



Case Presentation

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Introduction



68 years old veteran with a past medical history of hypertension, type II diabetes mellitus, hyperlipidemia, multiple myeloma, and basal cell carcinoma.



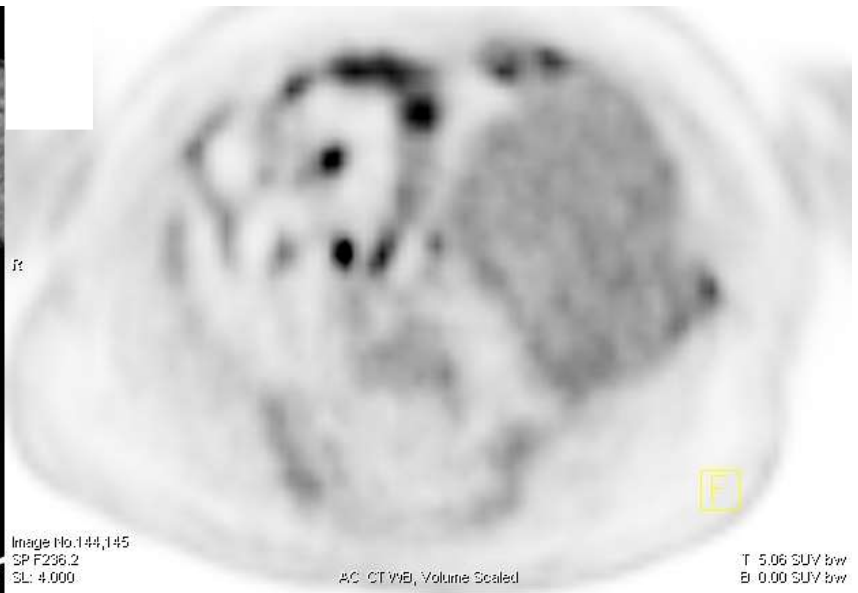
Patient complaining of weight loss, abdominal pain and constipation.




PET/CT performed to evaluate the disease course of multiple myeloma.



T: 28%
B: 1%







What is your
differential
diagnosis?

PATHOLOGY REPORT

Accession No. SP 17 7118

DIAGNOSIS

- Part # 1 - Left pelvic mass, CT-guided core needle biopsy:
 - Amorphous eosinophilic material suggestive of amyloid, pending further characterization.
 - Comment: Results of Congo Red stain will be interpreted in a supplemental report when available.

PATHOLOGY REPORT

Accession No. SP 17

1546

Gross description:

The specimen is received in formalin labeled with patient name, social security number and further designated on the container and the requisition as "left pelvic mass". The specimen consists of multiple needle core biopsies measuring 0.1-1.2 cm in length and 0.1 cm in diameter. Specimen is wrapped in lens paper and entirely submitted in cassette 1A.

Also received are four labeled, stained slide(s) made at the time of specimen collection.

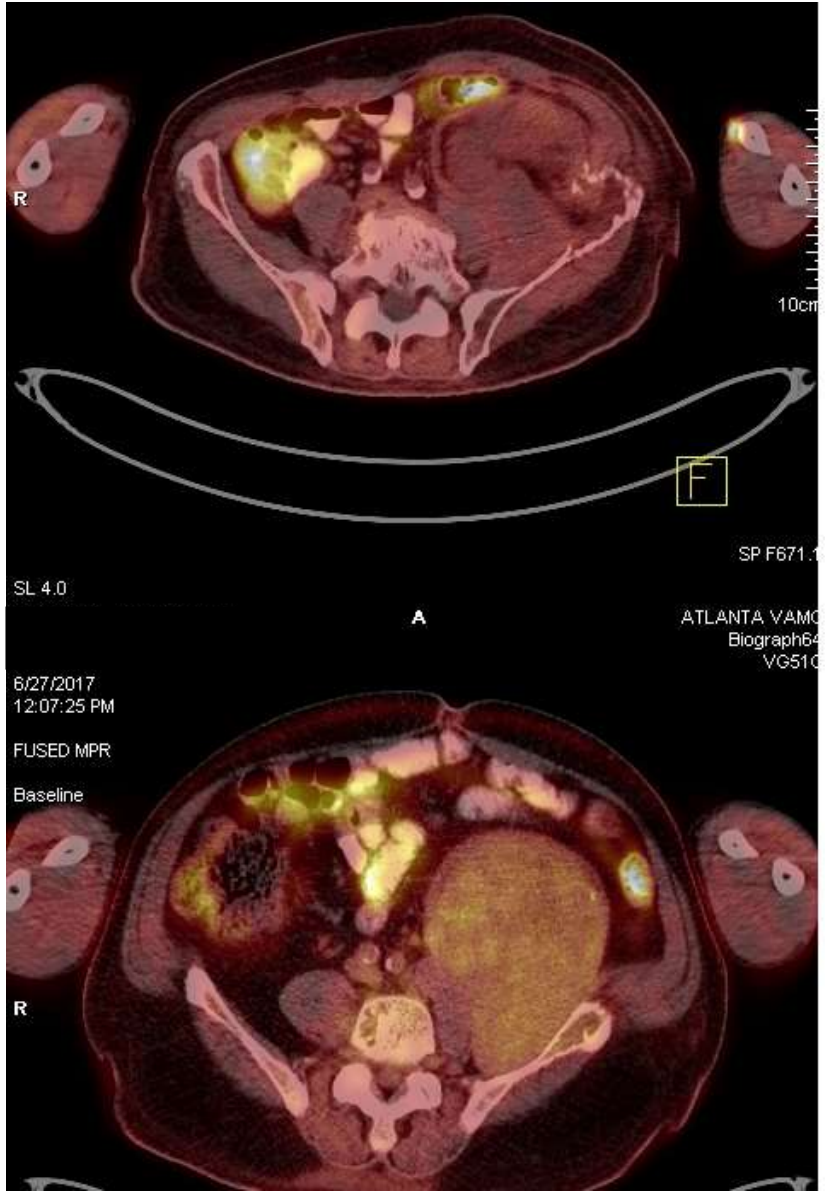
DIAGNOSIS

- Part # 1 - Left pelvic mass, biopsy:
 - Congo Red-positive material suggestive of amyloid.
 - Associated foreign body-type giant cell reaction, plasma cells, and lymphoid cells.

Follow-up

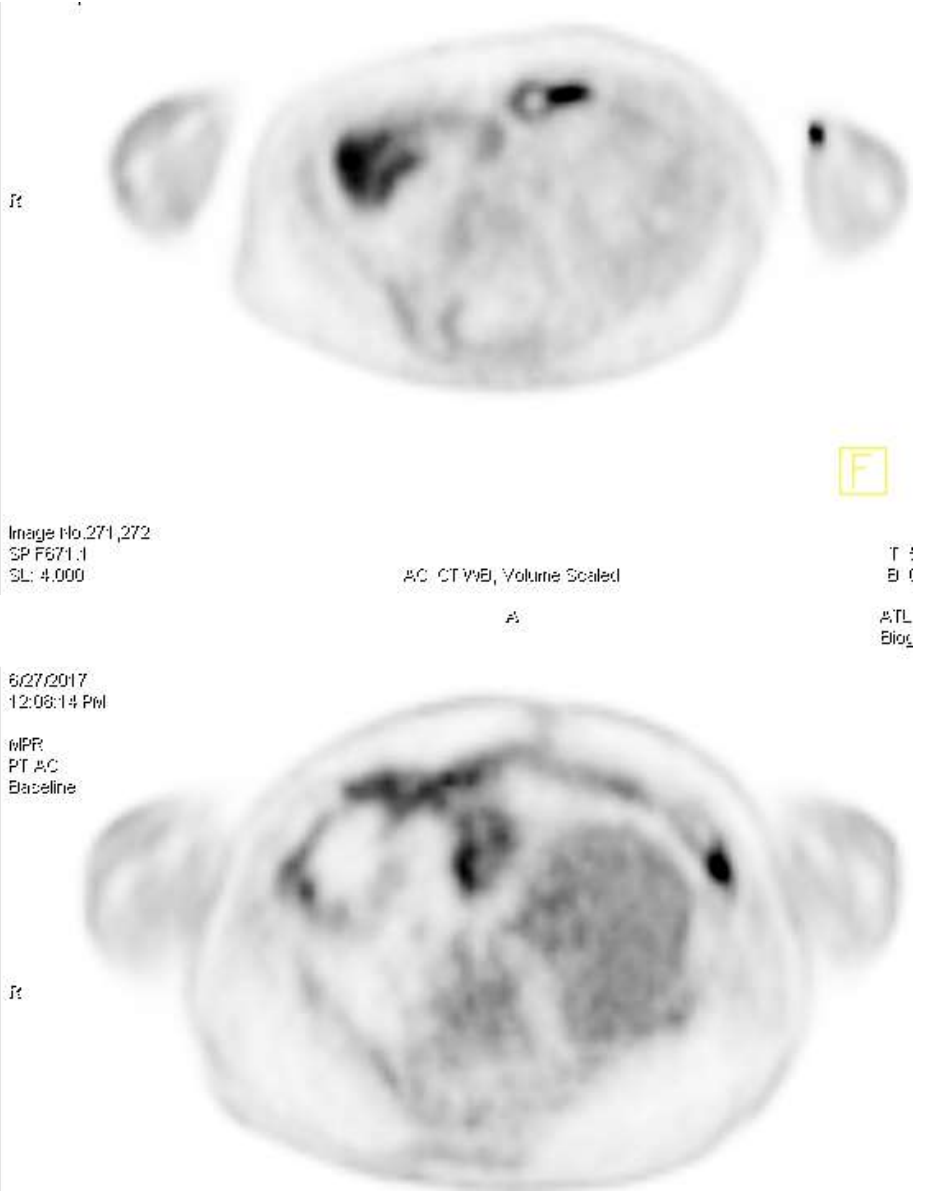
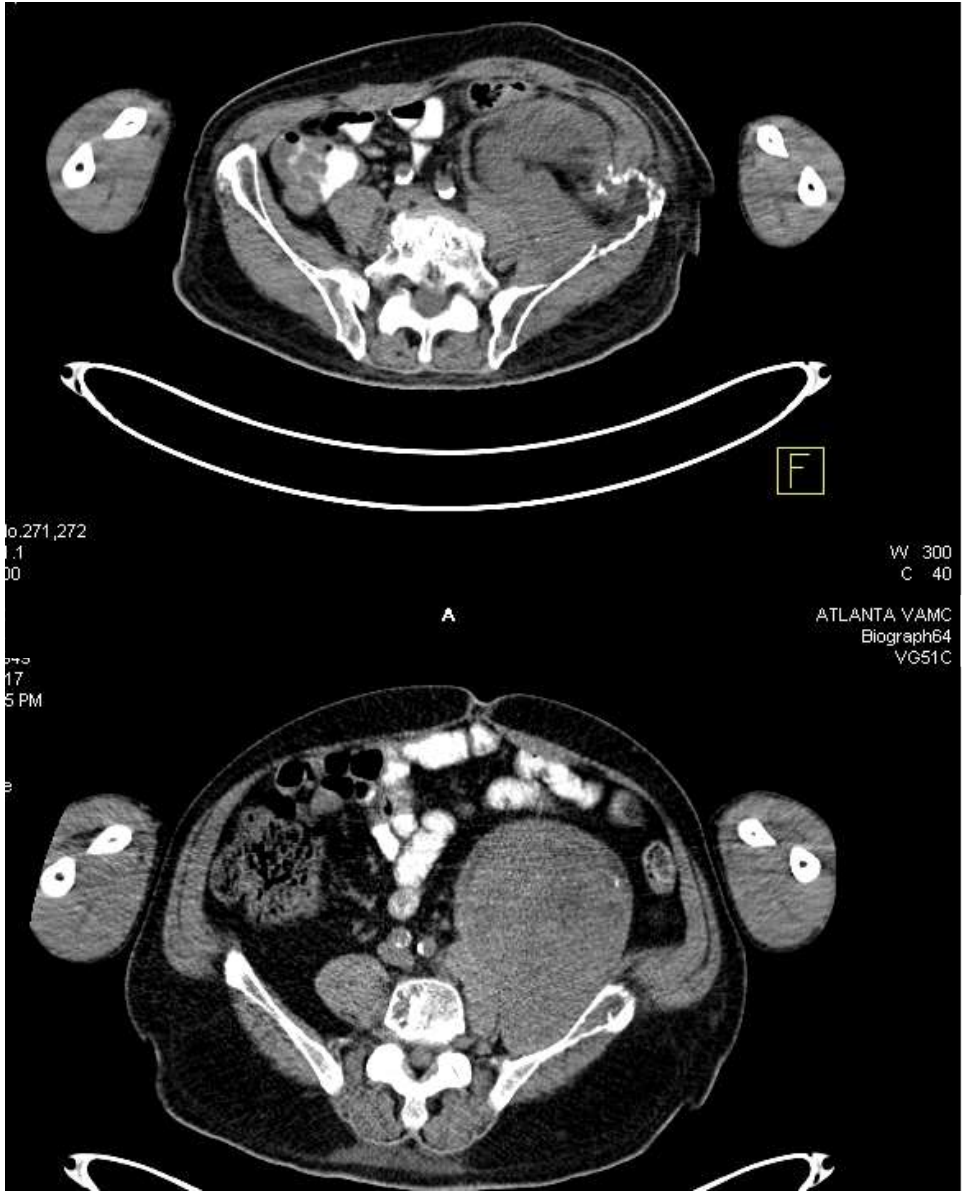
Patient received radiotherapy.

Followed by
Revlimid/Velcade/Dexamethasone



T: 28%
B: 1%






Amyloidoma

Defined as a solitary localized tumor-like deposit of amyloid, in the absence of systemic amyloidosis.

Is the least common presentation of amyloidosis.

Characterized by extracellular deposits of abnormally folded amorphous, insoluble and proteinaceous material that interferes with tissue function.



Amyloidoma of soft tissues is extremely rare and occurs mainly in the mediastinum and abdomen.

Associated with multiple myeloma, lymphoma, and other lymphoproliferative disease.

Some other cases of amyloidoma have been associated with local trauma, surgery, infection, or peripheral vascular disease, and a few patients have been diabetic.

Management

Primary localized amyloidoma possesses a relatively good prognosis.

Surgical debulking for limited disease.

Radiotherapy is used to decrease the size of bone-occupying lesions.

Aggressive chemotherapy with underlying lymphoproliferative disorders.