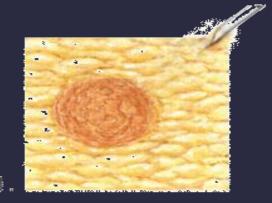




Αρχές και αναγκαιότητα της κλειστής κατευθυνόμενης βιοψίας

Dimitrios K Filippiadis MD, PhD, MSc, EBIR Associate Professor of Diagnostic and Interventional Radiology 2nd Radiology Dpt, University General Hospital "ATTIKON" Medical School, National and Kapodistrian University of Athens





ΔΙΑΔΕΡΜΙΚΕΣ ΒΙΟΨΙΕΣ



- Διαδερμική βιοψία διά βελόνης: εισαγωγή βελόνας εντός αλλοίωσης αγνώστου υποβάθρου με σκοπό τη λήψη κυττάρων ή ιστικού τεμαχίου
 - Αναρρόφηση διά λεπτής βελόνης (<20-22G):
 απόκτηση κυττάρων προς κυτταρολογικό έλεγχο

Ιστική βιοψία: απόκτηση ιστοτεμαχιδίου με βελόνη διαμέτρου >20G προς παθολογοανατομική εκτίμηση



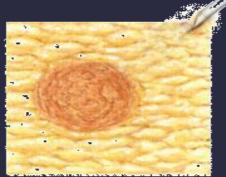
20-22 G NEEDLES











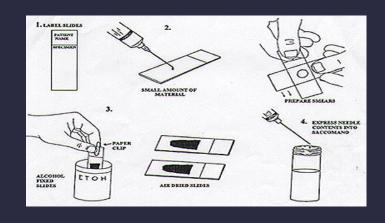












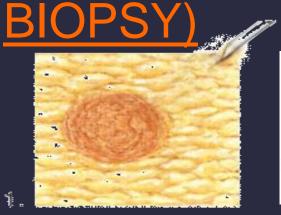


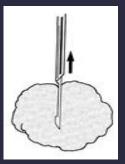


BIOΨΙΑ ΜΕ ΒΕΛΟΝΑ ΜΕ TEMNON AKPO (CORE NEEDLE



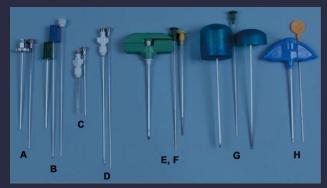
Βελόνες cut 10-18 G





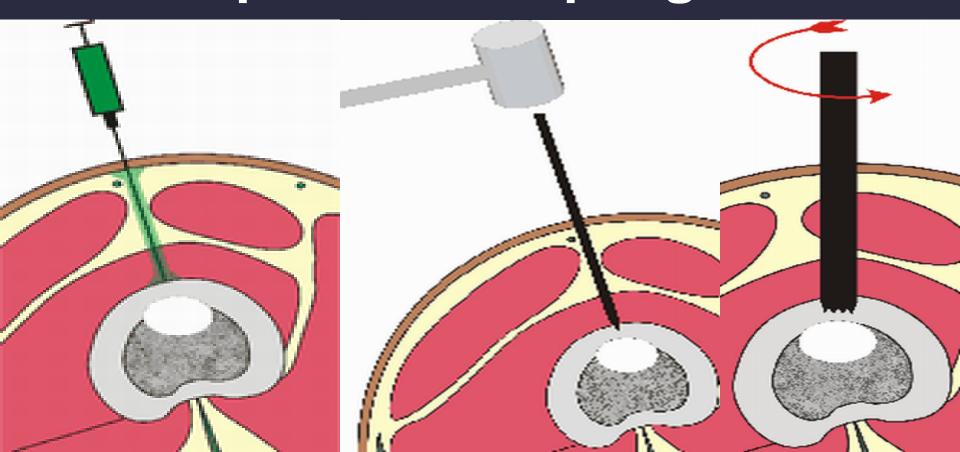








Techniques for sampling bone

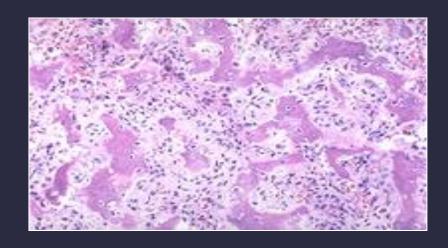
















 More than 1.4 million patients are diagnosed with cancer annually in the United States

Skeletal system: third most important filter for cancer metastases after lungs and liver





- Spine: the most common site of osseous metastatic disease
- WHY SPINE?
 - Presence of vascular red marrow in adult vertebrae
 - Communication of deep thoracic and pelvic veins with valveless vertebral venous plexuses





- Robertson and Ball (mid 1930s) posterolateral percutaneous approach
- Fluoroscopically guided needle biopsy of the spine:
 1949 -
- CT guided biopsy of the spine: 1981-





- Favourable risk-benefit ratio
- Technically feasible
- Sampling from different parts of the tumor
- Adequate for both histopathologic and molecular evaluation
- Potentially result in change of therapy







Table 1 Sensitivity and specificity of imaging modalities in bone metastasis

Imaging modality	Sensitivity (%) ^[12]	Specificity (%) ^[12]
18F NaF-PET/CT	100	97
MRI	95	90
SPECT	87	91
18F FDG-PET	98	56
CT	74	56
Bone Scintigraphy	78	48

Imaging of bone metastasis: An update

World J Radiol 2015 August 28; 7(8): 202-211 ISSN 1949-8470 (online)

Gerard J O'Sullivan, Fiona L Carty, Carmel G Cronin





ISSUES TO CONSIDER PRIOR TO BIOPSY:

- Review recent cross sectional imaging (x rays, CT, MRI, PET/CT)
- Lesion shape / location / lytic vs blastic
- SINS
- Vascularity
- Close by sensitive structures

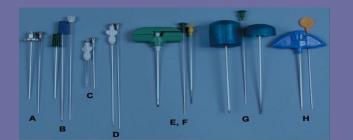




- Select the image guidance method optimal for the biopsy
- Assess the appropriate needle trajectory that would likely give the greatest diagnostic yield
- Choose the most appropriate biopsy system and type of anesthesia















- U/S (real-time imaging)
- Fluoroscopy (real-time imaging)
- CT (CT fluoro) (high-resolution images, 3D)
- Flat panel cone-beam CT (more accurate and faster)
- MR (thermal monitoring capacity)





MAGING GUIDANCE









Table 1 Advantages and Disadvantages of the Imaging Modalities Used to Guide Celiac Plexus Block			
Modality	Advantages	Disadvantages	
Fluoroscopy	Simple to perform	Anatomic structures overlap; does not allow distinction between the pancreas, blood ves- sels, tumors, and lymph node metastases, thus increasing risk for complications; diffusion of neurolytic solution is not clearly displayed	
US	Simple and inexpensive; the aorta, celiac artery, and SMA are clearly identified; diffusion of neurolytic agent may be seen without contrast medium	Operator dependent, retroperitoneal organs (including the pancreas) are not clearly identified	
Multide- tector CT	High contrast and spatial resolution; clearly depicts retroperitoneal structures and extent of tumor; celiac plexus may be directly identified; needle puncture site and needle course may be planned in advance; depicts the exact location of the needle tip and surrounding structures, helping avoid vital organ damage; accurately depicts diffusion of neurolytic agents; CT fluoroscopy allows real-time monitoring of the procedure	Risk for radiation exposure	
MR imaging	Superior soft-tissue resolution, contrast material not necessary, no ionizing radiation	Expensive, limited availability, requires MR imaging-compatible equipment	
Endoscopic US	Real-time monitoring of neurolytic agent in- jection, use of an anterior approach avoids neurologic complications	Operator dependent, invasive, risk for complica- tions (eg, gastric perforation, pancreatitis), snowstorm effect may hinder visualization of the celiac plexus	

Kambadakone A et al CT-guided Celiac Plexus Neurolysis: A Review of Anatomy, Indications, Technique, and Tips for Successful Treatment Radiographics 2011



IMAGING GUIDANCE

<u>Advantages</u>

- No radiation exposure
- Rapid image acquisition
- Widespread availability

Disadvantages

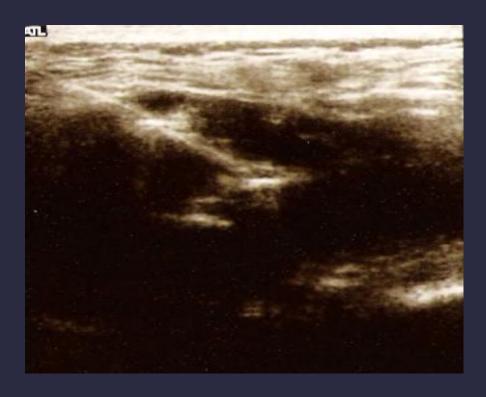
- Difficult imaging in some patients
- Image quality
- Difficult visualization (esp. of the iceball)



MAGING GUIDANCE













<u>Advantages</u>

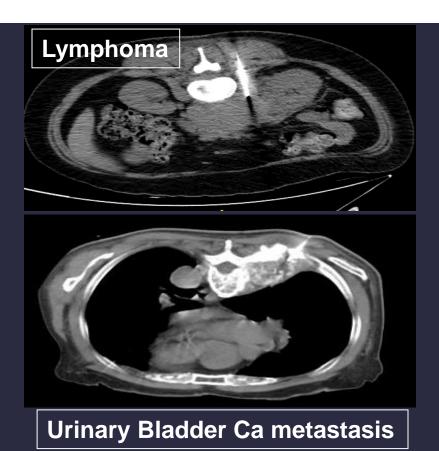
- Widely available
- Rapid image acquisition
- Better visualization of cryoprobe/RF electrode

Disadvantages

Radiation exposure







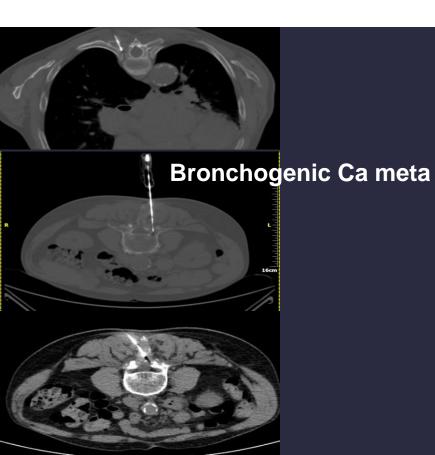


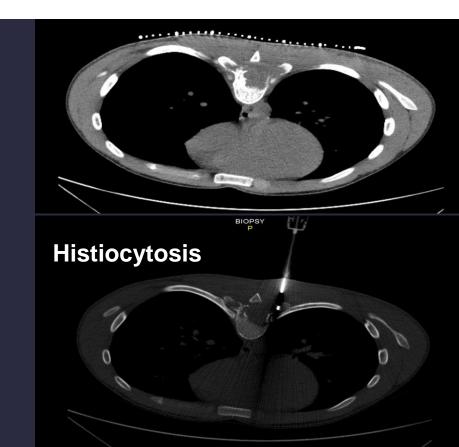


Multiple Myeloma















VR Advantages

- Best visualization of probe/RF electrode
- No radiation exposure
- Best visualization of iceball
- Image in sagital and coronal plane
- Thermal monitoring capacity

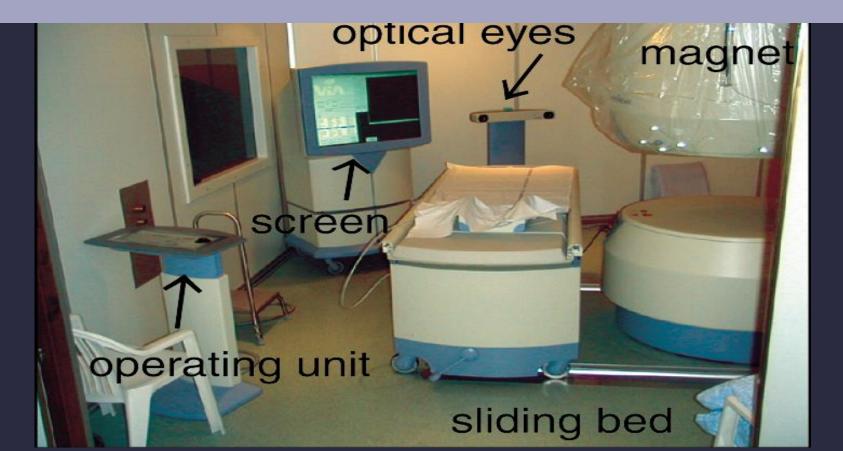
Disadvantages

- Small working area
- Limited availability
- Longer duration





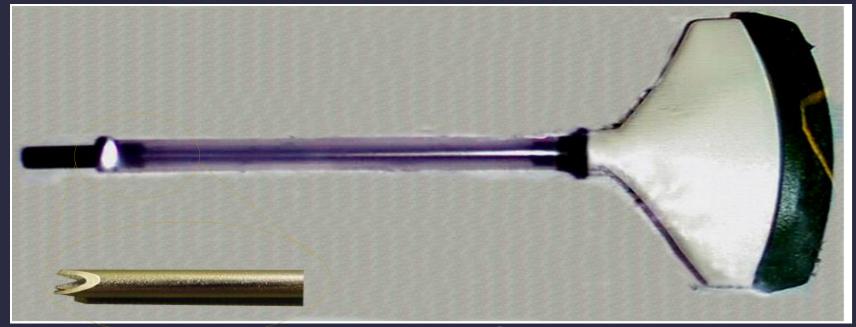






MAGING GUIDANCE



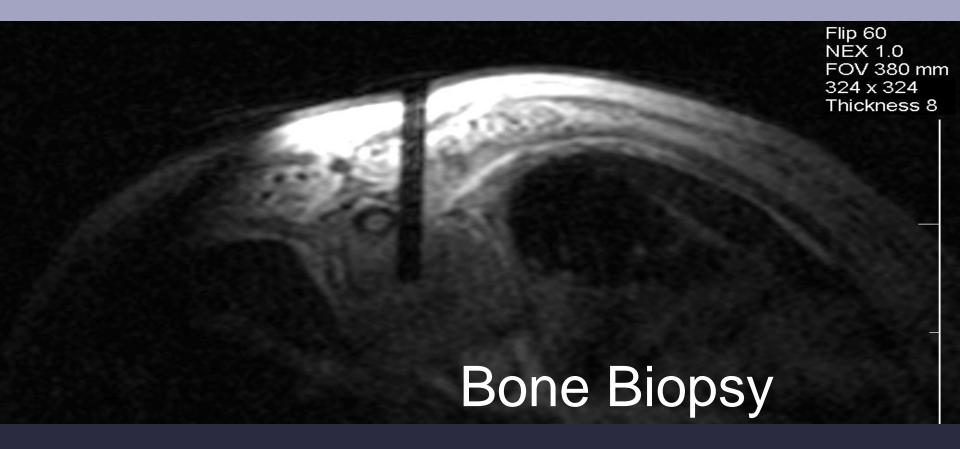


Titanium Biopsy Systems













Cardiovasc Intervent Radiol (2016) 39:290–295 DOI 10.1007/s00270-015-1216-y C RSE



TECHNICAL NOTE

Fluoroscopy-Guided Percutaneous Vertebral Body Biopsy Using a Novel Drill-Powered Device: Technical Case Series

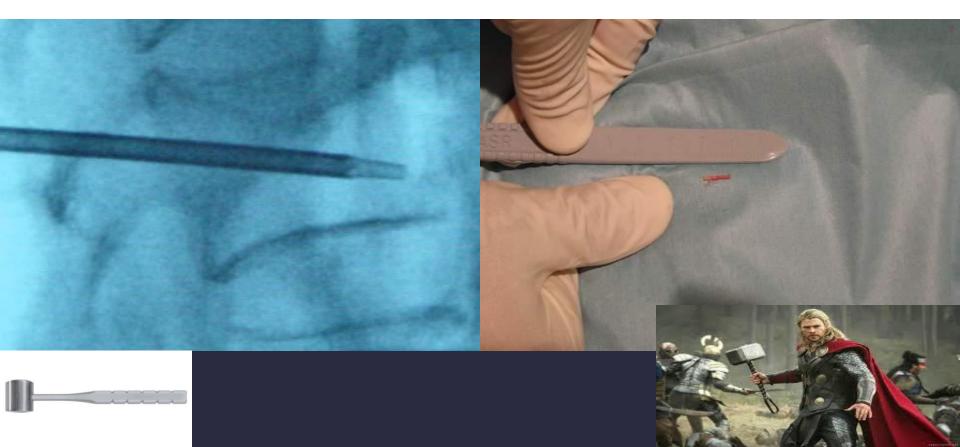
Adam N. Wallace¹ · Rafael A. Pacheco¹ · Anderanik Tomasian¹ · Andy C. Hsi² · Jeremiah Long¹ · Randy O. Chang³ · Jack W. Jennings¹

- Reduce procedure time and radiation exposure
- Easily cuts through even sclerotic bone lesions
- Obtain more bone core





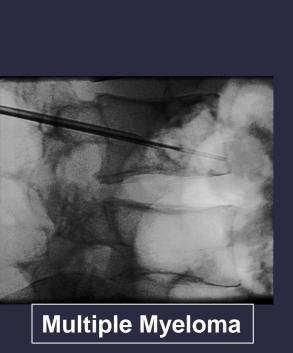




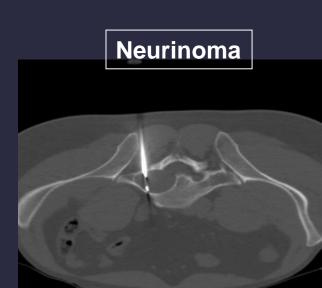


MSK













Targeting of metabolic and progressing lesions

Target the periphery of the lesion

After each core, aspirates or bleeding back through the large needle

CONSIDER PATIENT SAFETY





- Percutaneous biopsy of intramedullary lytic lesions that do not have a soft tissue component may only yield blood clots
- Specimens containing principally blood clots were obtained in 20.8% of cases with percutaneous needle biopsy providing diagnosis for a diagnostic yield of 75%

Harish S, Hughes RJ, Saifuddin A, Flanagan AM. Image-guided percutaneous biopsy of intramedullary lytic bone lesions: utility of aspirated blood clots. Eur Radiol. 2006





- Coagulopathy (INR>1.5, PT>1.5 times control, platelets<50.000/mm³)
- Systemic infection skin infection at puncture site
- Significant spinal cord compromise at the biopsied level (theoretical risk of causing or aggravating a myelopathy)
- Epidural tumor in the spinal canal (risk of tissue swelling/bleeding and further spinal cord compromise)





 Confirm metastatic tumor involvement of the spine in a patient with a known primary neoplasm - Confirm diagnosis of multiple myeloma

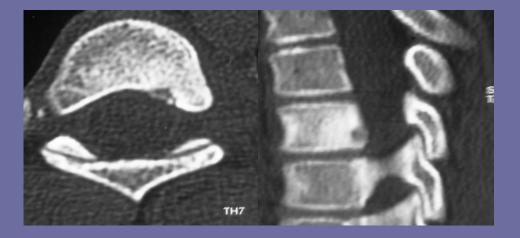
Determine a lesion with intermediate or aggressive imaging features

Assess benign vs pathologic compression fracture





routine biopsy at osteoid osteoma ablation







Overall accuracy 77-97%

Higher positive recovery rates for osteolytic lesions

Lower positive recovery rates for sclerotic lesions

 Lower accuracy rates does not necessarily indicate suboptimal technique or invalid procedure







Lesion-related or Technical Factor	C	umulative No. of	No. of Specimens to Reach Diagnostic				
	1	2	3	4	5	6	Yield Plateau
All lesions (n = 151)	65	71	75	77	77	77	4
General lesion subtype							
Soft tissue (n = 63)	67	71	73	76	76	76	4
Bone (n = 88)	64	70	77	77	77	77	3
Bone lesion subtype							
Sclerotic (n = 28)	39	46	57	57	57	57	3
Lytic (n = 60)	75	82	87	87	87	87	3
Lesion size (cm)							
≤2 (n = 24)	42	50	54	54	54	54	3
>2 To 5 (n = 57)	67	72	74	75	75	75	4
>5 (n = 70)	71	77	84	86	86	86	4
Biopsy needle gauge							
14 (n = 6)	67	67	67	83	83	14/41	4
15 (n = 68)	54	63	72	72	72	72	3
16 (n = 30)	67	73	73	77	77	77	4
18 (n = 47)	79	81	83	83	83	83	3
Imaging guidance modality							
CT (n = 133)	65	71	76	77	77	77	4
US (n = 18)	67	72	72	78	78	78	4

Wu JS et al. Bone and soft tissue lesions: What factors affect diagnostic yield of image guided core needle biopsy. Radiology. 2008; 248: 962-970.



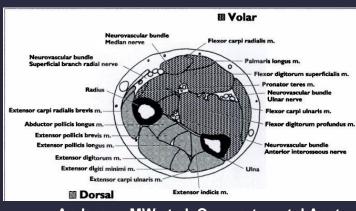


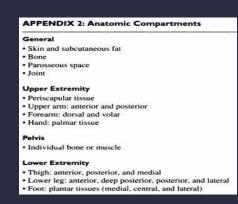
BONE BIOPSY: respect compartmental anatomy

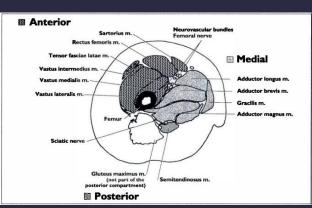
Perspective

Compartmental Anatomy: Relevance to Staging and Biopsy of Musculoskeletal Tumors

Mark W. Anderson 1.2, H. Thomas Temple 2.3, Robert G. Dussault 1.2, Phoebe A. Kaplan 1.2



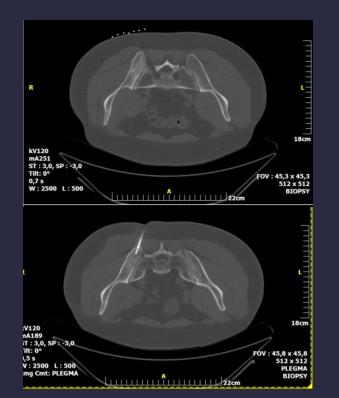


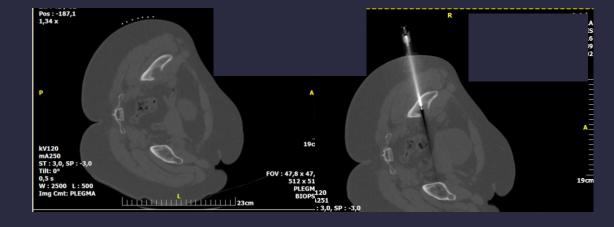




BONE BIOPSY: respect compartmental anatomy









BONE BIOPSY: respect compartmental anatomy





STEPS TO A SOLUTION:

Talk to your surgeon to determine the ideal approach route



SPINE BIOPSY



SPINE BIOPSY TARGET

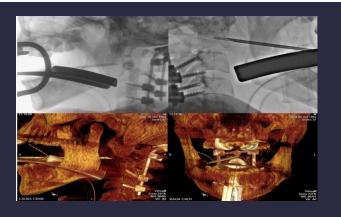


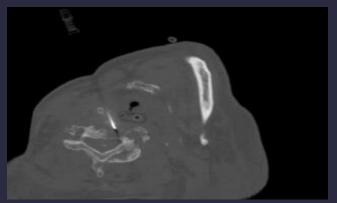


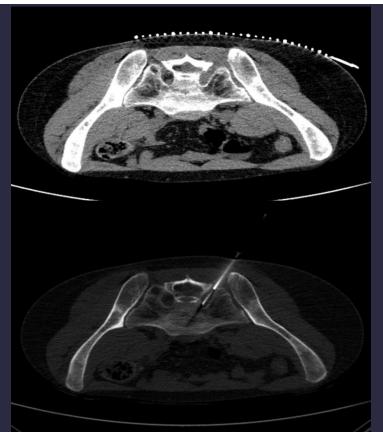
- 1. Vertebral body
- Posterior vertebral arch
- Intervertebral disc
- 4. Facet joint
- Paraspinal mass
- 6. Epidural space neural foramen
- Spinal cord

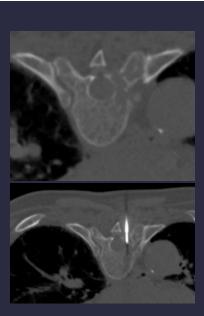






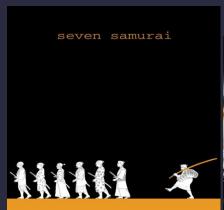








SPINE BIOPSY TARGET





- 1. Vertebral body
- 2. Posterior vertebral arch
- Intervertebral disc
- 4. Facet joint
- Paraspinal mass
- 6. Epidural space neural foramen
- Spinal cord

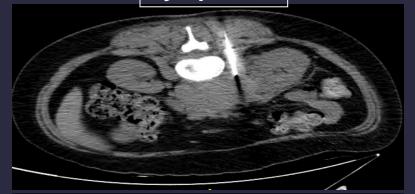


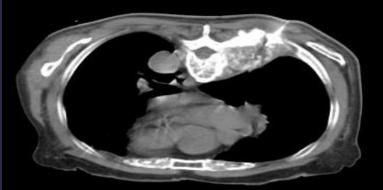
SPINE BIOPSY TARGET CONSIDERATIONS

PARASPINAL MASS



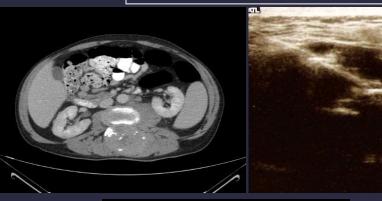
Lymphoma





Urinary Bladder Ca metastasis

Gastric Ca metastasis





Multiple Myeloma



SPINE BIOPSY TARGET CONSIDERATIONS

VERTEBRAL BODY



CERVICAL SPINE APPROACH

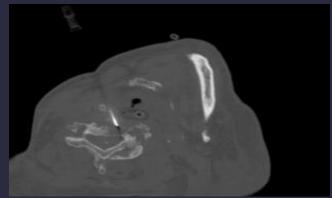
TRANS-ORAL





ANTERO-LATERAL

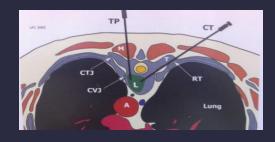


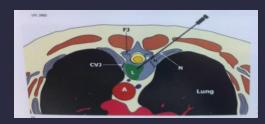


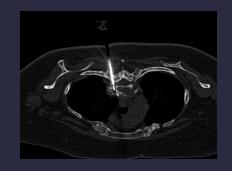


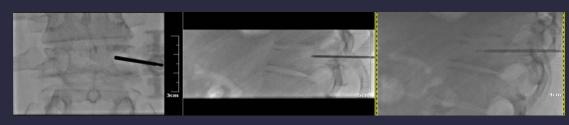
THORACIC SPINE APPROACH

COSTOVERTEBRAL
COSTOTRANVERSE JOINT
INTERCOSTAL
TRANS-PEDICULAR







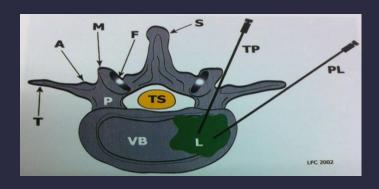




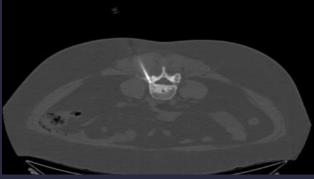
LUMBAR SPINE APPROACH

POSTEROLATERAL

TRANS-PEDICULAR



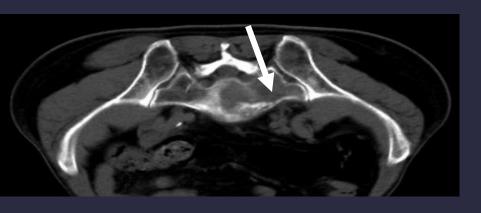


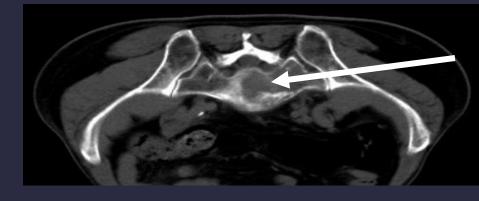






SACRAL SPINE APPROACH











Blood Clots

- Percutaneous biopsy of intramedullary lytic lesions that do not have a soft tissue component may only clots.
- Specimens containing principally blood clots were obtained in 20.8% of cases with percutaneous needle biopsy providing diagnosis allowing appropriate further management in 62 cases, for a diagnostic yield of 75%.

Harish S, Hughes RJ, Saifuddin A, Flanagan AM. Image-guided percutaneous biopsy of intramedullary lytic bone lesions: utility of aspirated blood clots. Eur Radiol. 2006



BIOPSY AND SPONDYLODISCITIS

- De Lucas et al (2009):
 - Diagnostic rates obtained in patients with previous antibiotic treatment were significantly lower (23% vs. 60%, p=0.013)..... this technique yields a lower diagnostic rate than previously reported biopsy of neoplastic vertebral lesions, especially if performed in patients with previous antibiotic treatment
- Pupaibool et al (2015):
 - The microbiological yield of image-guided needle biopsy varies between 36% and 91%... Image-guided spinal biopsy had a sensitivity of 52.2% and a specificity of 99.9%



Culture or Histological sample?

The results of our study support the conclusion that both microbiologic and histologic evaluation of bone biopsy specimens should be performed in cases of suspected osteomyelitis.

We found the sensitivity of culture alone in the diagnosis of osteomyelitis to be 42%, and the sensitivity of combined culture and histologic examination to be 84%.



BIOPSY AND SPONDYLODISCITIS

Downloaded from http://pmj.bmj.com/ on December 22, 2014 - Published by group.bmj.com

607

ORIGINAL ARTICLE

Therapeutic impact of percutaneous spinal biopsy in spinal infection

J J Rankine, D A Barron, P Robinson, P A Millner, R A Dickson

Postgrad Med J 2004;80:607-609. doi: 10.1136/pgmj.2003.017863

See end of article for authors' affiliations

Correspondence to: Dr James J Rankine, Department of Clinical Radiology, St James's University Hospital, Beckett Street, Leeds LS9 7TF, UK; james.rankine@leedsth. nhs.uk

Submitted 8 December 2003 Accepted 16 January 2004 **Objective:** To investigate the therapeutic impact of percutaneous spinal biopsy in patients with suspected spinal infection.

Design and patients: A review of the case notes and imaging features of 36 patients who underwent percutaneous spinal biopsy was performed. From this group 20 patients with a prebiopsy diagnosis of spinal osteomyelitis were identified. Management before biopsy was noted including the use of antimicrobial therapy. The results of the histology and microbiology were noted along with the subsequent diagnosis and management.

Results: Eight of the 20 patients (40%) had received antibiotics before the biopsy. An organism was isolated in 8/20 cases (40%). Of the eight patients on antibiotics, two grew an organism (25%), including one case of candida in a patient receiving flucloxacillin. Out of 12 patients not on antibiotics there were six cases where an organism was isolated (50%). The result of the biopsy led to a change in management in seven of the 20 patients (35%).

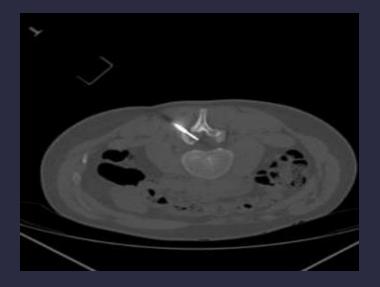
Conclusions: Many clinicians are treating spinal osteomyelitis empirically with antibiotics before biopsy, but this reduces the chance of isolating an organism and determining antibiotic sensitivity. Despite this biopsy led to a change in management in 35% of cases.

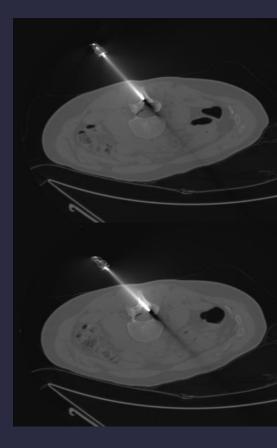


SPINE BIOPSY TARGET CONSIDERATIONS

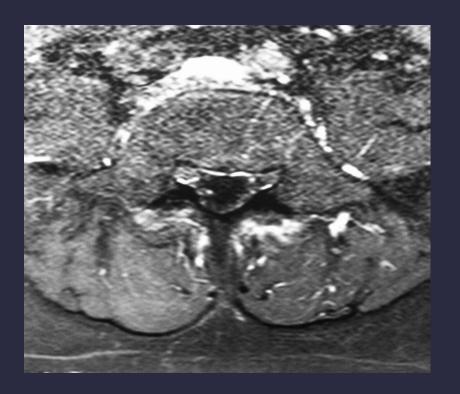
FACET JOINT

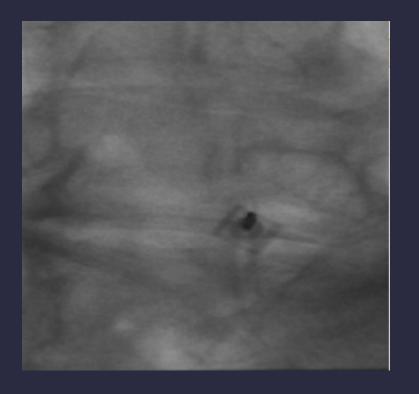












Septic arthritis of the facet joint

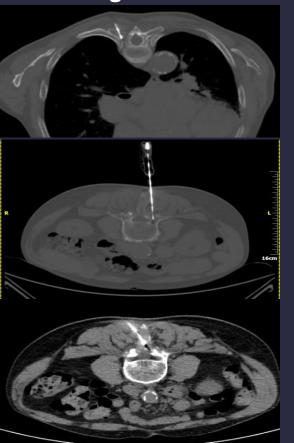


SPINE BIOPSY TARGET CONSIDERATIONS

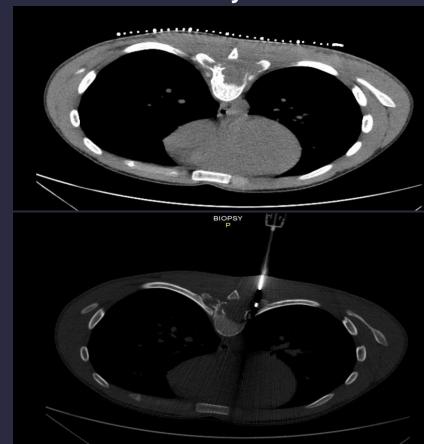
POSTERIOR VERTEBRAL ARCH



Bronchogenic Ca meta



Histiocytosis

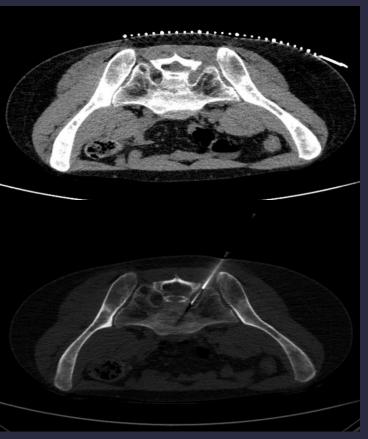


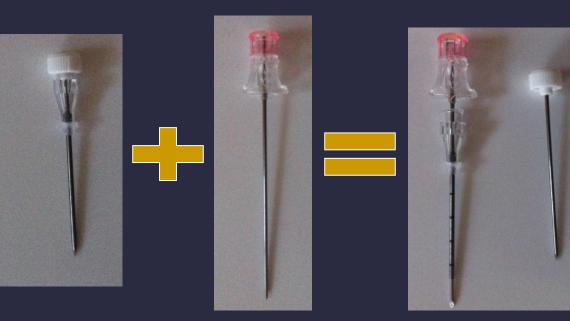


SPINE BIOPSY TARGET CONSIDERATIONS

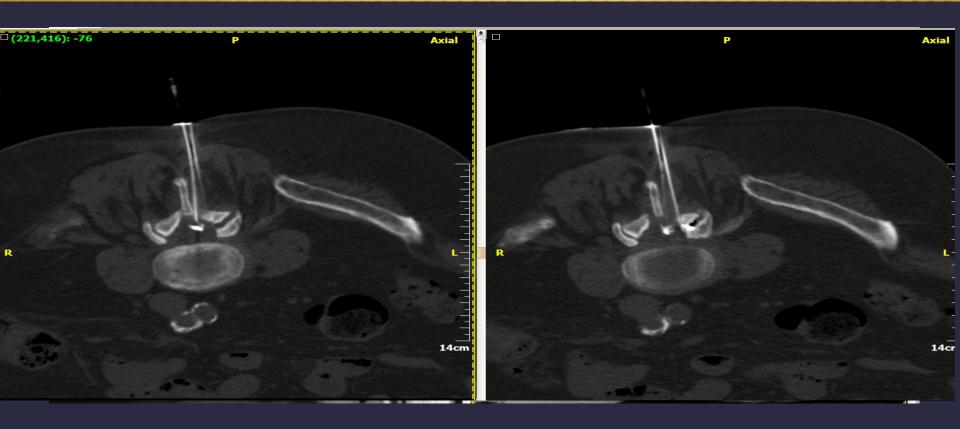
NEURAL FORAMEN
EPIDURAL SPACE











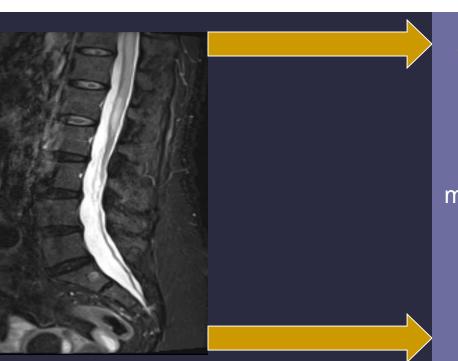


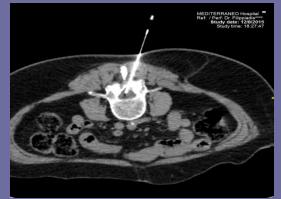
SPINE BIOPSY TARGET CONSIDERATIONS

SPINAL CORD













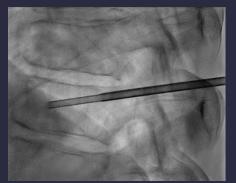


CLINICAL STUDY

CT-Guided Bone Biopsies in Metastatic Castration-Resistant Prostate Cancer: Factors Predictive of Maximum Tumor Yield

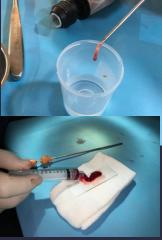
Michael G. Holmes, MD, Erik Foss, MD, Gabby Joseph, PhD, Adam Foye, BS, Brooke Beckett, MD, Daria Motamedi, MD, Jack Youngren, PhD, George V. Thomas, MD, Jiaoti Huang, MD, PhD, Rahul Aggarwal, MD, Joshi J. Alumkal, MD, Tomasz M. Beer, MD, Eric J. Small, MD, and Thomas M. Link, MD. PhD





Urology Case Reports 6 (2016) 45e46

c/o E. Eyheremendy Argentina



Growth 24 months	Accuracy
> 25%	81%
No change	42%





Decalcification

Hard: Freezing



- Optimal antigen preservation
- Poor morphological preservation



Cut slides



Soft: Decalcification



- Strong acids : RDO faster
- Chelating agent EDTA longer but do not damage Biomarkers







The accuracy and reliability of immunohistochemical testing is multifactorial:

- method of tissue fixation
- method of staining methods
- antigen retrieval
- subjective scoring

biopsies from different tissues require different preparation efforts leading to significant technical variations





Next-generation sequencing (NGS)

- Metastatic bone specimens : higher failure rate than nonbone specimens (36% vs 2.3%).
- Resection specimens, a higher failure proportion of which (6 of 10 specimens) were submitted for regular decalcification.
- FNA smear slides, alternative sources for mutational profiling of bone metastasis.

Core biopsy specimens should be grossly examined for the presence or absence of bones.

Clinical Mutational Profiling of Bone Metastases of Lung and Colon Carcinoma and Malignant Melanoma Using Next-Generation Sequencing

Gang Zheng, MD, PhD¹; Ming-Tseh Lin, MD, PhD¹; Parvez M. Lokhandwala, MD, PhD¹; Katie Beleri, MS¹; George J. Netto, MD¹³; Christopher D. Gocke, MD¹²; James R. Eshleman, MD, PhD¹²; Edward McCarthy, MD¹; and Peter B. Illei, MD¹².





- Vascular injury Hematoma myelopathy
- Infection
- Nerve root damage radiculopathy
- Thecal sac puncture headache arachnoiditis quadriparesis, quadriplegia
- Pneumothorax
- Vasovagal reaction
- Fracture
- Drug related allergy
- Tumor seeding

AJR 2009; 193:W407–W410. Eur Spine J (2008) 17:975–981. Skeletal Radiol (2002) 31:349–353 Tehranzadeh et al Acta Radiol 2007 Le et al Semin Intervent Radiol 2010 Huang et al Radiol Clin N Am 2011

0-7.4%





- Identify new targets
- Identify optimal treatment
- Predict tumor response in advanced stage
- Identify tumor recurrence
- Predict rate of recurrence
- Predicting recurrence in curative cases could help in treatment stratification, identification and validation of new targets







- Pathologic analysis:
 - Histology
 - Biomarkers (prognostic, predictive)
 - Molecular profiling
 - Genetic sequencing

- Pathologic assays:
 - Immunohistochemistry
 - Polymerase chain reaction
 - Fluorescence in situ hybridization
 - Direct gene sequencing





Tumor Type	Tumor Marker	Molecular Test	Clinical use
NSCLC	EML4-ALK EGFR mutation	FISH PCR	Targeted drug therapy
Colon Adenocarcinoma	Microsatellite instability KRAS BRAF	PCR	Prognostic,HNPCC screening Targeted drug therapy
Melanoma	BRAF	PCR	Targeted drug therapy
Pancreas	SMAD-4	IHC	Differentiate Pancreatic adenocarcinoma from Metastasis
HCC	HepPar-1 Glypican 3	IHC	Differentiate HCC from Metastasis
Renal	RCC monoclonal antibody	IHC	Differentiate Renal cyst from RCC





 Use the largest possible needle biopsy that can safely maximize the diagnostic yield - Perform all the necessary tests for the obtained sample

 Effective communication with pathologist, molecular diagnostic laboratory and oncologist treating the patient



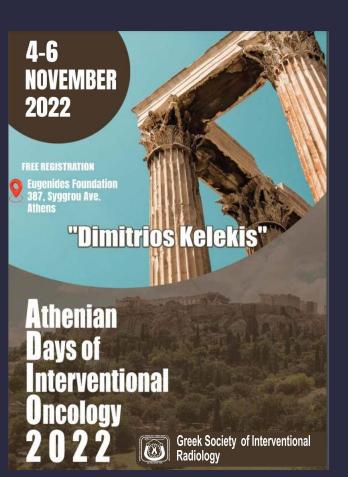


 Increasingly important as recent advances in molecular profiling and tissue biomarkers require adequate tumor sample

 Must be aware of the expanded indications, tissue yield and expectations related to biopsy: adapt to the new changes and alleviate possible challenges







16:30 - 18:00

MSK - A. KELEKIS & A. GANGI

Surgical approaches for MSK mets - P. Papaggelopoulos

Ablation in the spine and peripheral skeleton: which technique, which target - J. Jennings

Percutaneous augmentation in the spine: is there room for improvement? - S. Marcia

Percutaneous augmentation in peripheral skeleton: is there room for improvement? - D. Filippiadis

Trans-arterial embolization for MSK lesions: indications, technique, results - A. Ryan

Radiation therapies in the spine and peripheral skeleton - M. Trichas Combined approaches - M. Callstrom





