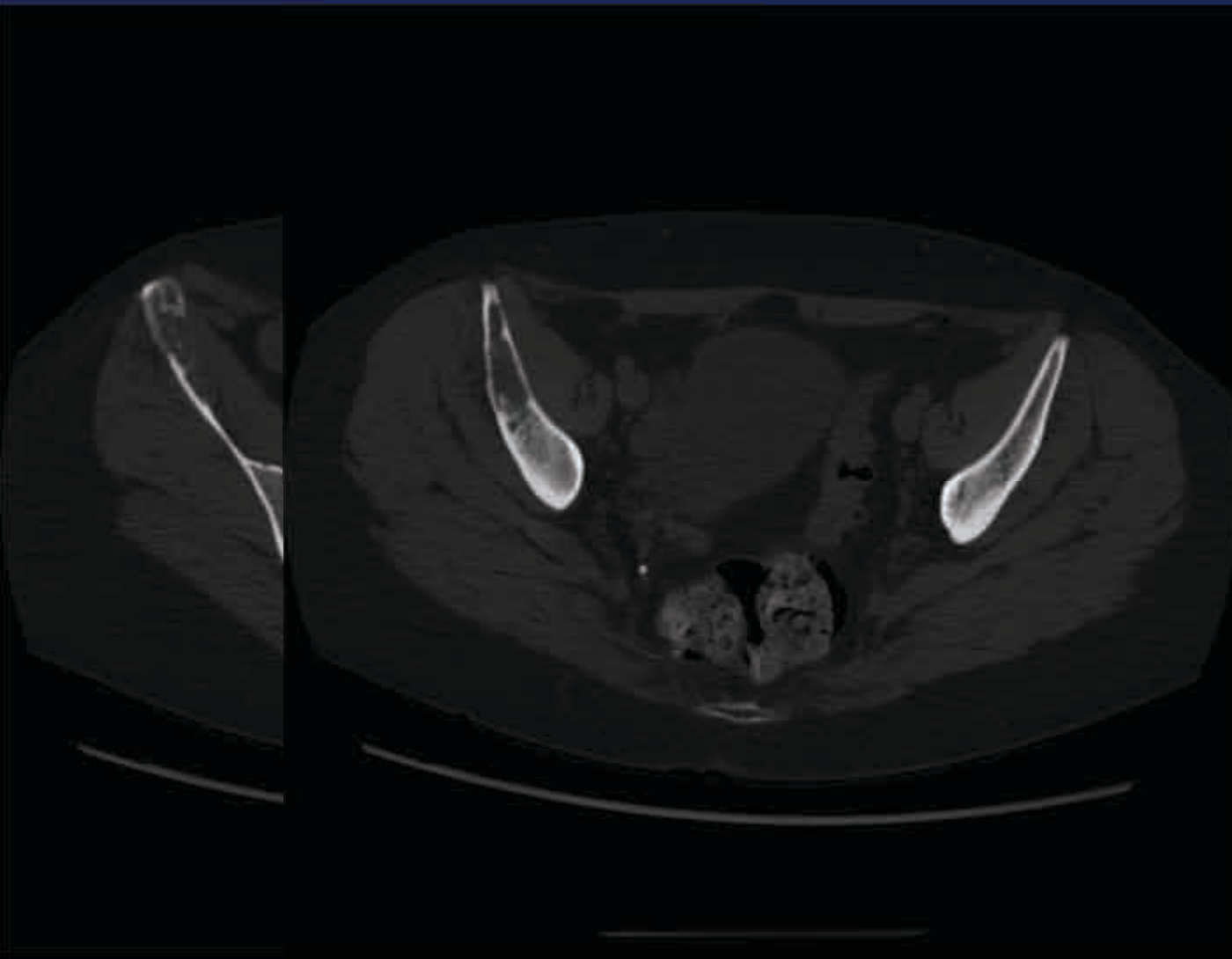


4/20/2009

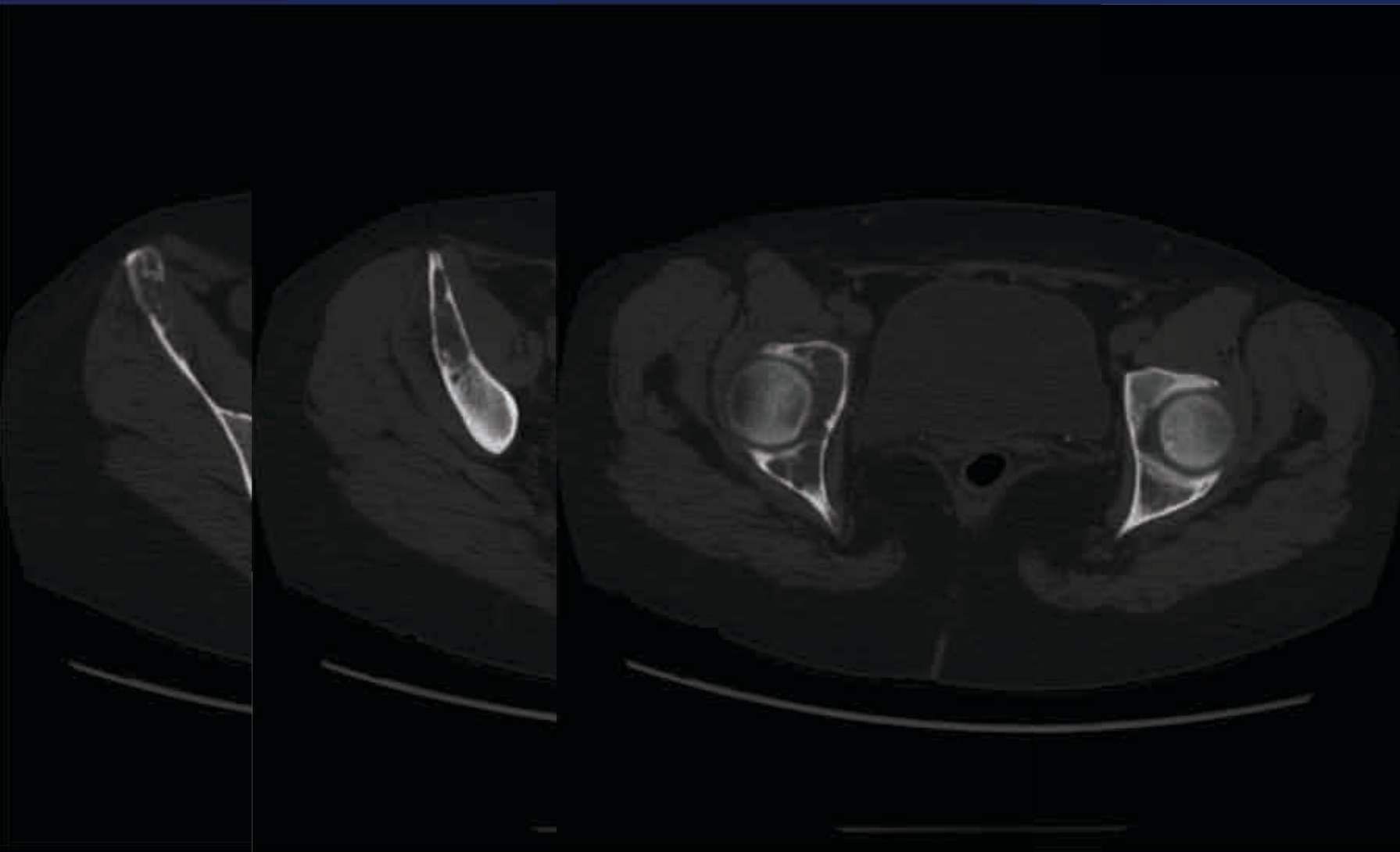




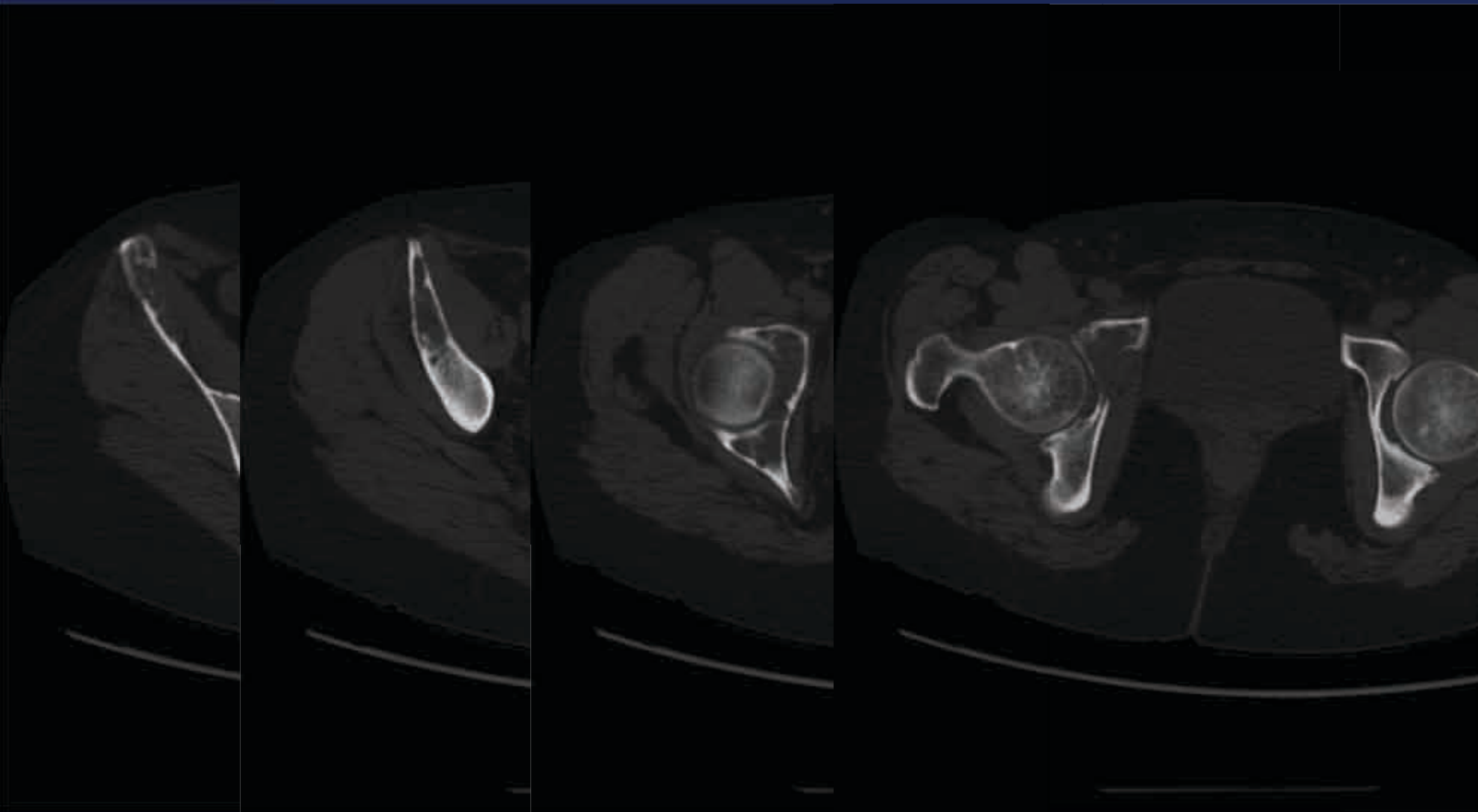
4/21/2009



4/21/2009

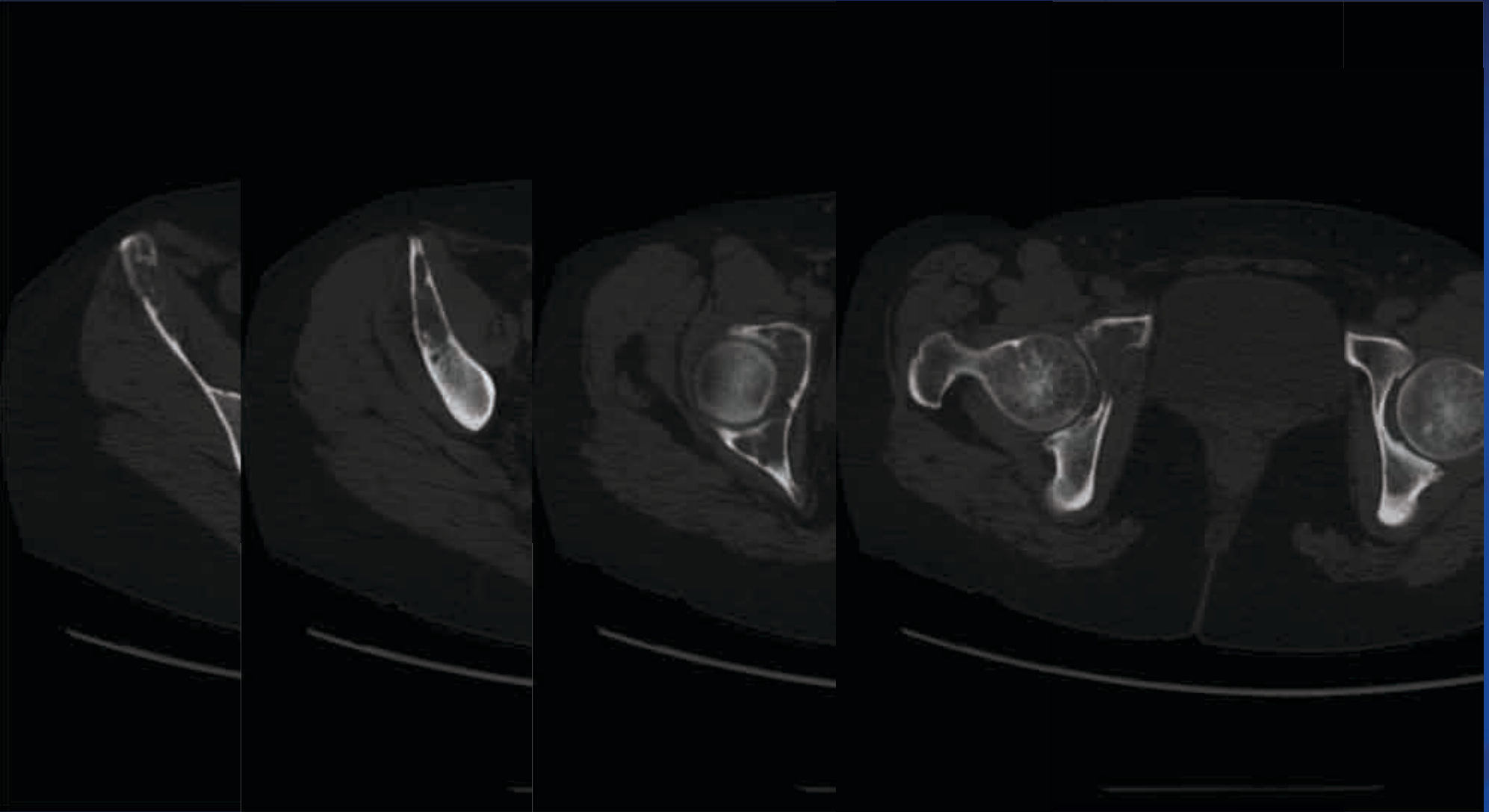


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4/21/2009

## Pelvic bone metastases.

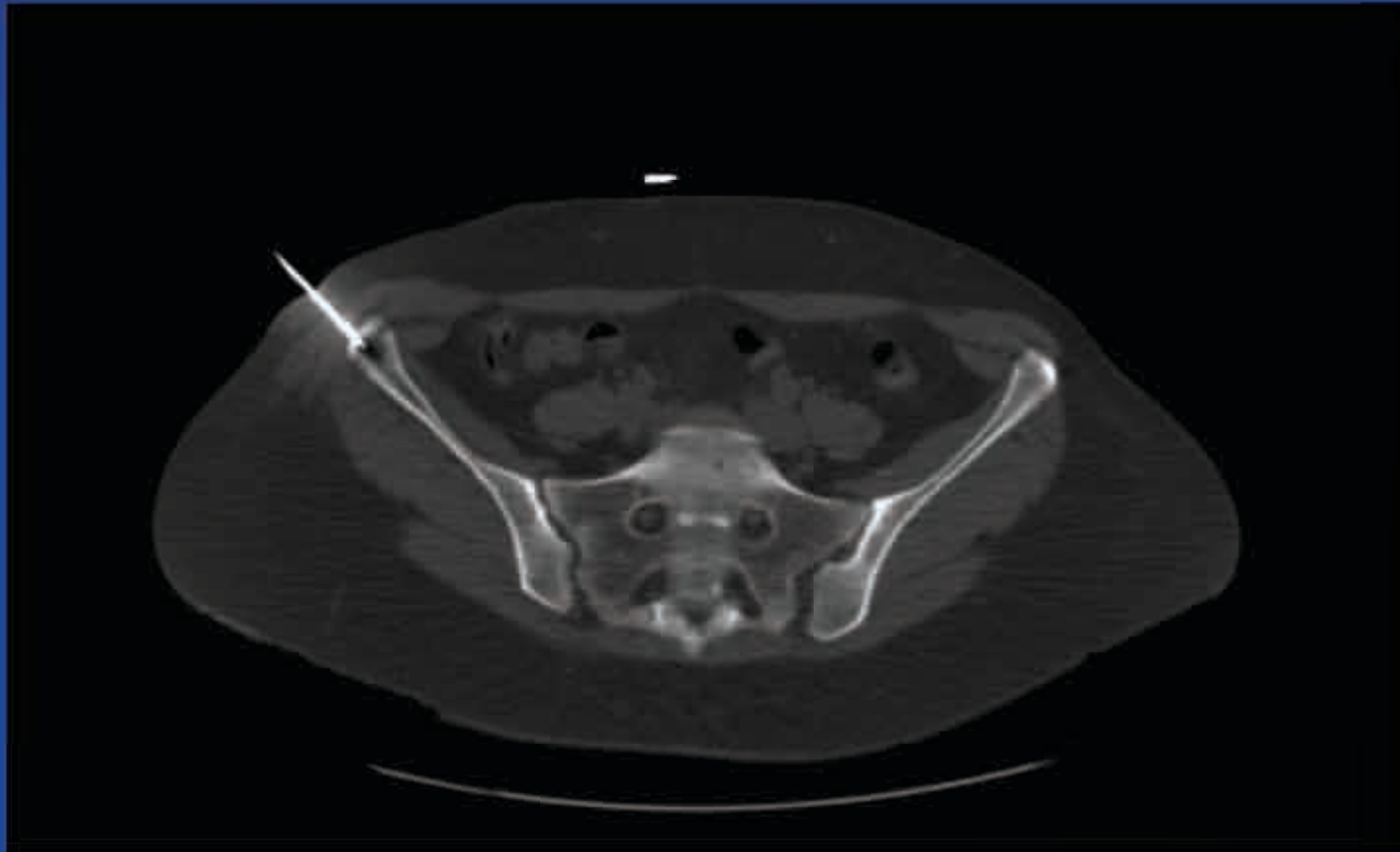


4/21/2009

- Since extensive mets, ortho suggested to first undergo radiation treatment to decrease her pain and for disease control given that there are no imminent fractures.
- Heme/Onc requested a biopsy to confirm not only breast cancer but more so to confirm receptor status and particularly HER2/neu positivity



Fine needle aspiration and core biopsy of right anterior iliac wing lesion submitted to pathology.



4/29/2009



# Radiology / Pathology Conference

Stacie Canacci, M.D.

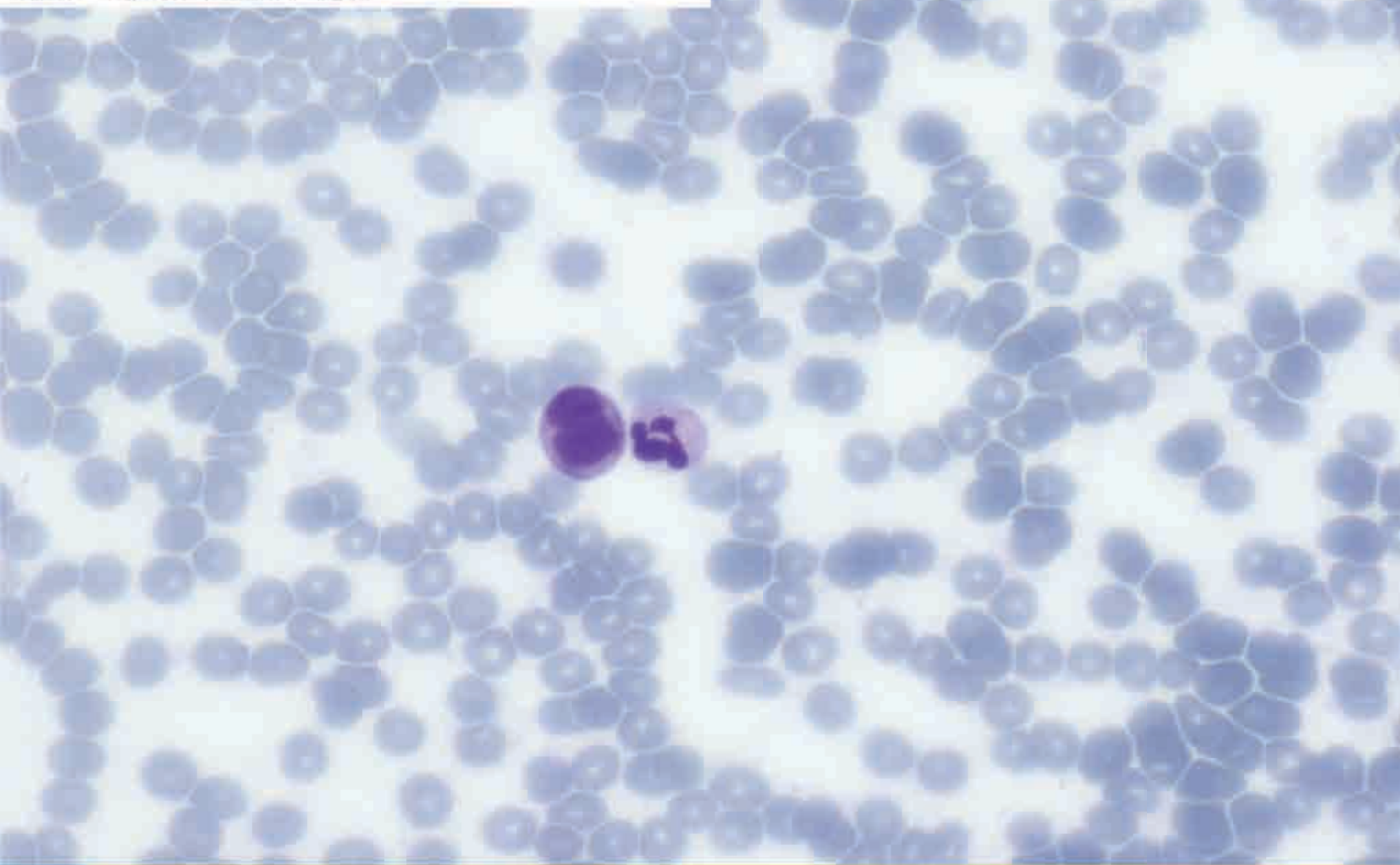
MEDICINE *of* THE HIGHEST ORDER



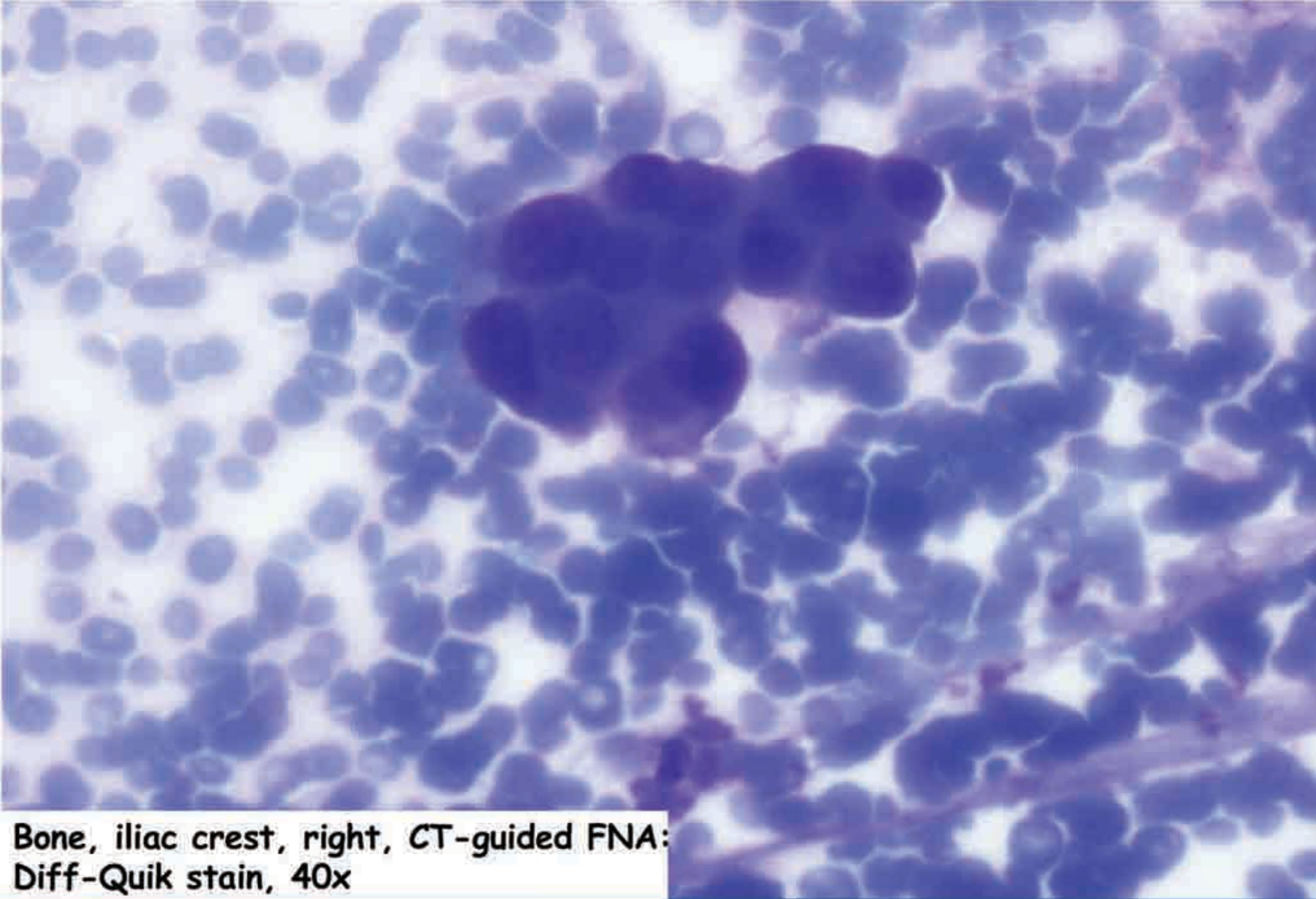
UNIVERSITY *of*  
**ROCHESTER**  
MEDICAL CENTER

# Case 1

**Bone, iliac crest, right, CT-guided FNA:  
Diff-Quik stain, 40x**

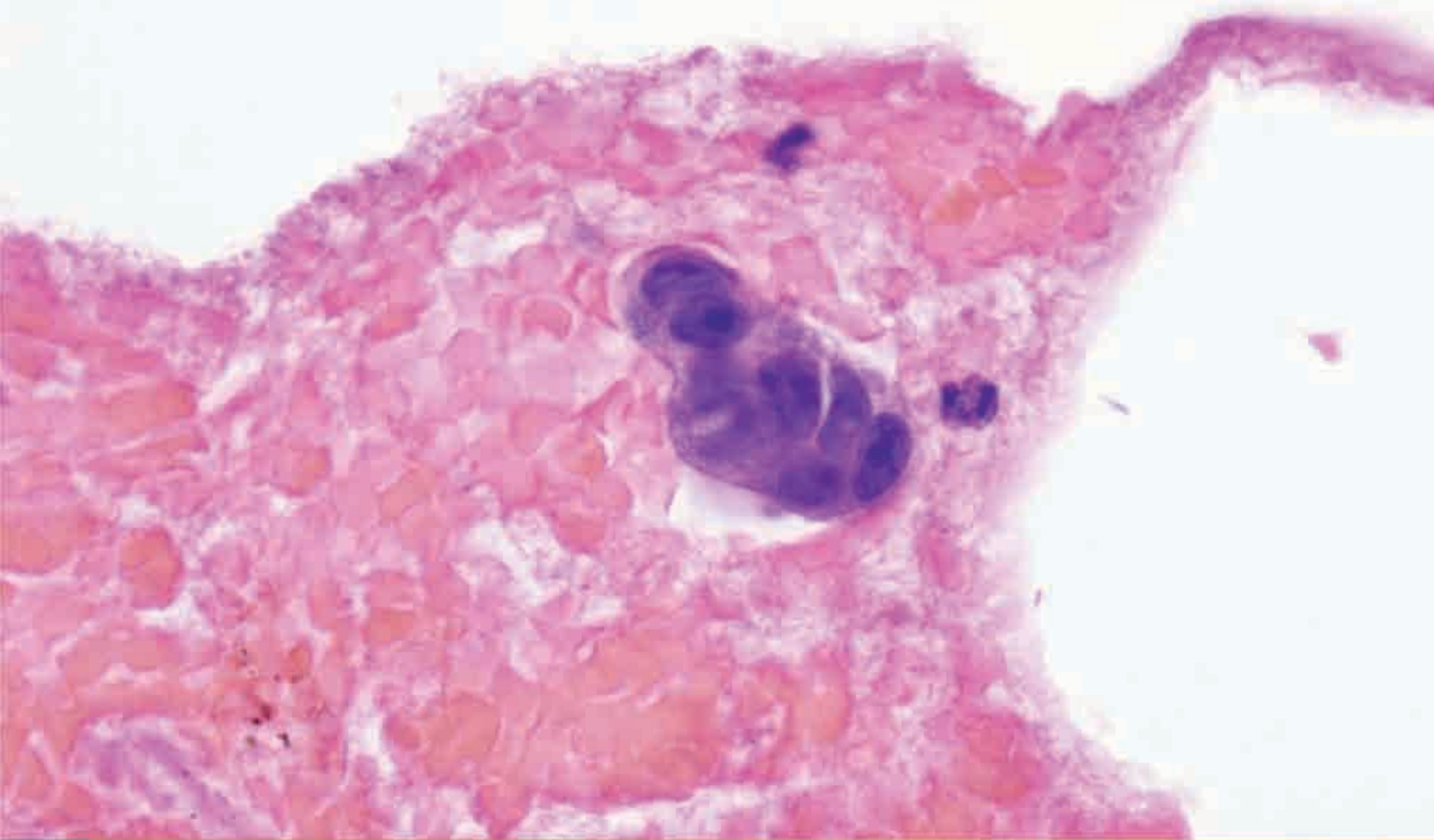






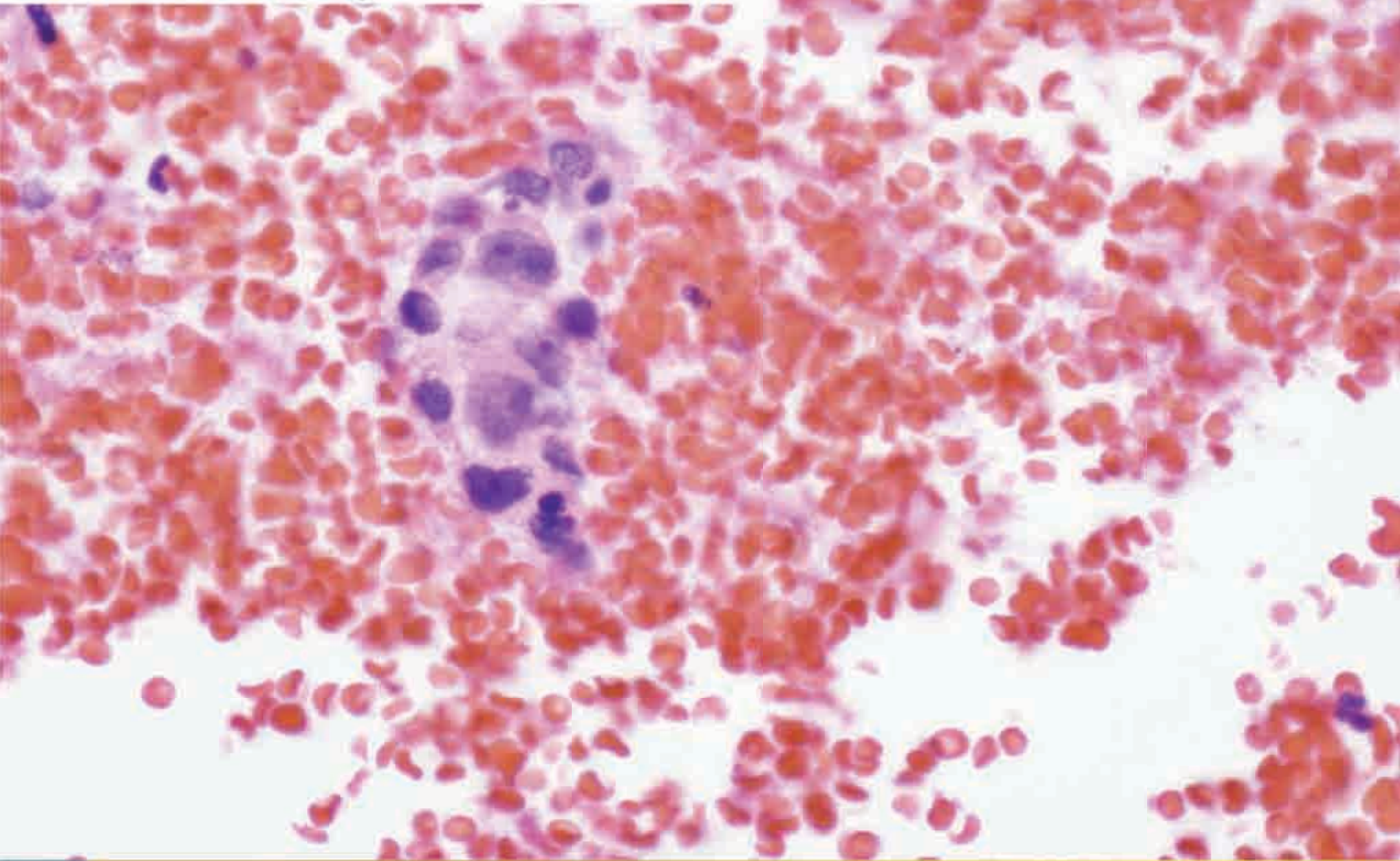
**Bone, iliac crest, right, CT-guided FNA:  
Diff-Quik stain, 40x**

**Bone, iliac crest, right, CT-guided FNA:  
Cell Block, Hematoxylin and eosin stain, 40x**





Bone, iliac crest, right, CT-guided FNA:  
Cell Block, Hematoxylin and eosin stain, 40x



## Bone, iliac crest, right, CT-guided fine needle aspiration:

**Malignant tumor cells present derived from  
adenocarcinoma consistent with origin from breast.**

**Cell block and cytologic preparations examined.**

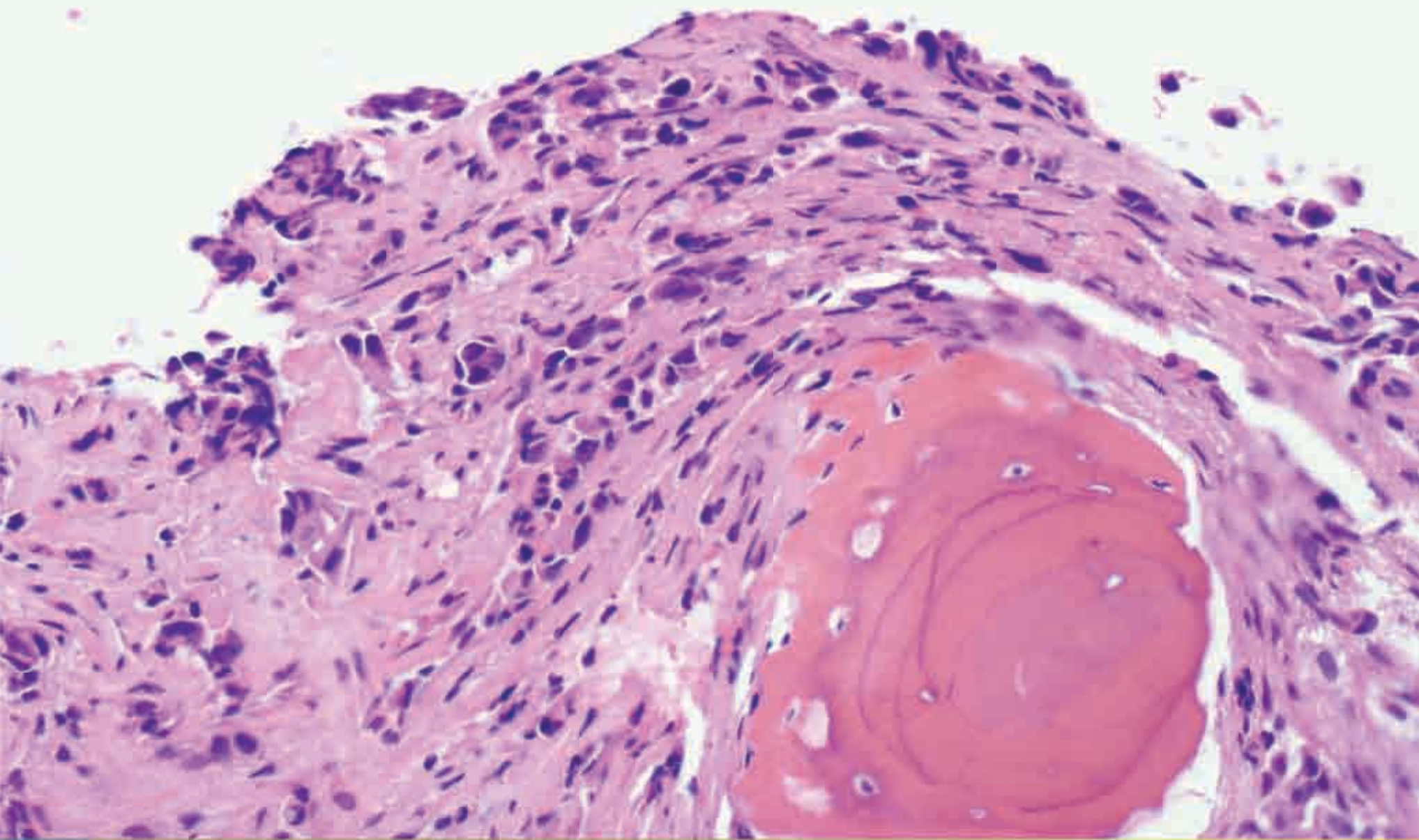


# Bone right anterior iliac, core needle biopsy:

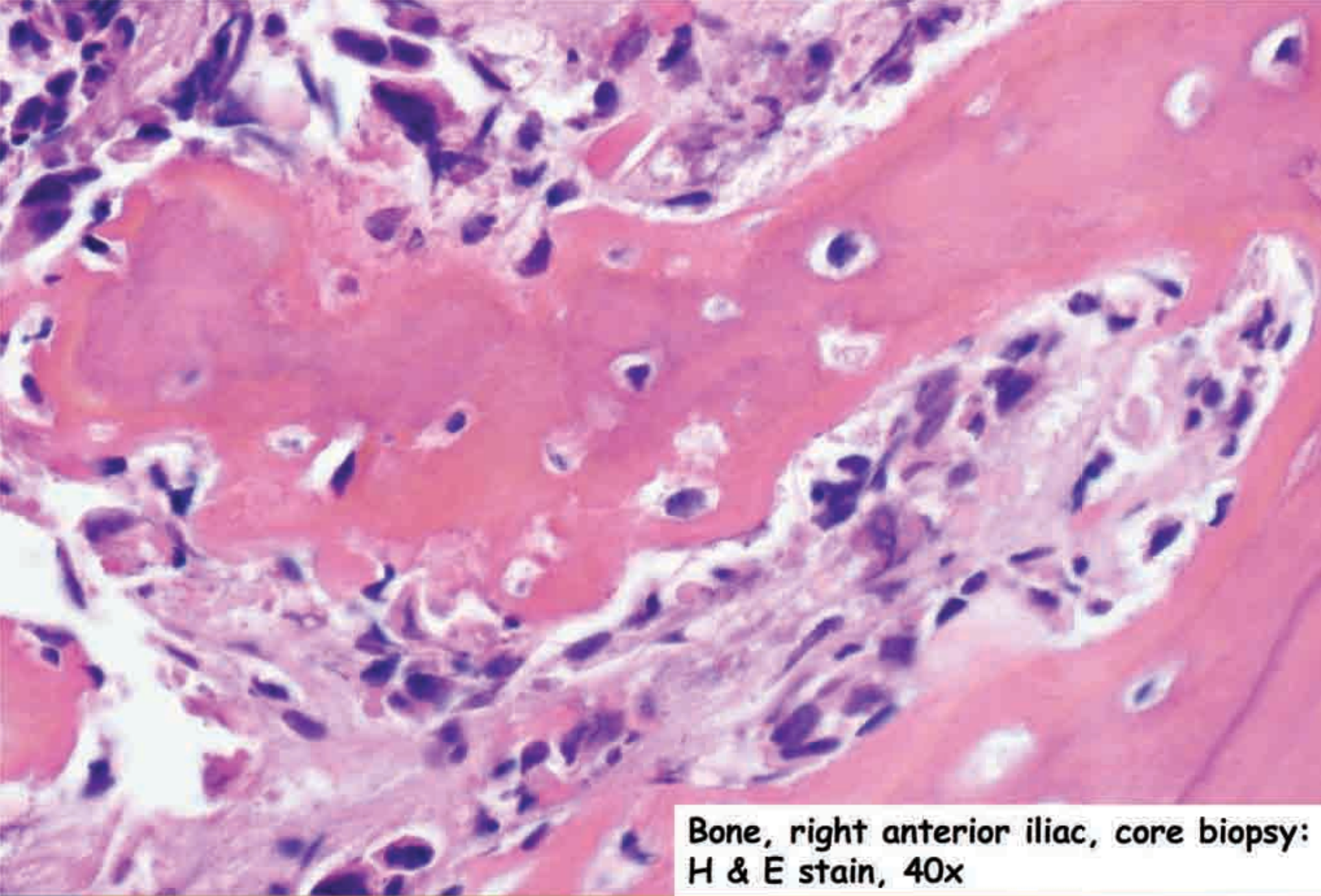
**Metastatic carcinoma, consistent with breast primary.**

**Immunohistochemical stain for Estrogen receptor is moderately positive and a few cells are positive for Progesterone receptor. Her2 staining is negative.**

**Bone, right anterior iliac, core biopsy:  
H & E stain, 20x**







**Bone, right anterior iliac, core biopsy:  
H & E stain, 40x**

# Restaging Breast Cancer

- SUV measurements helpful in determining adequate response to chemotherapy:
  - Greater than 50% SUV reduction at 9 weeks accurately defined good clinical response
  - Greater than 55% response after one cycle also defined good response

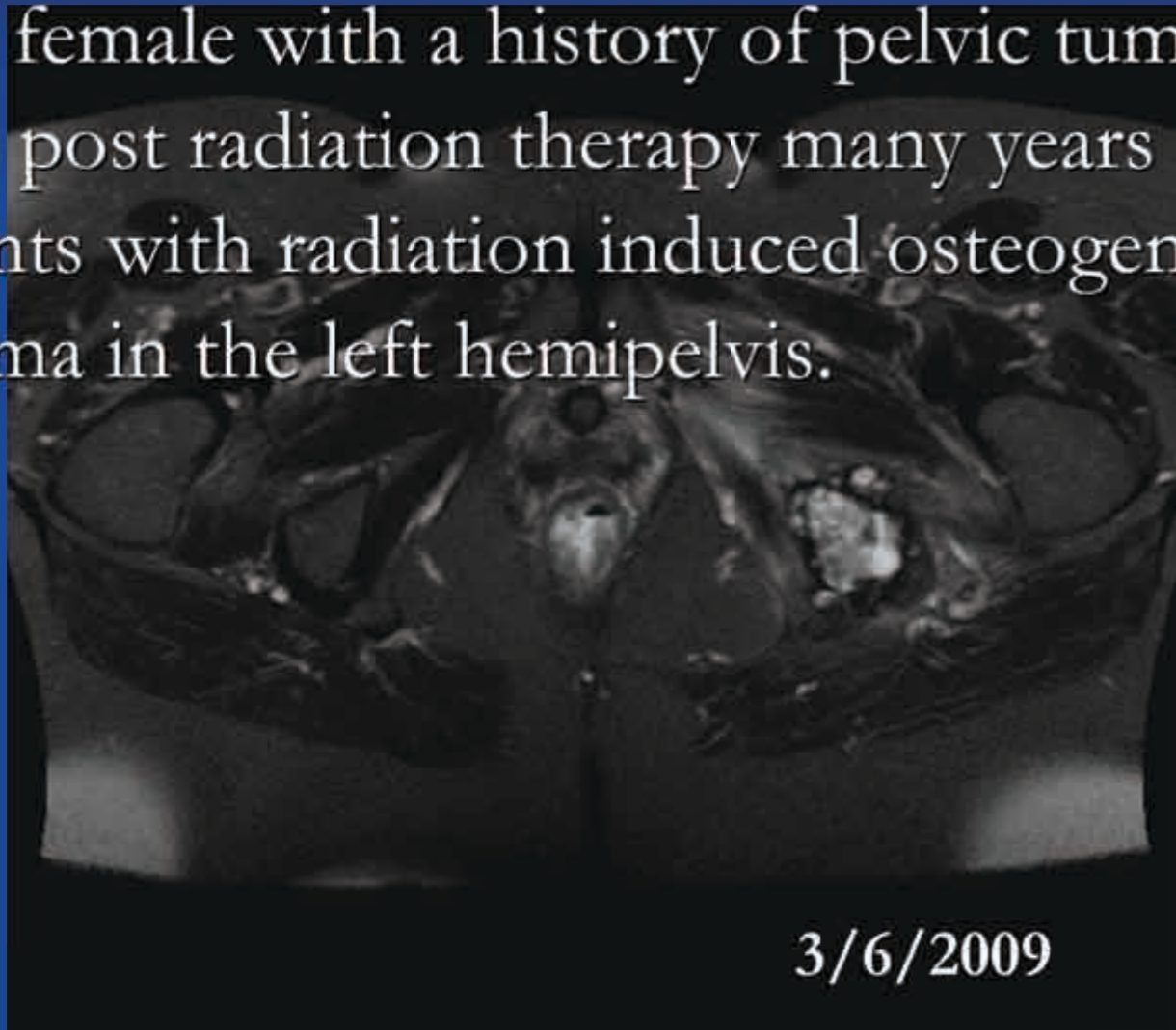


## Case 2

- 63 yo female with a history of pelvic tumor status post radiation therapy many years ago presents with radiation induced osteogenic sarcoma in the left hemipelvis.

## Case 2

- 63 yo female with a history of pelvic tumor status post radiation therapy many years ago presents with radiation induced osteogenic sarcoma in the left hemipelvis.

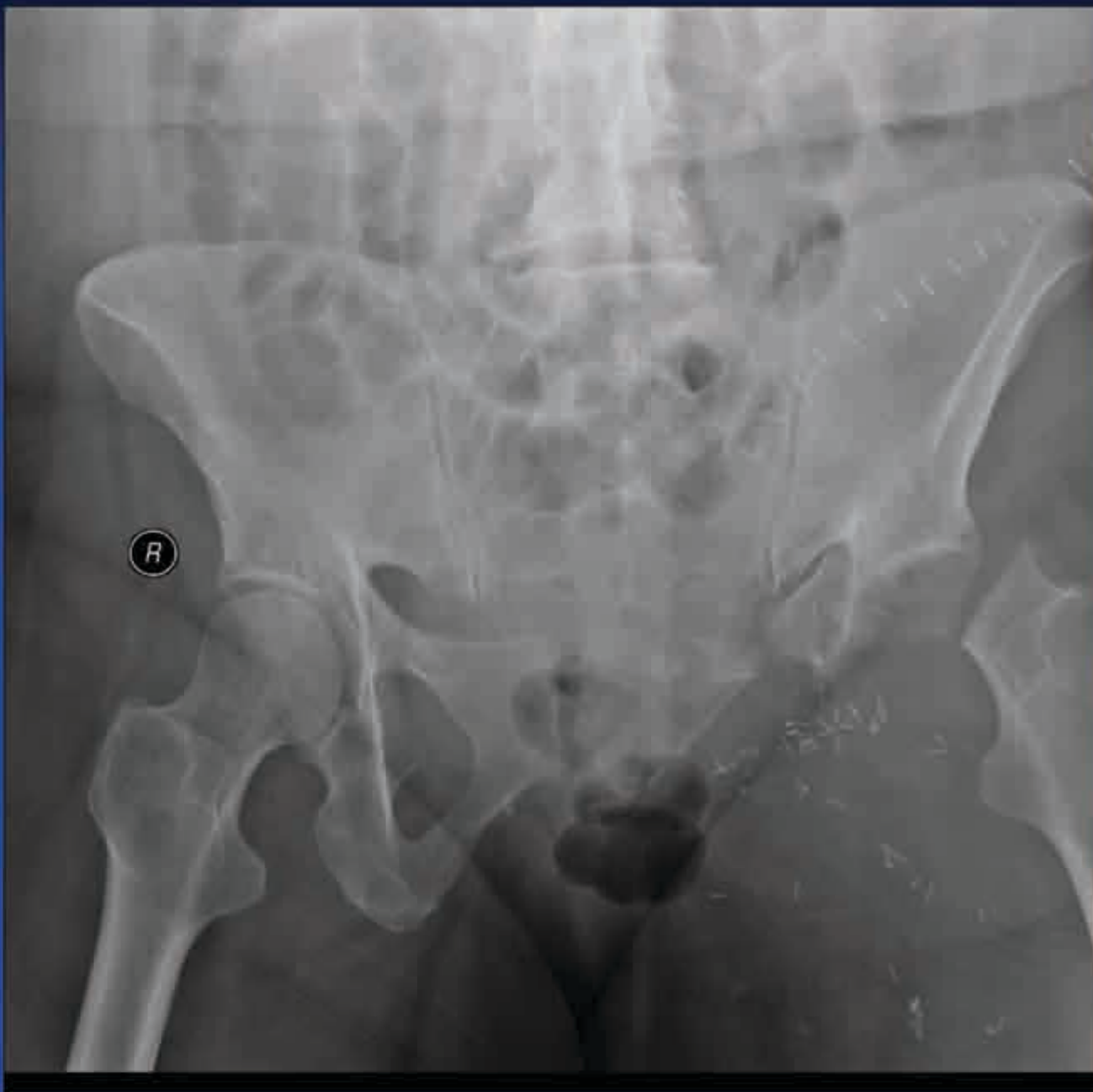


3/6/2009

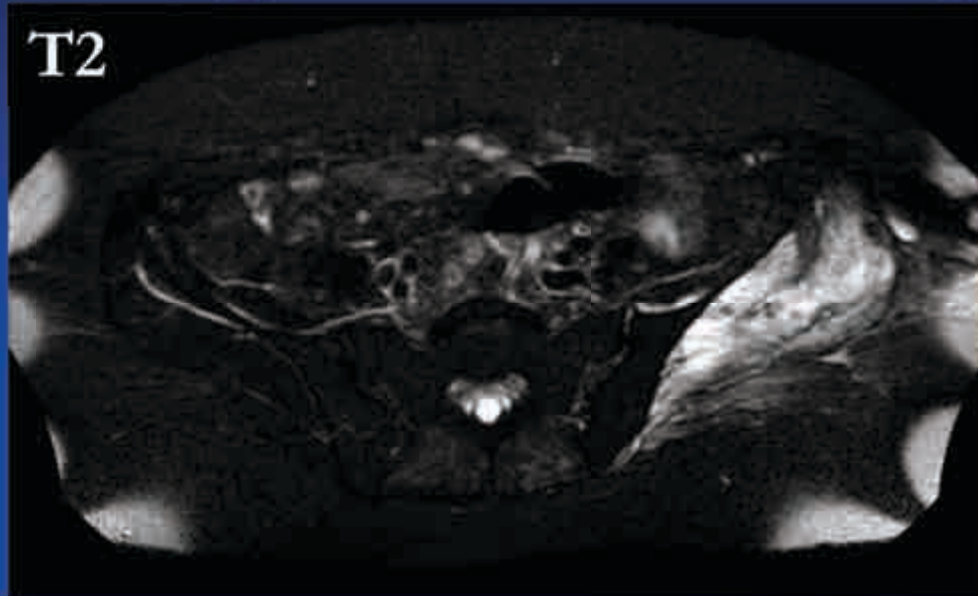
## Case 2

- 63 yo female with a history of pelvic tumor status post radiation therapy many years ago presents with radiation induced osteogenic sarcoma in the left hemipelvis.
- Initial treatment performed at RGH consisted of chemotherapy with a poor response. She was transferred to Strong and underwent two surgical resections on 3/17/2009 and 3/27/2009.





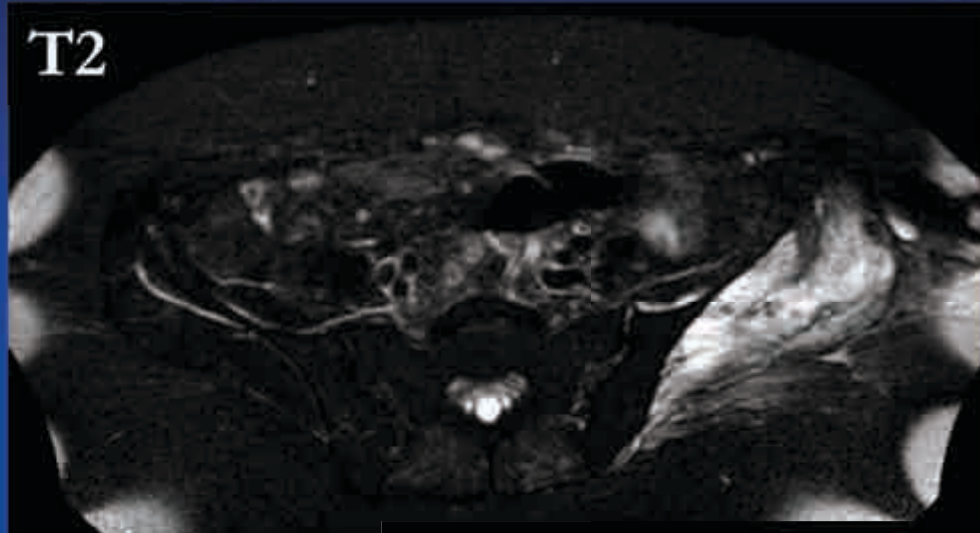
# Follow-up MRI of the pelvis



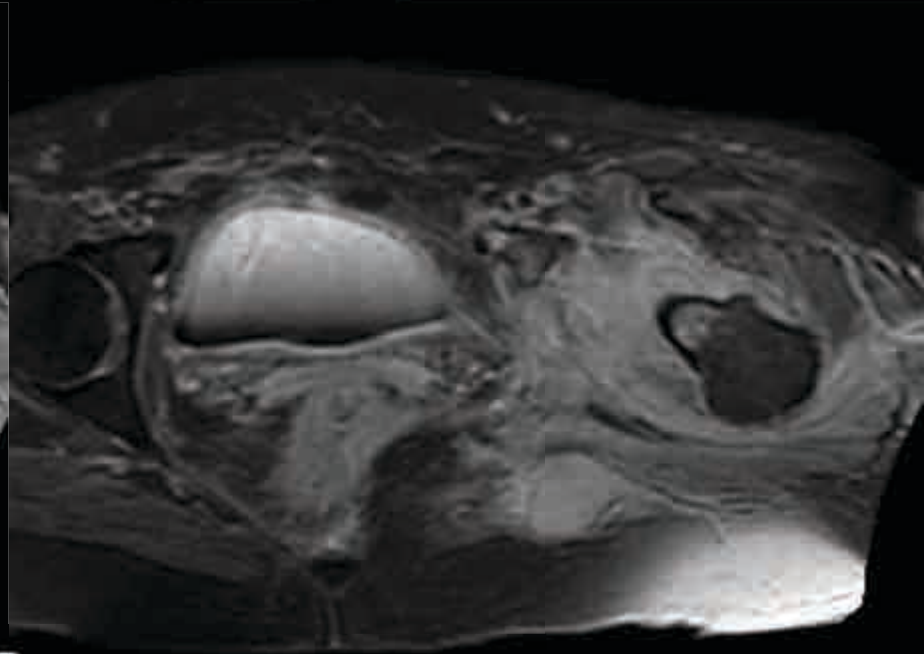
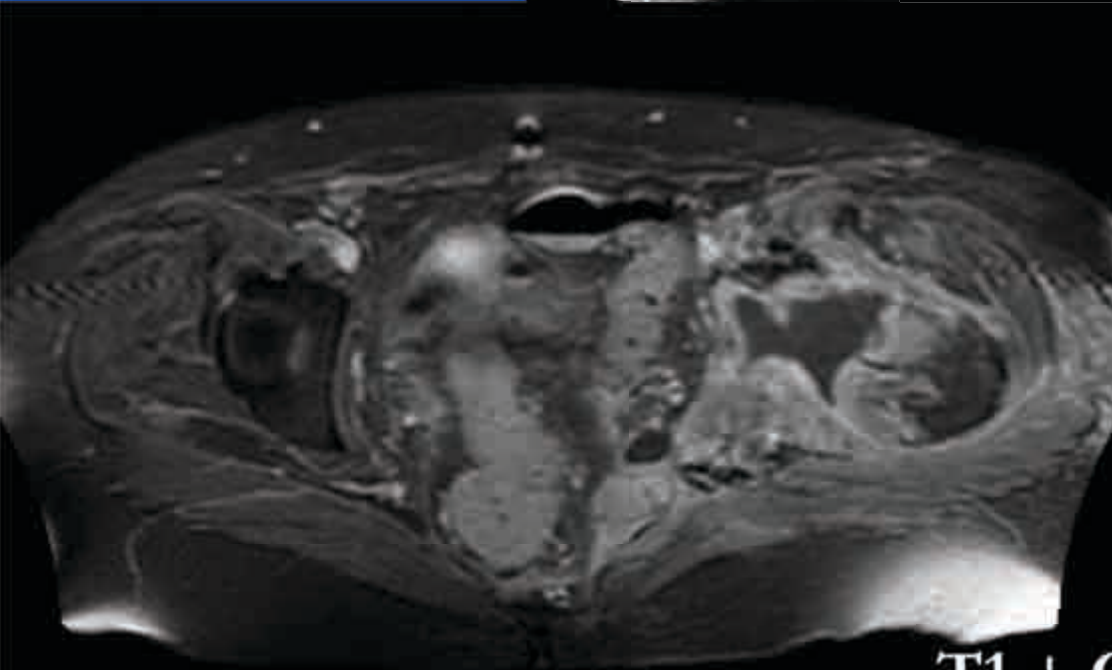
5/1/2009

# Follow-up MRI of the pelvis

T2



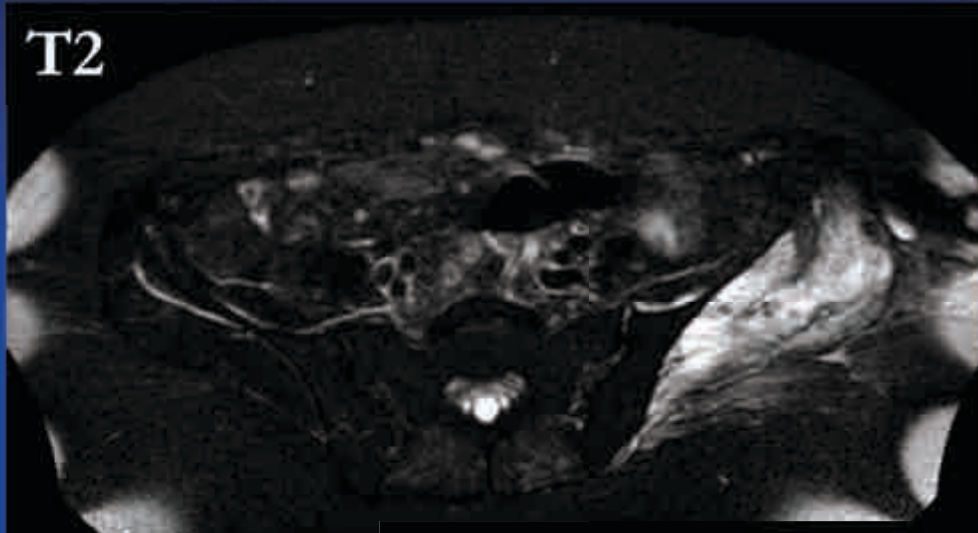
5/1/2009



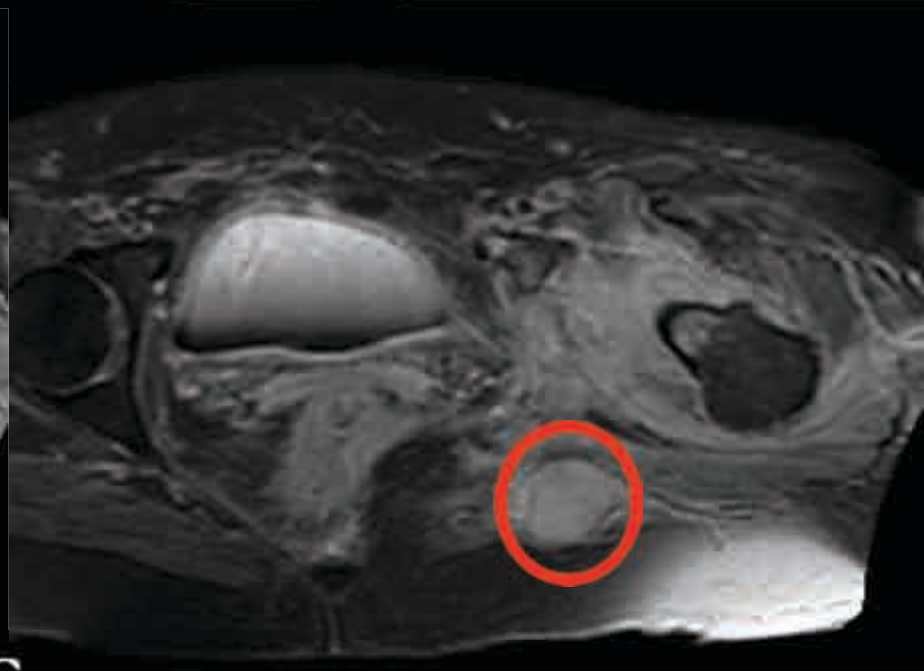
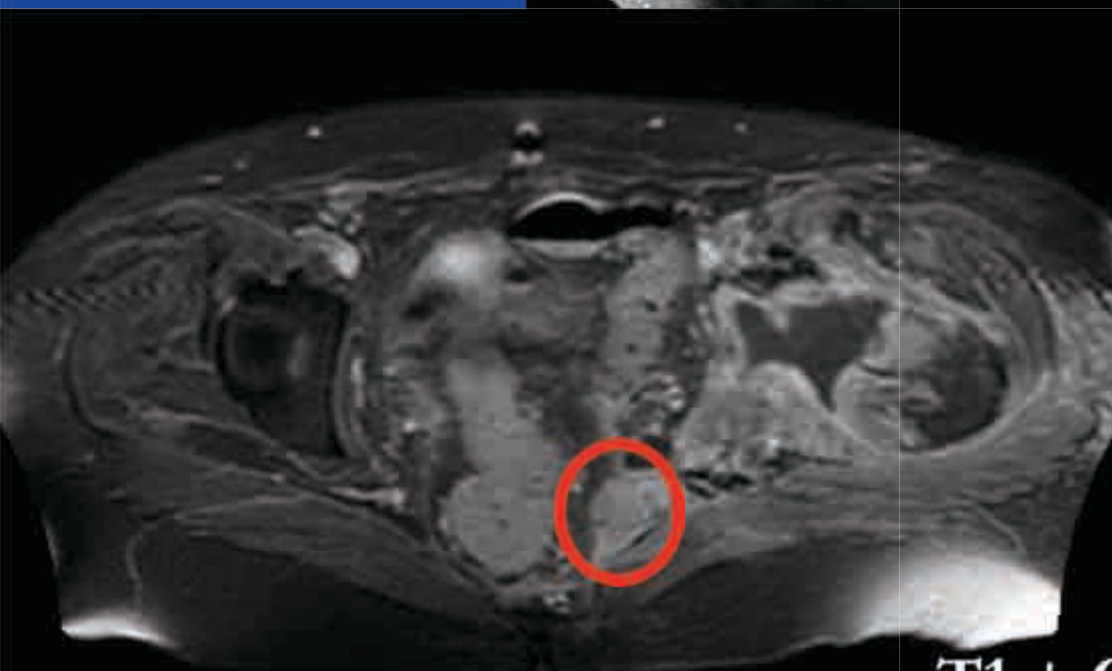
T1 + C

# Follow-up MRI of the pelvis

T2



5/1/2009

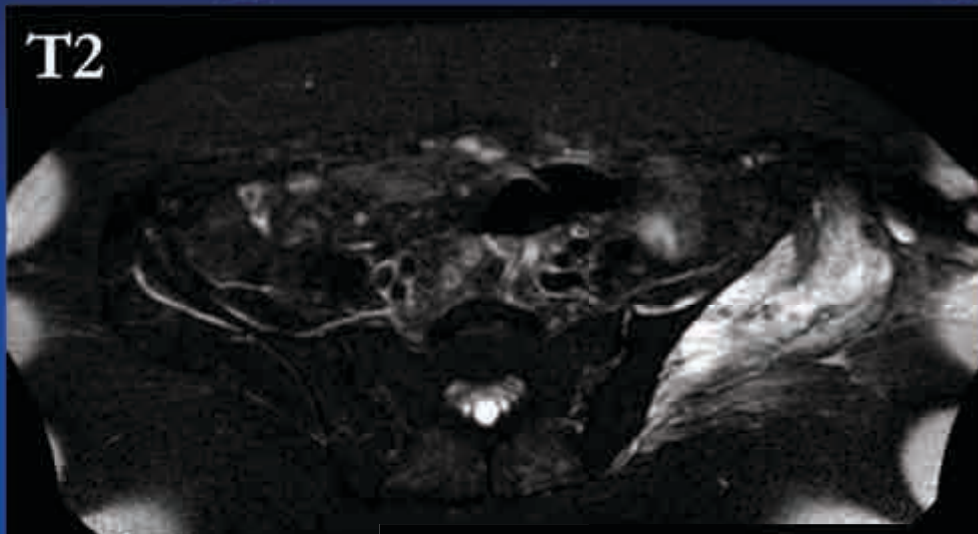


T1 + C

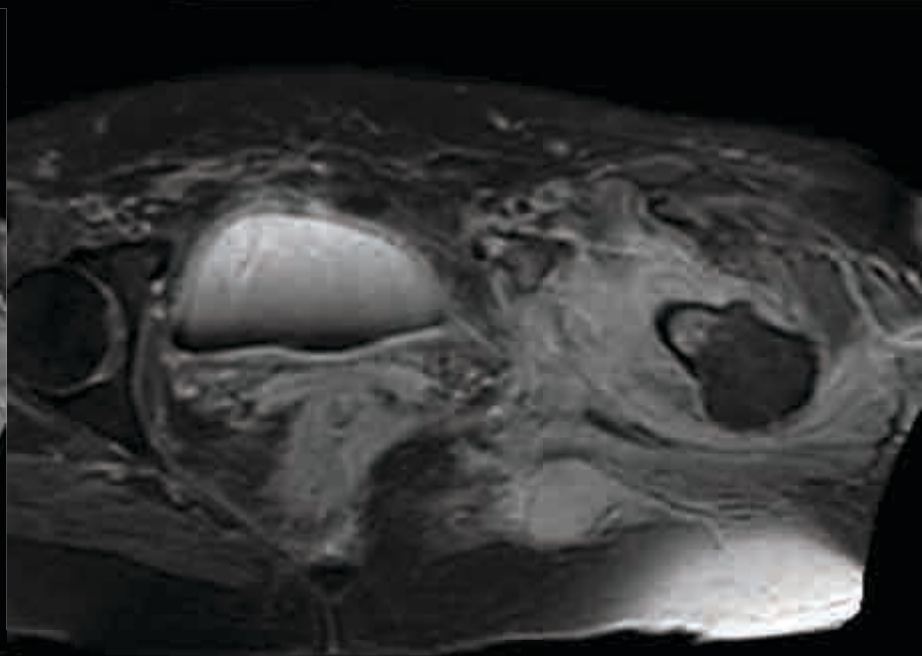
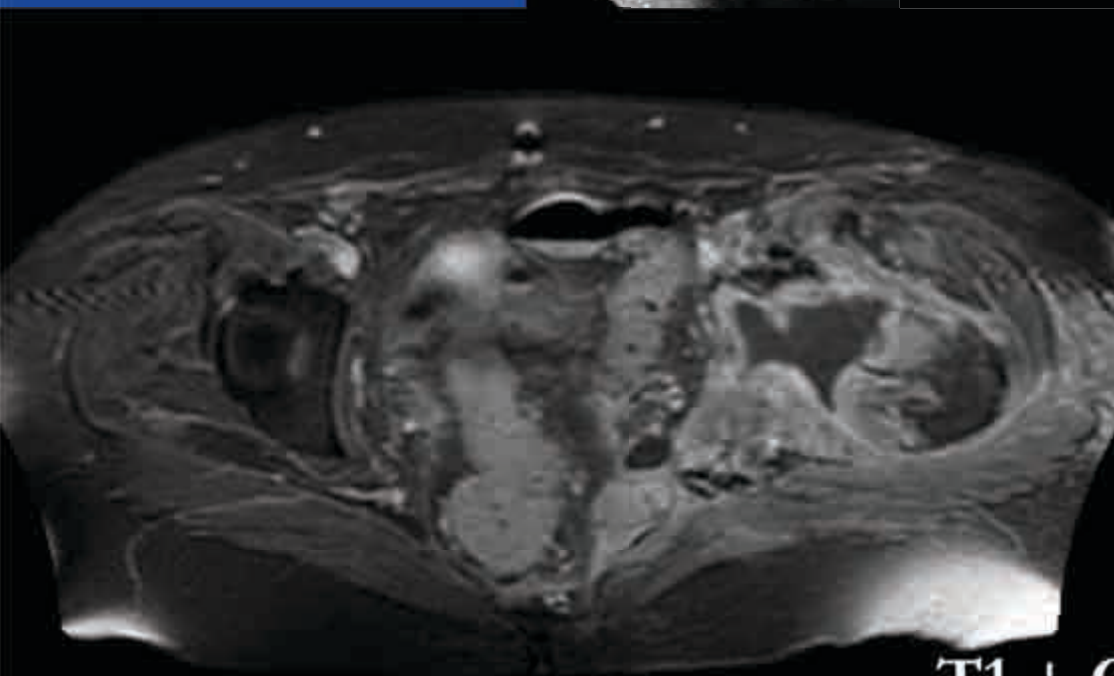


# Follow-up MRI of the pelvis

T2



5/1/2009

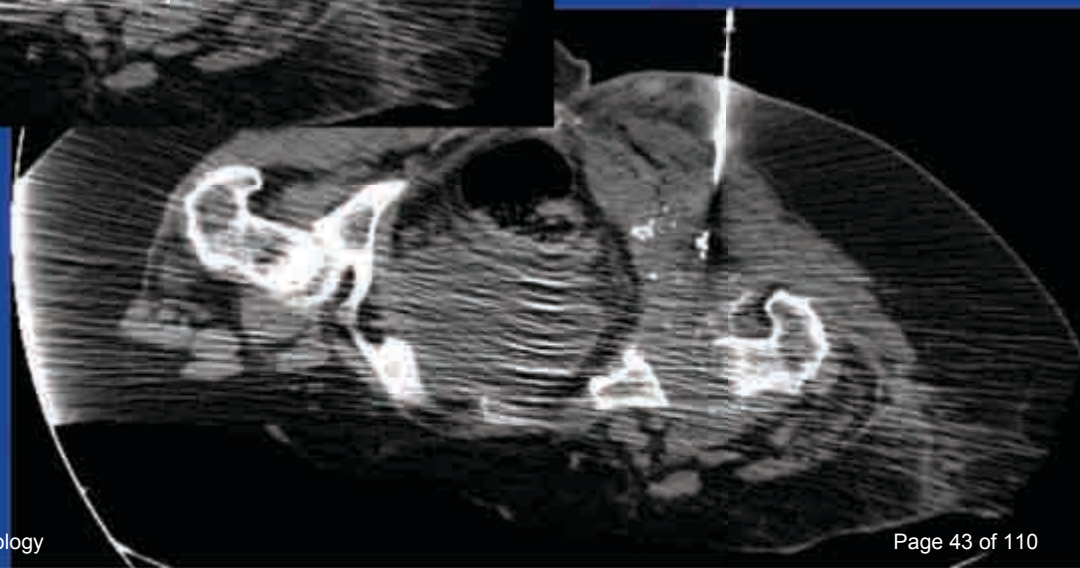
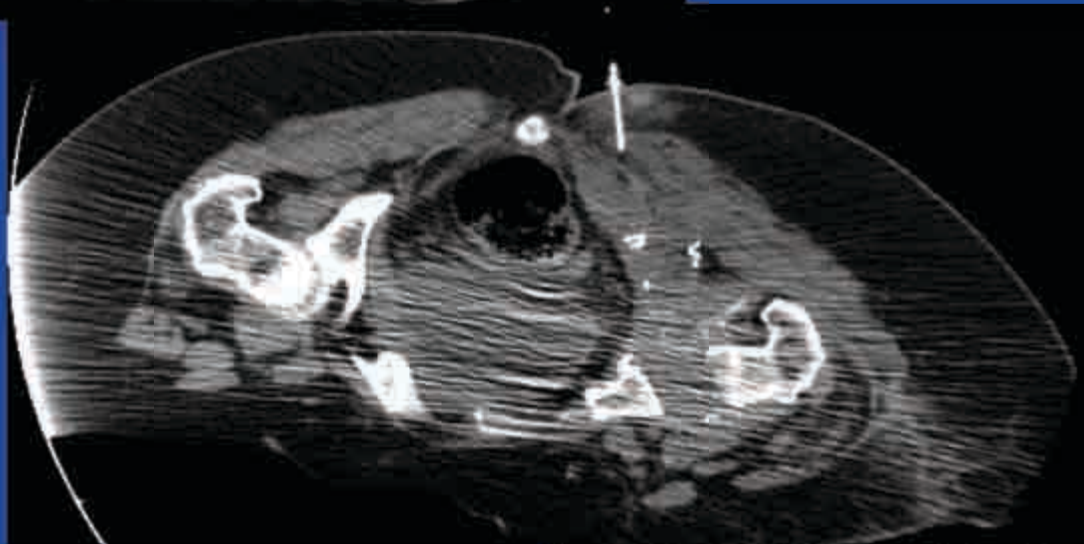
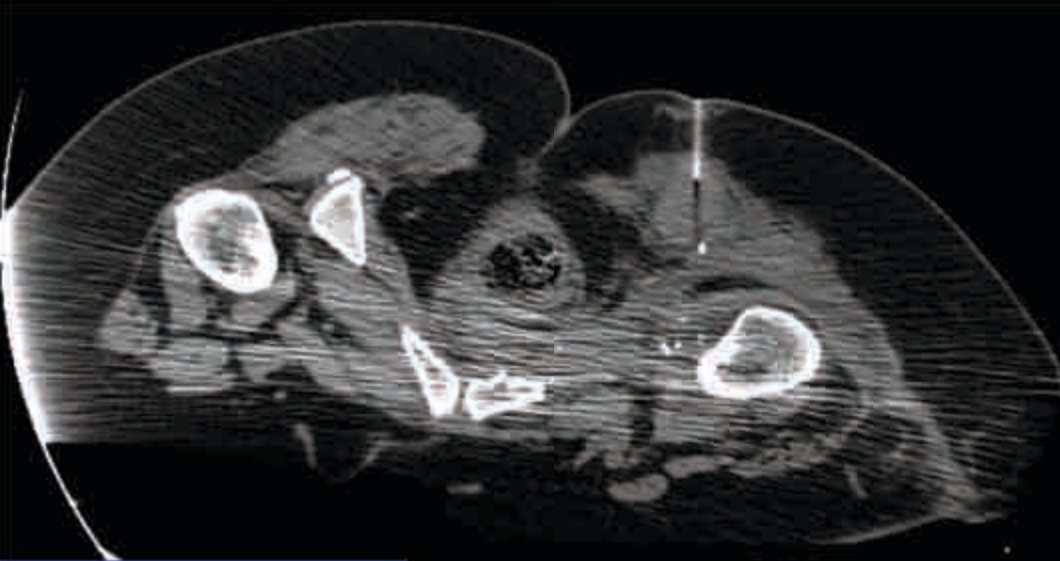


T1 + C

# Differential Diagnosis

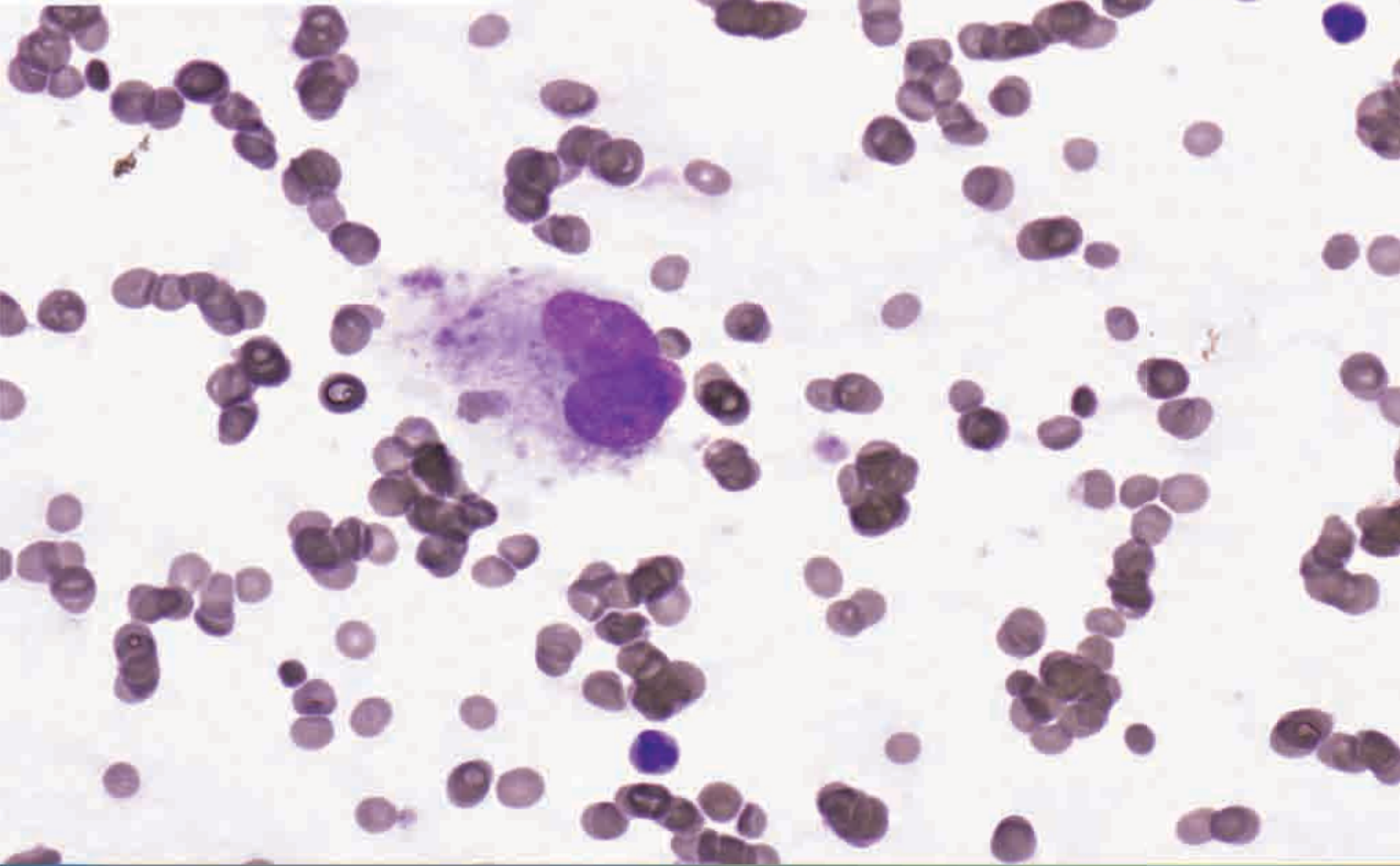
- Metastasis
- Atypical hematoma





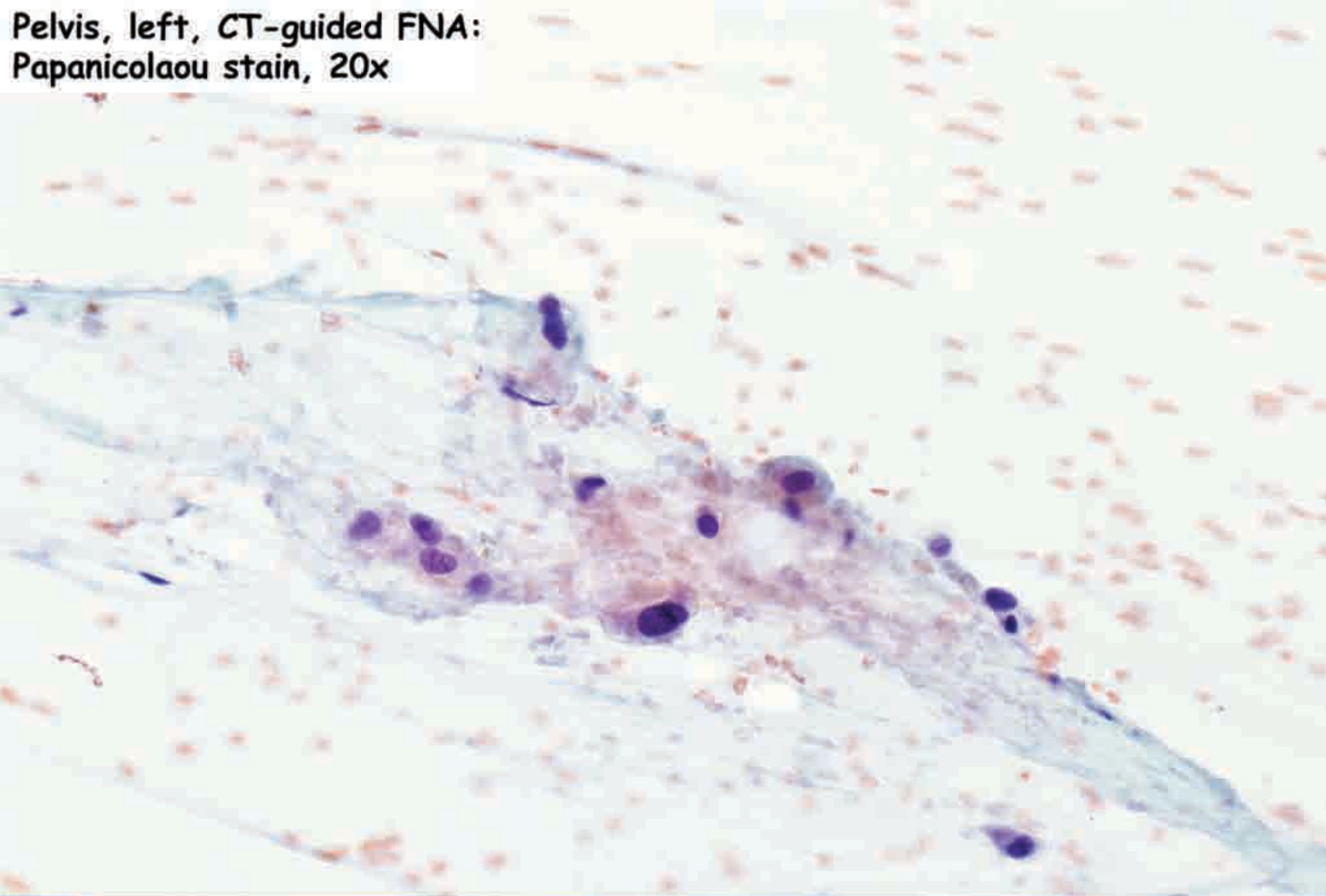
# Case 2

**Pelvis, left, CT-guided FNA:  
Diff-Quik stain, 40x**



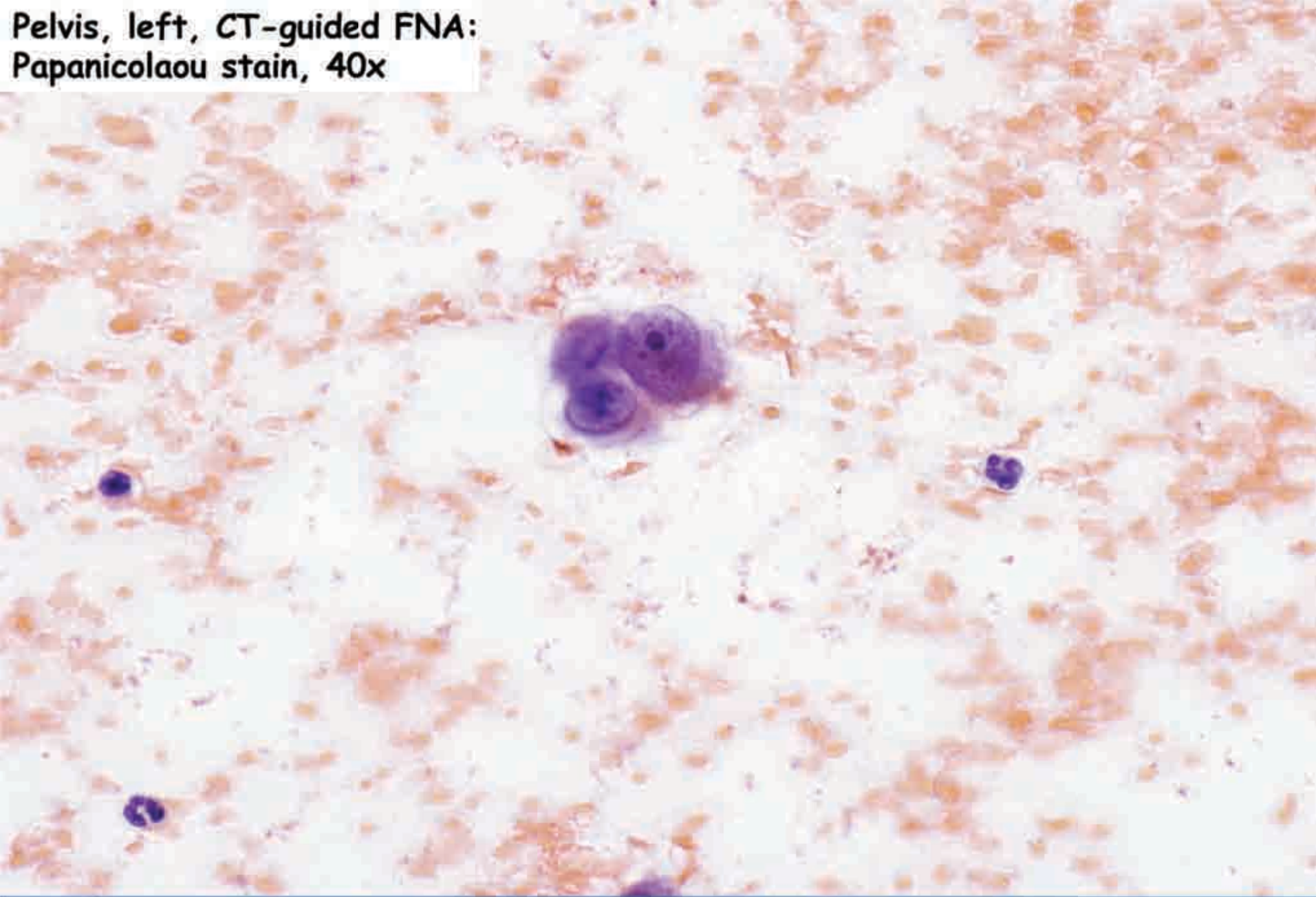


**Pelvis, left, CT-guided FNA:  
Papanicolaou stain, 20x**

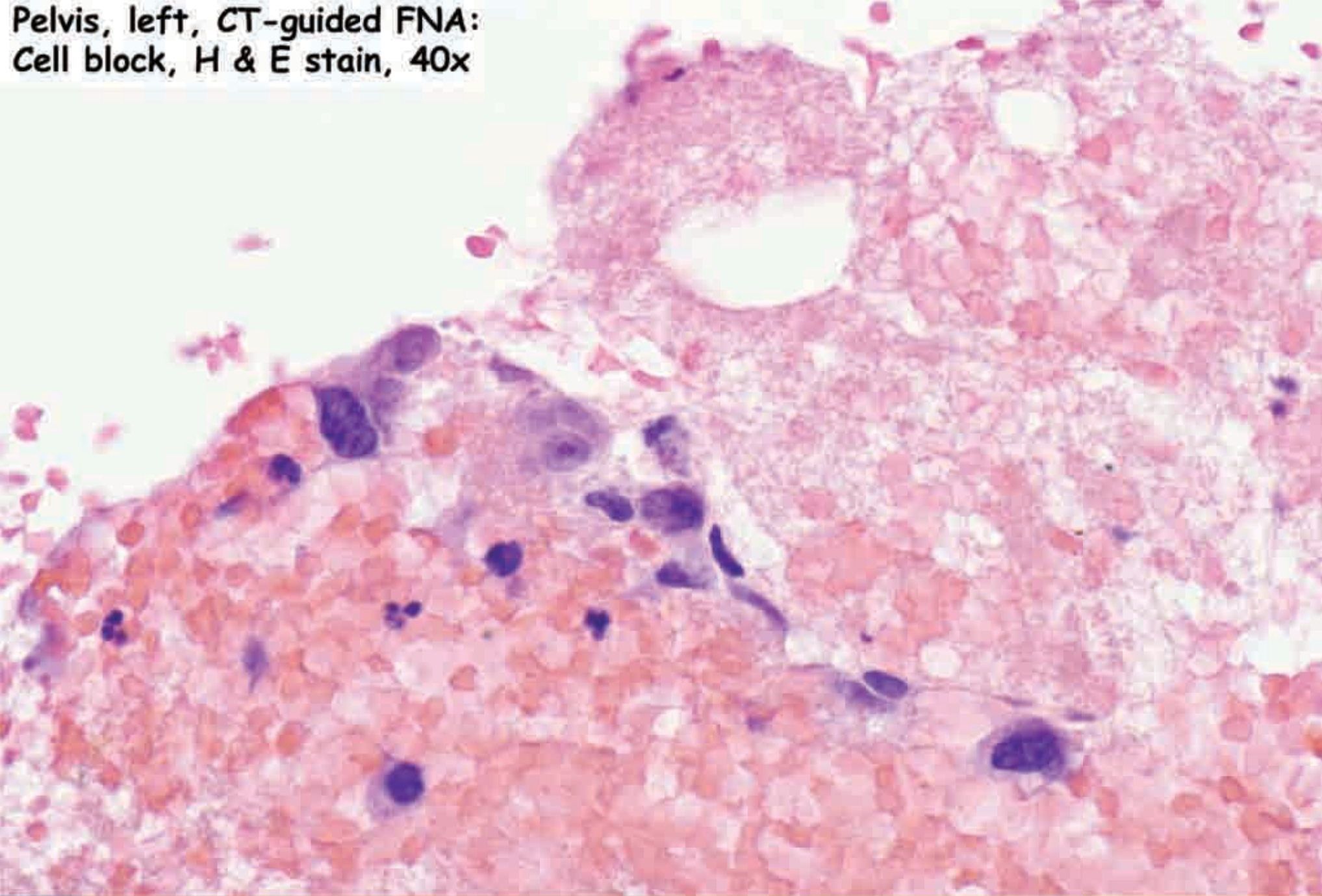




Pelvis, left, CT-guided FNA:  
Papanicolaou stain, 40x



**Pelvis, left, CT-guided FNA:  
Cell block, H & E stain, 40x**





# Pelvis, left, CT-guided fine needle aspiration:

**Malignant tumor cells present consistent with osteogenic sarcoma.**

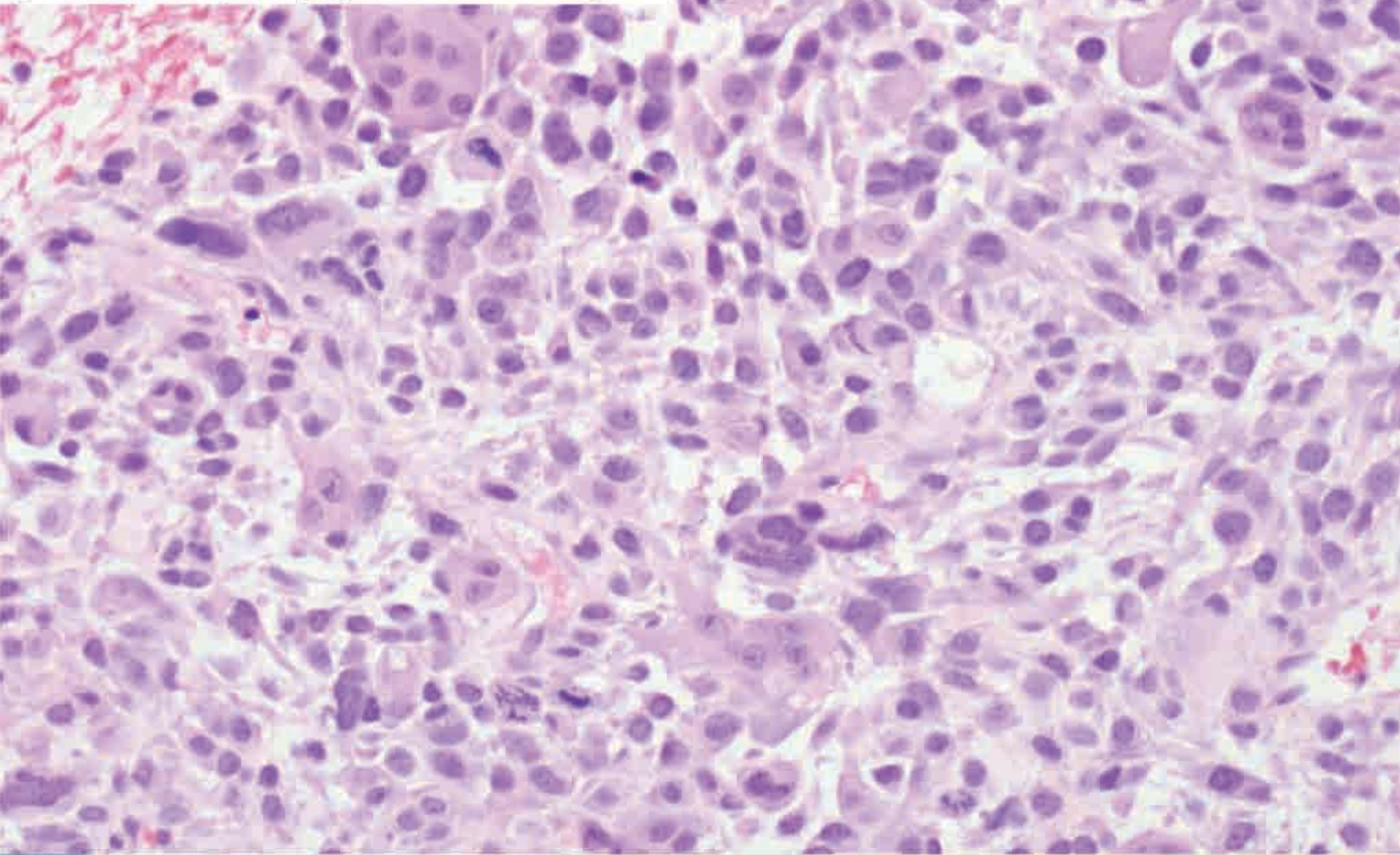
**Cell block and cytologic preparations examined.**

**Soft tissue, left piriformis and gluteal mass, biopsy:**

**High grade osteogenic sarcoma (clinically recurrent)**

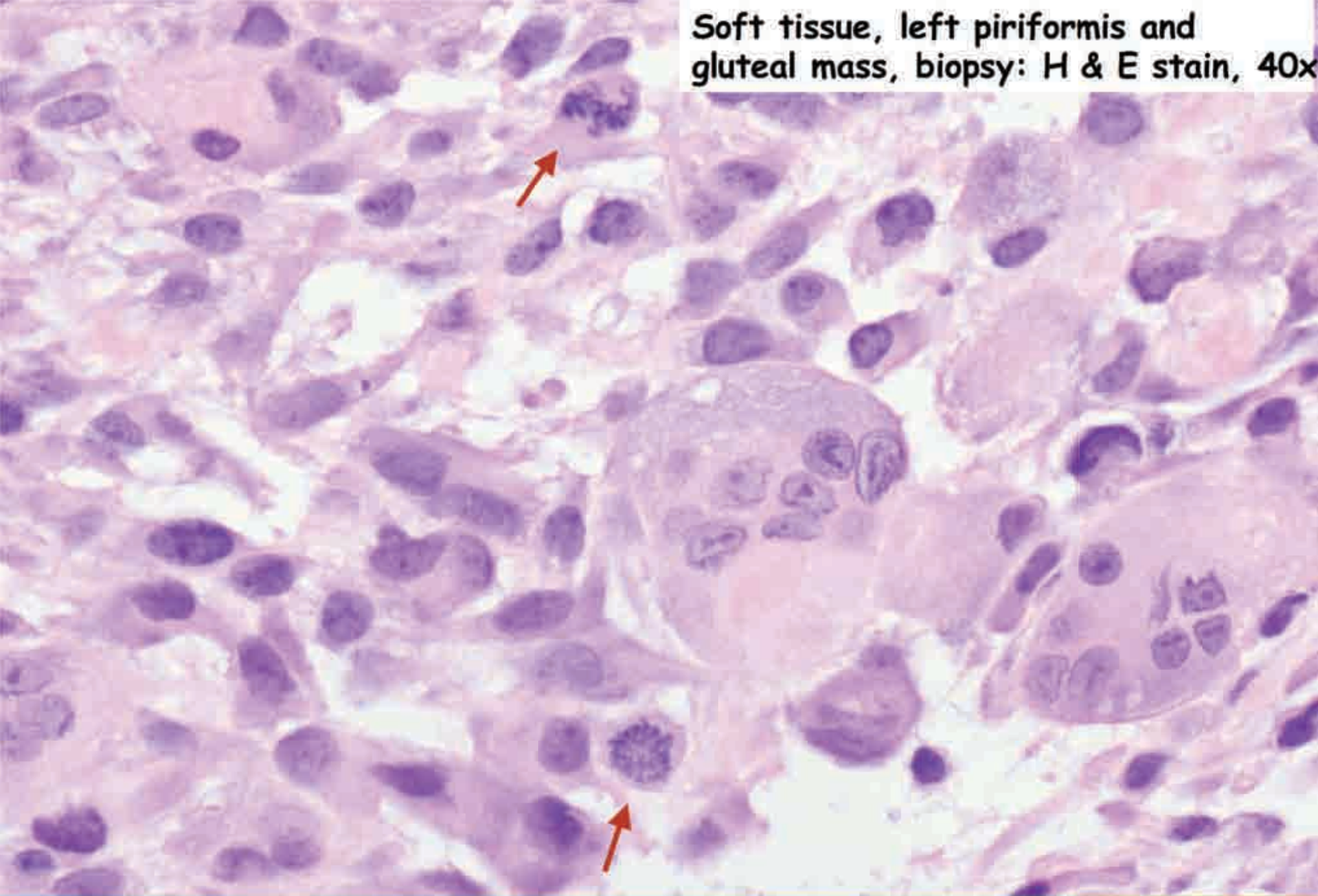


Soft tissue, left piriformis and  
gluteal mass, biopsy: H & E stain, 20x





Soft tissue, left piriformis and  
gluteal mass, biopsy: H & E stain, 40x



# Osteogenic Sarcoma

- Malignant tumor with ability to produce osteoid directly from neoplastic cells
- Best diagnostic clue: Bone destruction with associated tumor bone formation and aggressive periosteal reaction.
- Multiple types:
  - Conventional, telangiectatic, multicentric, parosteal, periosteal, and
  - Secondary Osteosarcoma (5%) – in association with preexisting bone lesion such as prior radiation, bone infarct, Pagets



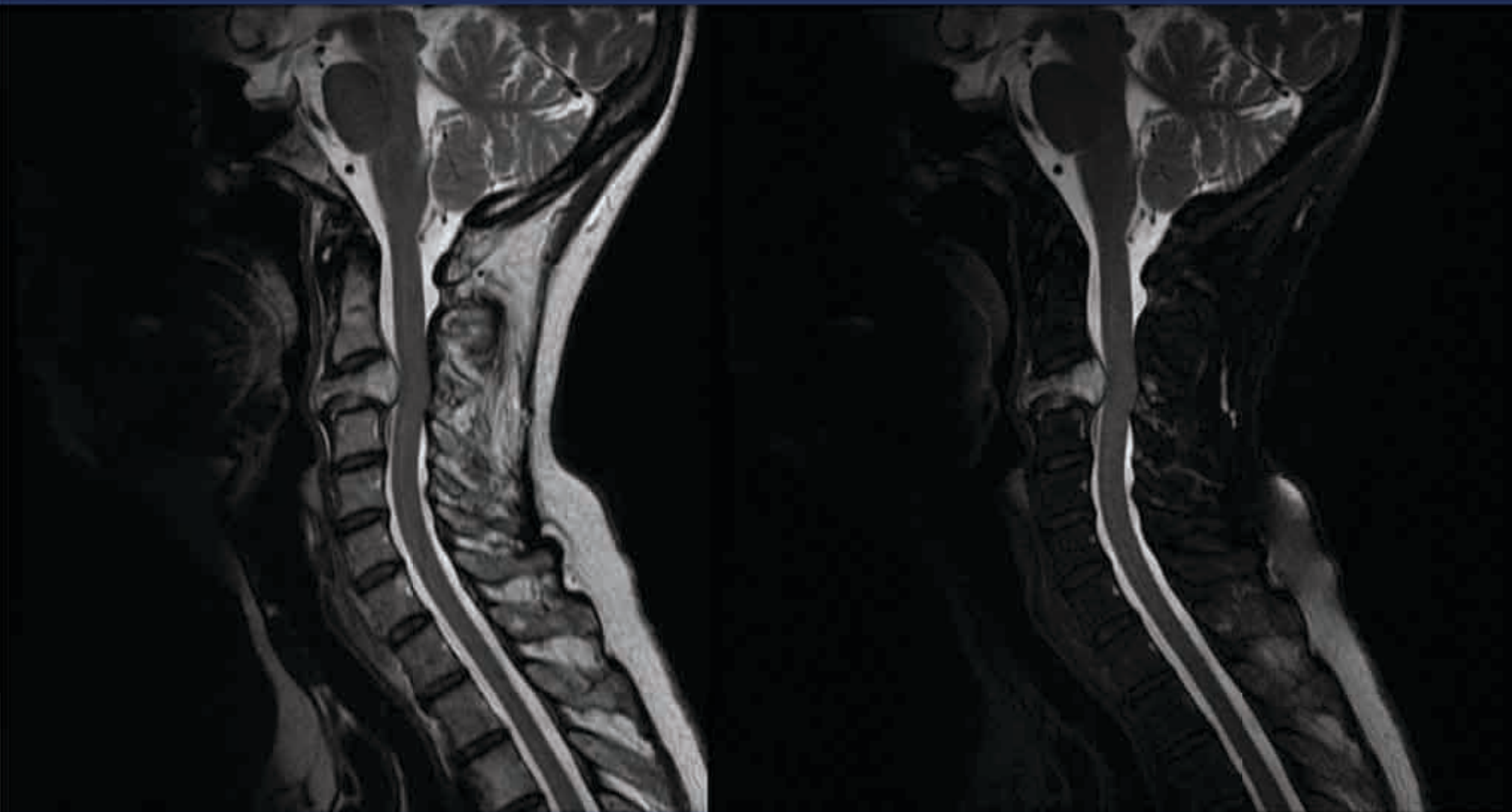
# Osteogenic Sarcoma - Clinical Issues

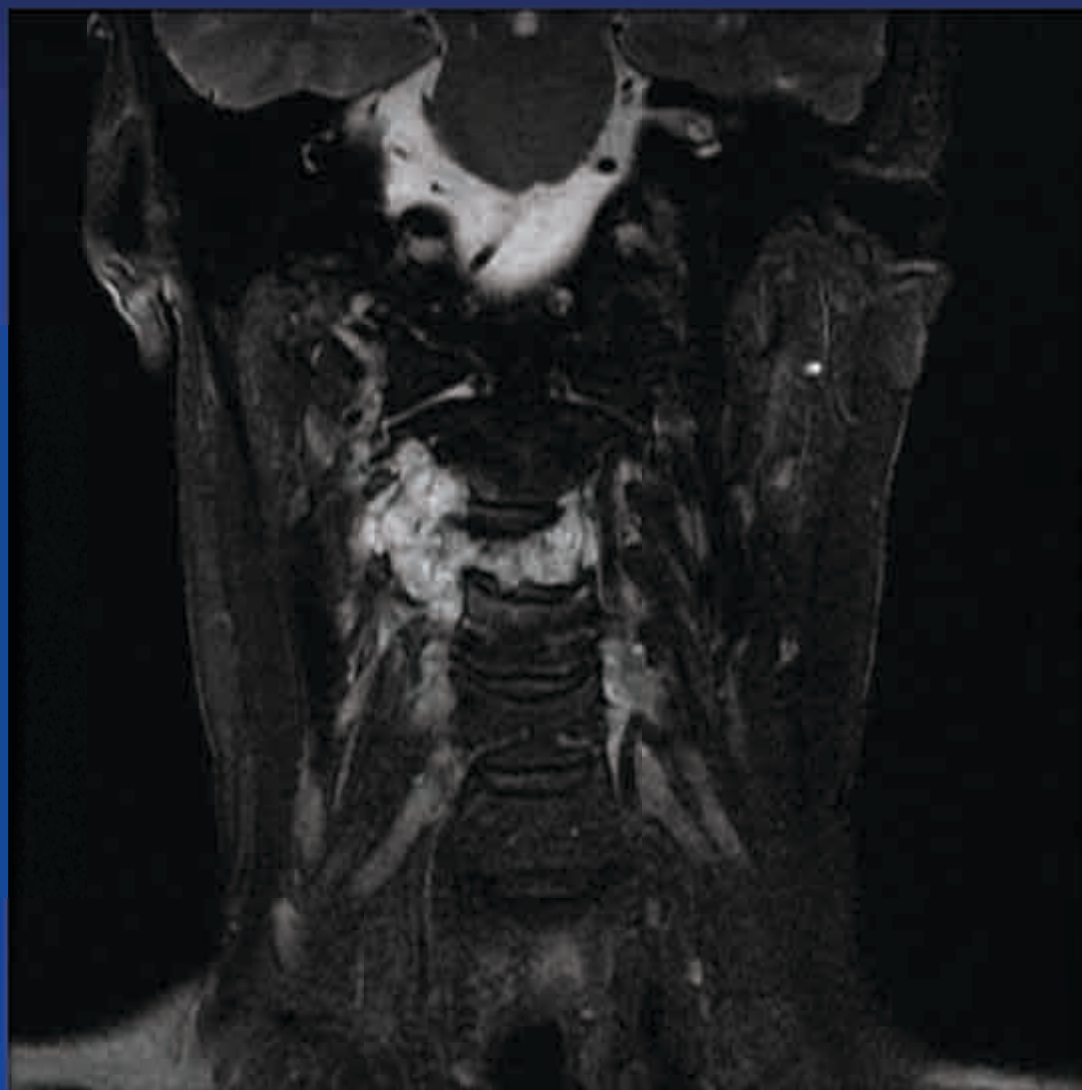
- Progressive pain, soft tissue mass/swelling
- Bimodal age (2<sup>nd</sup>-3<sup>rd</sup>, >6<sup>th</sup> decade)
- Very aggressive tumor
- Surgical resection with wide margins
- Adjuvant and neoadjuvant chemo
- 5 year survival: 41%
- 5 year survival (if resected with no mets): 60-70%

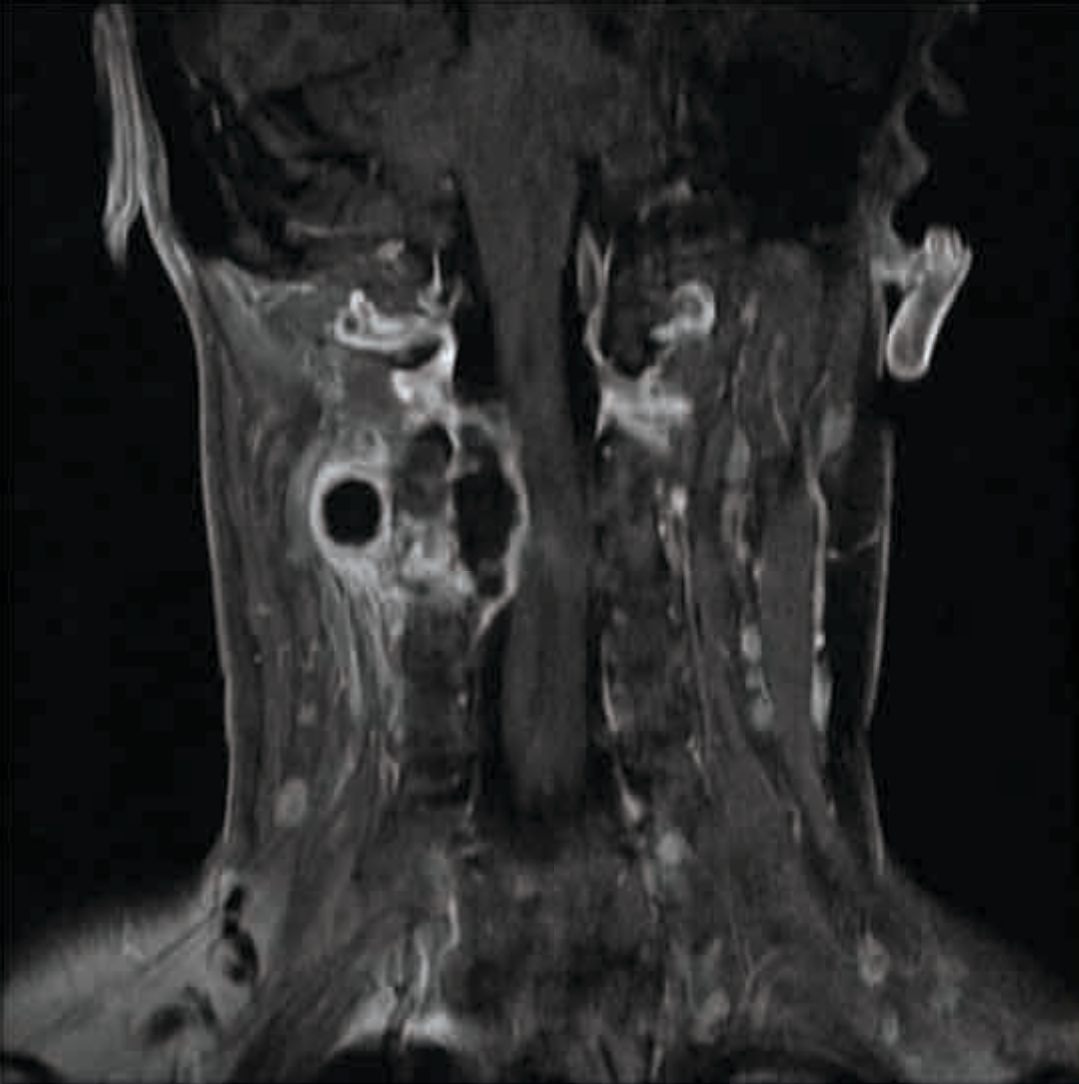


# Case 3

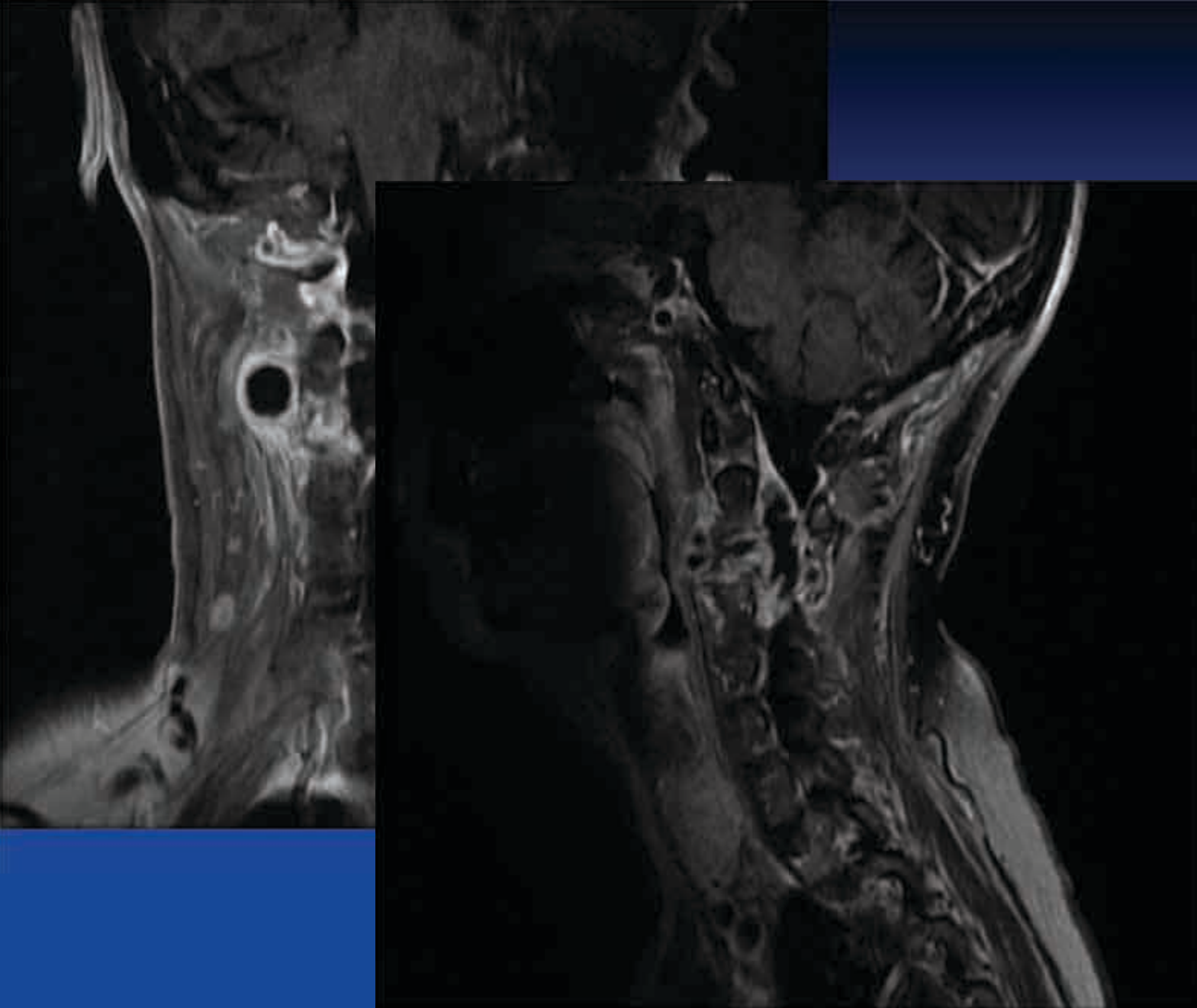
- 60 year-old female with neck pain

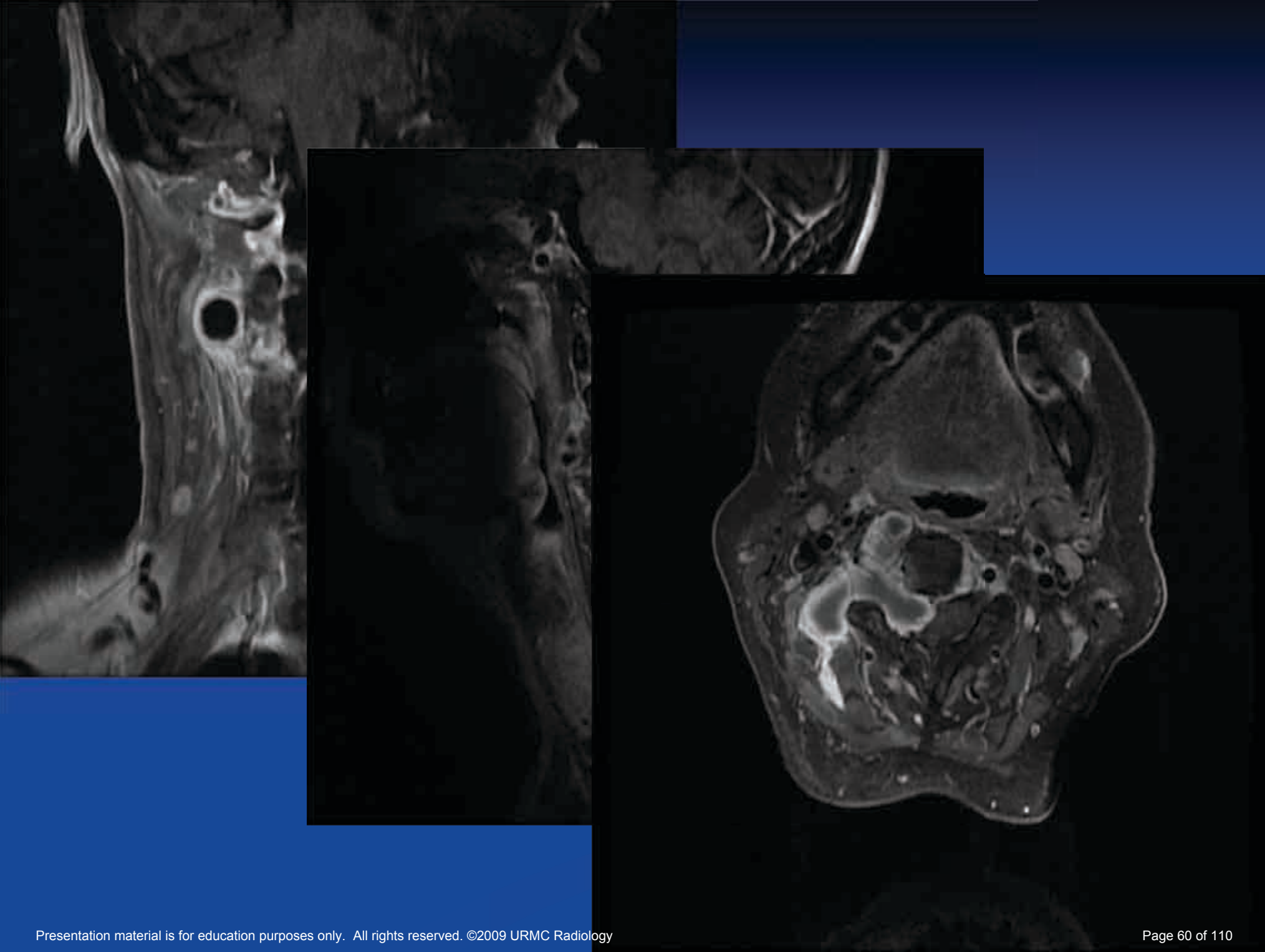












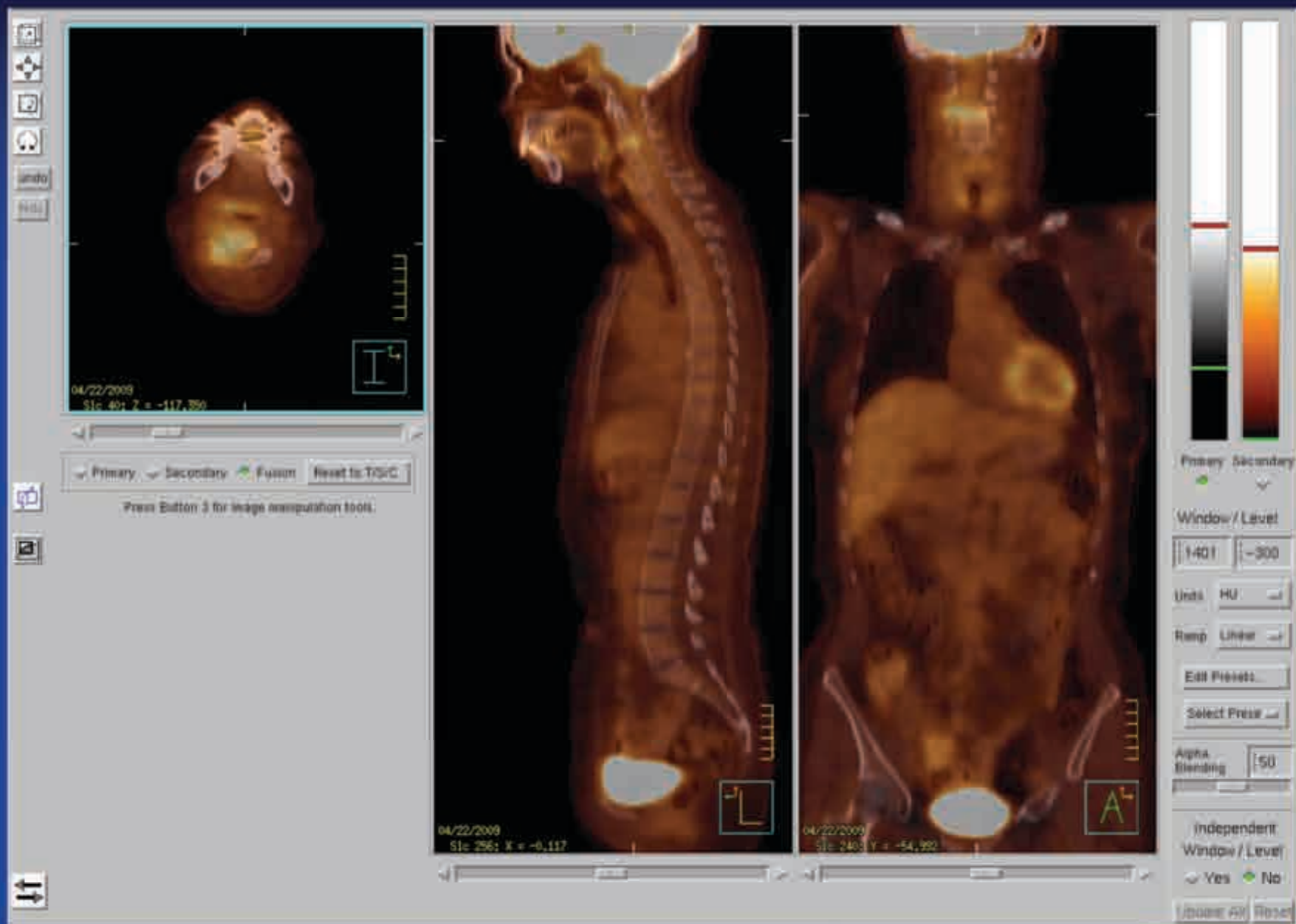
# Findings

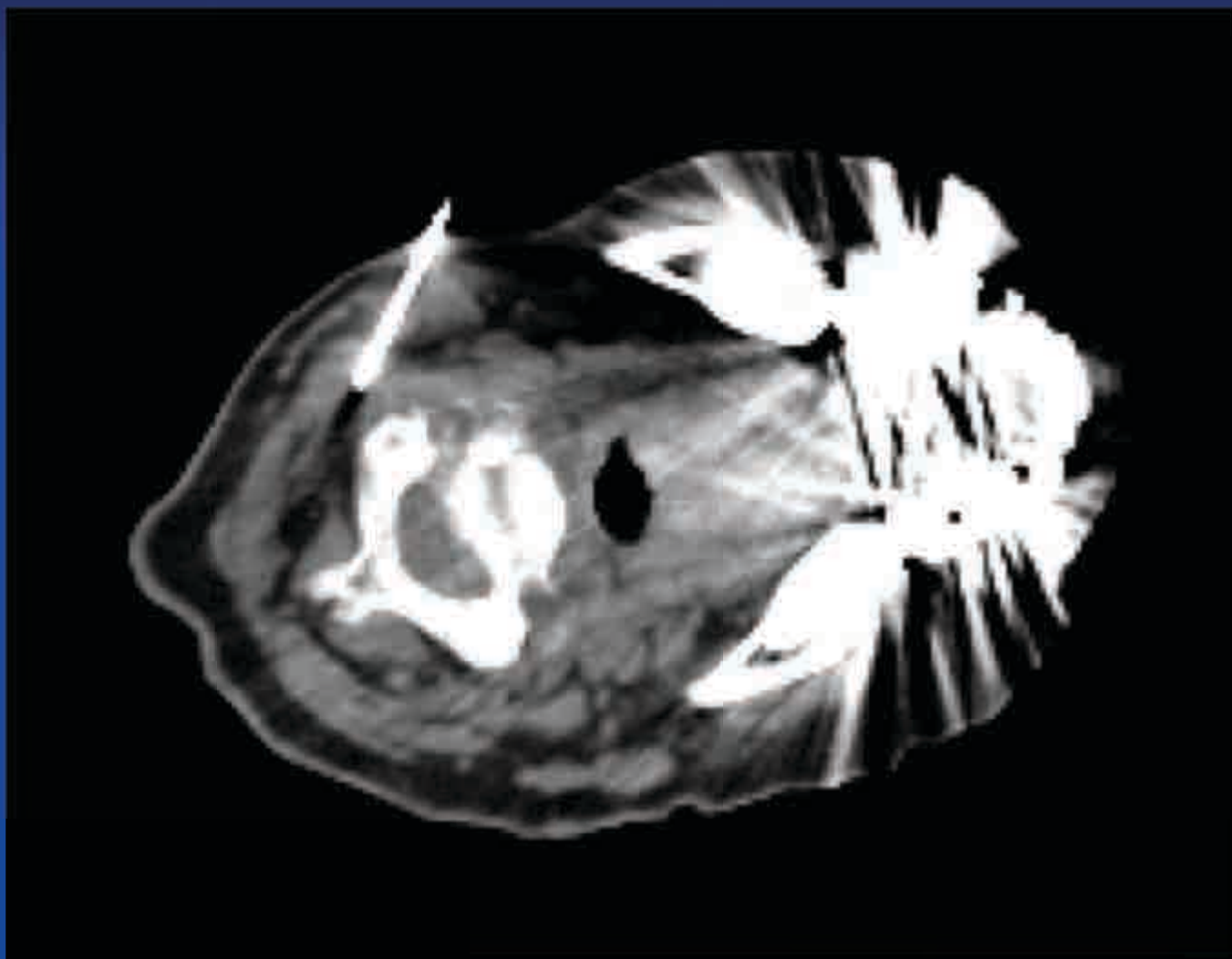
- Diffuse abnormal high T2 signal and enhancement of C3 with pathologic compression fracture.
- Large peripherally enhancing soft tissue component which extends into and expands the right neural foramen and thecal sac.

# Differential Diagnosis

- Chondrosarcoma
- Multiple myeloma
- Lymphoma
- Metastatic disease

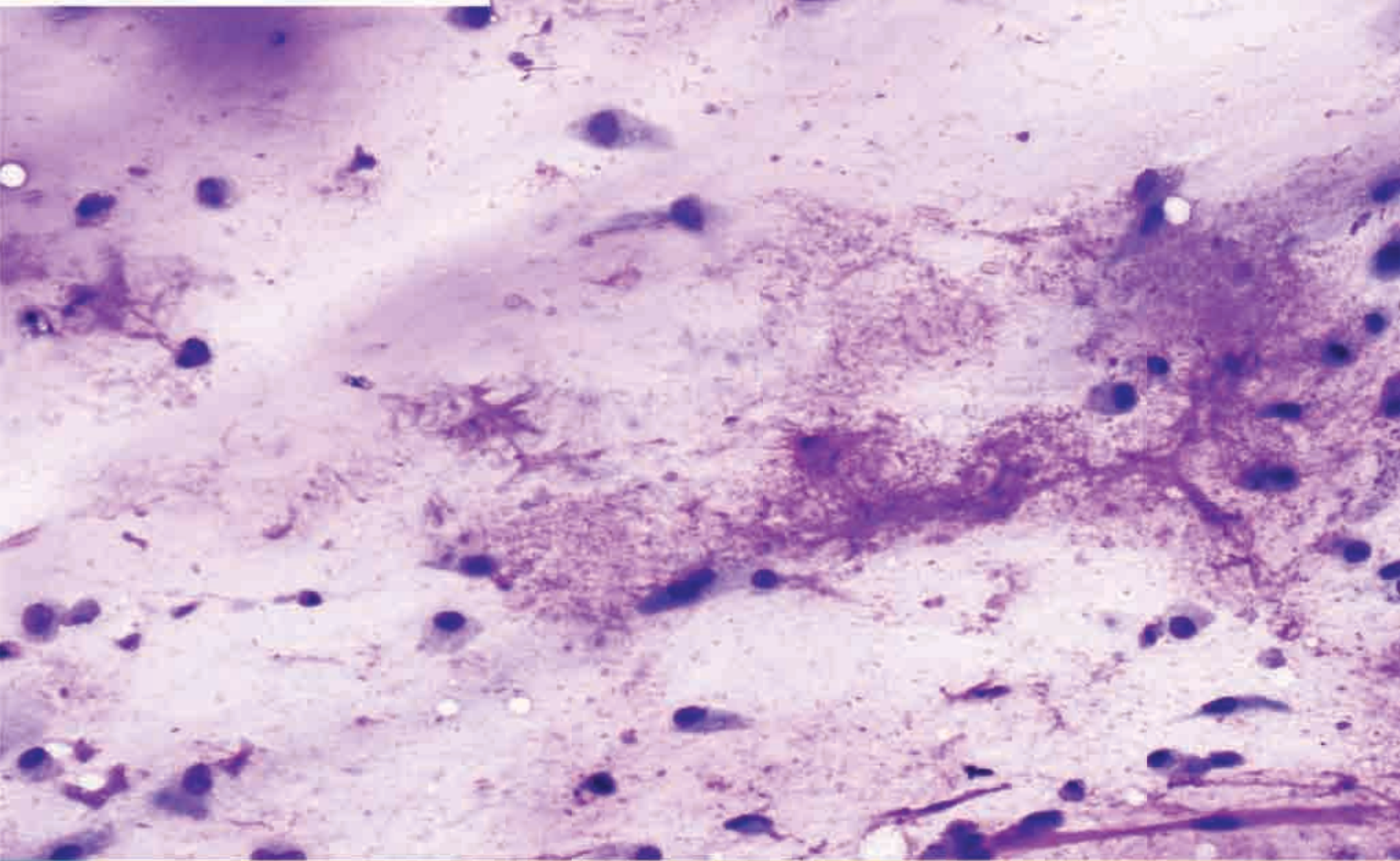






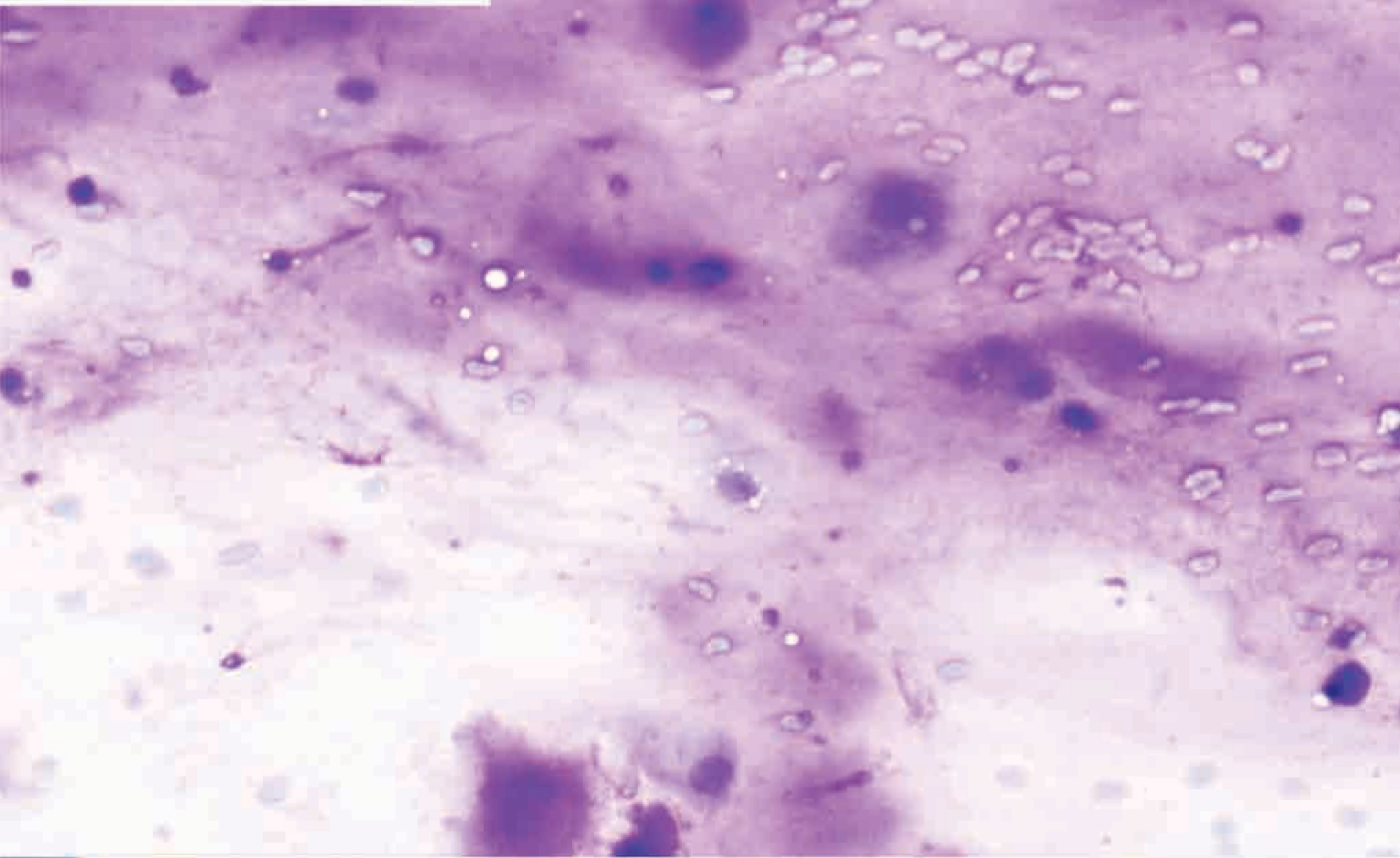
# Case 3

**Spine, C3, CT-guided FNA:  
Diff-Quik stain, 20x**





**Spine, C3, CT-guided FNA:  
Diff-Quik stain, 40x**



**Spine, C3, CT-guided FNA:  
Diff-Quik stain, 40x**



# Spine, C3, CT-guided fine needle aspiration:

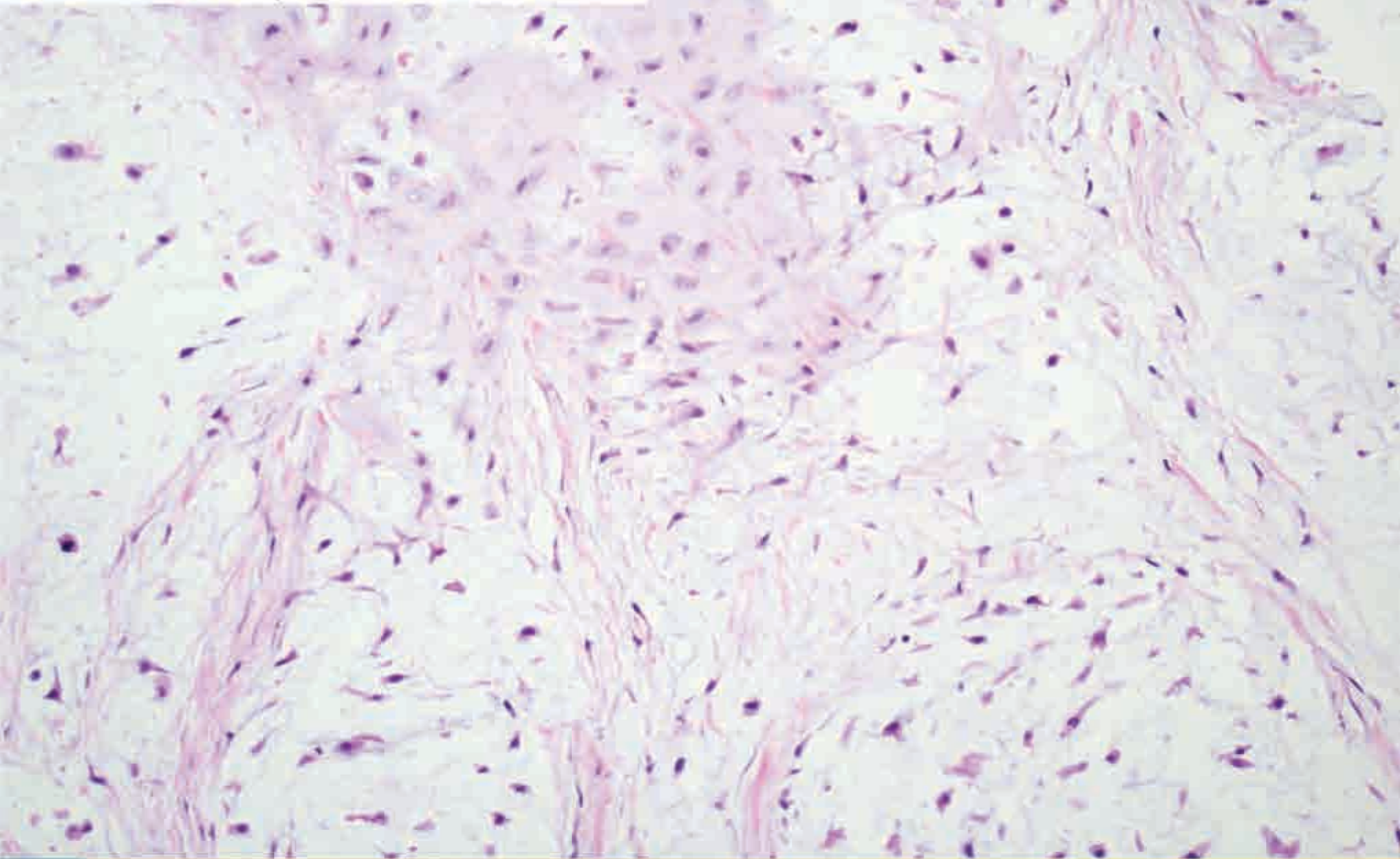
**Malignant tumor cells present consistent with chondrosarcoma.**

**Soft tissue, C3, right paravertebral body,  
core needle biopsy:**

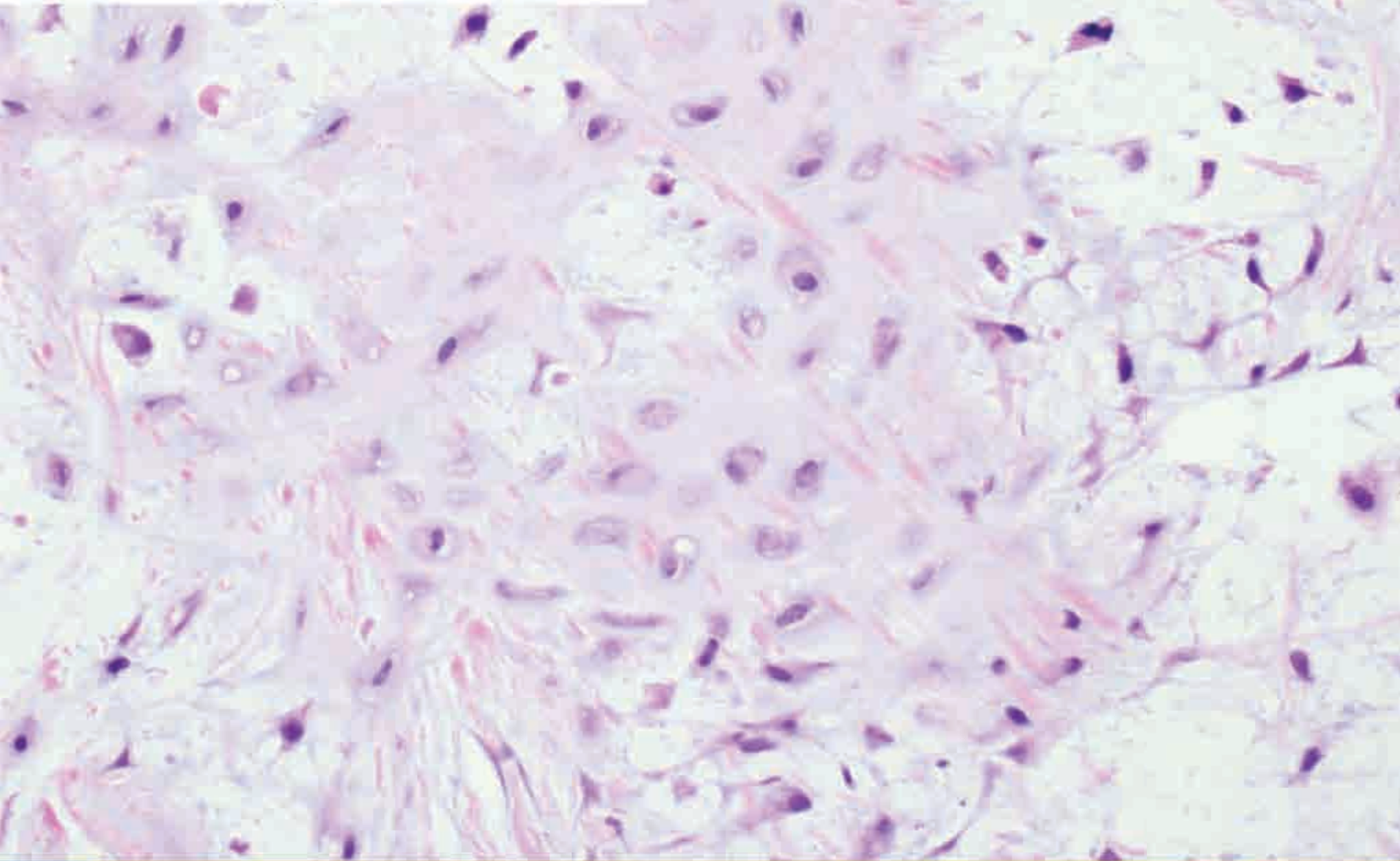
**Chondrosarcoma, grade 2.**



Soft tissue, C2, paravertebral body,  
core needle biopsy: H & E stain, 10x



Soft tissue, C2, paravertebral body,  
core needle biopsy: H & E stain, 20x





# Chondrosarcoma

- Malignant tumor in which neoplastic cells form cartilage
- Best diagnostic clue: Lytic mass with or without chondroid matrix, cortical disruption, and extension into soft tissues
- Chondroid matrix mineralization of “rings and arcs”
- Multiple types:
  - Medullary, exostotic, dedifferentiated, mesenchymal, clear cell, and extraskeletal



# Chondrosarcoma - Clinical Issues

- Dull aching pain at rest, soft tissue mass/swelling
- Age is 20-90 years, peak at 40-60 years
- Surgical resection with wide margins
- Limited role for chemo or radiation therapy
- 5 year survival: 48-60%
- 5 year survival: 90% (Grade 1), 81% (2), 29% (3)

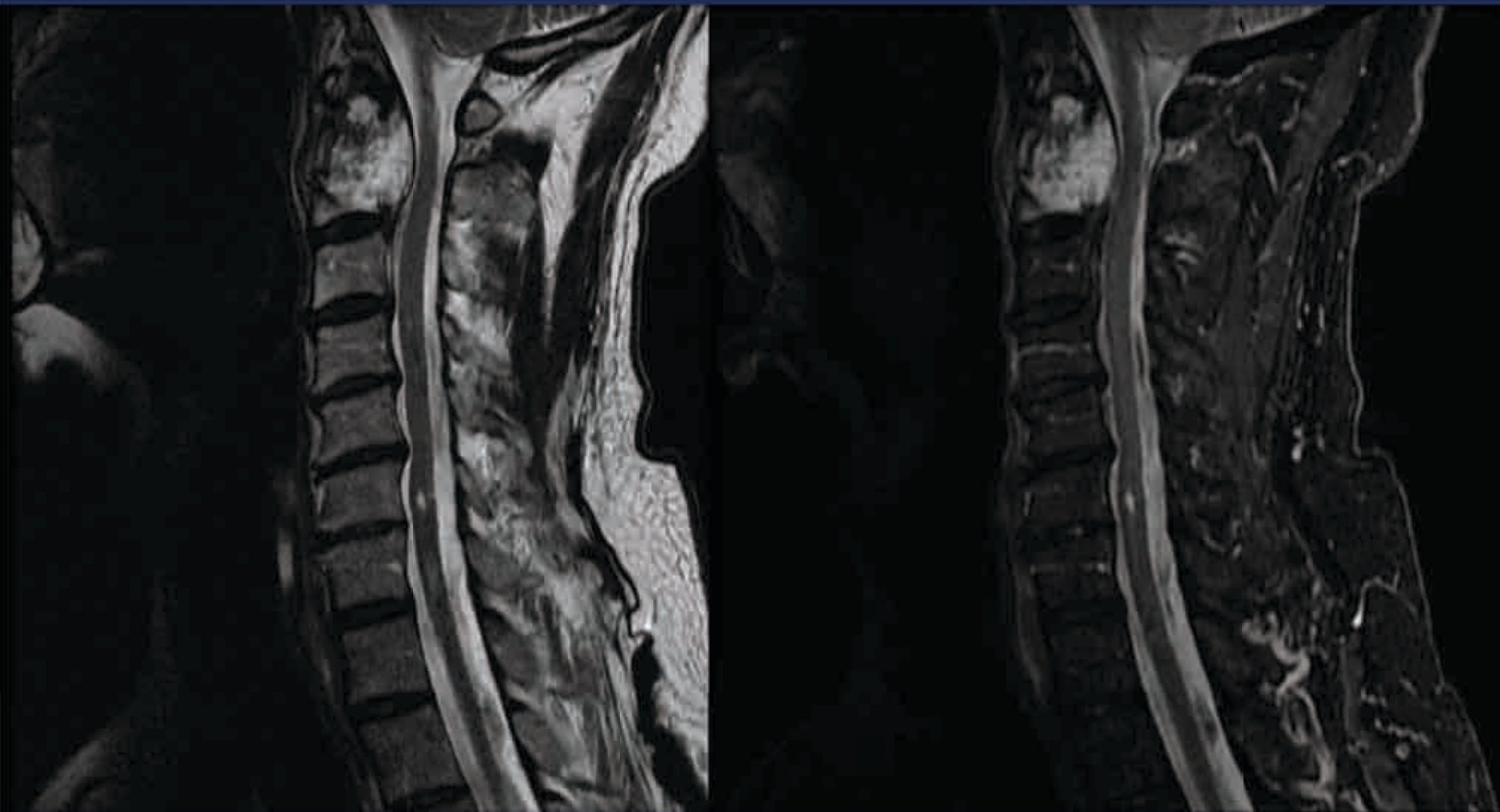
# Chondrosarcoma - Grading

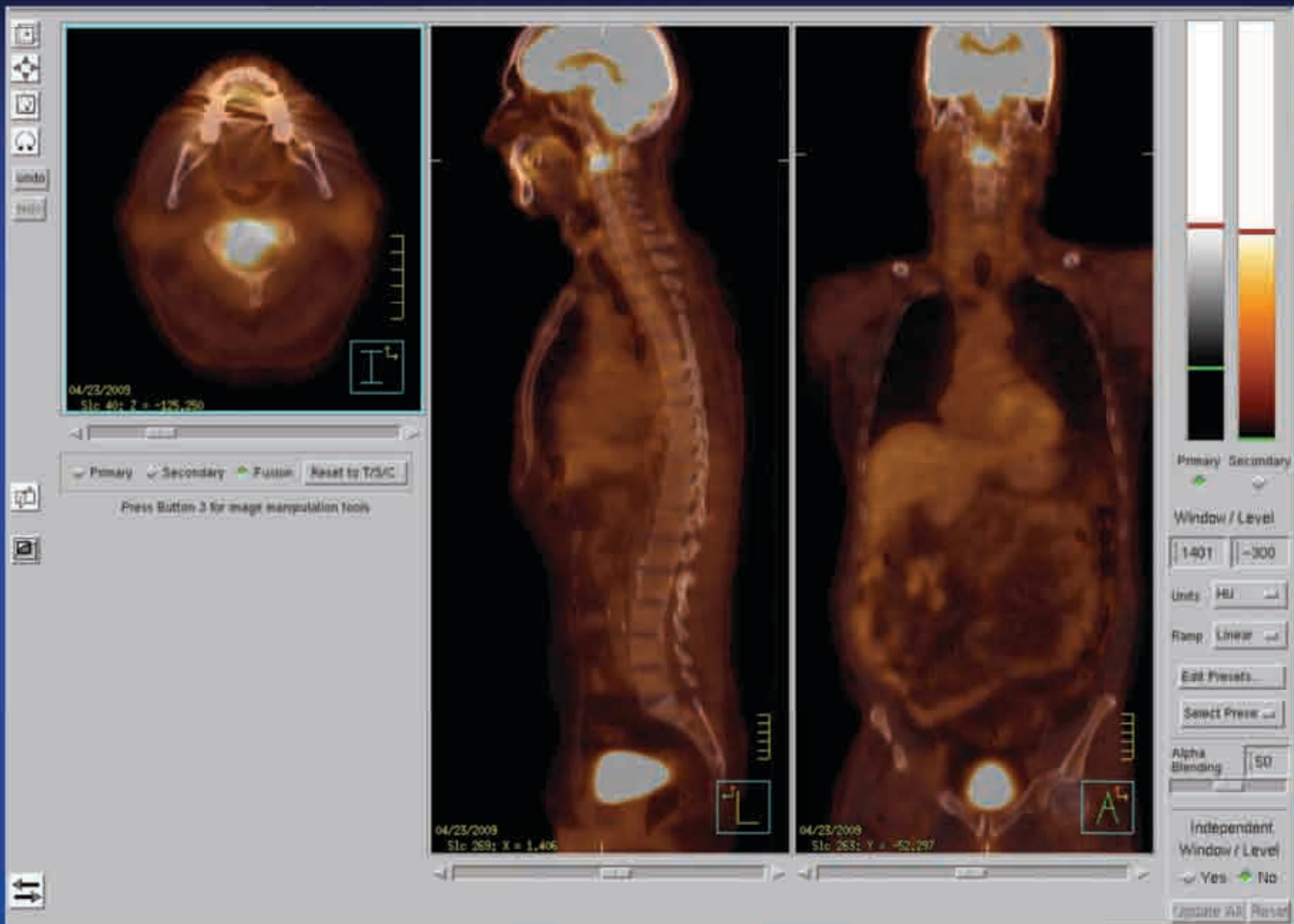
- Grade 1: Slowly growing, locally aggressive, indolent course
  - Recurrent growth potential, no metastatic spread
- Grade 2: Locally aggressive tumor with great potential for local recurrence
  - Mets in 10-15%
- Grade 3: Highly aggressive, rapidly growing
  - Mets in over 50%

# Case 4

- 66 year old male with neck pain







# Findings

- Hypermetabolic focus corresponding to a destructive lytic lesion at C2. This lesion has markedly increased T2 signal.

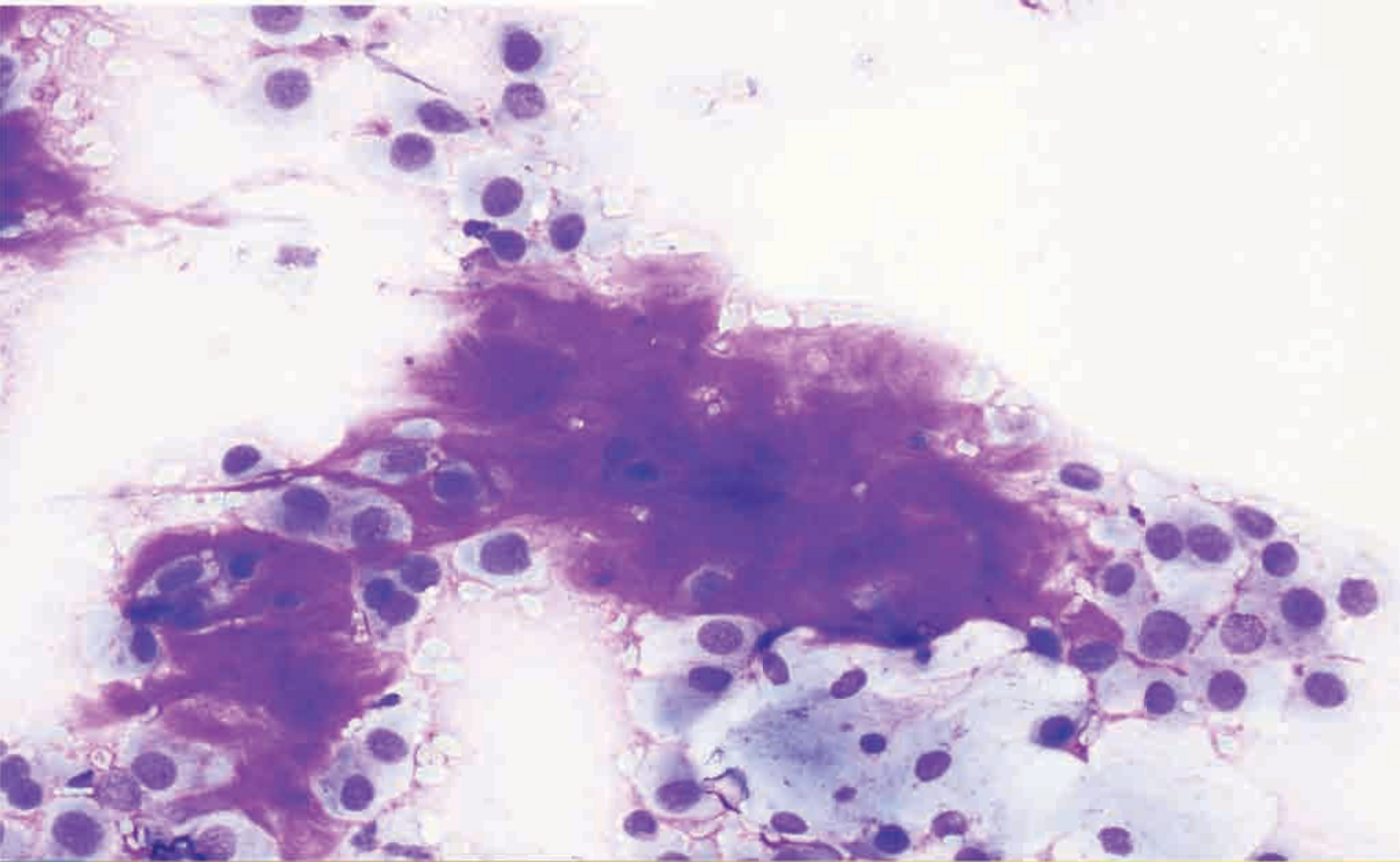


# Differential Diagnosis

- Chondrosarcoma
- Metastases
- Multiple myeloma
- Lymphoma
- Chordoma

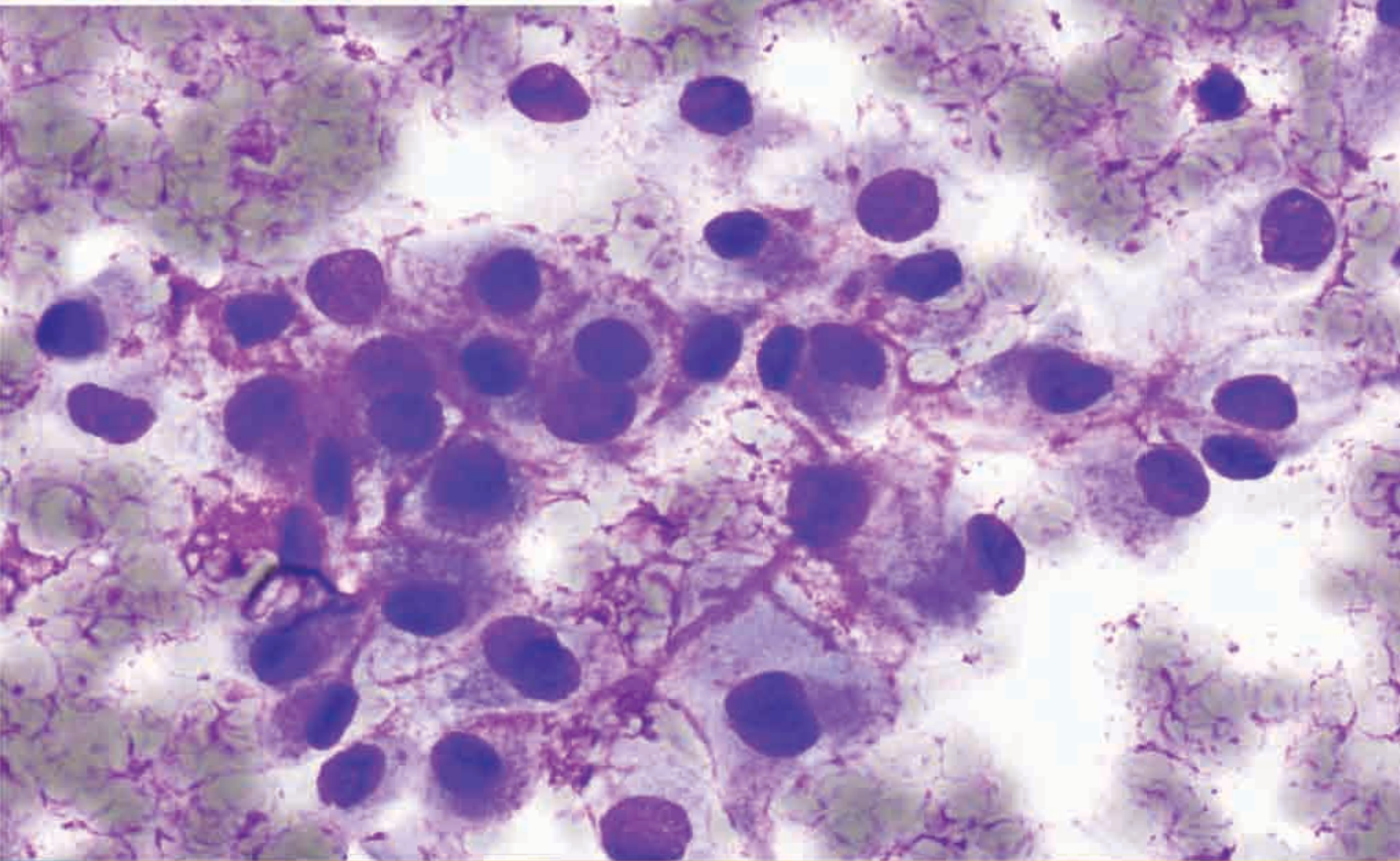
# Case 4

**Bone, vertebra, C2, CT-guided FNA:  
Diff-Quik stain, 20x**



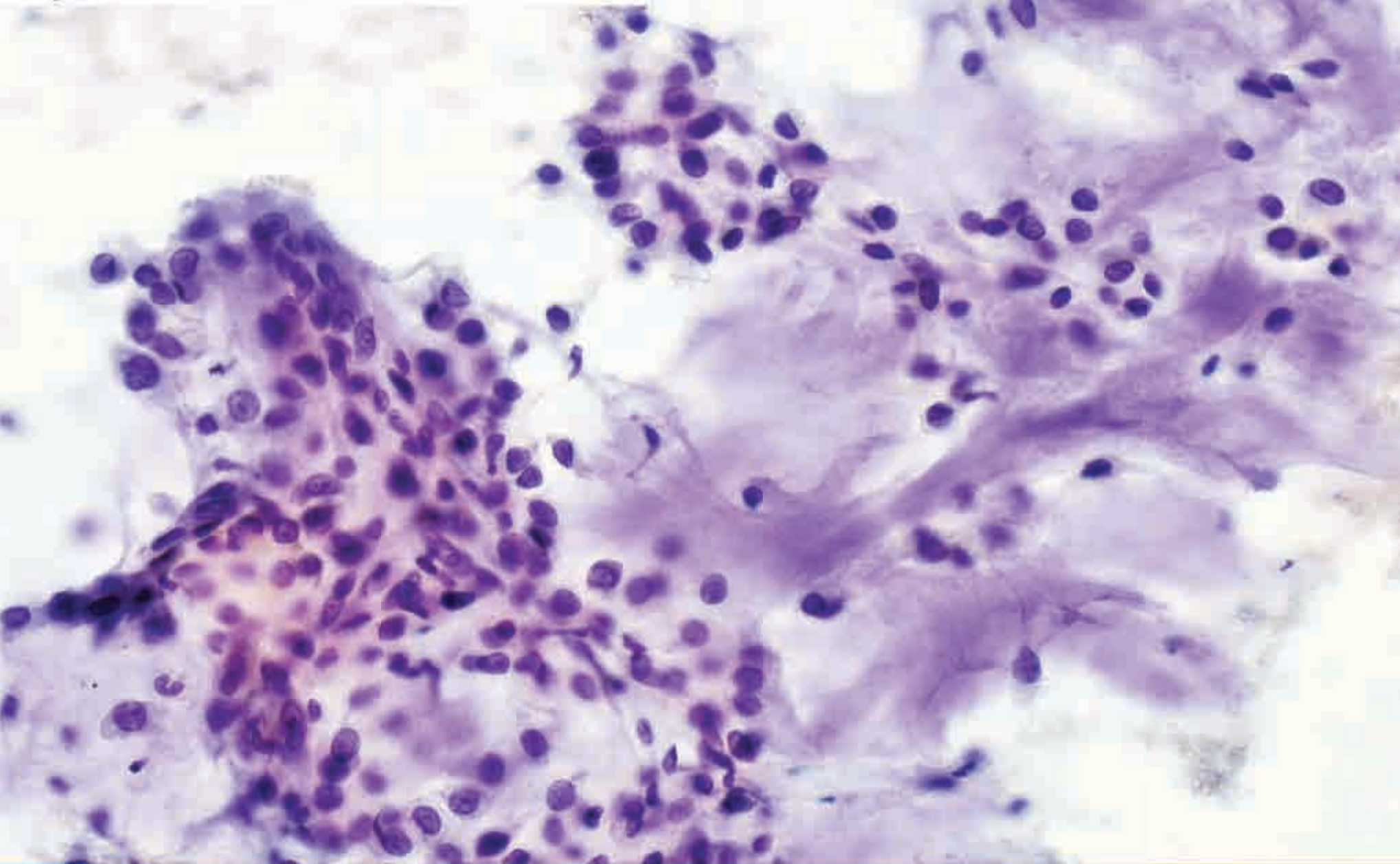


**Bone, vertebra, C2, CT-guided FNA:  
Diff-Quik stain, 40x**

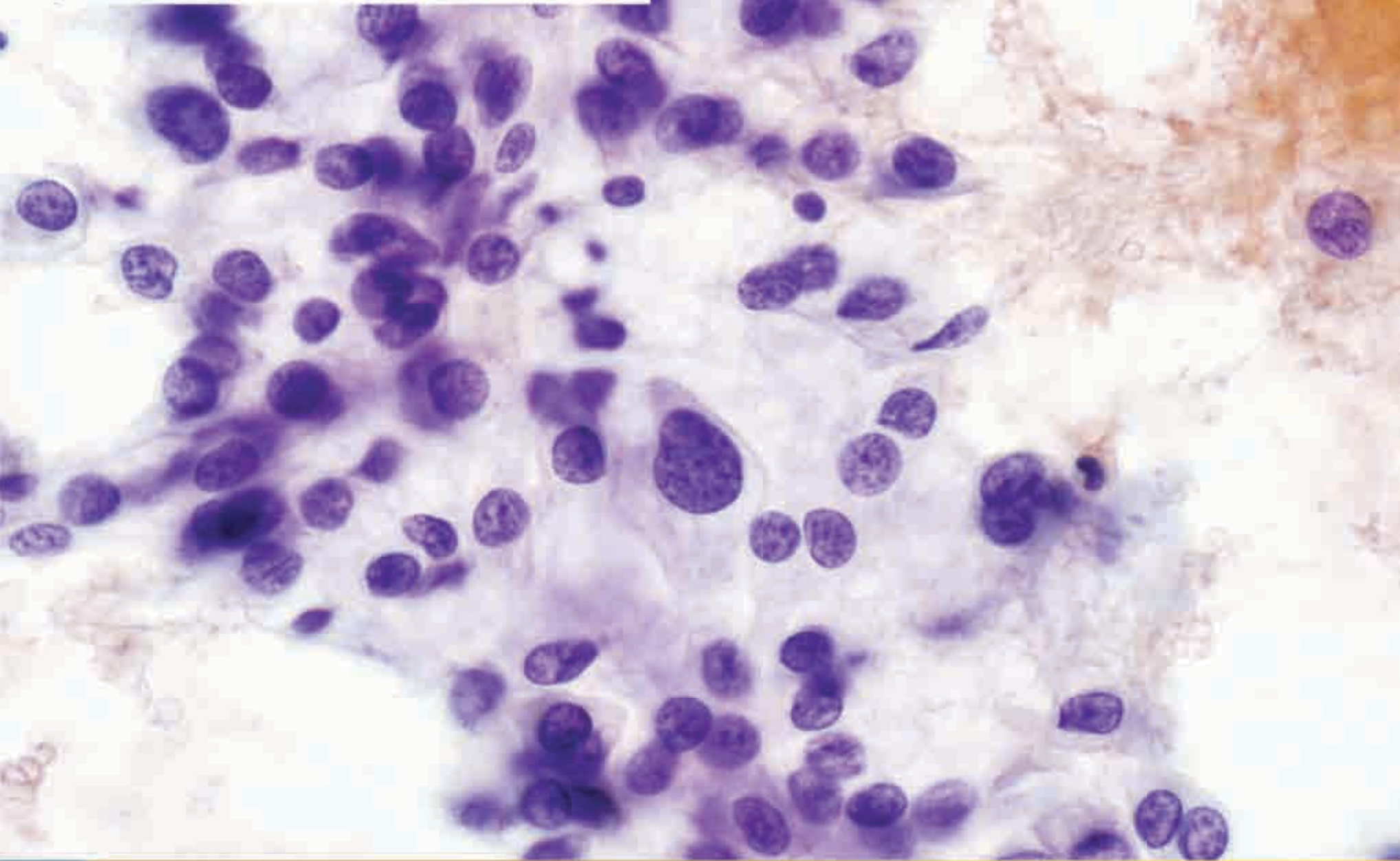




**Bone, vertebra, C2, CT-guided FNA:  
Papanicolaou stain, 20x**

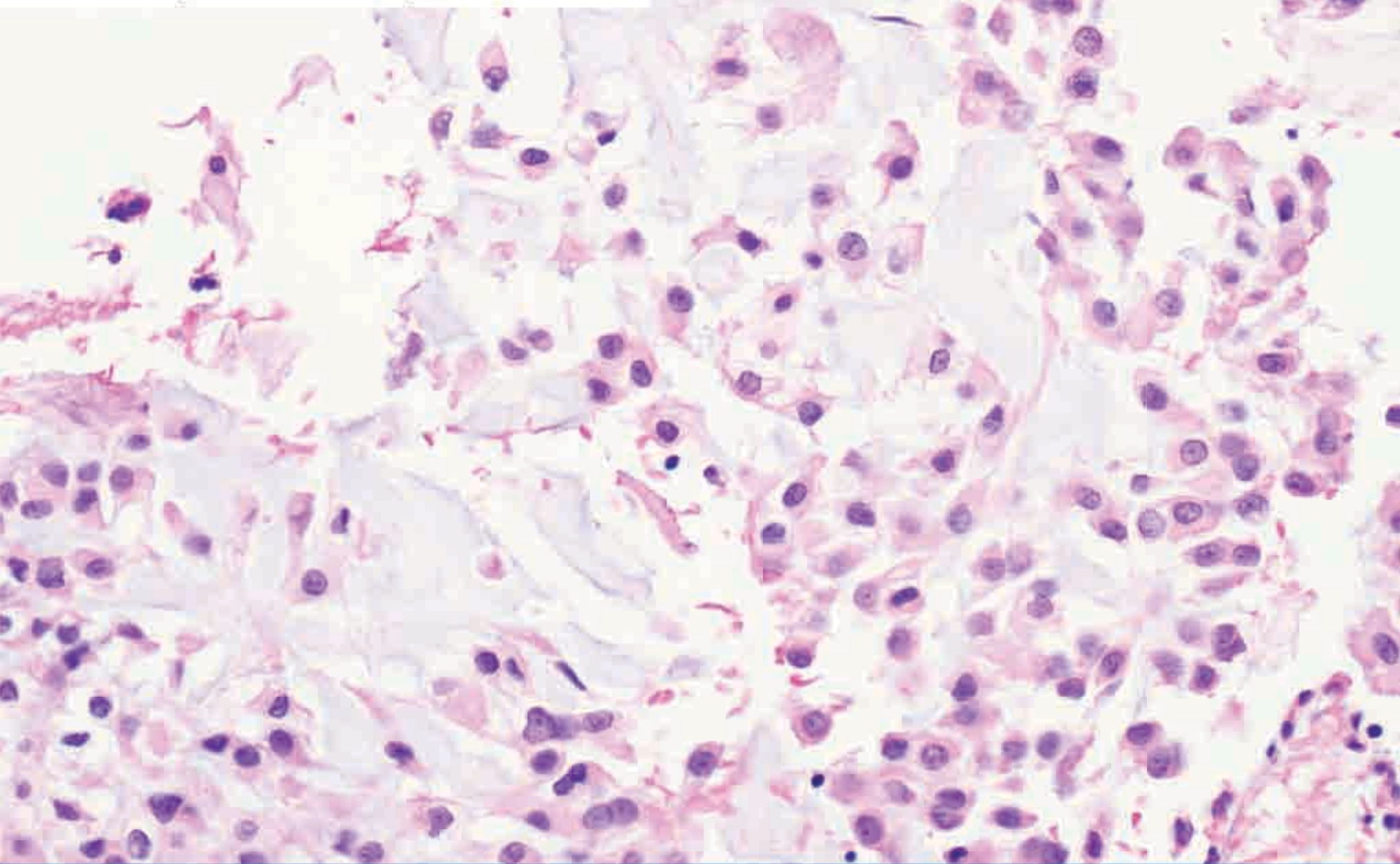


**Bone, vertebra, C2, CT-guided FNA:  
Papanicolaou stain, 40x**

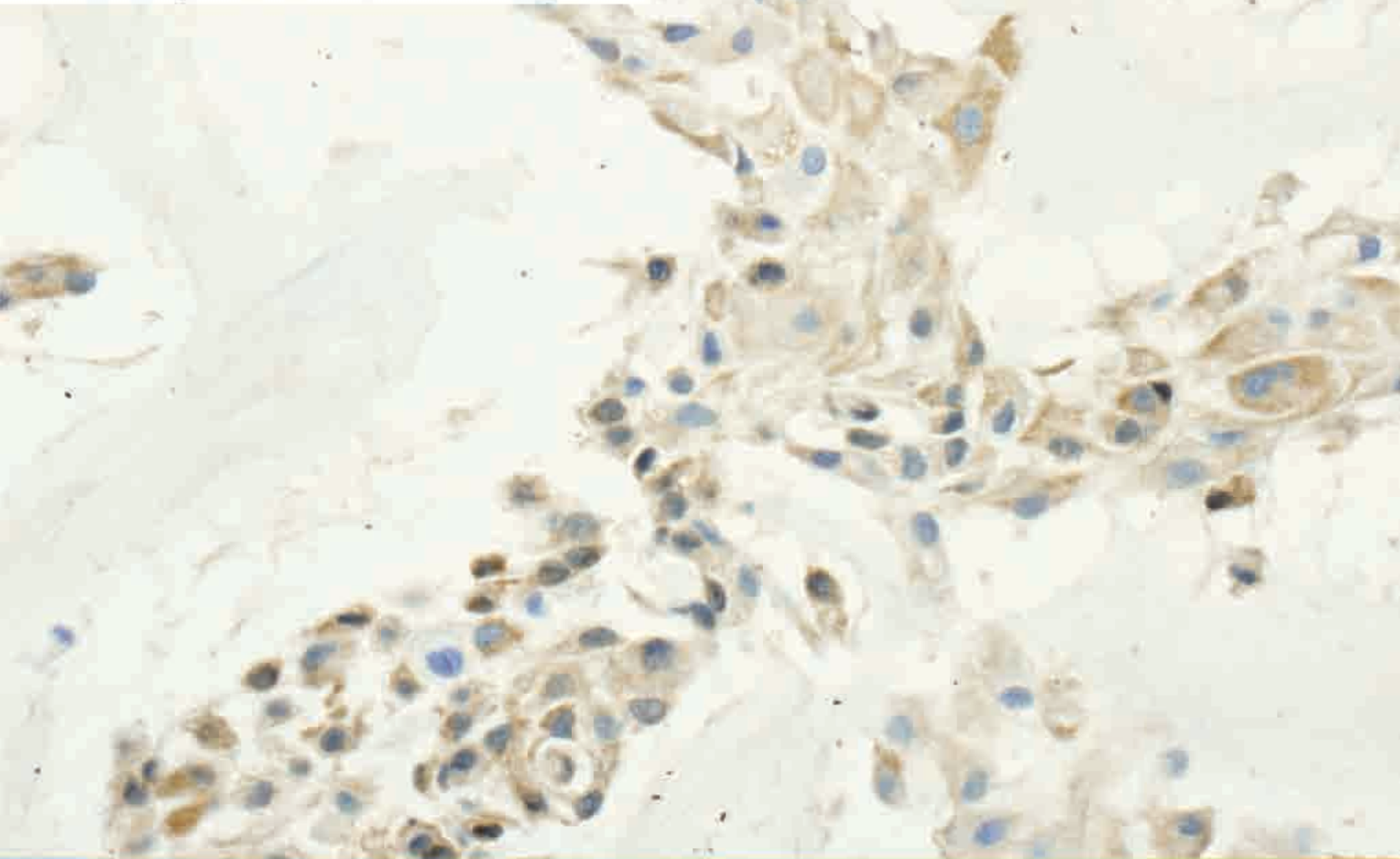




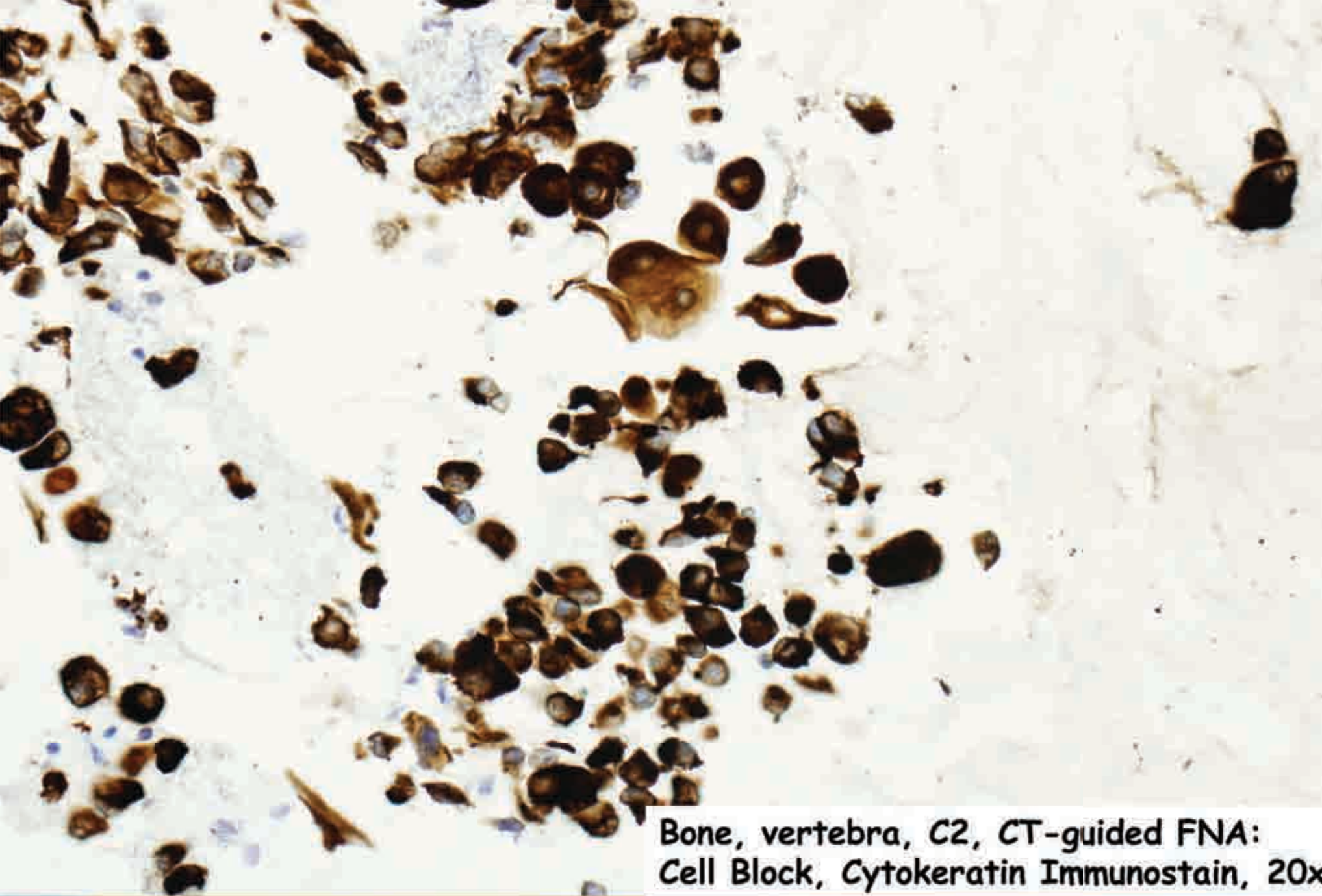
**Bone, vertebra, C2, CT-guided FNA:  
Cell Block, H & E stain, 20x**



**Bone, vertebra, C2, CT-guided FNA:  
Cell Block, S-100 Immunostain, 20x**







**Bone, vertebra, C2, CT-guided FNA:  
Cell Block, Cytokeratin Immunostain, 20x**



# Bone, vertebra, C2, CT-guided fine needle aspiration:

Cellular evidence of chordoma.

Cell block and cytologic preparations examined.

Immunohistochemical stains were performed to aid in interpretation. Stains for pancytokeratins and S-100 are positive. Stains for chromogranin, synaptophysin and CD56 are negative.

# Chordoma

- Malignant tumor arising from notochord remnants
- Midline:
  - sacrococcygeal > spheno-occipital >> vertebral body
- Hyperintense to discs on T2WI, with multiple septa
- Histologic identification of physaliphorous cells confirm diagnosis

# Hyperintense to discs on T2WI, with multiple septa



**Case 3: Chondrosarcoma**



**Case 4: Chordoma**



# Chordoma – Clinical Issues

- Location dependent presentation
  - pain, numbness, weakness, incontinence
- Age: 5<sup>th</sup>-6<sup>th</sup> decade
- Slow growing
- Surgical resection with adjuvant XRT
- Recurrence is common
- 5 year survival: 67 to 84%
- 10 year survival: 40%

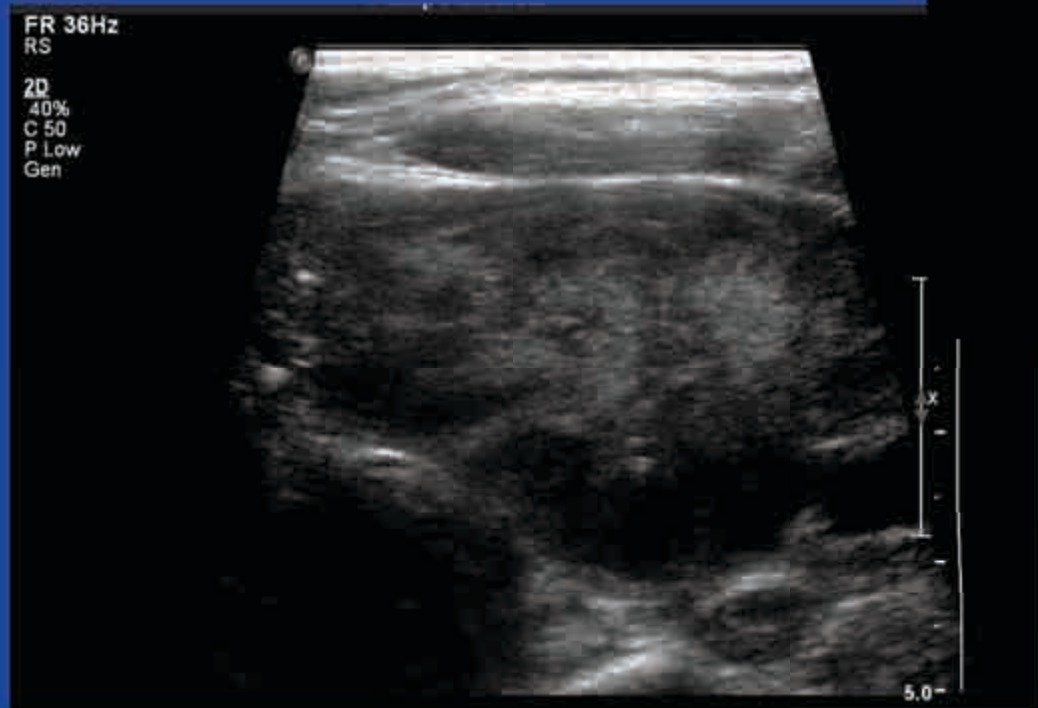
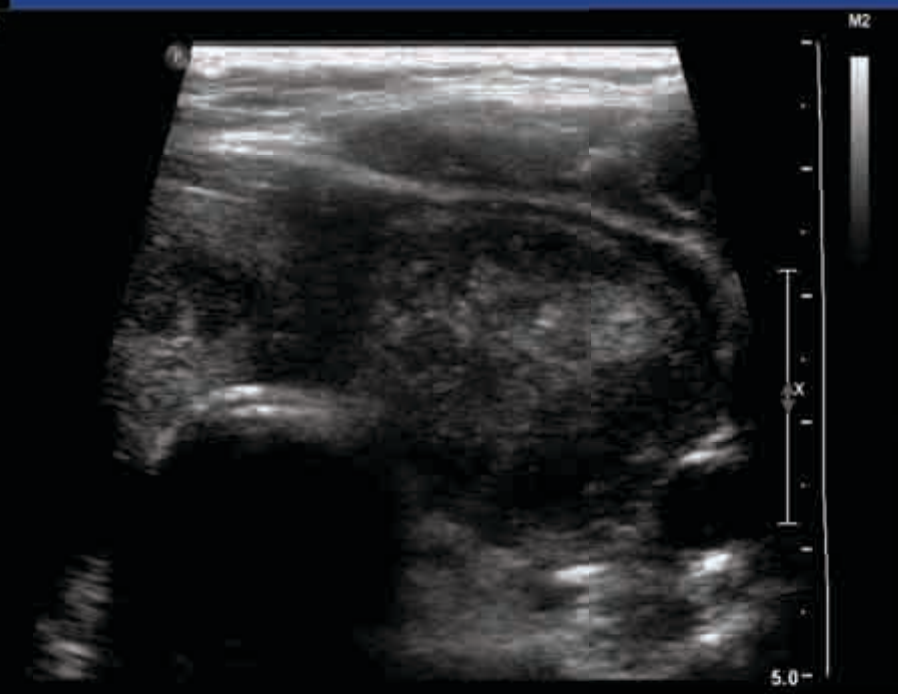
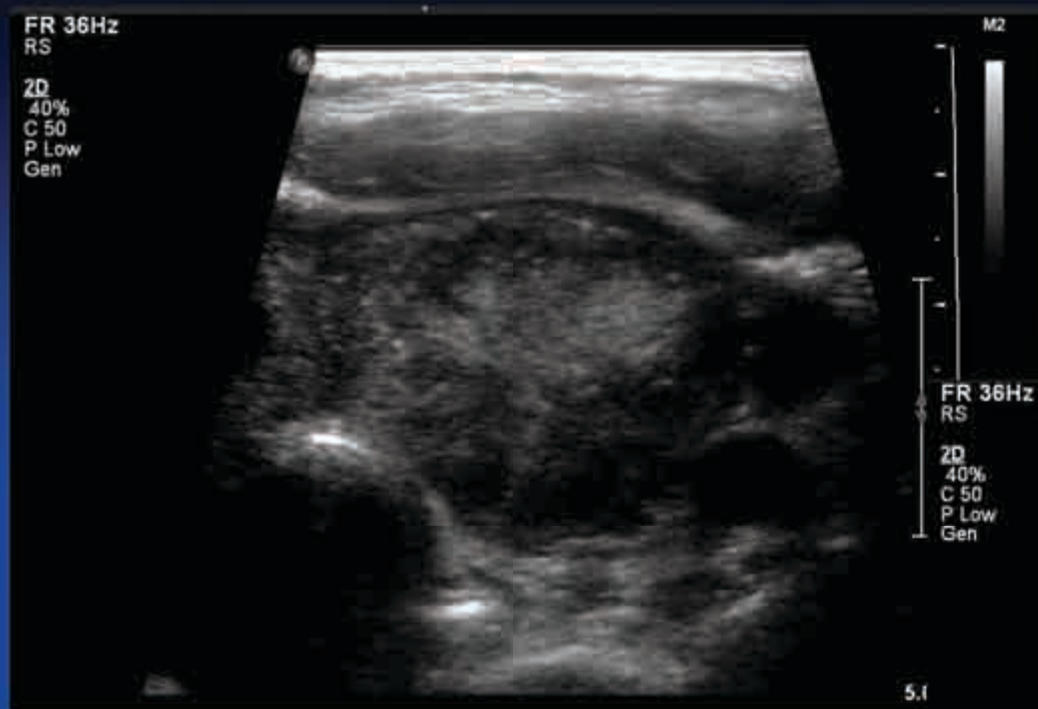
# Follow-up

- Complex case requiring highly specialized team:
  - Maxillofacial surgeon
  - ENT surgeon
  - Orthopedic oncologist
  - Neurosurgeon or an orthopedic complex spinal surgeon
- 3 centers in the nation perform this procedure:
  - Johns Hopkins, University of Pittsburgh, MD Anderson
- The referral for initial consultations at these hospitals has been denied by his insurance company

# Case 5

- 41 year old female with thyroid nodules. The left lobe is mostly replaced with a nodule.





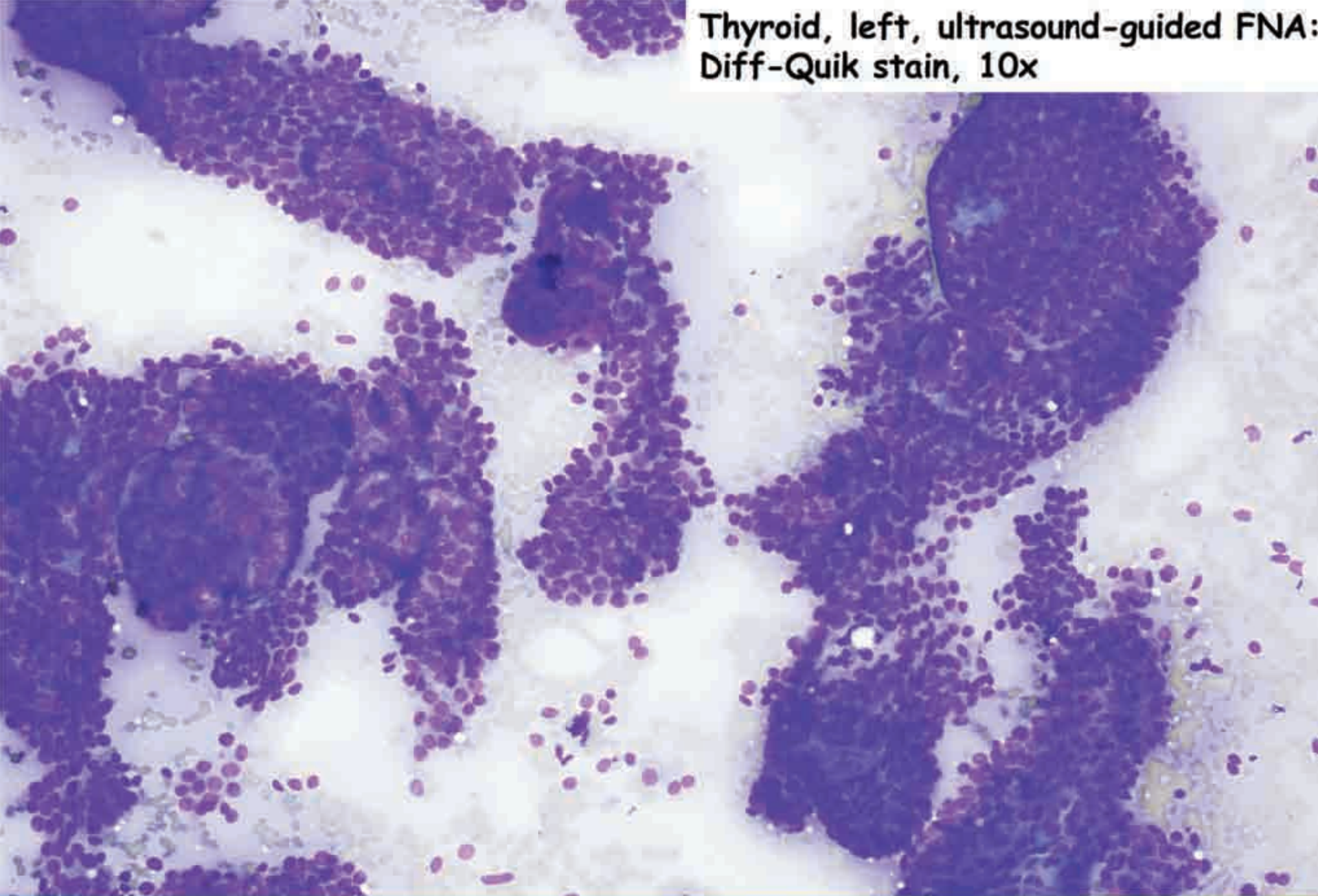
# Differential Diagnosis

- Thyroid Cancer
  - Differentiated thyroid carcinoma
    - Papillary (80%) and Follicular (10%)
  - Medullary carcinoma (5-10%)
  - Anaplastic carcinoma (1-2%)
- Hemorrhagic colloid cyst
- Multinodular goiter
- Follicular adenoma
- Non-Hodgkin lymphoma

# Case 5

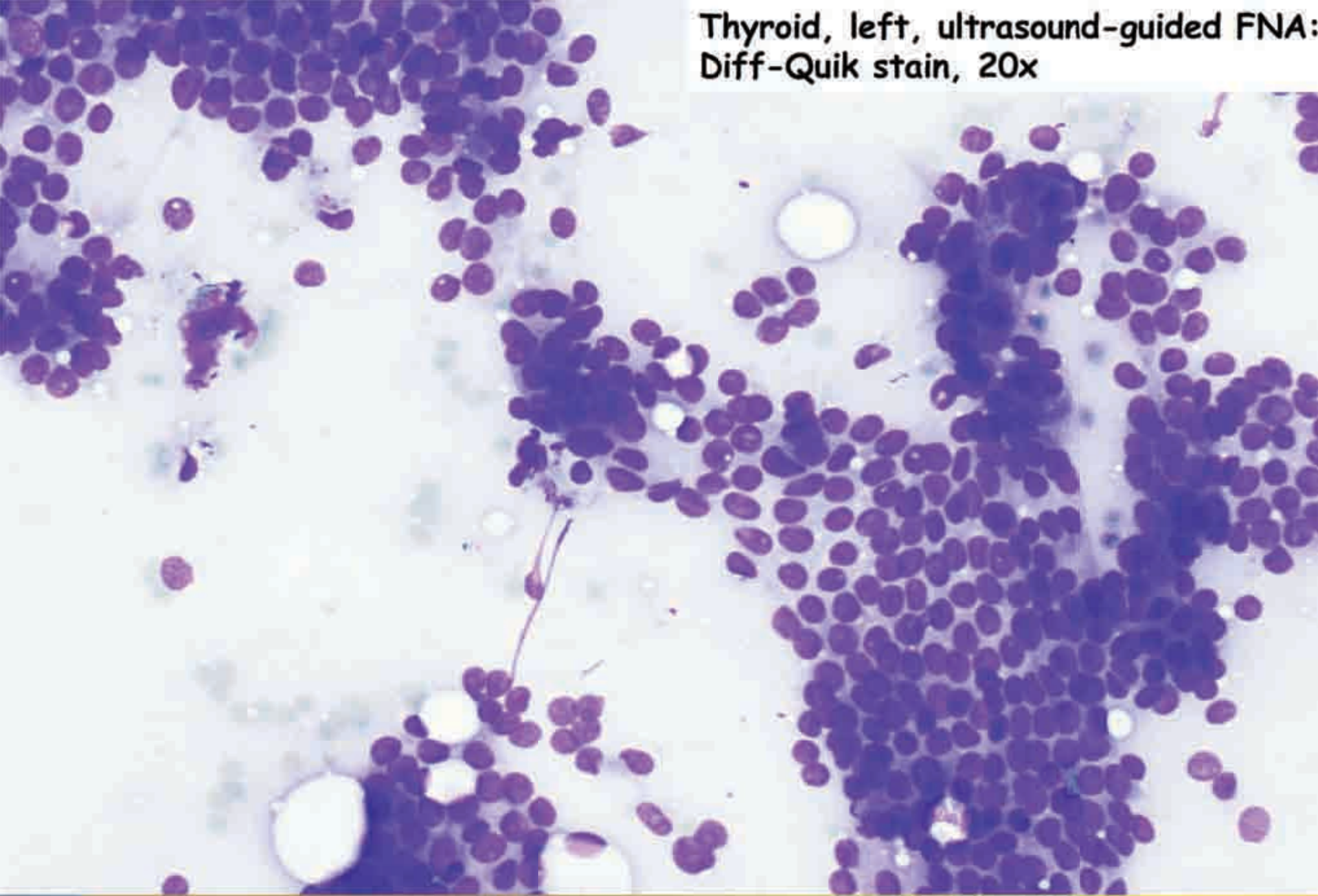


**Thyroid, left, ultrasound-guided FNA:  
Diff-Quik stain, 10x**

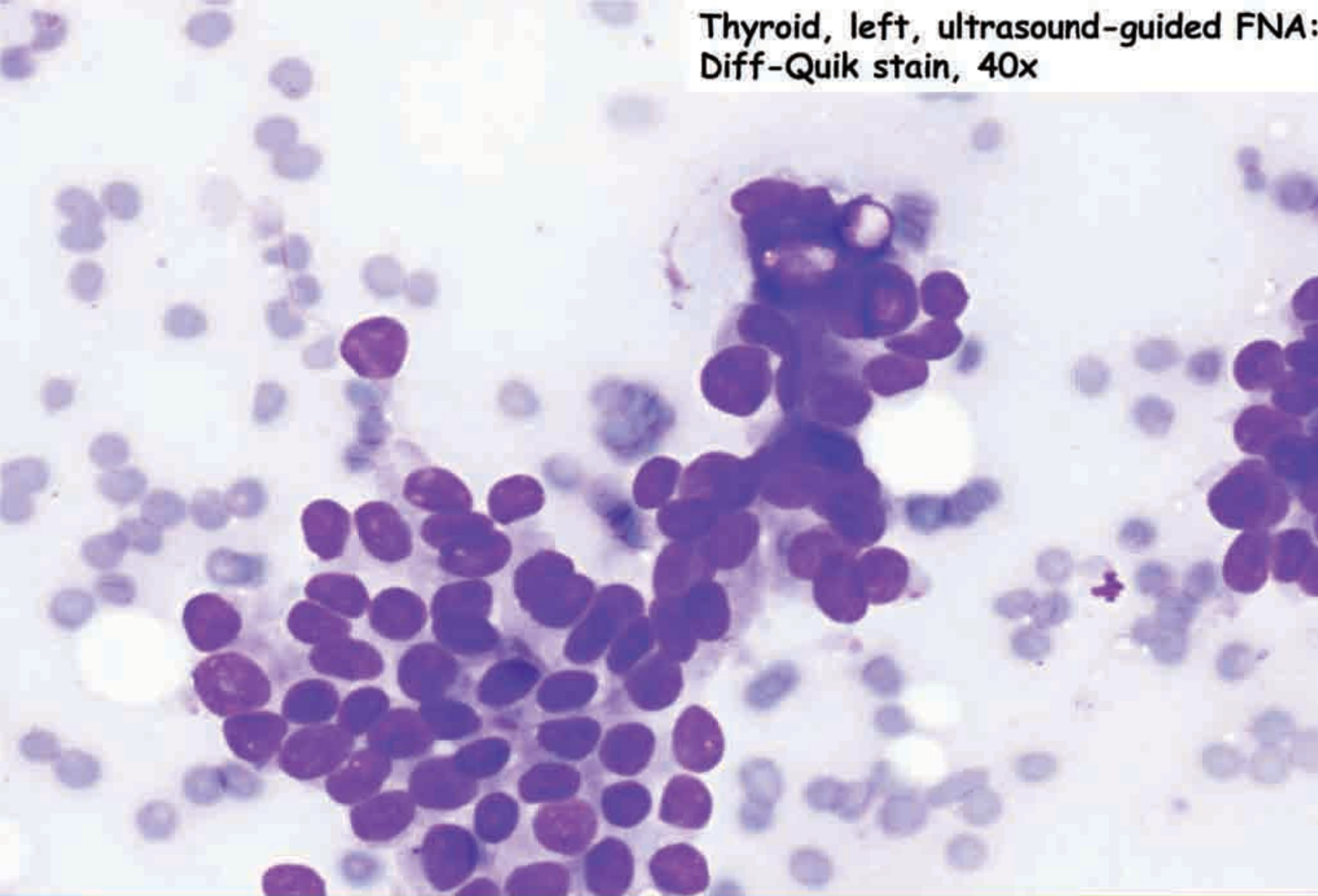




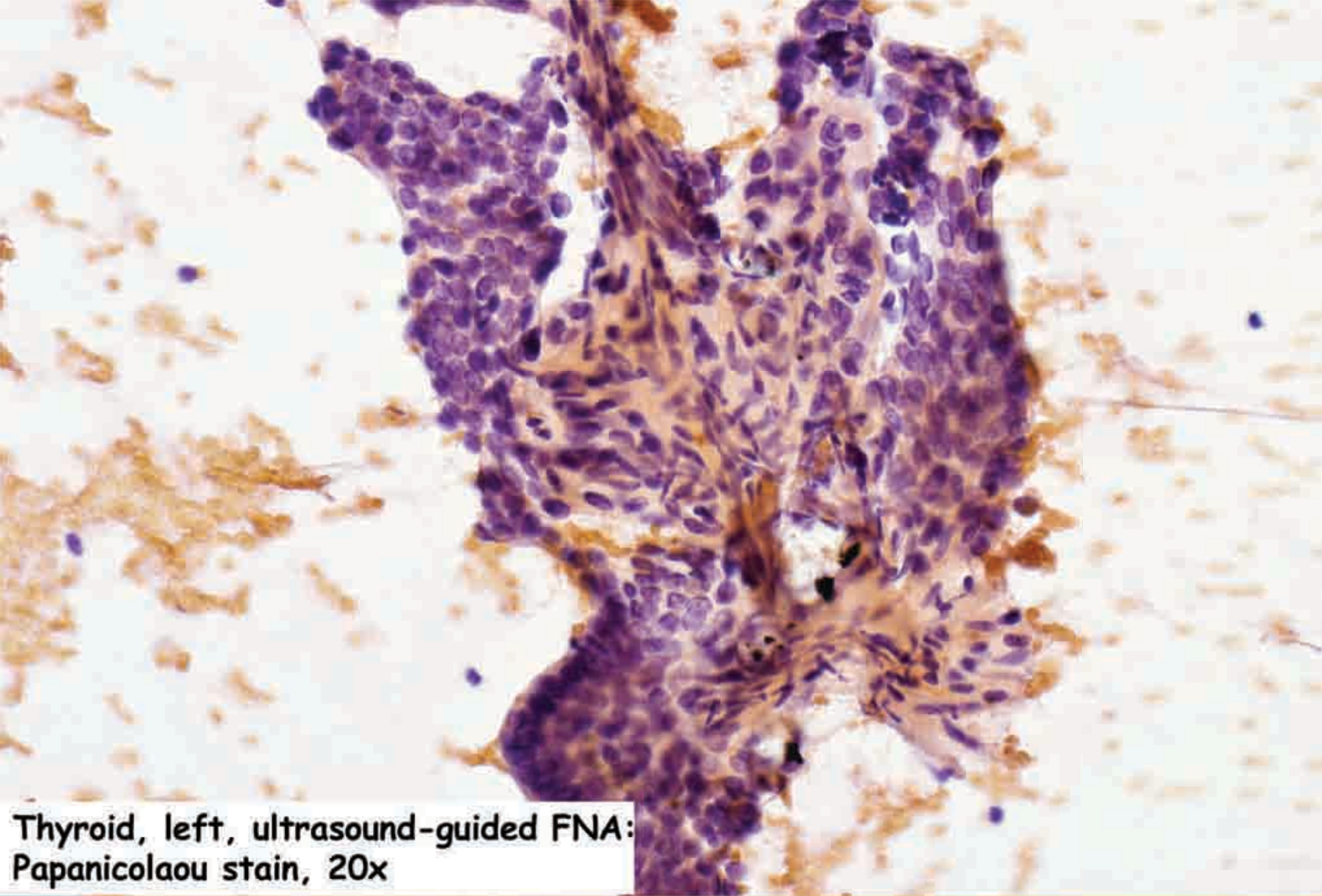
**Thyroid, left, ultrasound-guided FNA:  
Diff-Quik stain, 20x**



**Thyroid, left, ultrasound-guided FNA:  
Diff-Quik stain, 40x**

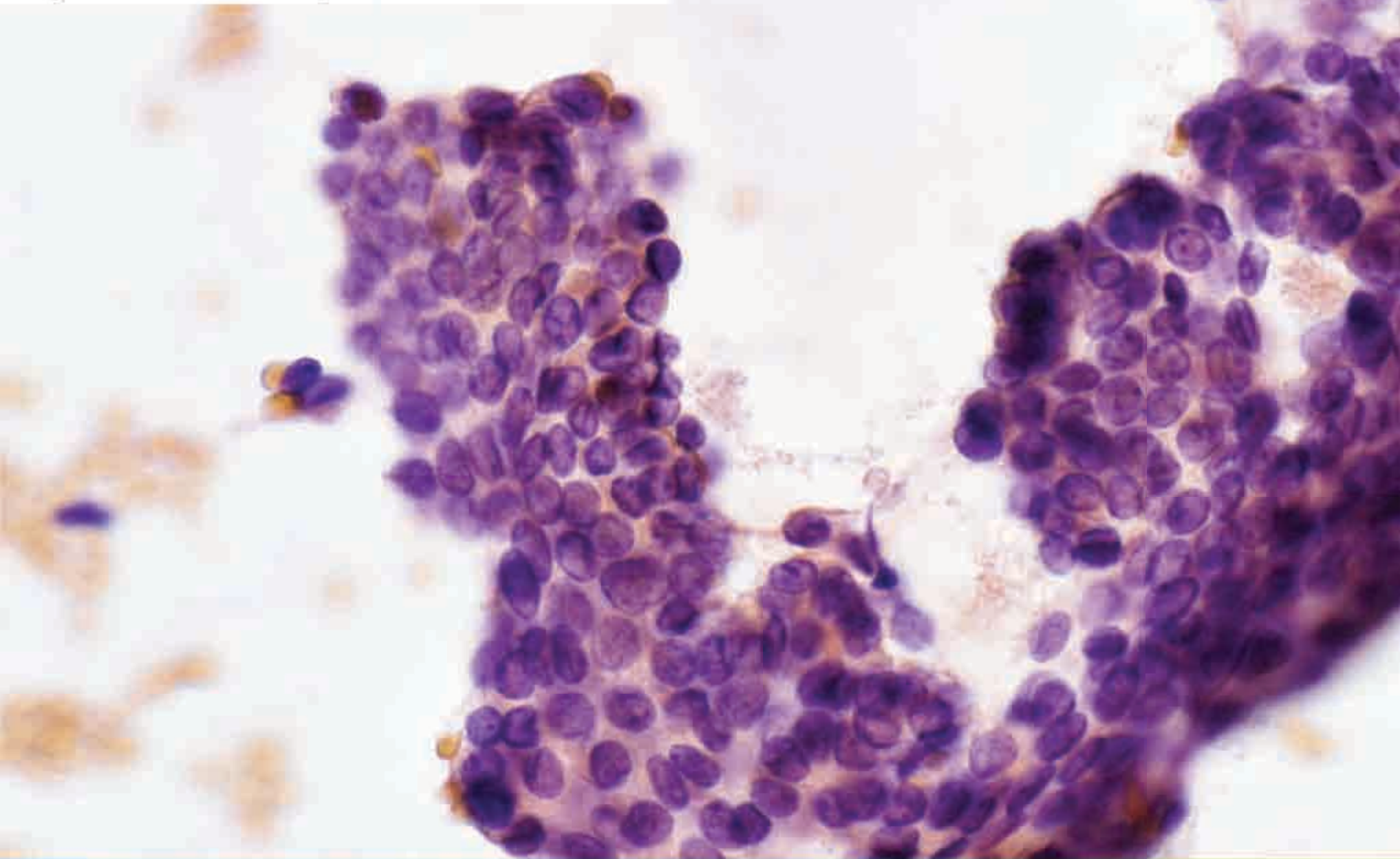






**Thyroid, left, ultrasound-guided FNA:  
Papanicolaou stain, 20x**

**Thyroid, left, ultrasound-guided FNA:  
Papanicolaou stain, 40x**





# Thyroid, left, ultrasound-guided fine needle aspiration:

**Malignant tumor cells present derived from papillary carcinoma.**



# Thyroid and neck dissection, excision:

Papillary carcinoma, well differentiated.

Tumor necrosis: Not present

Tumor size:

Right lobe 1.7 x 1.5 cm

Isthmus and left lobe 7 x 5 cm

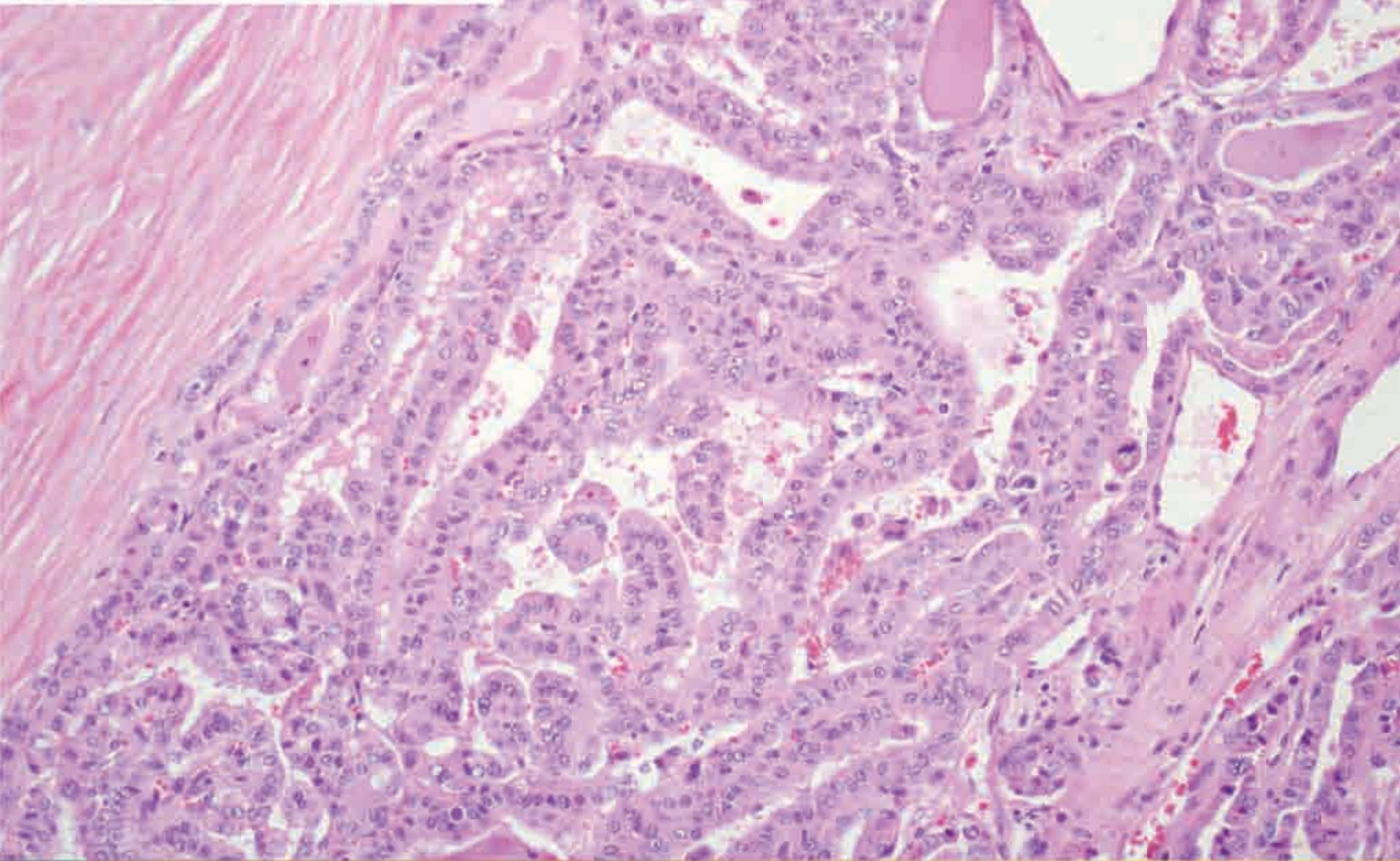
The tumor is partially encapsulated.

Other tumor features Psammomatous calcification.

Lymph node: Nine out of twenty four lymph nodes are + for metastatic disease.

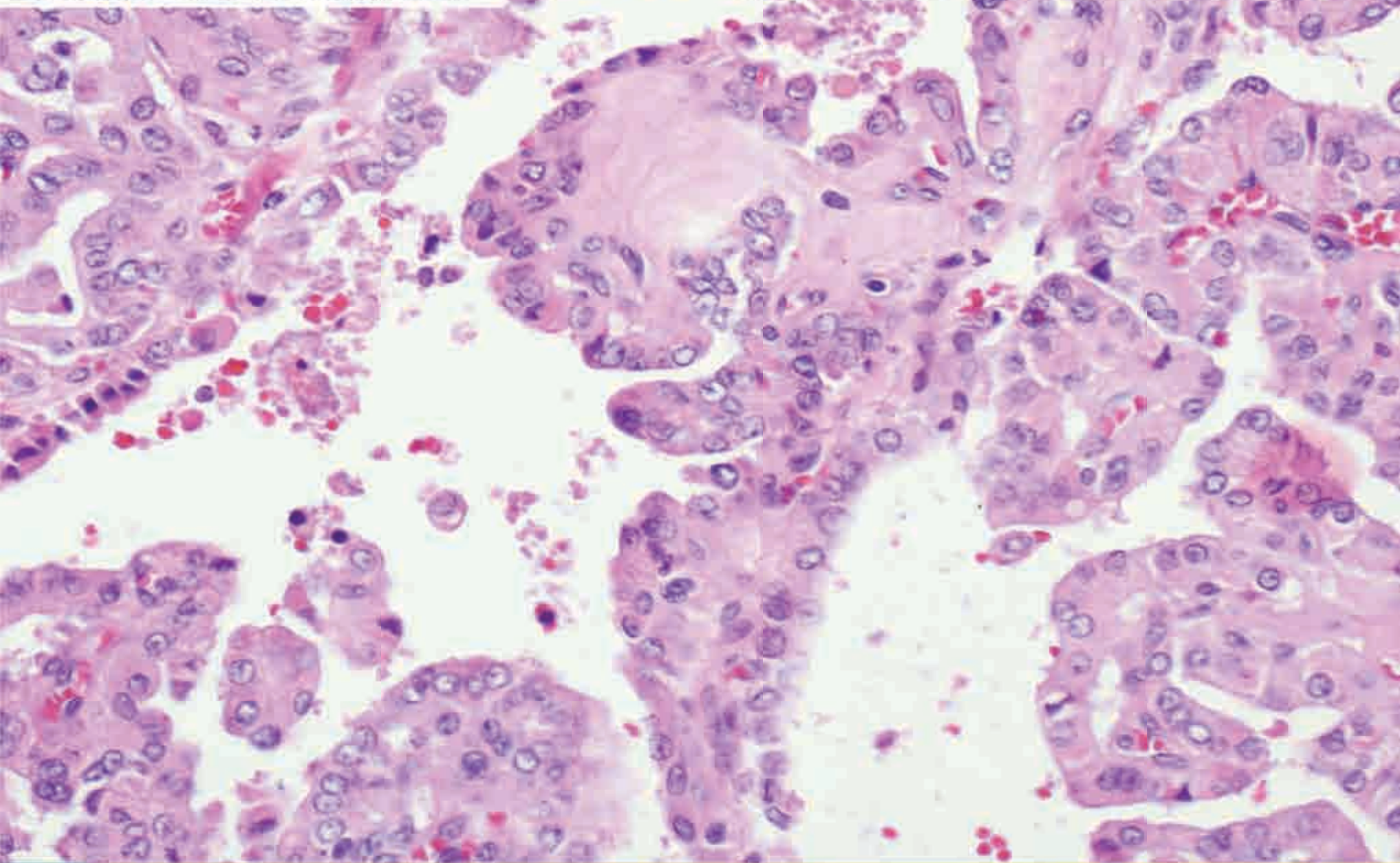


**Thyroid and neck dissection,  
excision: H & E stain, 10x**



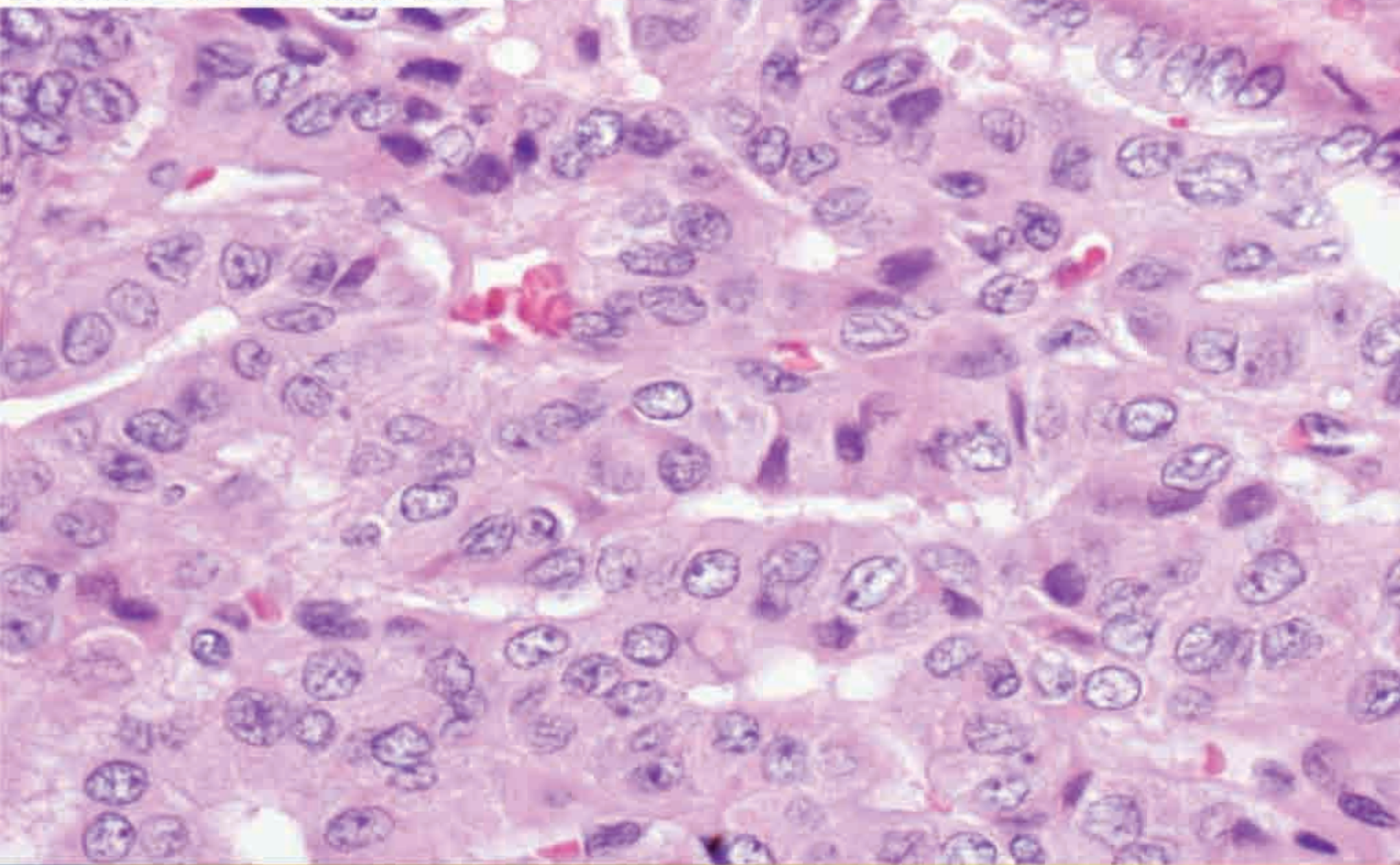


**Thyroid and neck dissection,  
excision: H & E stain, 20x**





**Thyroid and neck dissection,  
excision: H & E stain, 40x**



# Papillary Thyroid Carcinoma

- Best diagnostic clue: Focal, intrathyroidal mass with associated extracapsular invasion +/- metastatic nodes
- Highly variable ultrasound appearance with no definite features to differentiate benign from malignant
- Most common appearance is a solid hypoechoic or isoechoic mass +/- calcification



# Papillary Thyroid Ca – Clinical Issues

- Painless, palpable, solitary thyroid nodule
- Clinical profile: female with firm thyroid nodule
- Age: Peak incidence in 3<sup>rd</sup> and 4<sup>th</sup> decades
- 20 year survival rate:
  - 90% (papillary), 75% (follicular)
- Surgical resection: Total thyroidectomy and regional lymphadenectomy
- Followed by I-131 ablative therapy if indicated



# Follow-up

- Patient underwent total thyroidectomy and had two avid areas of radiotracer uptake in the neck on subsequent I-131 total body scan.
- She underwent thyrogen protocol and received 148 mCi of I-131.
- There was good radiotracer uptake in the thyroid surgical bed and therapeutic benefit can be expected.