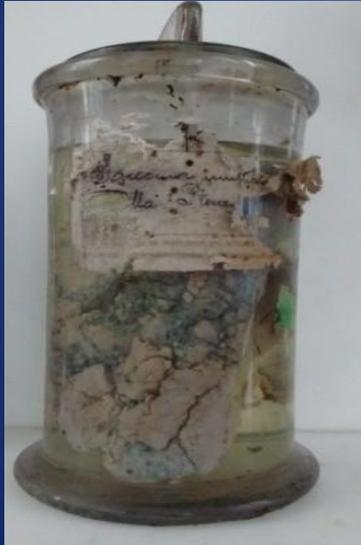




31st European Congress
of Pathology
Pathology is Nice
7–11 September 2019, Nice, France



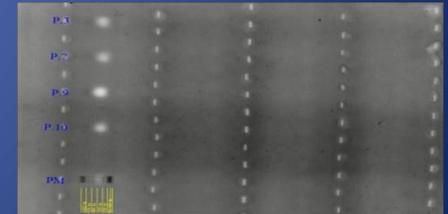
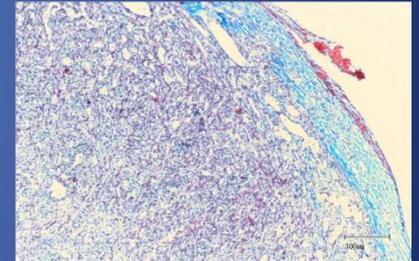
Histopathology and DNA evaluation of wet specimens from the Pathology Collection of Turin



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Salvatore Hospital L'Aquila

The authors declare no conflict of interest



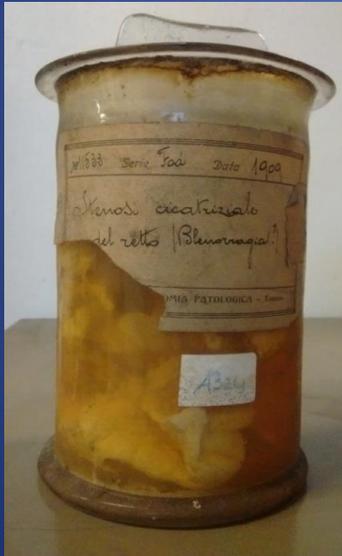
**The Pathology Collection of Turin houses
around 300 wet specimens dating back to
XIX-XX century**



**Most of these specimens are in their original jars
with labels describing year, necropsy number
and original old diagnosis.**



Due to their original conditions these specimens
are an actual
biological archive
and may represent a valid source for research on
molecular features of ancient diseases.



Four cases originally diagnosed as

lung cancer

uterine myosarcoma

pleural sarcomas (2)

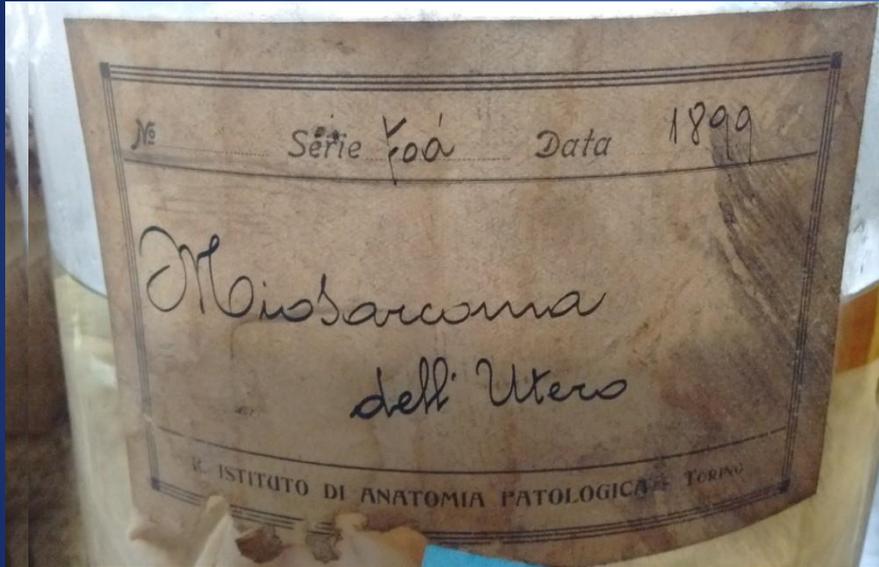
underwent modern diagnostic revision and

DNA evaluation by conservative sampling.

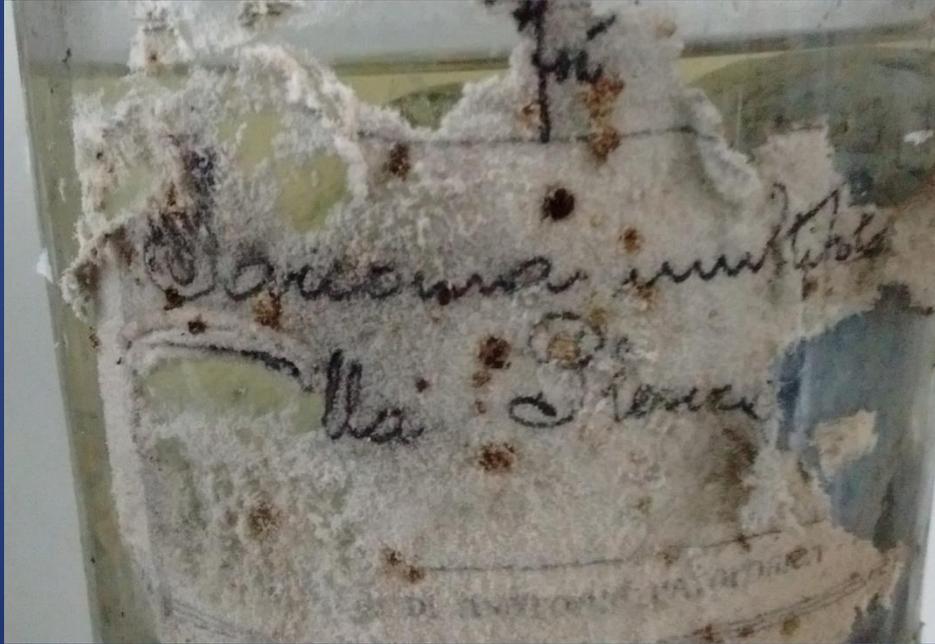
Case 1: Lung cancer



Case 2: Uterine Myosarcoma



Case 3: Pleural sarcoma (1896)



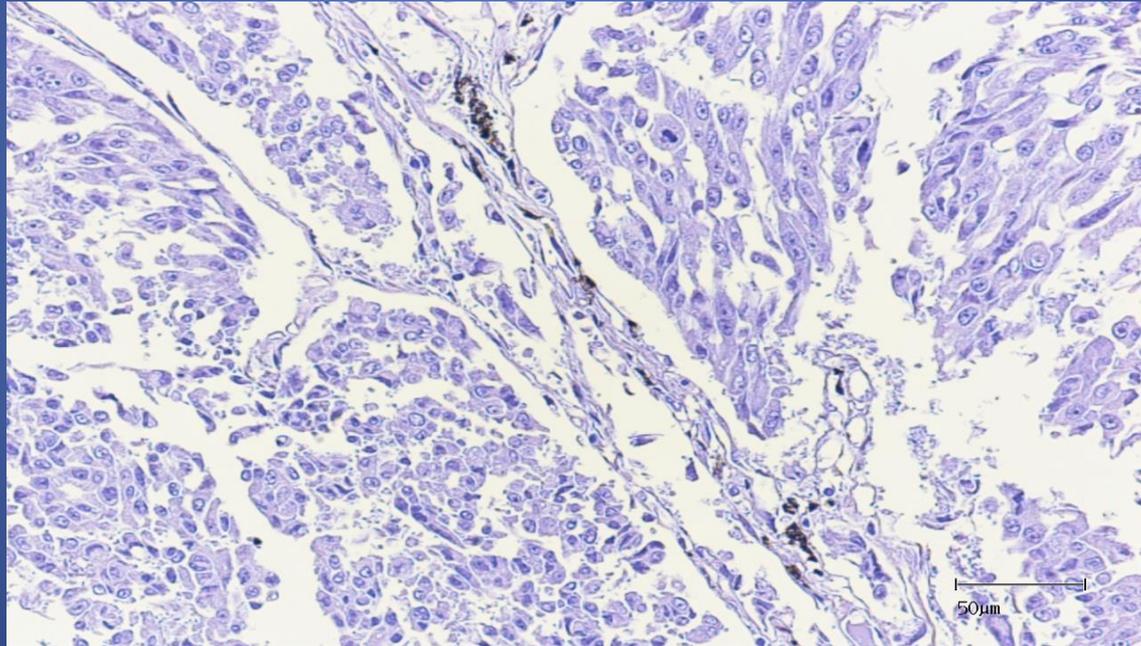
Case 4: Pleural sarcoma(1898)



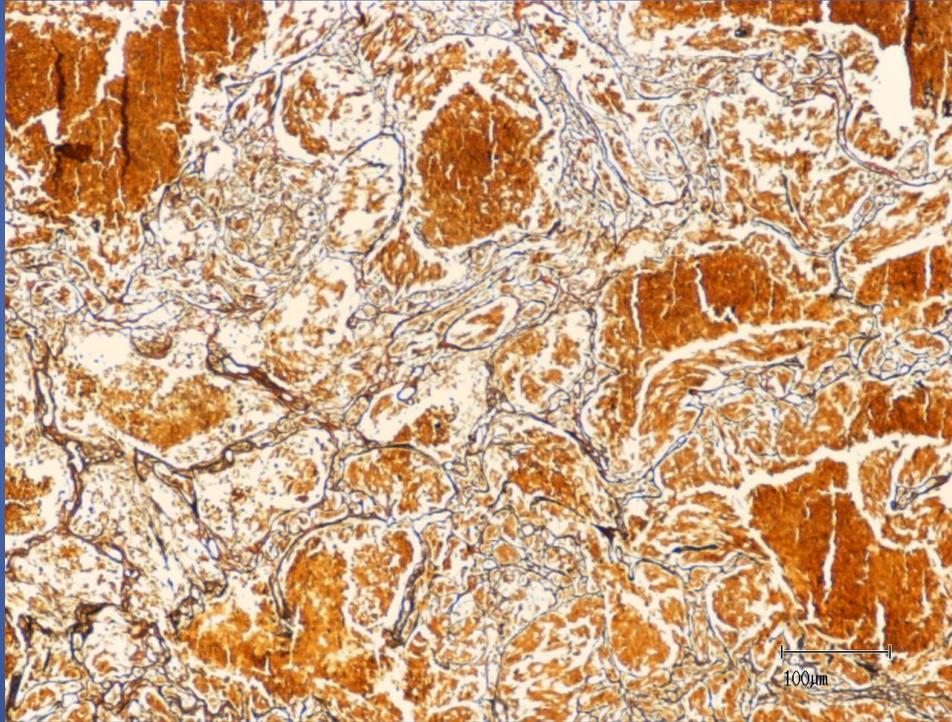
The revised diagnoses were:

- necrotic lung carcinoma**
- uterine leiomyosarcoma**
- lung metastases from squamous carcinoma of unknown primary**
- lung metastases from uterine leiomyosarcoma**

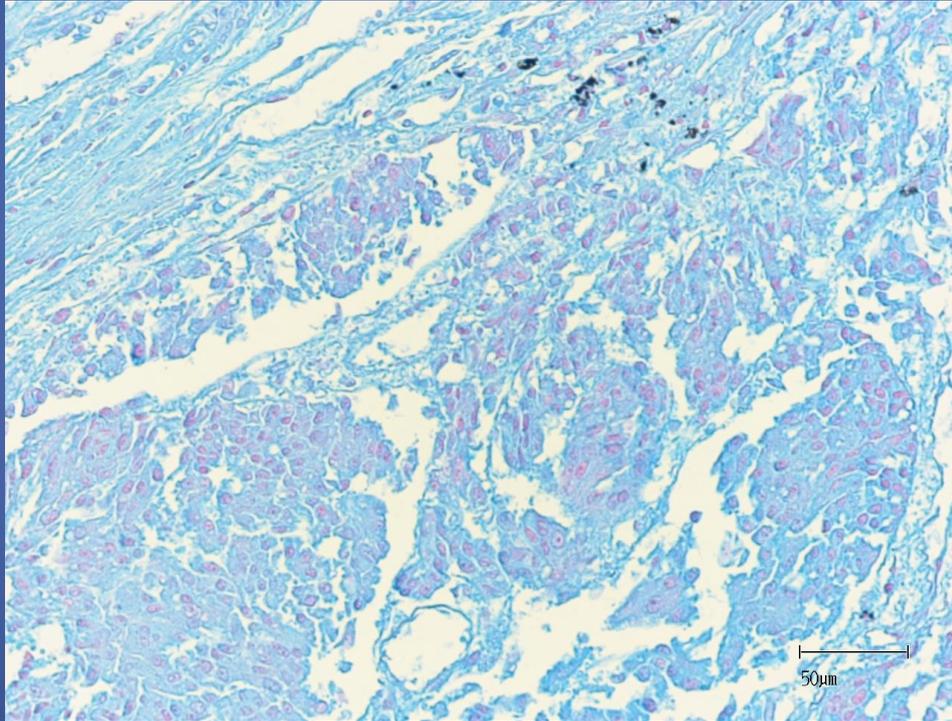
Lung metastases of squamous carcinoma of unknown primary



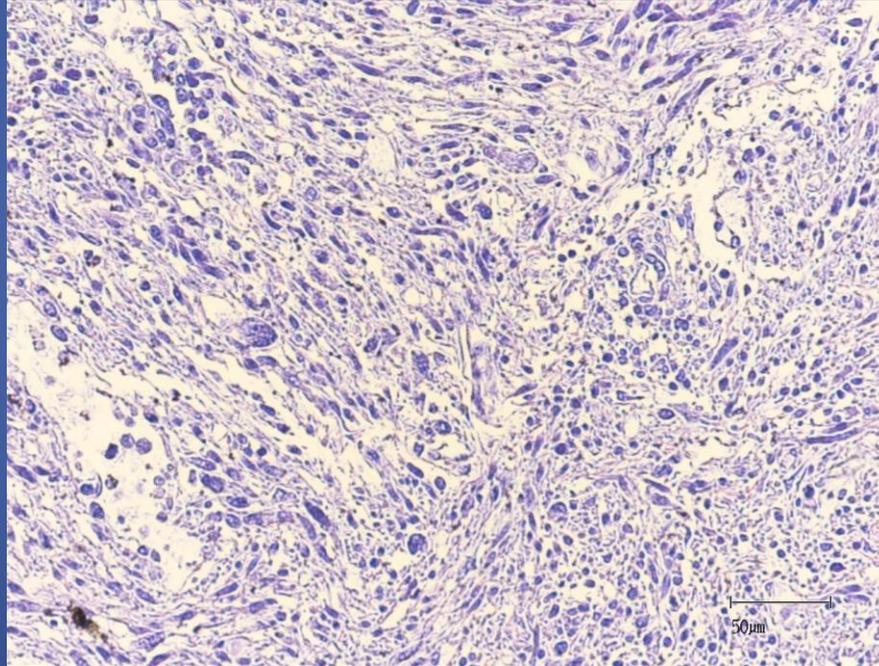
Reticular fibers staining



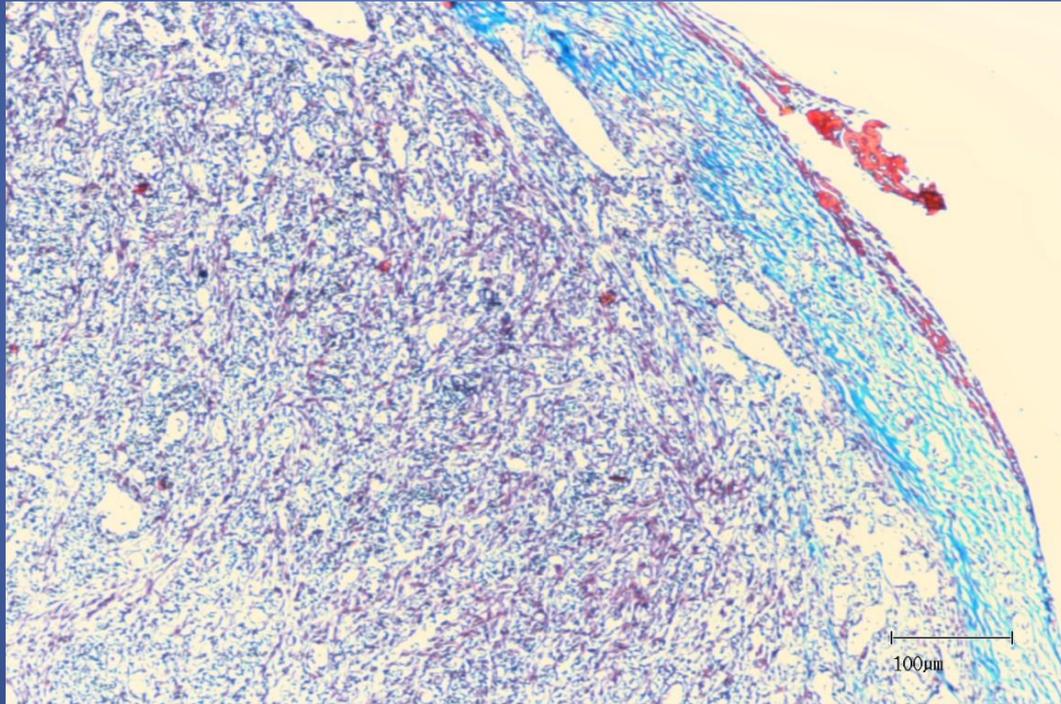
Alcian blu staining



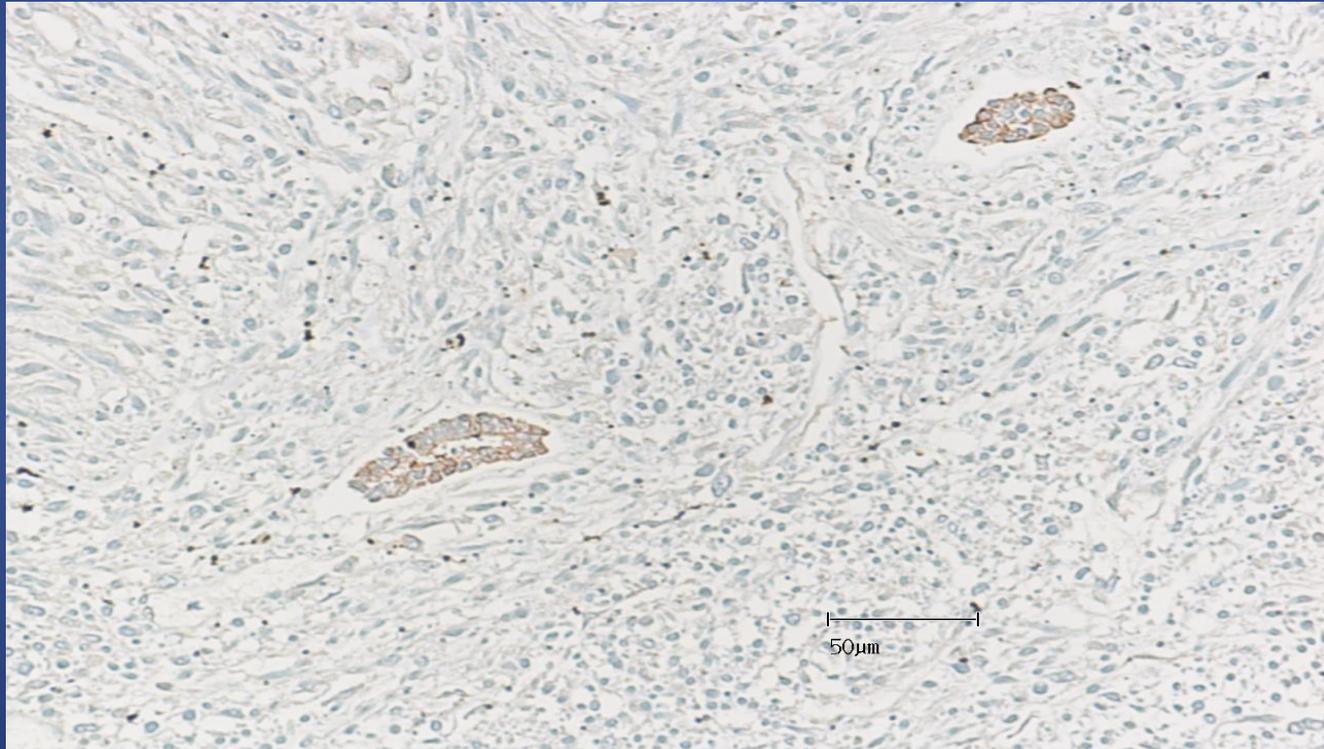
Lung metastases from uterine leiomyosarcoma



Trichrome stain



Cytokeratin MNF116



**Additional tiny samples underwent
DNA extraction and analysis by
spectrophotometry and
electrophoretic run in agarose gel.**

**As the chemical composition of the
storage fluids is unknown,
pH value
was measured in each specimen.**

pH values were

- **2.56** necrotic lung carcinoma
- **3.15** uterine leiomyosarcoma
- **4.45** lung metastases from squamous carcinoma of unknown primary
- **4.65** lung metastases from uterine leiomyosarcoma

The samples were taken by conservative method and cryostatic sections (-20 degrees C) were obtained 10 microns thick.

**Digestion of the section with solution of 75 mM NaCl, 10 mM tris, 0.5 mM EDTA to pH 8.0 and 100 ml of K Proteinase solution.
(18 mg/ml)**

**The samples were incubated at 56 degrees C
for 48 hours**

**50 ml of fresh K Proteinasi solution was
added for 72 hours**

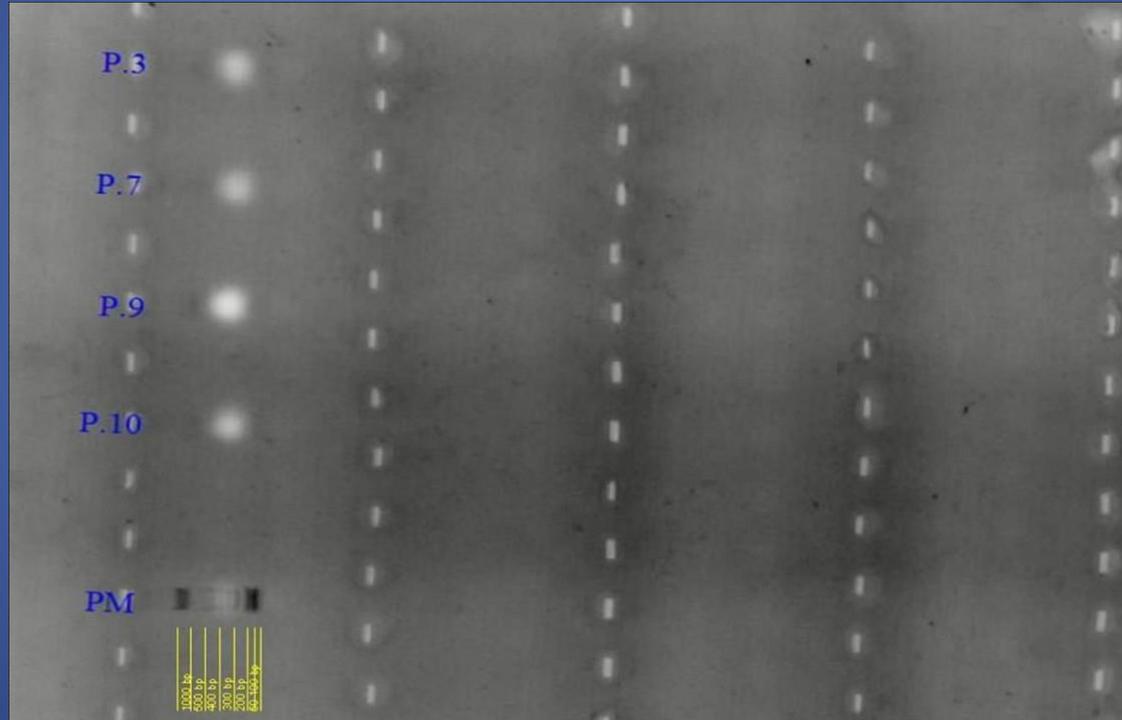
**400 ml of solution was then extracted
(magnetic beads -Roche MAGNA PURE
COMPACT instrumentation)**

DNA quantity were assessed using the entire absorption spectrum (220/340 nm) obtained from the Nanophotometer P 300 spectrophotometer.

The concentration of DNA in ng/ml and absorption at 260/280 nm were evaluated on 4 ml samples.

Gel electrophoresis

agarose at 1.3%



RESULTS

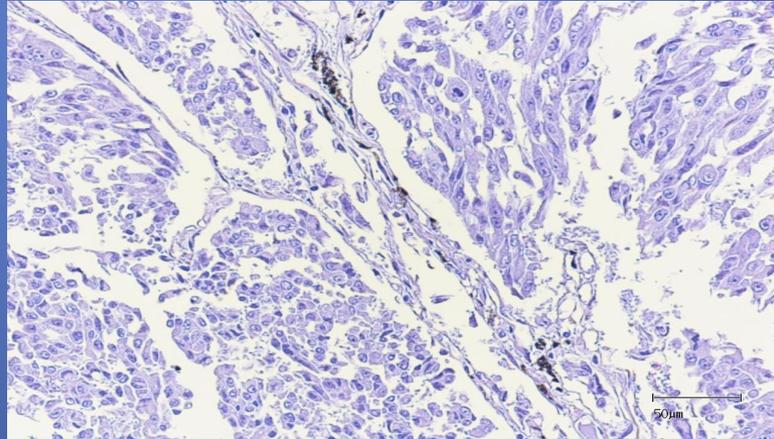
The first two samples gave negative results on both spectrophotometric and electrophoretic analysis



Case 3 – pH 4.45

spectrophotometric analysis absorbance 1.53 6ng/ml

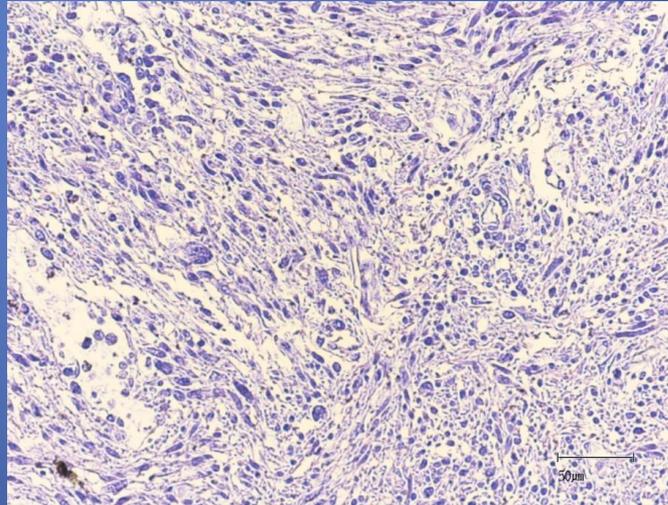
Electrophoretic analysis DNA band with molecular weight about 400bp



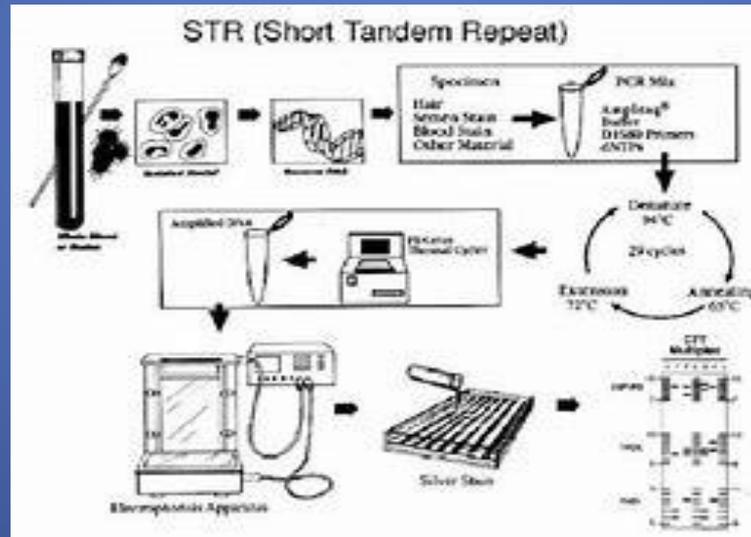
Caso 4 - pH 4.65

Spectrophotometric analysis absorbance 1.50 7ng/ml

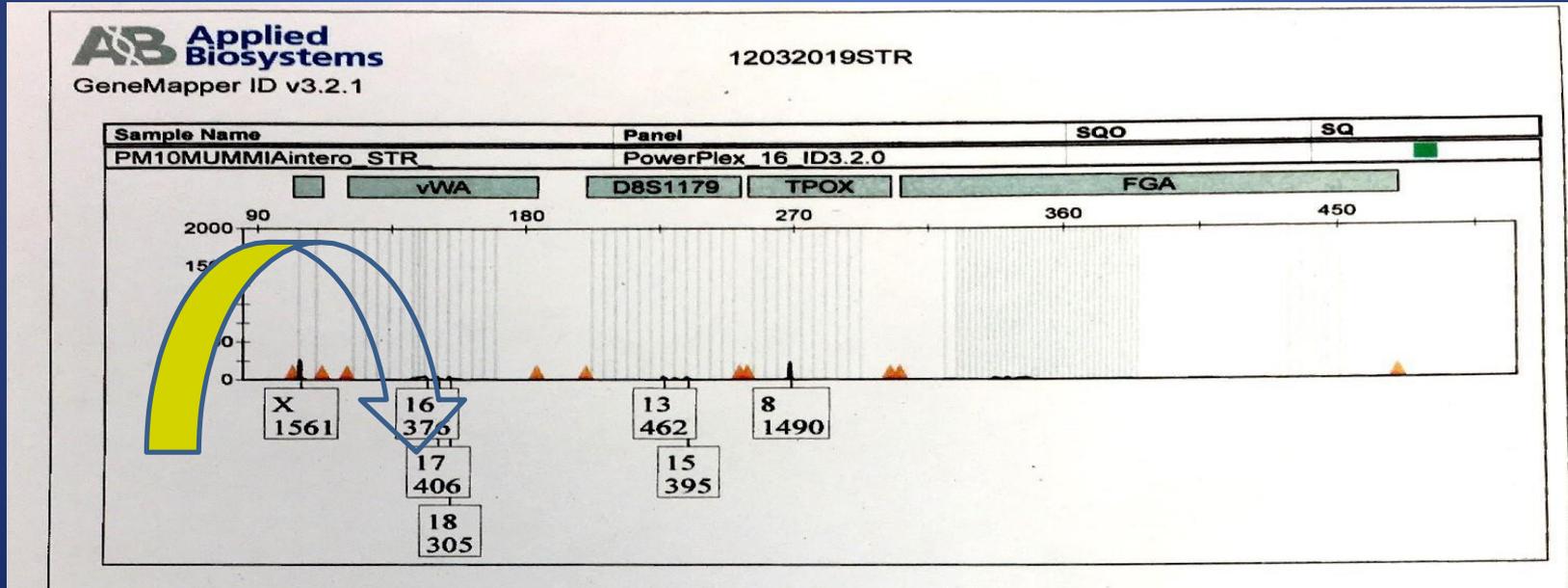
Electrophoretic analysis DNA band with molecular weight about 500bp



To assess DNA integrity,
short tandem repeat (STR) analysis
with PowerPlex 16 HS (PROMEGA) panel
employed for personal identification was used



The amplification of amelogenin STRs of chromosome X demonstrates the probable presence of a female subject in case 3



CONCLUSIONS

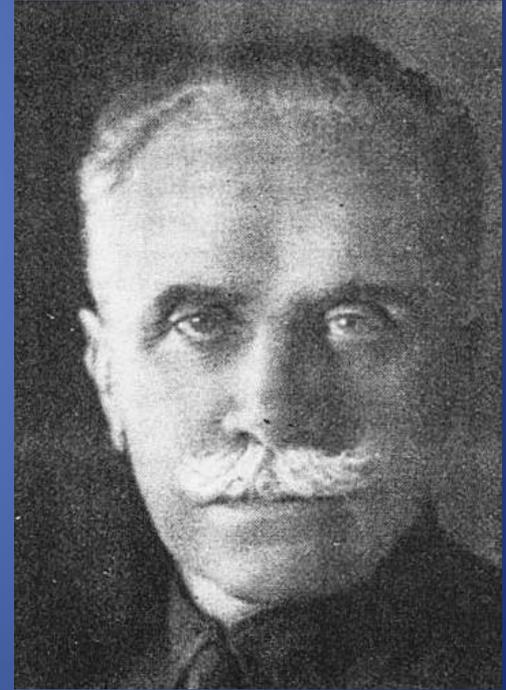
This study confirms the importance of
Pathology Collections as a

historical archive of

diseases that no longer exist
or with natural
course unmodified
and also as

biological archive

Pathology Museums must be preserved as a precious historical and biological heritage!!



Thanks for your attention!

