MUSCLE DISORDERS

KG  4/15/2010
“The Quizmaster”

- 12 unknowns
- 1 best diagnosis
51 yo F with right shoulder pain
54 yo F with thigh pain
54 yo M with foot pain
35 yo M with limited neck extension
47 yo F with history of trauma
23 yo M with calf pain and inability to walk
12 yo M with knee pain and fevers
41 yo F who recently underwent bowel surgery
37 yo M in MCA several months ago
60 yo M with weakness and muscle pain
Middle aged man with calf pain
TYPES OF MUSCLE INJURIES

- Denervation
- Laceration
- Strain
- Contusion
- Fascial Tear
- Avulsion
- Fatigue
- Iatrogenic
MUSCLE DENERVATION

- Acutely denervated muscle does not show signal abnormality
- High signal on T2WI is usually does not become evident for 2-4 weeks after denervation has occurred

- If innervation is restored, MR findings eventually return to normal. If innervation is not restored, fatty atrophy occurs over a period of months, indicating irreversible changes in the muscle

- Unlike strained muscles:
  - No perifascial edema
  - Specific nerve territory
48 yo M with right shoulder pain and weakness after MCA 8 months ago
Suprascapular nerve compression by HO
MUSCLE LACERATION

- Produced by penetrating injury, such as a knife
- In the acute setting, these are rarely imaged with MR
- Show focal, sharply margined discontinuity of fibers and high T2 signal intensity caused by hemorrhage and edema

MUSCLE STRAIN
**MUSCLE STRAIN**

- Most common pathology we see
- Occur in muscles that cross 2 articulations
- Usually occurs with eccentric contraction (contract during elongation)
  - Hamstrings
  - Rectus Femoris
  - Gastrocnemius
First Degree/Mild/Low grade injury

• No significant loss of strength or range of motion
• Heals completely with appropriate rest from aggravating activities
• (+) Increased signal on T2 and STIR
  • Edema and hemorrhage at the MTJ that extends into adjacent muscle fascicles
  • Feathery appearance on MRI
• (-) No architectural distortion of muscle or tendon
Connell, David A. et al: Longitudinal study Comparing sonographic and MRI assessments of acute and healing hamstrings injuries
Second Degree/Moderate grade tear

- Some loss of strength
- Pain relief within 2 weeks and return to sports after about 3 weeks
- Increased signal on T2 and STIR
- Hematoma at MTJ
- Perifascial fluid
- Architectural distortion
Moderate grade strain
Third Degree/Severe/High grade tear

- Complete musculotendinous disruption, with or without retraction
- Loss of strength
- Prompt surgery may be required
- Diagnosis usually made on clinical grounds but MR is helpful:
  - For preoperative assessment of the extent of retraction
  - To pinpoint hematomas which require percutaneous drainage
CHRONIC CHANGES AFTER TEARS

*Tendon thickening or attenuation
*Muscle atrophy

Normal Example

Our Patient

47 yo F 8 months after complete hamstrings origin avulsion
MUSCLE CONTUSION

• Imaging similar to a low grade muscle strain
  – High signal on T2 and STIR: diffuse or geographic pattern, no fiber discontinuity
    • *with contusion, overlying edema within subcutaneous fat
• History is different than muscle strain
  – Direct trauma, usually a blunt object
• Larger in size than strain injury but shorter recovery times
SEQUELAE OF MUSCLE INJURY
14 yo M with persistent left thigh swelling and pain

No response to physical therapy
MYOSITIS OSSIFICANS

• Trauma, paralysis, burns

• Radiographic findings:
  – Zonal pattern of mineralization
  – Ossifies from the outside in
  – May merge with the underlying bone and resemble an osteochondroma
  – May be resorbed over a period of 1-5 years
MYOSITIS OSSIFICANS

• MRI findings
  – EARLY: heterogenous muscle edema
  – SUBACUTE: mass like region of high T2 signal during first days to weeks after the injury (difficult to distinguish from sarcoma)
  – LATE: older lesions develop peripheral calc: peripheral low signal intensity and central fat signal intensity.

• Recognition of peripheral calcification pattern is important for making the correct diagnosis because biopsy, particularly of the central portion, may lead to a false diagnosis of osteosarcoma due to the presence of abundant osteoid and mitotic spindle cells

*Workup should ALWAYS begin with plain films
*Short term follow up in 3-4 weeks with X-ray or CT scan is necessary to confirm suspected MO
*Allows postponement of a biopsy or surgical procedure until diagnostic imaging features have declared themselves
37 year old male several months after motorcycle accident
Chronic Degloving Injury

• Skin and subcutaneous fatty tissue abruptly separating from the underlying fascia
• Disrupted capillaries may continuously drain into the perifascial plane, filling up the virtual cavity with blood, lymph, and debris
• An inflammatory reaction commonly creates a peripheral capsule
32 yo M with increasing left leg mass after MCA one month prior
Injury disrupts segmental perforating vessels and results in a hematoma composed of hemolymphatic fluid with a mixture of viable and necrotic fat.
41 yo M with anterior lower leg pain
MUSCLE HERNIATION

• Protrusion of muscle through a focal fascial defect
  – **Traumatic**: tear of the fascial sheath due to penetrating wounds or violent impact associated with fractures.
  – **Constitutional**: Muscular overuse or hypertrophy may lead to fascial rupture at weak spots such as those traversed by vessels and nerves

• Rarely, a familial cause: congenital weakness in the fascia in some persons
MUSCLE HERNIATION

• Usually occur in the middle to lower portion of the leg
• **Anterior tibialis muscle** is most commonly involved
• May be multiple and bilateral

MUSCLE HERNIATION

• Asymptomatic hernias are treated conservatively

• For severe symptoms or cosmetic complaints, fasciotomy is the preferred surgical technique

• Fascial repair (closing fascial defect) may result in compartment syndrome and is no longer performed

MUSCLE ISCHEMIA AND NECROSIS

• Compartment syndrome
• Myonecrosis and Rhabdomyolysis
• Diabetic Muscle Infarct
Causes of ACS

• Blunt or penetrating trauma
  – fracture
  – soft tissue contusion
  – crush injury
  – gunshot wound
  – vascular injury

• Thermal or electrical injury

• Iatrogenic insult
  – Reperfusion edema
  – Arthroscopy, osteotomy, arthrodesis, THA
  – Pt positioning during long operations
  – Extravasation of contrast material or other parenteral drugs in the forearm
  – Anticoagulation
  – Hematoma formation after transaxillary arteriography
  – Prolonged application of an excessively tight cast, constrictive dressing, blood pressure monitor, or pneumatic antishock garment
Compartment Syndrome

- Fibular fracture
- Increased capillary permeability
- Swelling, edema, pain, erythema, and heat

AJR
Compartment Syndrome

- Serious complication of fractures of the tibial shaft
- Most important diagnostic feature is the presence of inappropriate pain even after stabilization of the fracture by a cast or by internal or external fixation
- Pain is made worse by passive stretching of the muscles involved and there is an associated sensory disturbance
  - 4 Ps: paresthesia, pallor, paresis, lack of pulse
- It has been shown that muscle can tolerate only 4 hours of ischemia without injury
- Treatment: Surgical fasciotomy and decompression

Hyder, N. Compartment syndrome in tibial shaft fracture missed because of a local nerve block. JBJS 1996 78-B: 499-500.
66 yo M referred for CT angiogram
• 4 hours after the injection, the patient was brought to the operating room for fasciotomy, washout, and placement of vacuum drainage device.

• At surgery, subfascial edema was found in the biceps compartment but the tissues appeared healthy and viable.

• The biceps fascia was divided longitudinally and the tissues were irrigated.
54 yo M with foot pain
Compartment syndrome and subsequent myonecrosis

**Sheet-like calcification or ossification of abductor hallucis and quadratus plantaris**
23 year old man with bilateral calf swelling and inability to walk
Supplements:

Glucosamine
Antiestrogen
Protein powder
Dicalcium phosphate
Stearic acid
Caffeine
• Compartment pressures in calves 35-45 mm Hg (nl 0-15)

• Operative findings:
  – Left side:
    • Immediate bulging of purple-gray muscle from medial calf
    • Massively swollen soleus and gastroc
    • Posterior compartment with poorly contractile musculature
  – Right side:
    • Similar but less severe findings, preserved contractile function

• Procedure:
  – Right and left calf four-compartment fasciotomies
T1: Patient

T1: Normal comparison
Rhabdomyolysis, compartment syndrome and myonecrosis
Rhabdomyolysis

- Striated muscle dissolution or disintegration
- Damages the sarcolemma, leading to muscle necrosis and release of toxins (potassium, phosphate, myoglobin, creatine kinase, urate)
- This leads to myoglobinemia, which often results in myoglobinuria
- Locally, the released products result in microvascular damage, capillary leak, and increased intracompartmental pressure, reduced tissue perfusion and ischemia
- Early complications: hyperkalemia, cardiac arrhythmia/arrest
- Late complications: acute renal failure and DIC
- **MRI is the method of choice in patients with clinical diagnosis of rhabdomyolysis to evaluate the distribution and extent of muscle lesions, especially when fasciotomy is considered for treatment; also to assess for muscle viability.**
35 yo M with limited neck extension
54-year old woman with type II diabetes mellitus presenting with acute thigh pain
3 months later
Diabetic Muscle Infarction

- Middle aged: mean 42 yo
- Poorly controlled or longstanding diabetes
- 77% have type 1 diabetes
- 94% have known microvascular diabetic complication (neuropathy, retinopathy, nephropathy)
- **ABRUPT ONSET OF SEVERE PAIN THAT OCCURS AT REST**
- THIGH muscles most commonly affected (80%)

Resnick. IDJ. 565-7.
Diabetic Muscle Infarction

- First described in 1965
- Caused by thrombosis of small and medium sized arterioles
- Occurs because of hypercoagulability and associated vascular endothelial damage
- Although histopathologic diagnosis is sometimes necessary, muscle infarction can often be diagnosed when the characteristic clinical presentation is combined with typical imaging findings

Resnick. IDJ. 565-7.
Diabetic Muscle Infarction

- Imaging DDx
  - Pyomyositis (insidious onset of symptoms; more than 1 month before clinical presentation)
  - Necrotizing fasciitis (fever, leukocytosis)
  - Non infectious myositis: focal myositis, proliferative myositis (biopsy is necessary in some cases)

Resnick. IDJ. 565-7.
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Fed “the washboard” Discepola
MYOSITIS

INFECTIONOUS ENTITIES AFFECTING MUSCLE:

– Pyomyositis
– Necrotizing fasciitis
55 yo F with 2 weeks lower extremity swelling, pain and cough
Diagnosis

- PET
  - Diffuse myositis or myonecrosis with secondary inflammation

- MR
  - Myositis with microabscesses in v. intermedius
  - Subacute hematomas in gastrocnemii

- US aspiration
  - R calf: S. aureus

- Bilateral Knee Aspirates
  - Elevated Neutrophils
  - S. aureus

- OR
  - Bilateral knee irrigation
  - Bilateral calf incision/irrigation
  - “cloudy fluid”

- BCx on admission: S. aureus

- CK < 25 (low)

- TTE: negative for veg
Final Diagnosis

Staph Aureus Bacteremia
Pyomyositis and Multiple intramuscular abscesses
Septic arthritis
Septic Emboli
Pyomyositis

• Skeletal muscle is particularly resistant to infection
  – Unless concurrent disease
    • Diabetes, HIV, chronic steroid use, connective tissue disorder, history of malignancy
    • Key elements in the development of PM are damaged muscle with varying degrees of immunosuppression and a source of bacteremia

• S. aureus is the most common agent (90%) in both tropical and temperate climates
  – Tends to occur in the large muscles of the lower extremities

12 yo M with knee and thigh pain and fevers
OPERATIVE PROCEDURE: 1. irrigation and debridement, right knee with arthrotomy.

2. irrigation and debridement, right femur (including bone).

FINDINGS: The knee was drained and approximately 5 cc of cloudy fluid was aspirated and was sent for culture. The dissection was carried out medially into the subvastus area and the abscess was entered. Purulent material was aspirated again, approximately 5 cc was aspirated although approximately 100 cc was drained and this was sent also for Gram stain and culture.

<table>
<thead>
<tr>
<th>Date</th>
<th>Time</th>
<th>Type of Sample</th>
<th>Culture Type</th>
<th>Gram Stain</th>
<th>Susceptibility Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>01/24/08</td>
<td>16:15</td>
<td>Fluid, Knee</td>
<td>Aerobic culture</td>
<td>Rare polymorphonuclear leukocytes seen. No organisms seen.</td>
<td></td>
</tr>
<tr>
<td>01/24/08</td>
<td>14:35</td>
<td>Fluid, Thigh, right</td>
<td>Aerobic culture</td>
<td>Abundant polymorphonuclear leukocytes seen. Moderate Gram Positive Cocci</td>
<td></td>
</tr>
<tr>
<td>01/24/08</td>
<td>00:18</td>
<td>Fluid, Knee, right</td>
<td>Aerobic culture</td>
<td>Rare Staphylococcus aureus</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>No evidence of inducible clindamycin resistance detected in this isolate of Staphylococcus aureus. Therefore, it can be considered susceptible to clindamycin.</td>
<td></td>
</tr>
<tr>
<td>01/23/08</td>
<td>22:10</td>
<td>Blood</td>
<td>Blood culture</td>
<td>Staphylococcus aureus in 1 of 2 bottles</td>
<td>For susceptibility results refer to accession 08:025-000650 on the Blood culture from 1/23/08.</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Test result called to Phoebe Lee, MD on 1/23/08 9:12:22 AM by C. Masadao.</td>
<td></td>
</tr>
</tbody>
</table>
Acute osteomyelitis
Subperiosteal abscess
Knee and hip septic arthritis
Myositis

• Taken back to OR 2 more times for I&D of knee and femur
45 yo M with hx of IV drug abuse
61 year old woman with diabetes

NECROTIZING FASCIITIS:
Surgical Emergency
Fascial Biopsy is the gold standard for diagnosis

Idiopathic Inflammatory Myopathies

- Unknown causes
- Characterized by nonsuppurative inflammation of muscle
  - Dermatomyositis
  - Polymyositis
  - Juvenile dermatomyositis
  - Inclusion body myositis
  - Focal Myositis
  - Idiopathic eosinophilic Fasciitis
40 yo F who recently underwent abdominal surgery
Dermatomyositis

• Hallmark: weakness and inflammatory infiltrates in proximal muscles
• Extent of calcification relates to severity of illness
• Intermuscular or fascial calcification is characteristic
• Median age at diagnosis is 51 yo
• Significant FEMALE predominance
• More likely than polymyositis to affect organ systems other than muscle:
  – Dysphagia
  – Pulmonary fibrosis
  – Cancer
27 yo F with generalized malaise
60 yo M with diffuse muscle pain and generalized weakness
FINDINGS

There is severe multifocal myositis characterized by muscle edema in a bilateral, patchy and slightly asymmetric fashion, with normal muscle bulk and no fatty atrophy. Virtually the entire pelvic musculature is involved. In the thighs, there is relative sparing of the rectus femoris and adductor muscles. The most inflammation is present in the vastus lateralis bilaterally, which would be a potential biopsy site with a high diagnostic yield. In the upper extremities, there is myositis of the pectoralis and latissimus dorsi muscles bilaterally as well as the serratus muscles and rotator cuff and deltoid muscles. There is a small amount of involvement of the brachialis muscles bilaterally and sparing of the biceps and triceps muscles.

There is mild skin edema overlying the vastus lateralis bilaterally.

The pattern of involvement is typical for the active phase of dermatomyositis or polymyositis.

In addition, there is evidence of hemosiderosis with intensely low signal intensity on the T2-weighted and STIR images within the lower lumbar spine, sacrum, and both proximal femora as well as the pelvis. Less hemosiderin deposition is seen in the humeri.

IMPRESSION:

1. Extensive active myositis of the pelvis, thighs, and both shoulder girdles in a pattern of dermatomyositis or polymyositis. Severe inflammation without fatty atrophy is present in the vastus lateralis muscles bilaterally, which if biopsy, would likely be of high diagnostic yield. 2. Marrow changes of hemosiderosis.
Muscle Biopsy

DIAGNOSIS:
SKELETAL MUSCLE, LEFT QUADRICEPS, BIOPSY
- MYOSITIS WITH PERIFASCICULAR ATROPHY (SEE COMMENT)

rxp/11/24/2006 17:00      By this signature, I attest that the above diagnosis is based upon my personal examination of the slides (and/or other material indicated in the diagnosis).

Robert E. Schmidt, M.D., Ph.D.
***Report Electronically Reviewed and Signed Out By Robert E. Schmidt, M.D., Ph.D.***

Microscopic Description and Comment:
Microscopic examination of the left quadriceps muscle biopsy material shows two pieces of striated muscle tissue with perifascicular atrophy and a patchy endomysial lymphoplasmacytic infiltrate with a minor eosinophil component. In addition to atrophy, a subset of the perifascicular fibers exhibits basophilia and plump nuclei with relatively open chromatin and prominent nucleoli, consistent with regeneration. There is no evidence of increase in the amount of endomysial/interstitial connective tissue, group atrophy, or angionecrosis. This pattern is that of a myositis.
4 months post-treatment

Polymyositis
How we can help

• MRI displays areas with the most edema and the least atrophy

• With MR imaging directing the site of biopsy, the sensitivity is 97%

• Accurate assessment of disease activity with serial studies is important when selecting and adjusting the effective dosage of steroids and other immunosuppressives
55 yo M with bilateral thigh muscle pain
Diagnosis

• Radiology IMPRESSION:
  – Eosinophilic fasciitis
  – Fascia superficial to vastus lateralis amenable to biopsy

• Labs
  – Absolute eosinophil count: 2.7 (nl 0.0 – 0.5)

• Surgical Biopsy:
  – Eosinophilic fasciitis
  – “mixed inflammatory infiltrate with readily identifiable eosinophils as well as lymphocytes and some plasma cells. The stromal background has a fibromyxoid appearance to indicate the involvement of the deep connective tissues.”
Eosinophilic Fasciitis

- Eosinophilic infiltration of muscle and fascia
- Late findings include fibrosis in the fascia that results in joint contractures
- No known cause for the persistent eosinophilia
- Treatment: STEROIDS
HEREDITARY DISORDERS OF HETEROTOPIC OSSIFICATION
51 yo F with shoulder pain
Myositis Ossificans Progressiva

• Since the 1800s—references in medicine describing people who “turn to stone”

• calcification/ossification of subcutaneous fat, skeletal muscle, tendons, aponeuroses, and ligaments.
  – Soft tissue masses coalesce leading to formation of “bony bridges” which cause restriction of respiration and ambulation and skeletal contractures

• Assoc with symmetric malformation of the digits, esp thumbs and great toes
Myositis Ossificans Progressiva

• No known treatment
  – Attempts at resection of heterotopic bone are considered futile because the soft tissue trauma induced by surgery can stimulate recurrent heterotopic ossification.

• Prognosis
  – Pts often develop compromised pulmonary and cardiac function, and premature death may result from respiratory failure caused by ankylosis of the thoracic cage.
HERITABLE DISORDERS AFFECTING MUSCLES

• Muscular Dystrophies
  – Heterogenous group of heritable disorders characterized by progressive muscle weakness and loss of muscle tissue
    • Most common MD is Duchenne
  – Muscles are edematous initially and then rapidly become atrophic
Normal

45 yo F with long standing MD
Muscular Dystrophies: Future MR Applications

• Potential for MRI in diagnosis of MD
  – Biopsy planning, limiting false negative biopsies
  – Distinguishing conditions with similar clinical phenotypes
  – To assess if muscle grossly normal or abnormal in cases of confusing clinical presentations

• Potential for MRI in management of MD
  – Marker for disease progression
  – Marker for response to therapy

Timberlake, J. Muscular Dystrophies: What the radiologist should know. URMC 2006.
• 31 yo F with diminutive gluteal musculature bilaterally
• i.e. “lacking junk in the trunk”
You’d just have to give up:
**Eating
**Drinking
**Buying new underwear
**Going to the movies
**Getting health insurance
**Buying bagels and coffee for Saturday morning conference
**Parking at Hillcrest and Thornton
THANK YOU FOR AN AMAZING YEAR!

- Drs. Chen, Chung, Fliszar, Gentili, Hiller, Hughes, Pathria, Resnick
- My co-fellows and research fellows
Overview of Imaging Techniques

- Radiography
- CT
- Sonography
- Scintigraphy
- MR imaging
Indications for MR imaging of Muscle

- To provide prompt diagnosis for initiating proper treatment
- To evaluate a soft tissue mass in a patient without a clear history of trauma
- To assess uncommon causes of muscle pain
- To investigate for an underlying structural cause of neuropathy
- To assess the extent and type of infection
- To evaluate the location and extent of myopathy, especially before biopsy
Causes of T1 Hyperintensity in Muscle

• Common causes
  – Fat deposition
  – Hematoma
  – Gadolinium contrast
  – MR artifacts

• Uncommon causes
  – Proteinaceous material
  – Melanin
Common causes of T2 Hyperintensity in Muscle

- Muscular exertion
- Trauma
- Vascular insufficiency
- Inflammation-infection
- Inflammation-autoimmune
- Subacute denervation
- Iatrogenic
- Infiltrative neoplasm
- Muscle cell death

IDJ Resnick
Causes of T2 hypointensity in muscle

- Calcification
- Foreign bodies
- Hemosiderin (old hemorrhage)
- MR artifact (flowing blood in a vessel)