

PRINCIPAIS FIXADORES, SUA IMPORTÂNCIA E ALGUMAS TÉCNICAS DE COLORAÇÕES ESPECIAIS

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FIXAÇÃO

A Fixação é a maneira de preservar as células e manter partes de suas estruturas que tinham no organismo vivo

FIXADOR UNIVERSAL– FORMALINA TAMPONADA

- Água destilada900 ml
- Formol 37% a 40%.....100 ml
- Fosfato de sódio Monobásico.....4,0 g
- Fosfato de sódio Bibásico..... 6,5 g

Volume do Líquido Fixador deverá ser no mínimo 20 vezes maior que o tamanho da peça

- Tempo ideal de Fixação de 01 a 24 horas
- PH ideal para boa Fixação – 7,0 ou PH neutro
- A FORMALINA TAMPONADA não precipita as proteínas, ótimo para os Lipídios, mantem o Glicogênio
- A FORMALINA **não** deve ser usada com cromatos, porque oxida-se facilmente

- A FORMALINA sem Tampão oxida, produzindo ácido fórmico-formando um composto FORMALINA hematina.
- Álcool como fixador, preserva o Glicogênio, altera o detalhe nuclear e comprime o Citoplasma.
- ACETONA- Boa para estudos histoquímicos, preserva Enzimas:
LIPASES / FOSFOTASES

LÍQUIDO DE BOUIN

- Água destilada 300 ml
- Formol 37 a 40% 100 ml
- Ácido Acético Glacial 20 ml
- Ácido Pícrico a Saturação..... 1,5g

Fixador rápido: até 24 horas

Desvantagens: Precipita as proteínas – perda da basofilia

FIXADOR DUBOSCQ - BRASIL

- Álcool 80% 150 ml
- Formol 37 a 40% 60 ml
- Ácido Acético Glacial 15 ml
- Ácido Pícrico 1,5 g

Fixador rápido: Fixa em 02 a 12 horas

Muito usado em biópsias renais e de testículos.





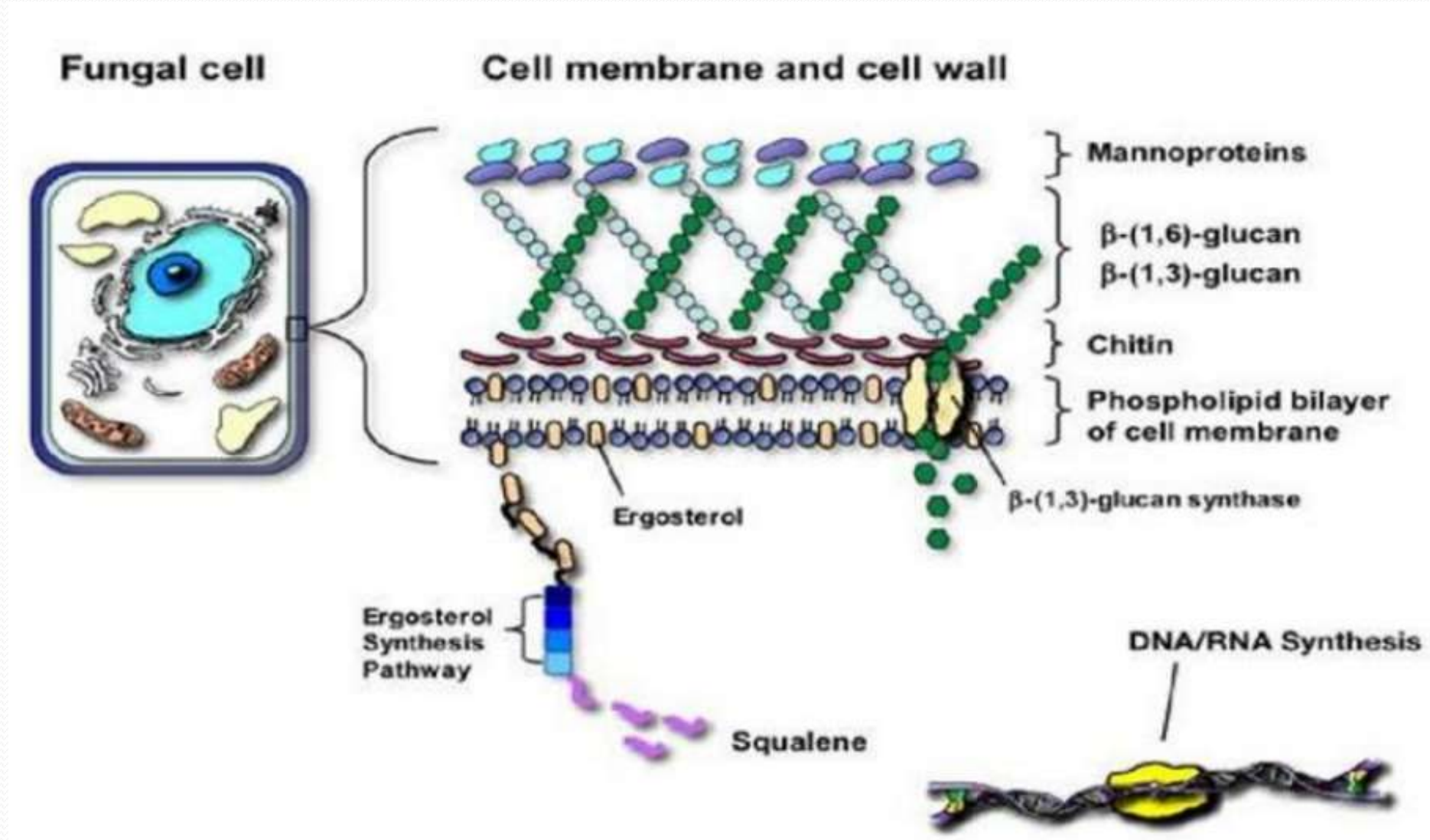




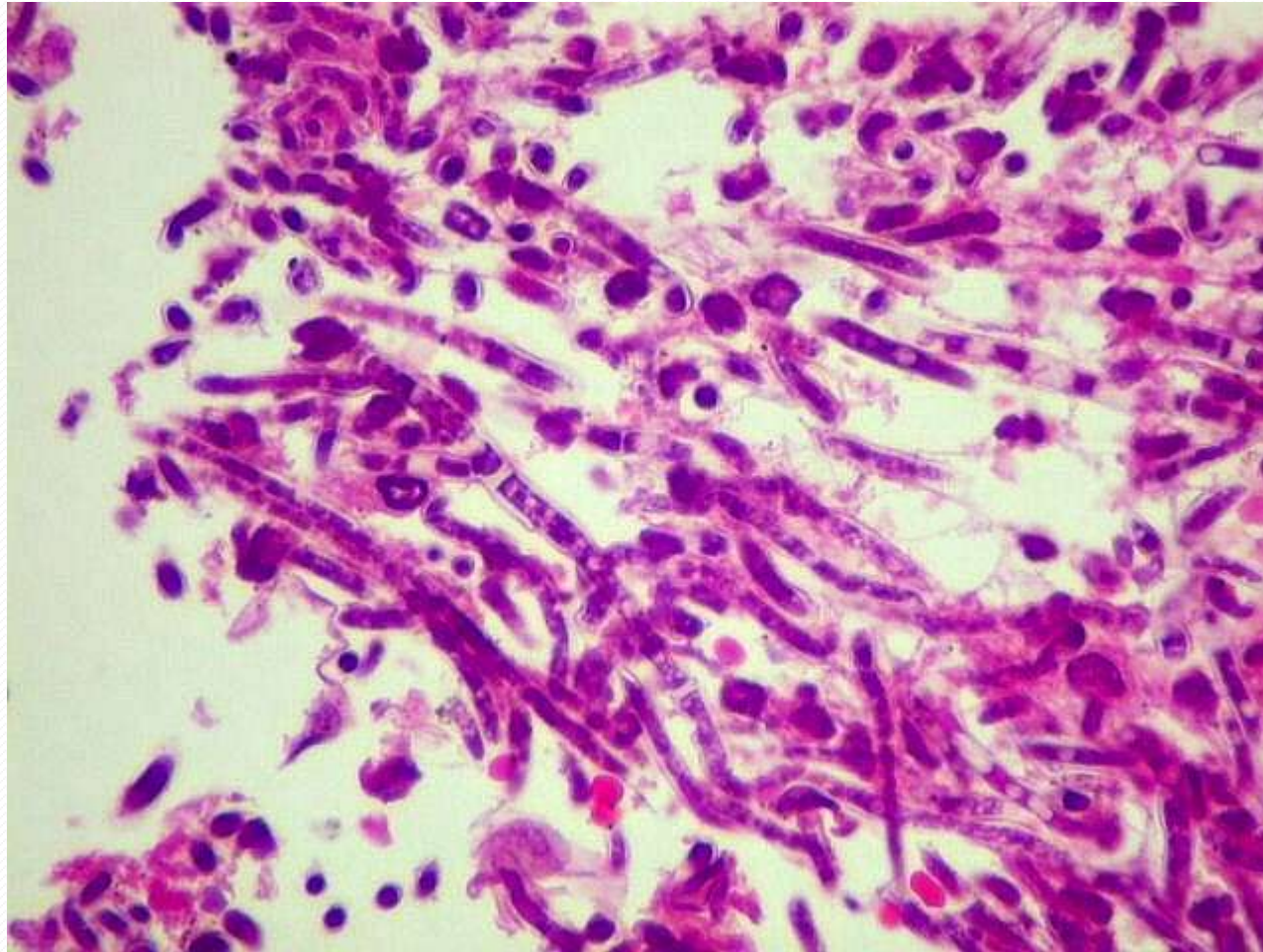
MÉTODO DE GOMORI - GROCOTT – COM METENAMINA - ARGÊNICA

- Membranas basais - pela redução da prata
- Paredes celulares - pela oxidação do ácido – crômico
- Composição das paredes celulares dos fungos – Quitina / Proteínas e Lipídeos
- Reação do ácido crômico com polissacarídeos

PAREDE E MEMBRANA DOS FUNGOS

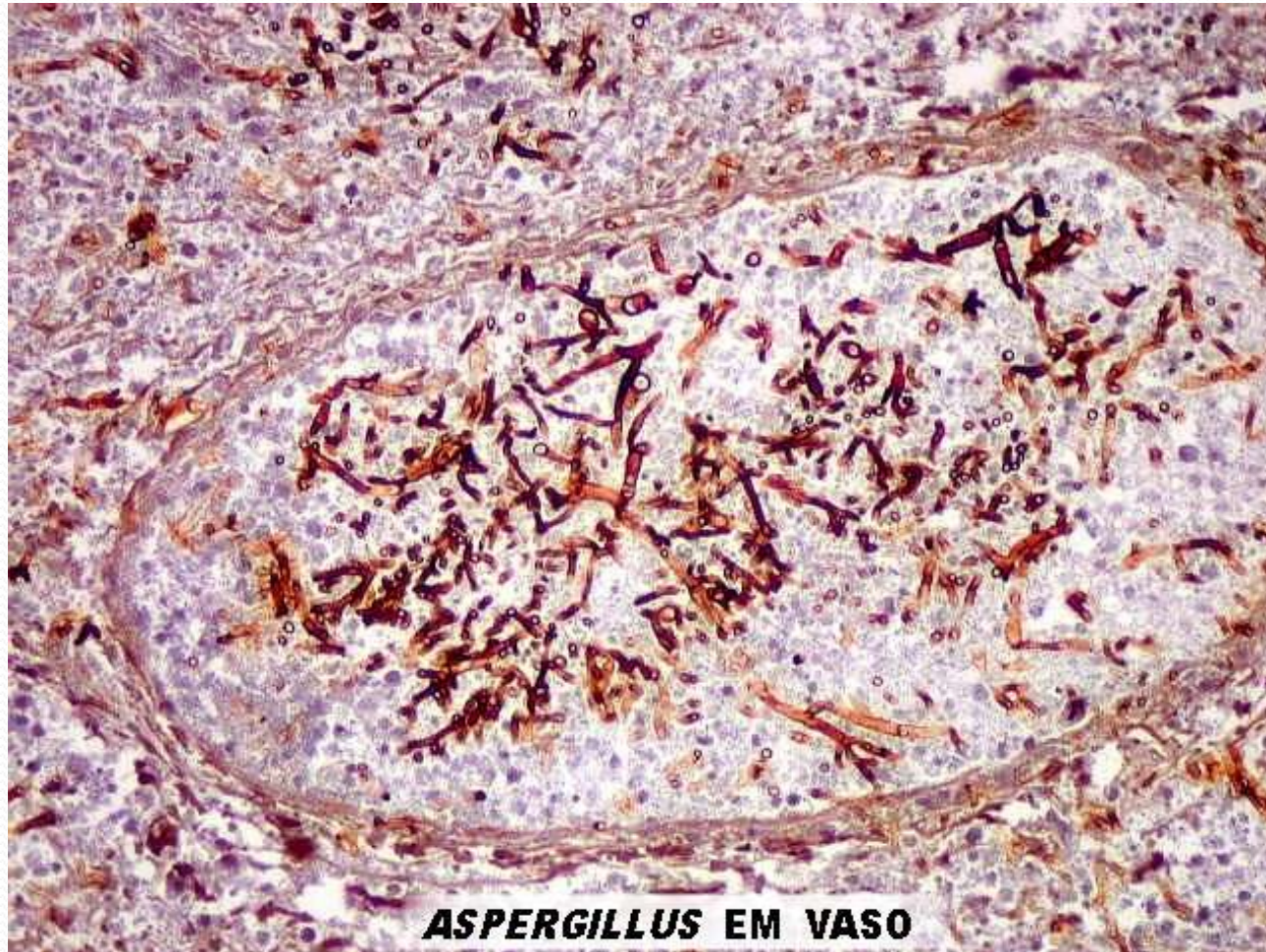


COLORAÇÃO DE HEMATOXILINA/EOSINA



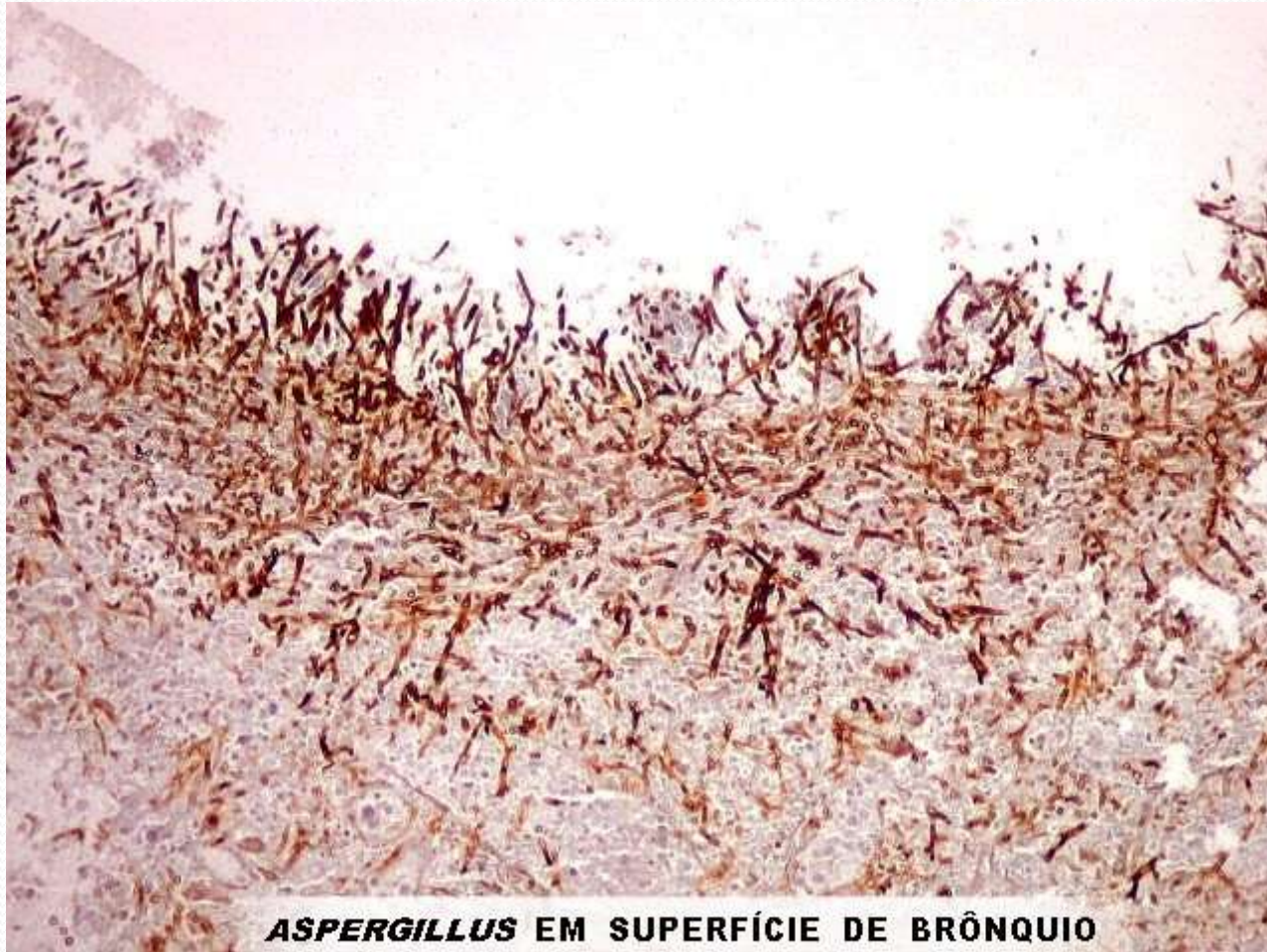
Hifas de *Aspergillus*

GROCOTT SEM COLORAÇÃO DE FUNDO



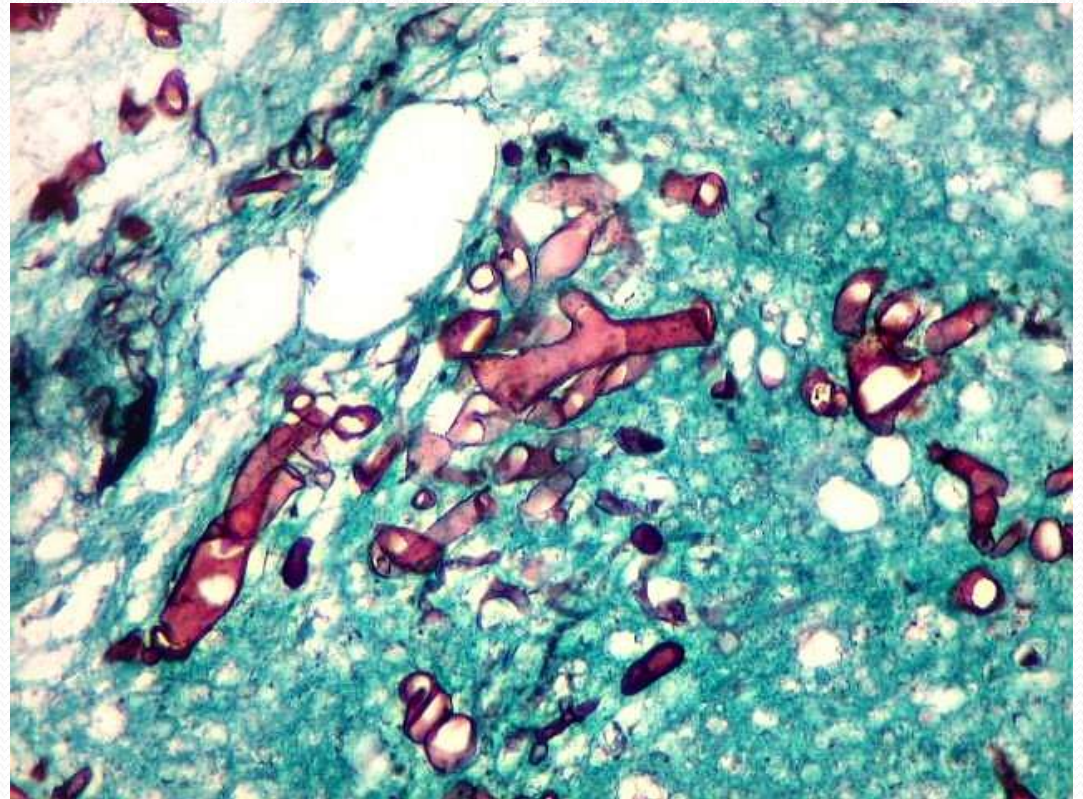
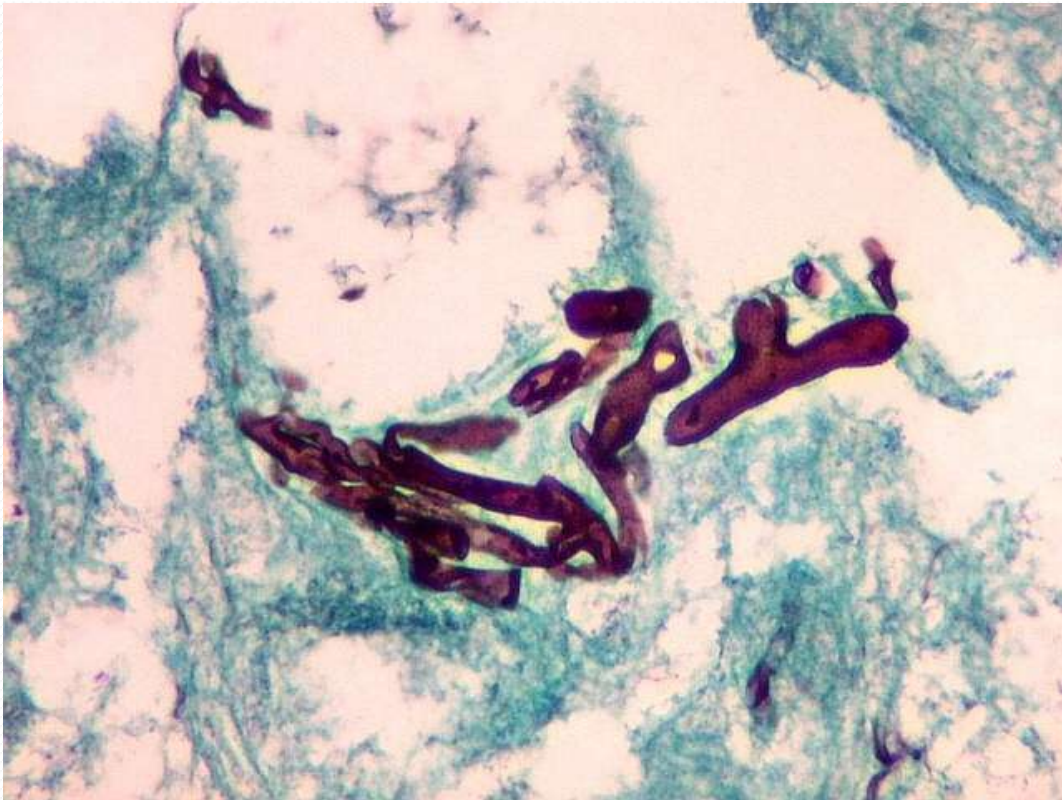
ASPERGILLUS EM VASO

GROCOTT SEM COLORAÇÃO DE FUNDO



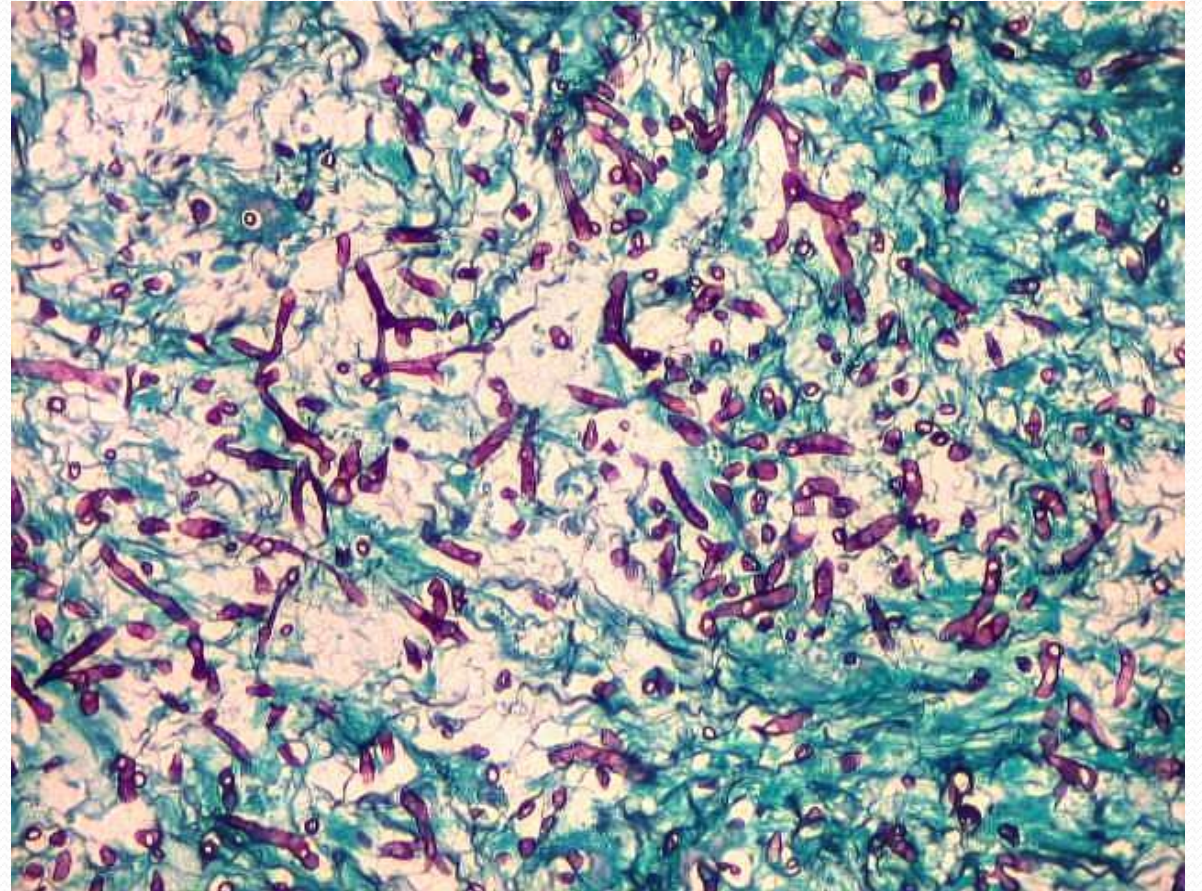
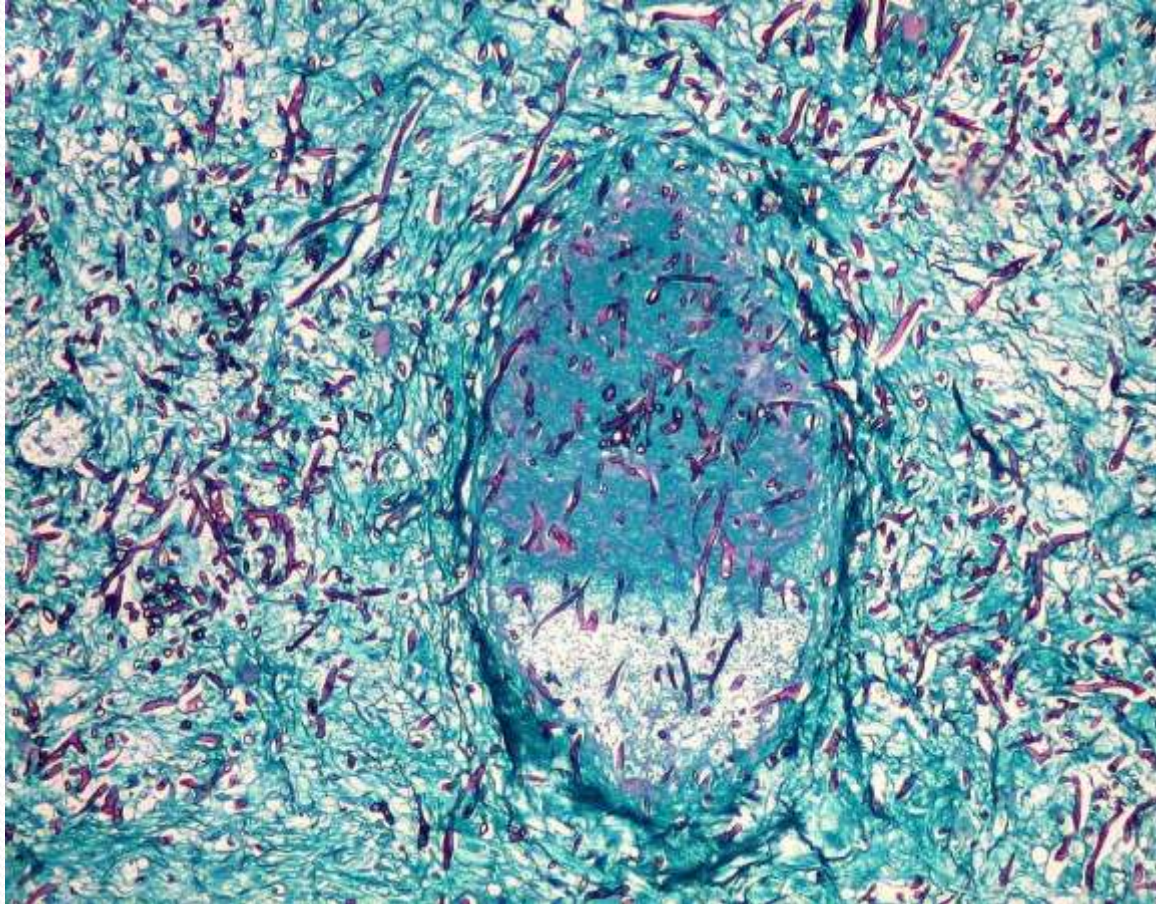
ASPERGILLUS EM SUPERFÍCIE DE BRÔNQUIO

GROCOTT + VERDE LUZ

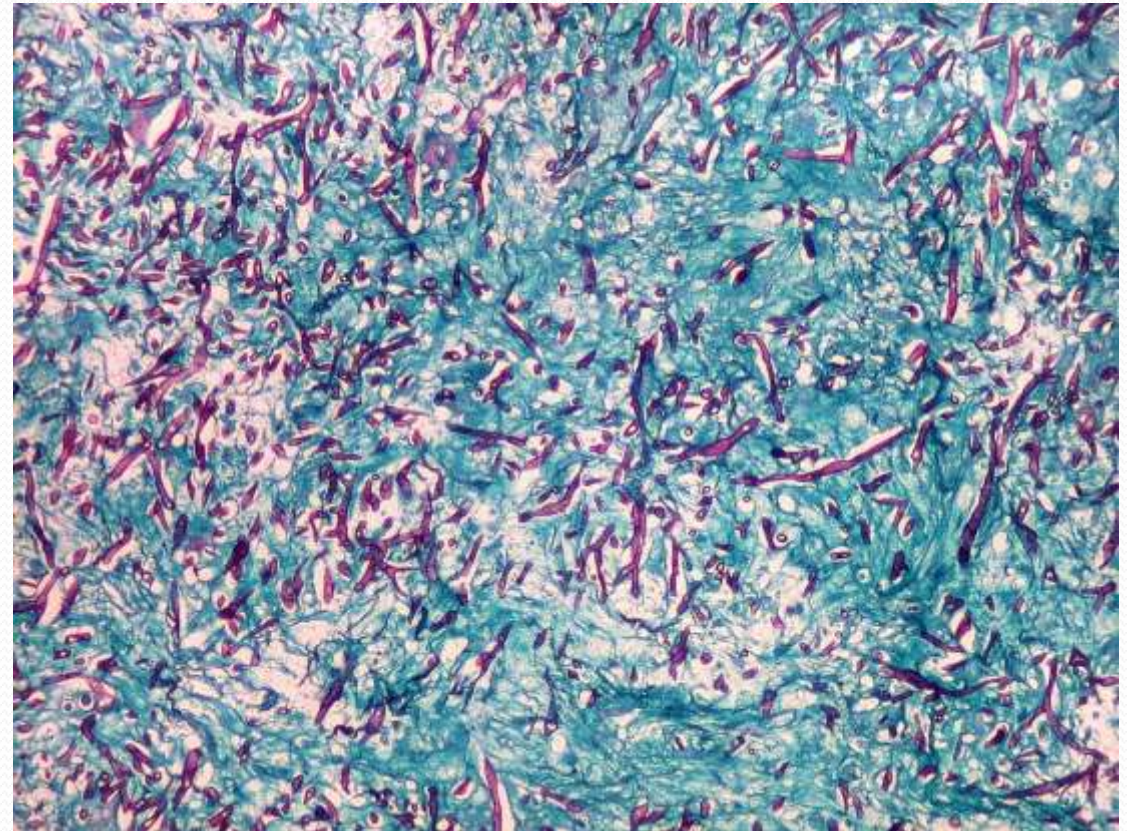
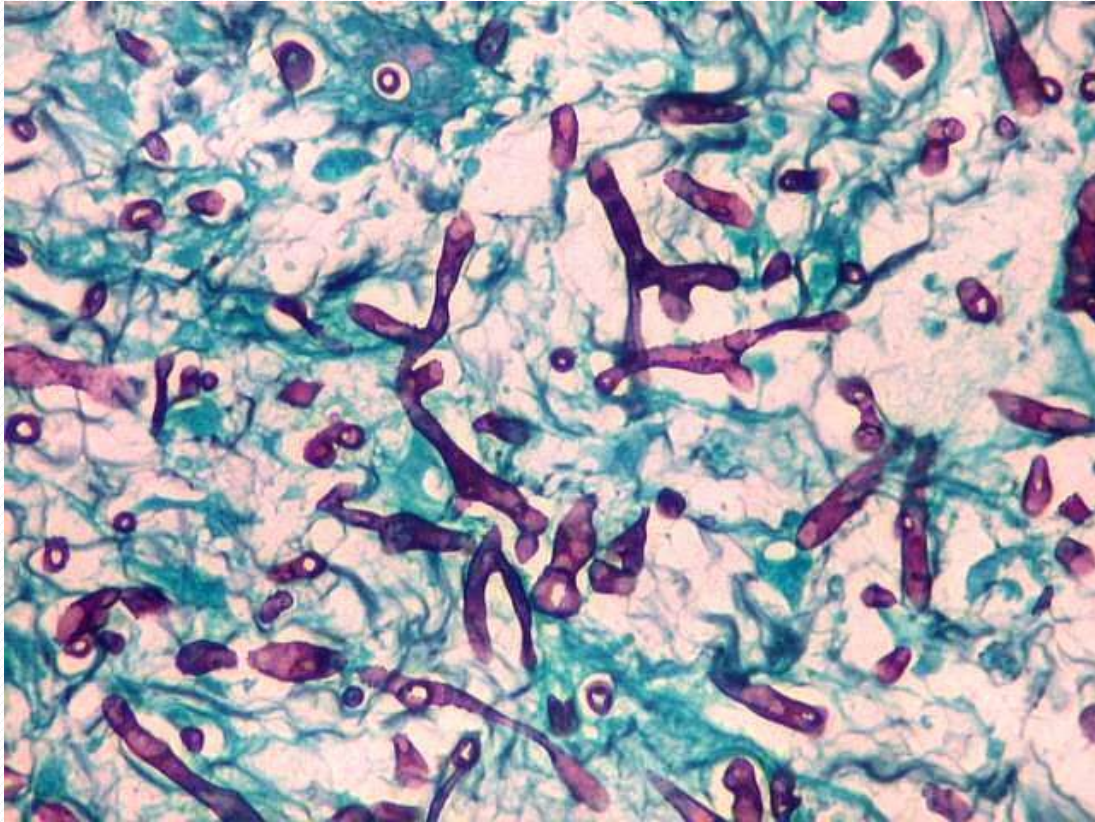


Mucormicose rino-órbito-cerebral

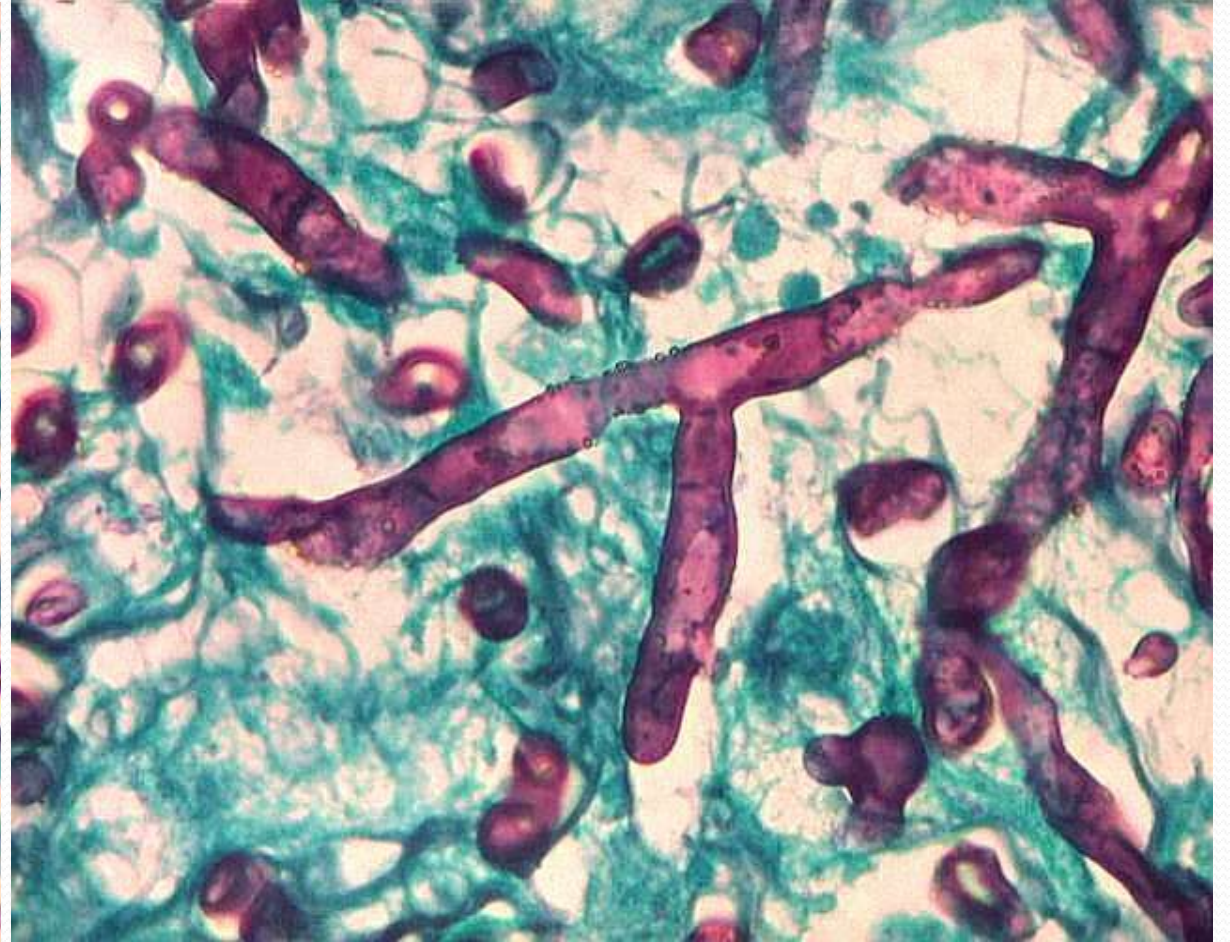
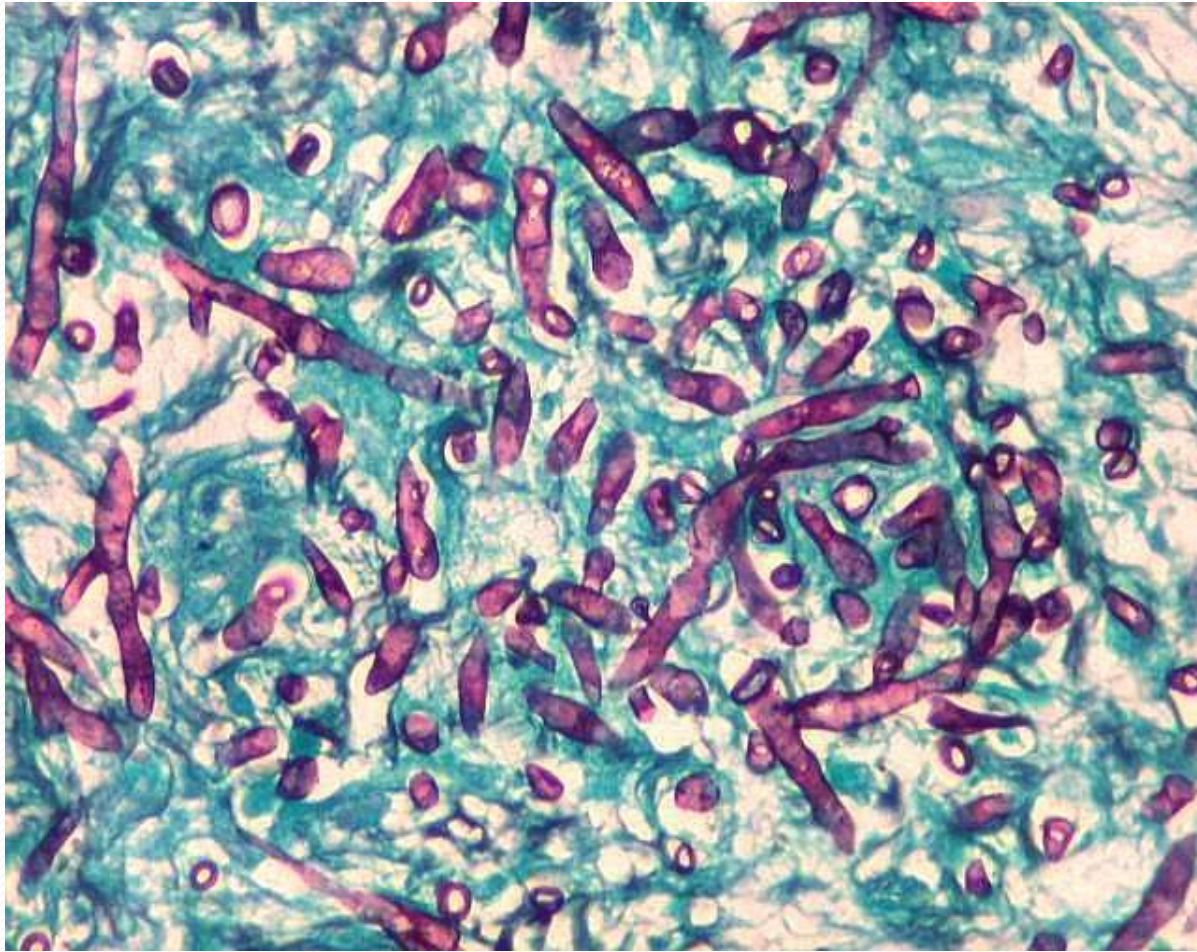
GROCOTT+VERDE LUZ(ASPERGILOSE)



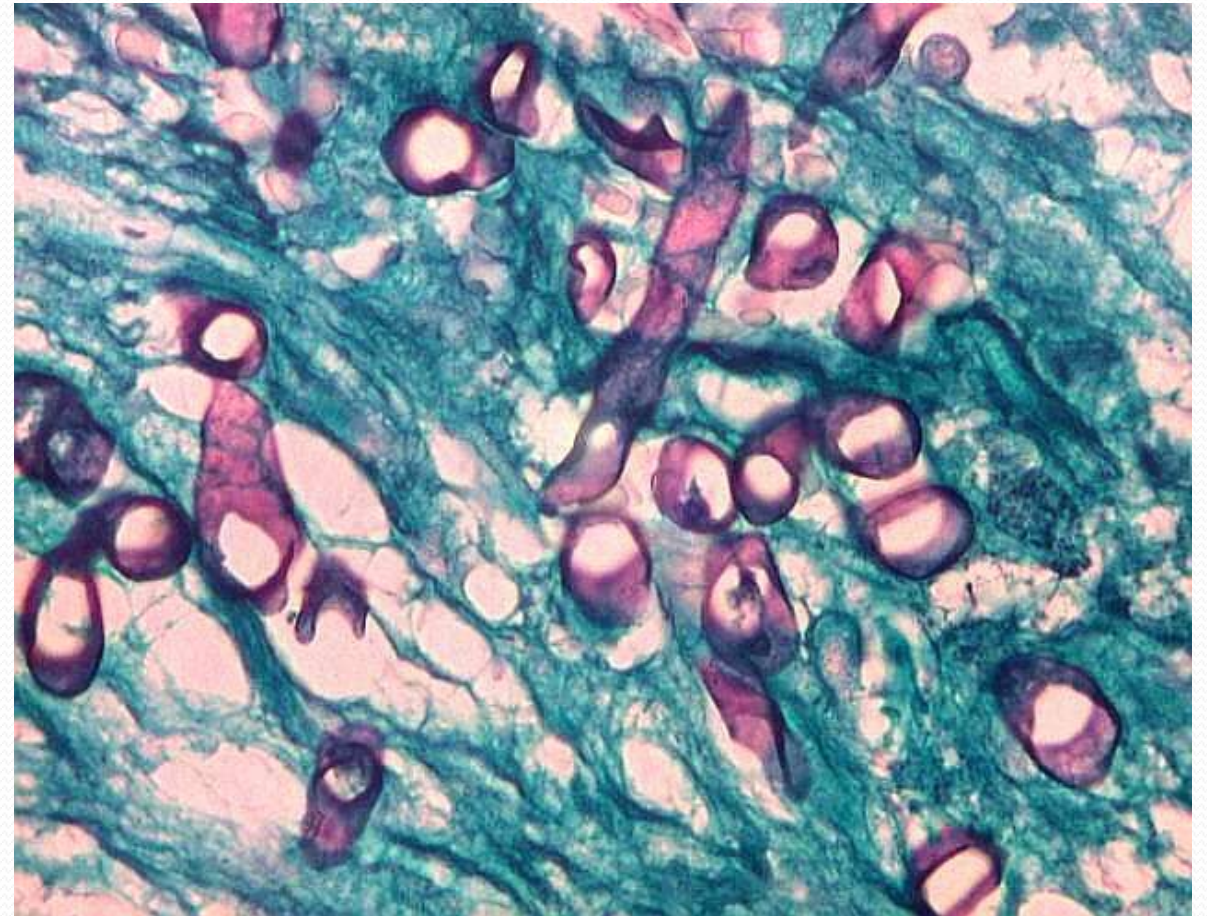
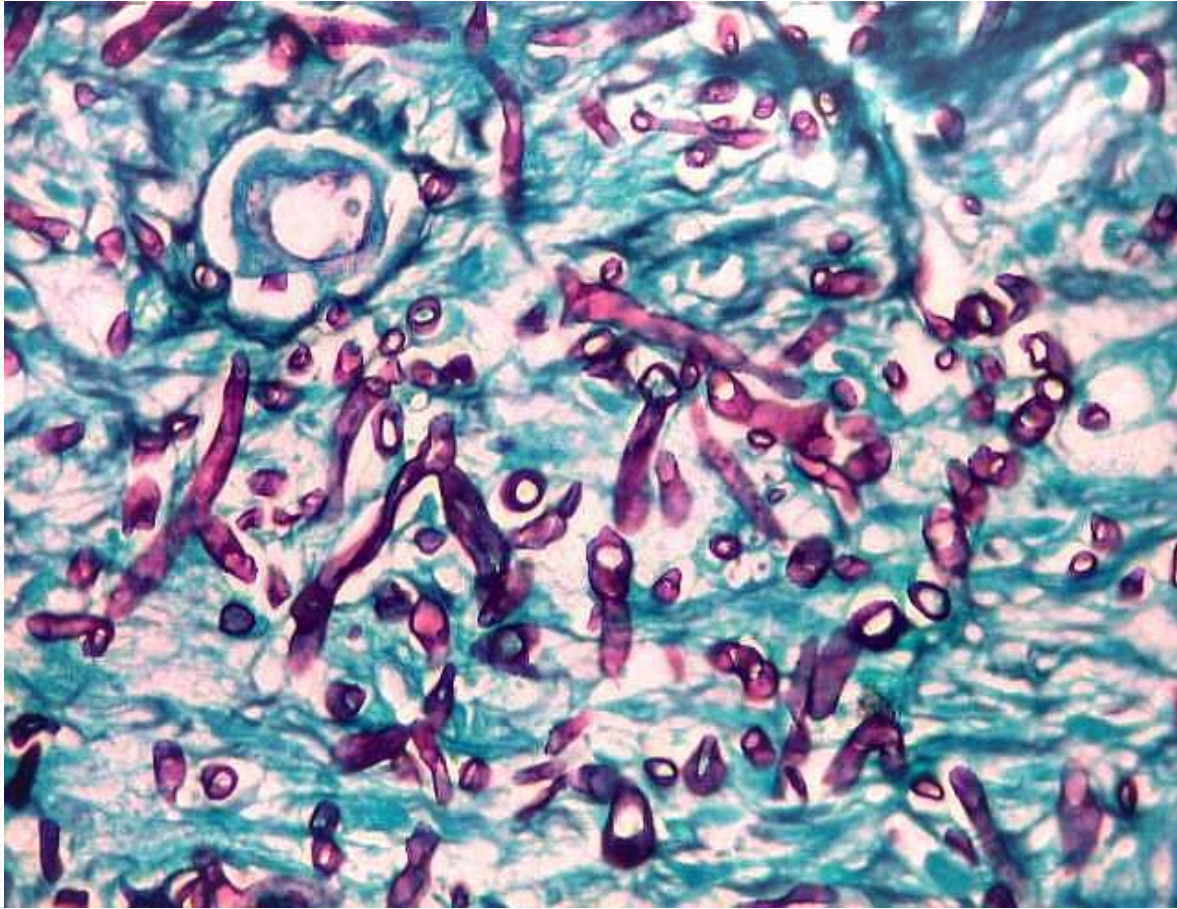
GROCOTT+VERDE LUZ(ASPERGILOSE)



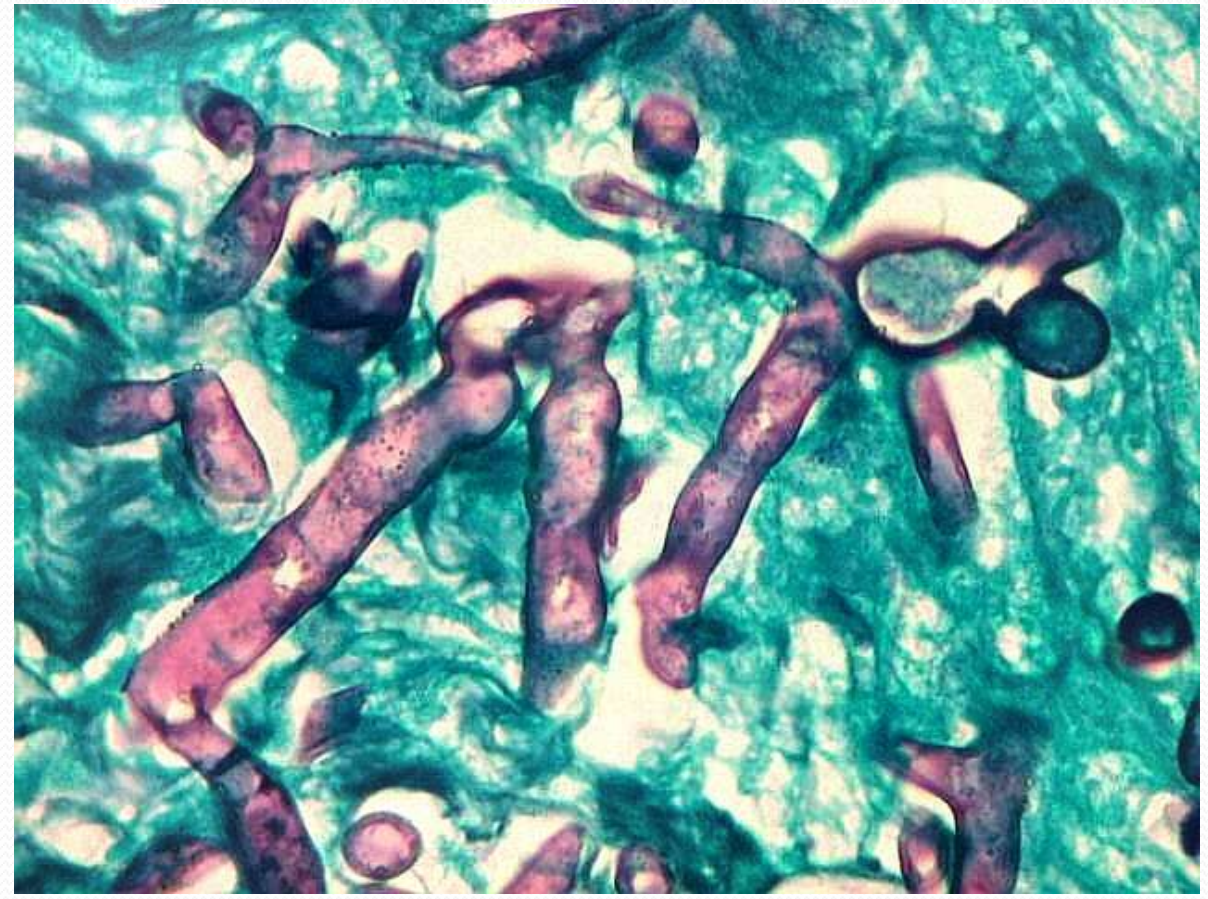
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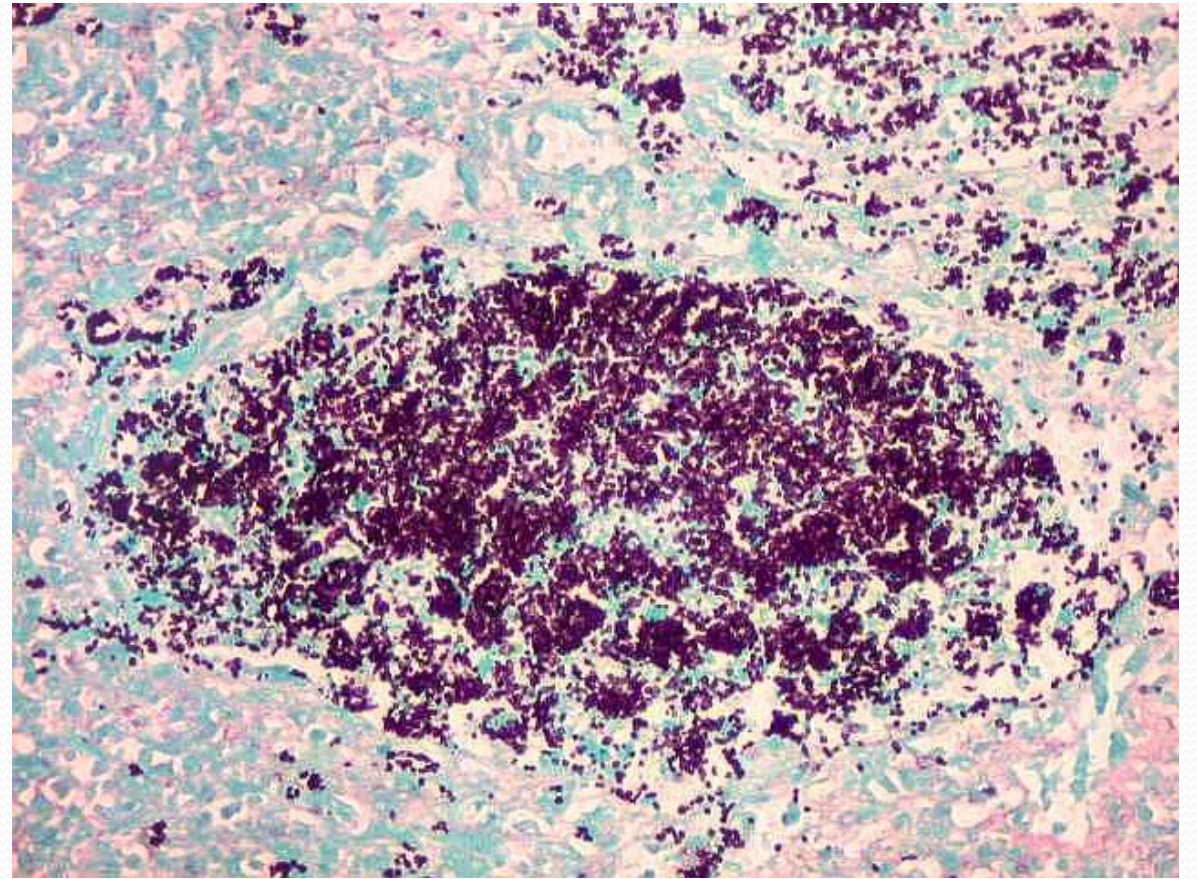
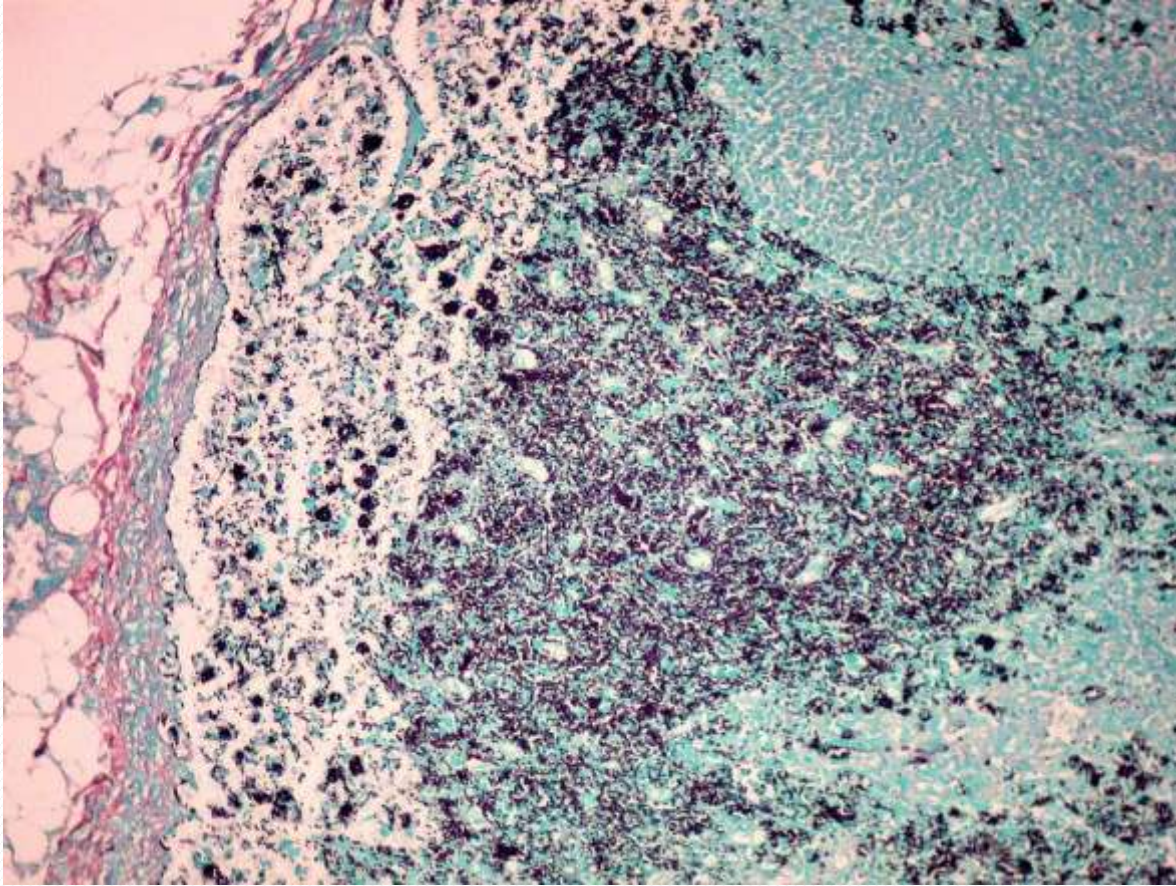
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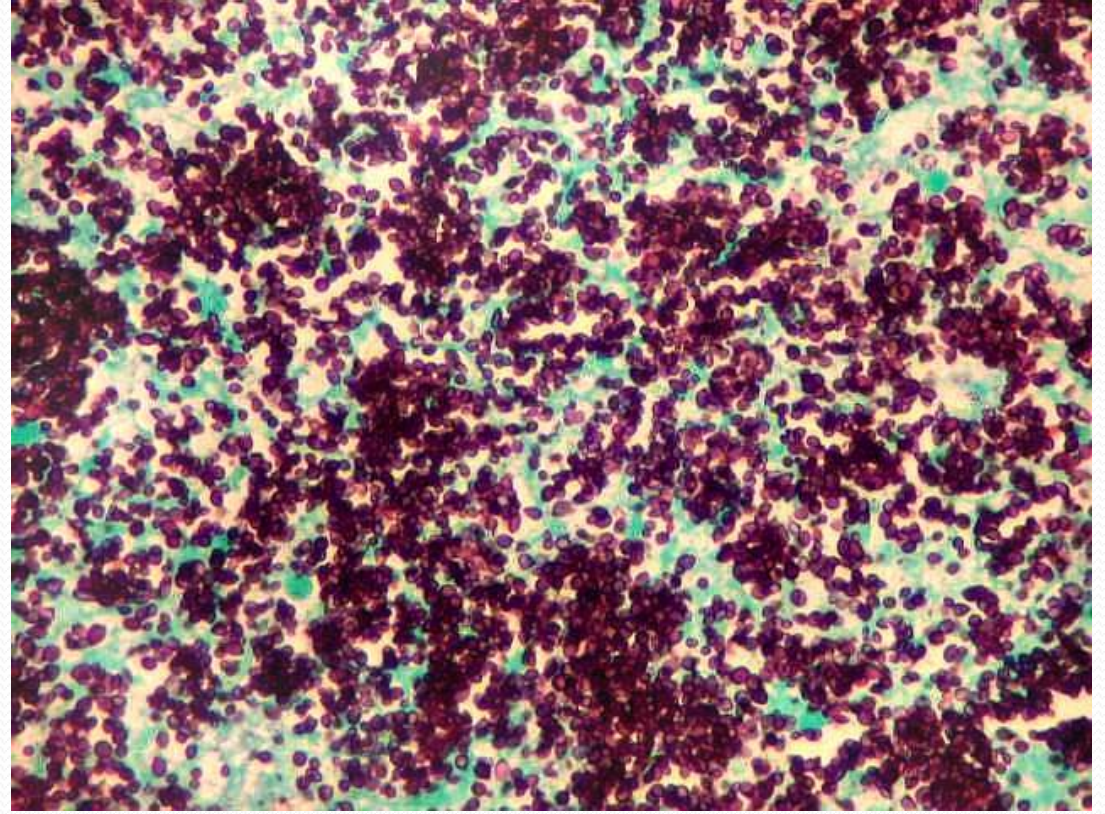
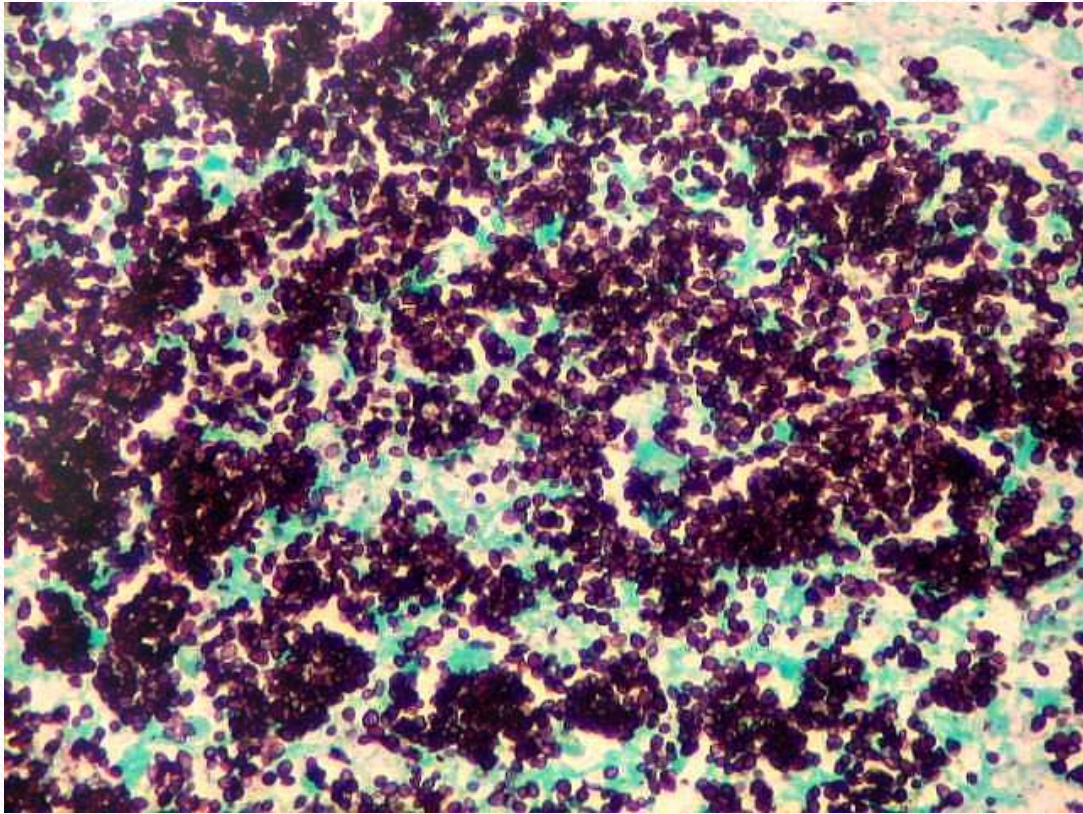
GROCOTT+VERDE LUZ(ASPERGILOSE)



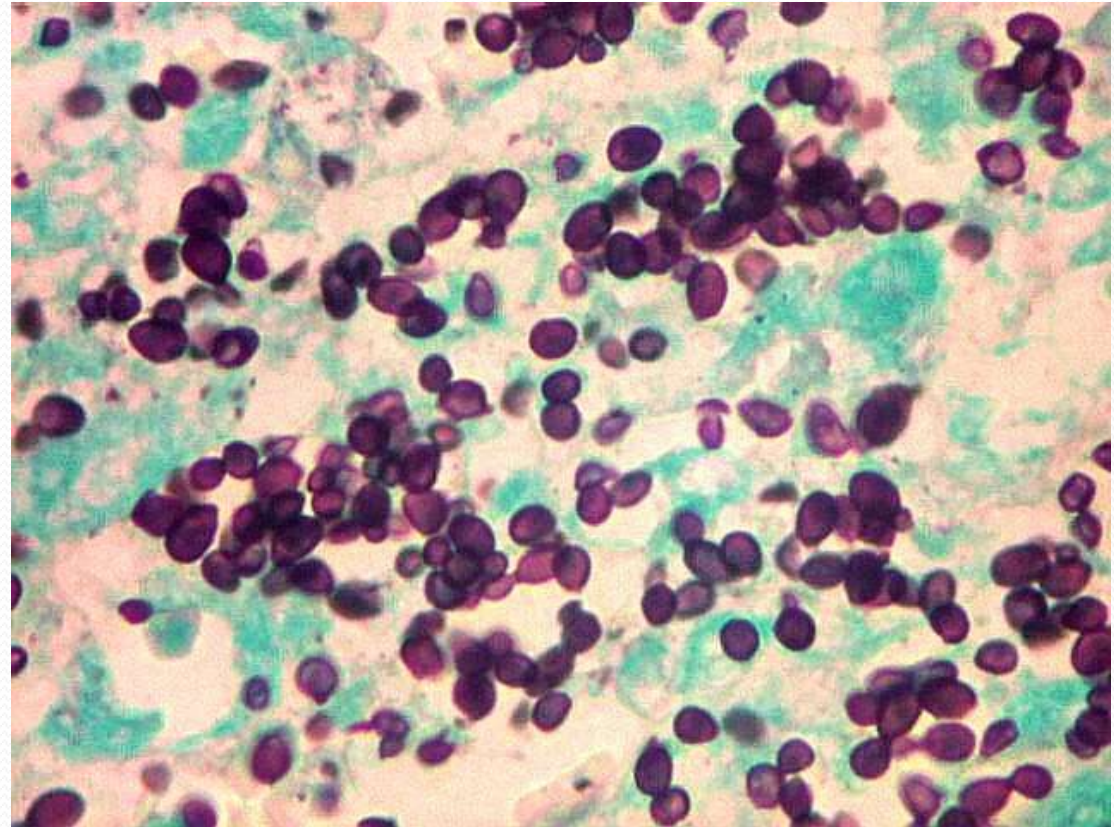
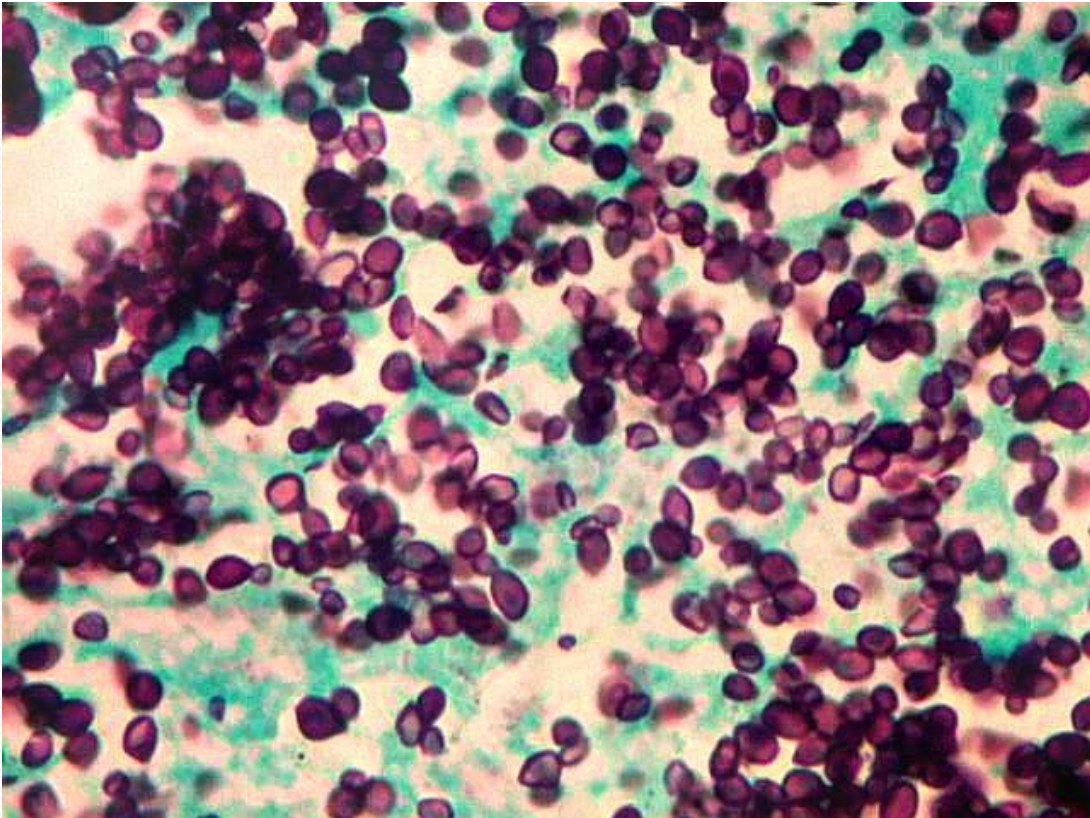
GROCOTT+VERDE LUZ(HISTOPLASMOSE)



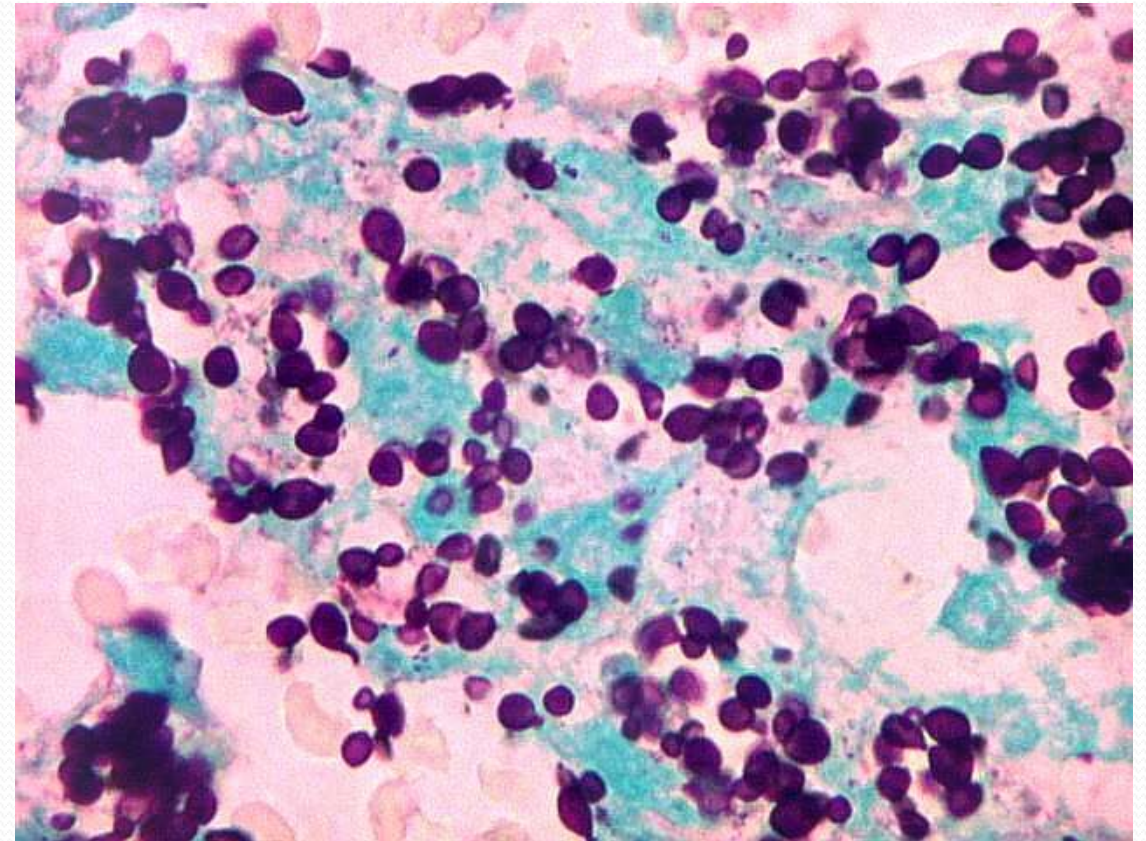
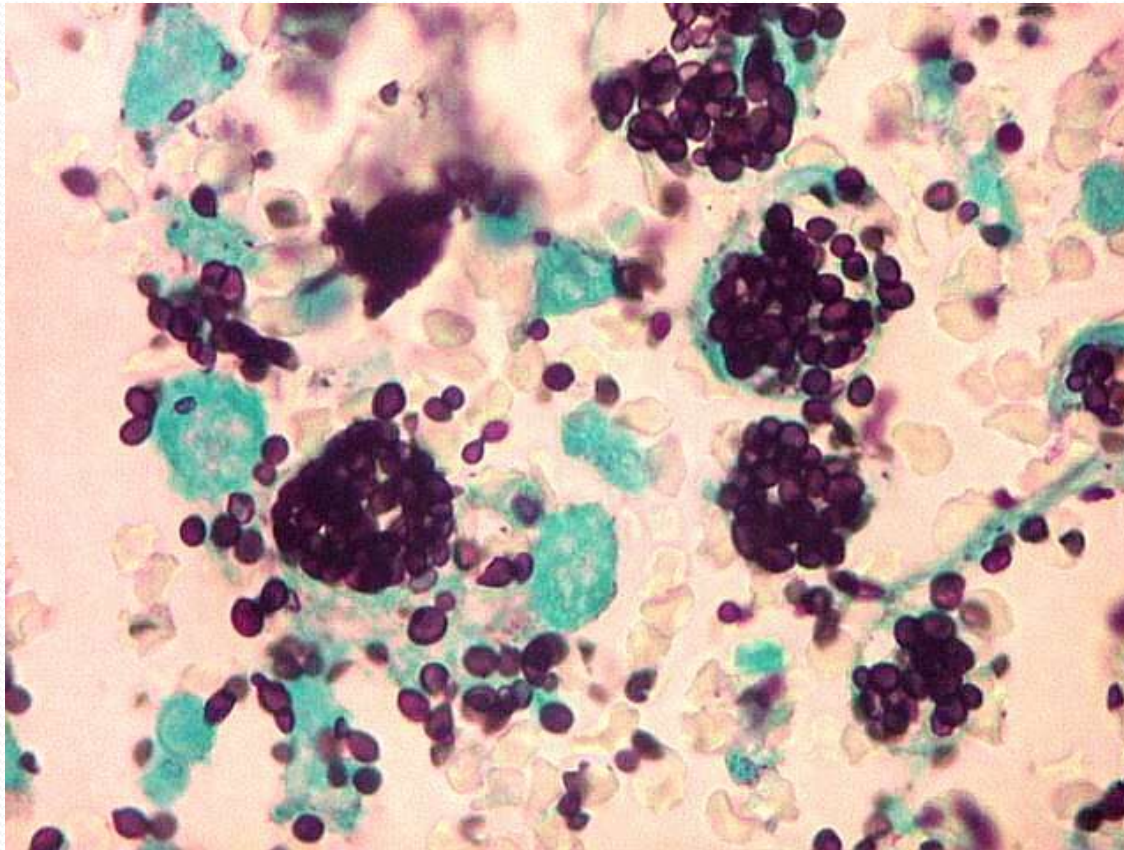
GROCOTT+VERDE LUZ(HISTOPLASMOSE)



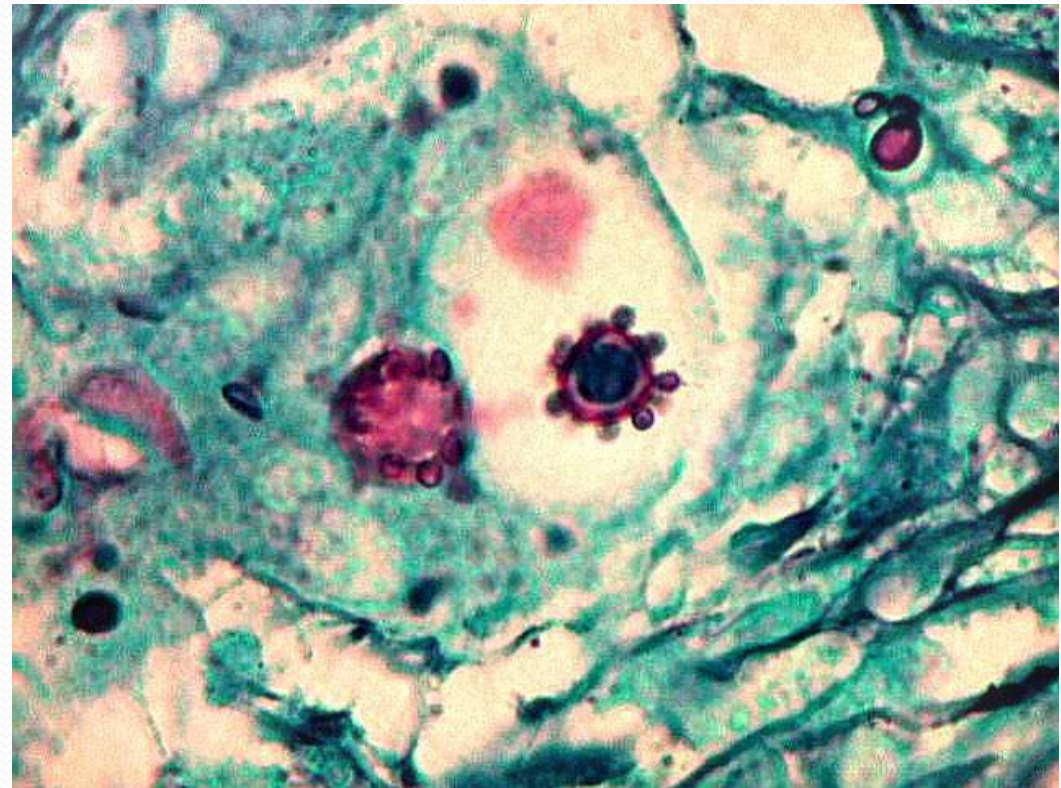
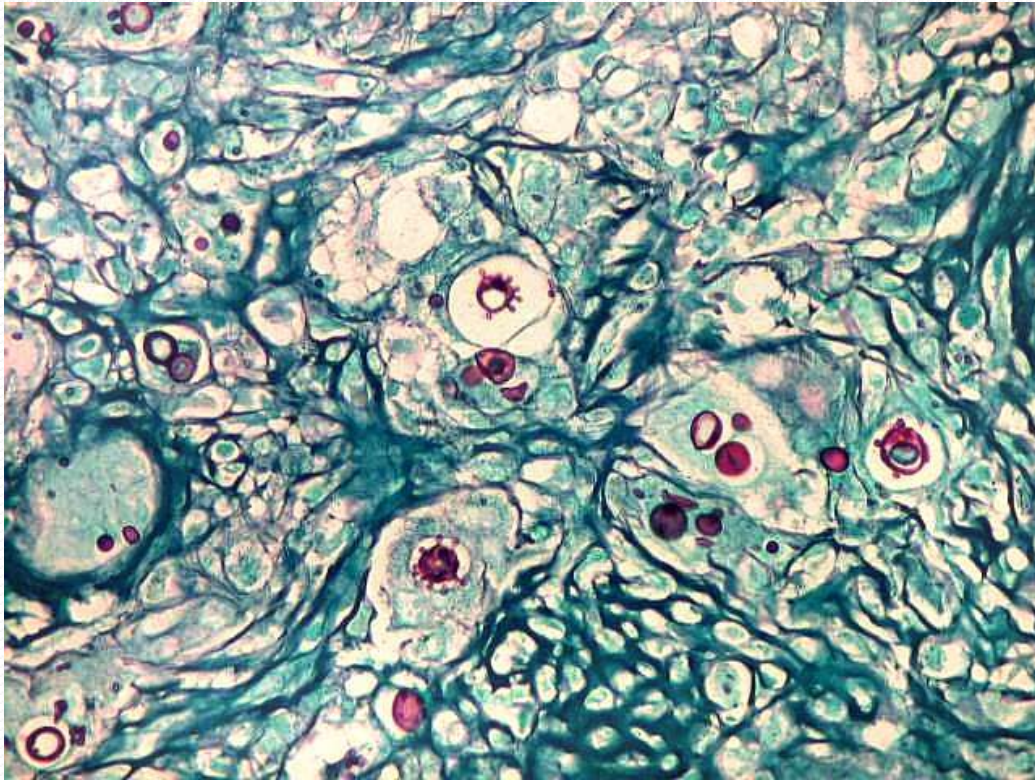
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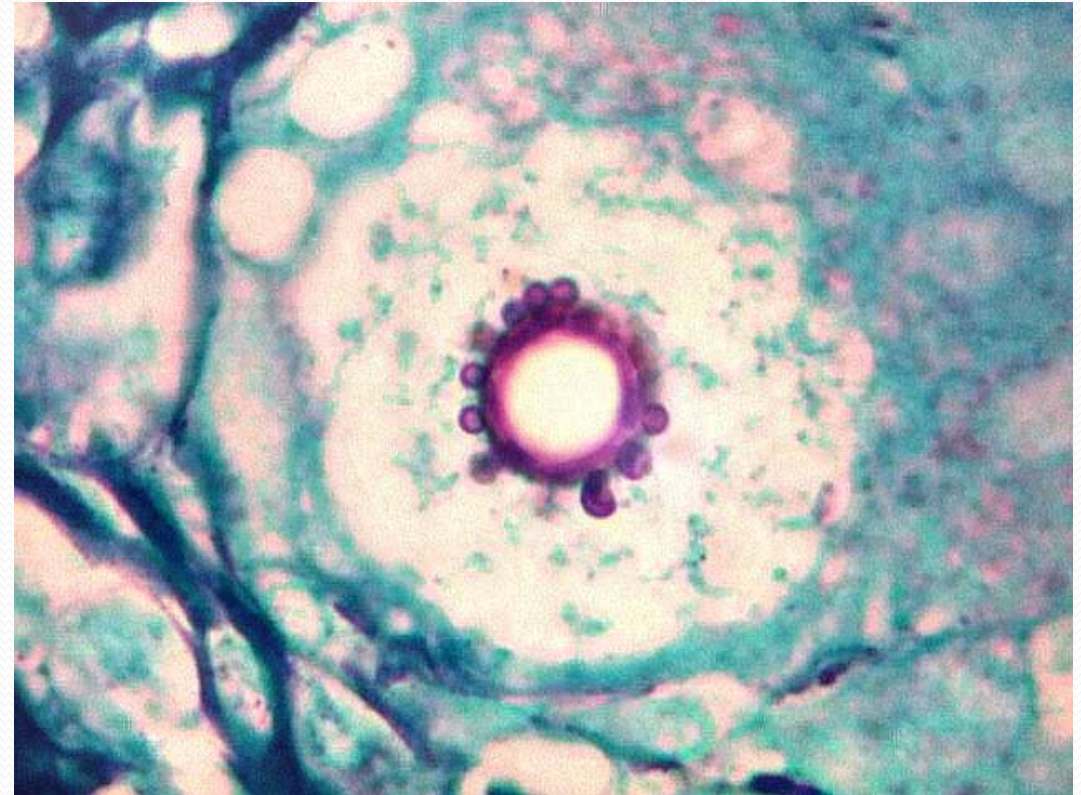
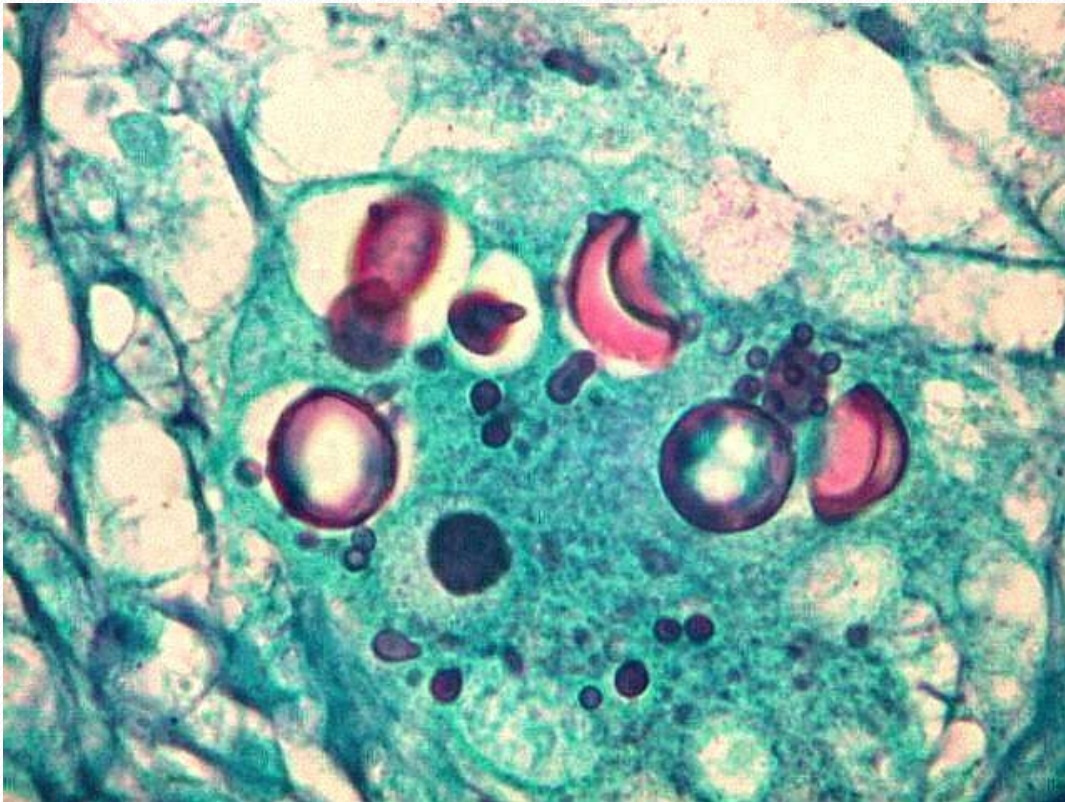
GROCOTT+VERDE LUZ(HISTOPLASMOSE)



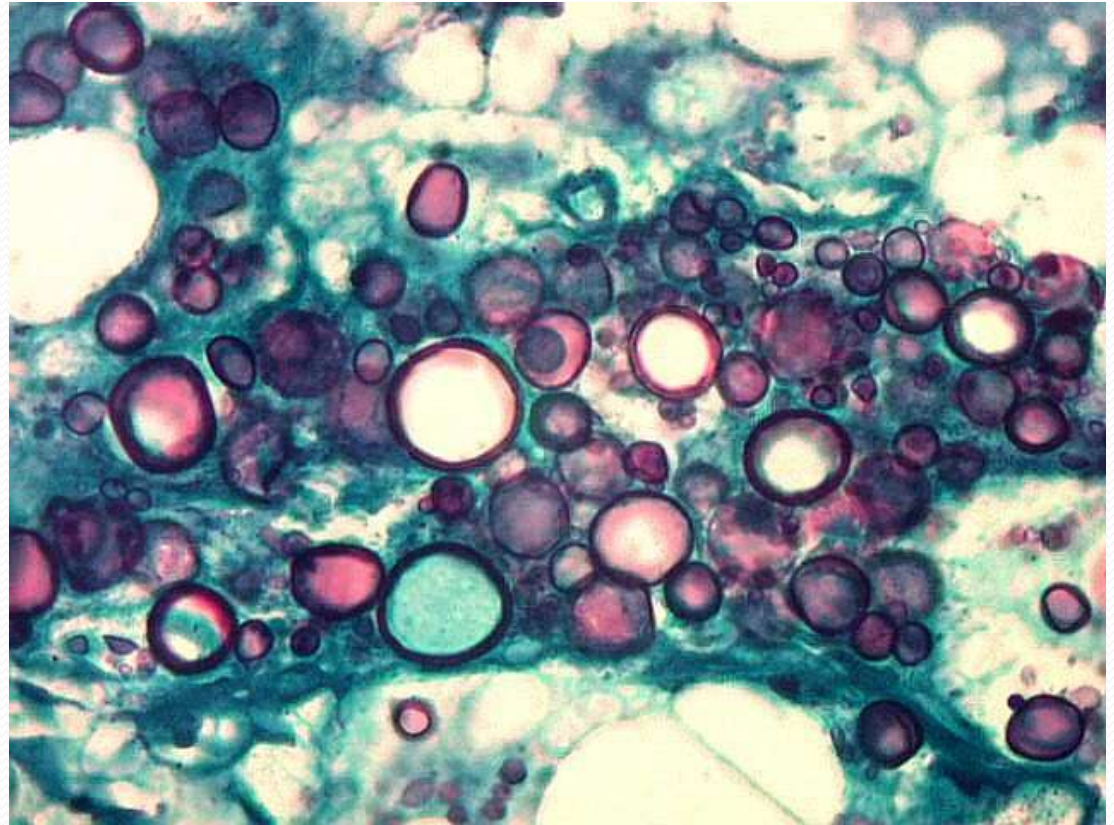
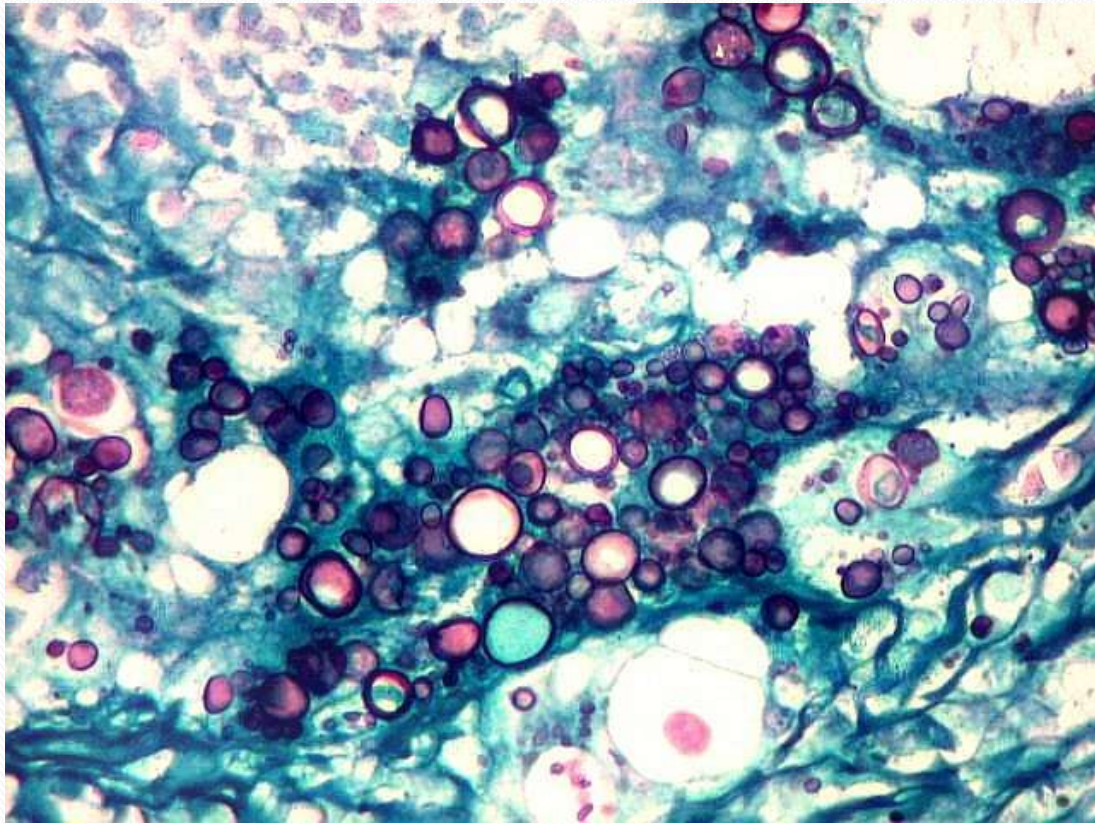
GROCOTT+VERDE LUZ(PARACOCO)



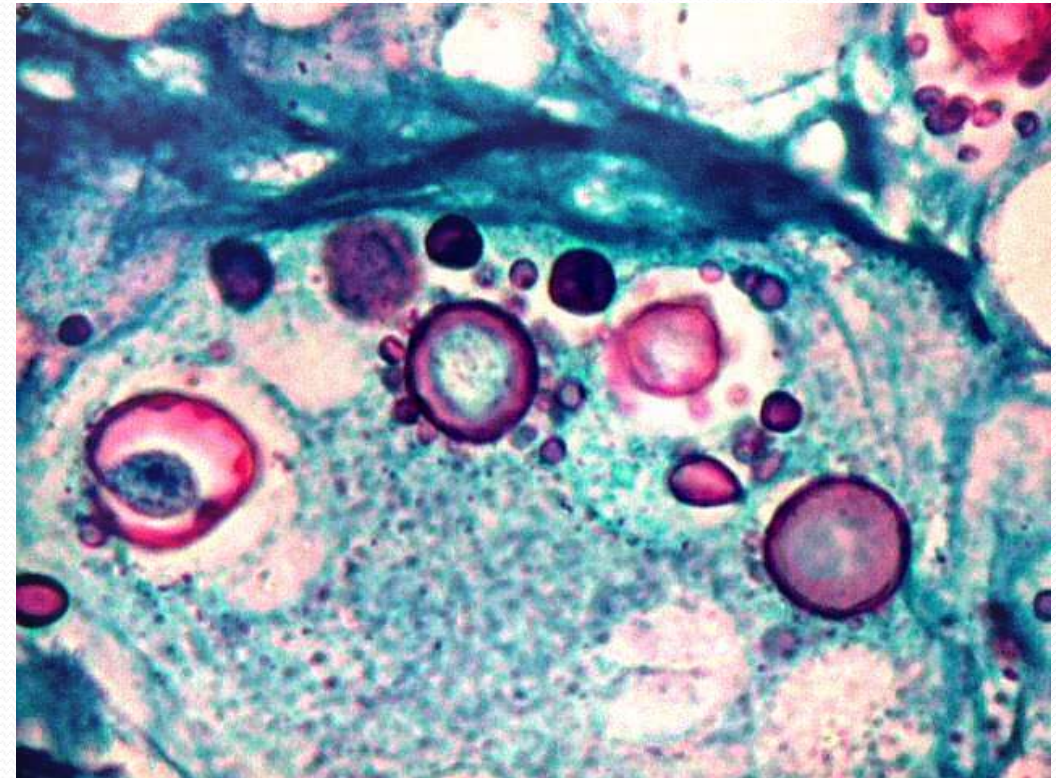
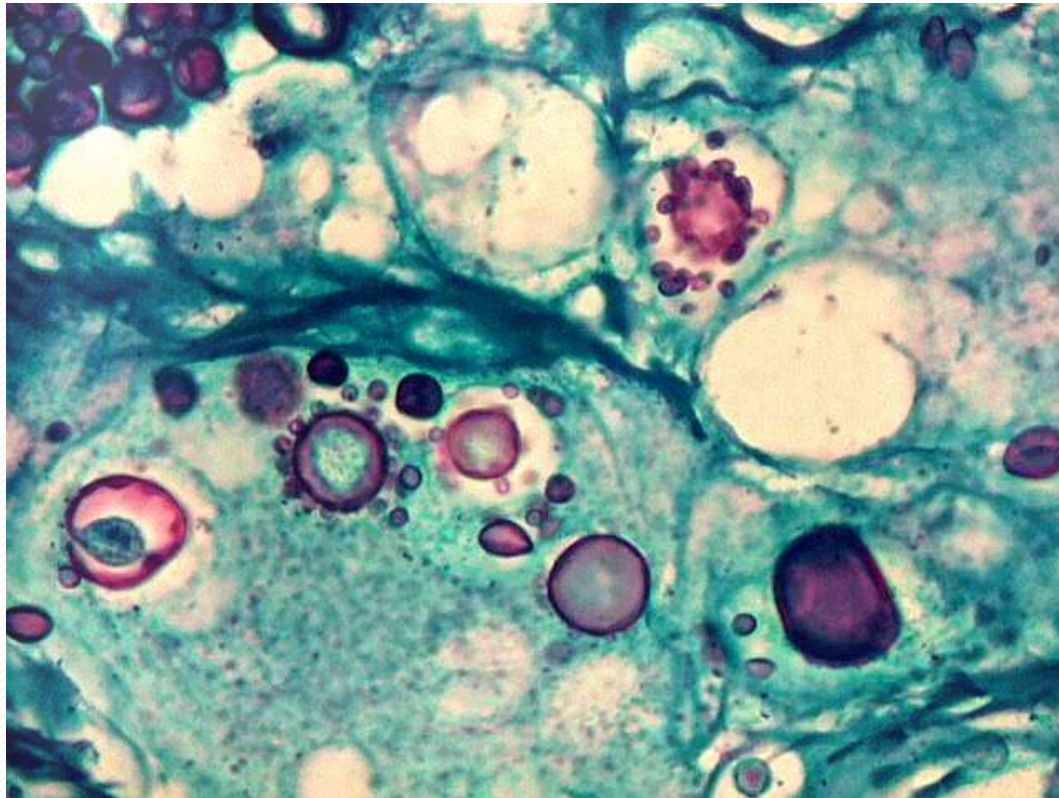
GROCOTT+VERDE LUZ(PARACOCO)



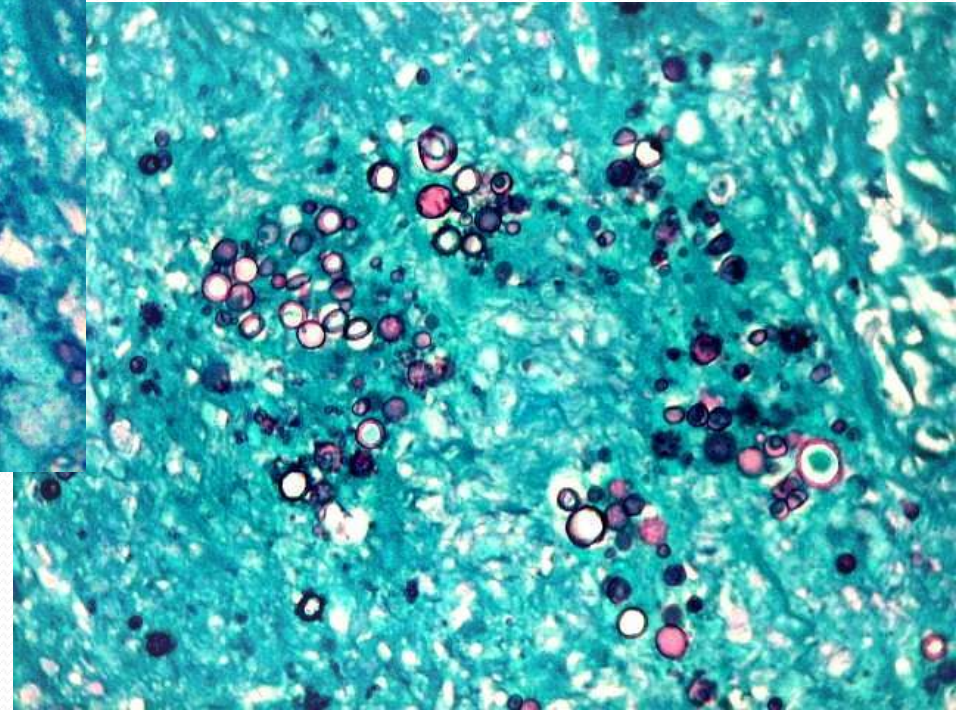
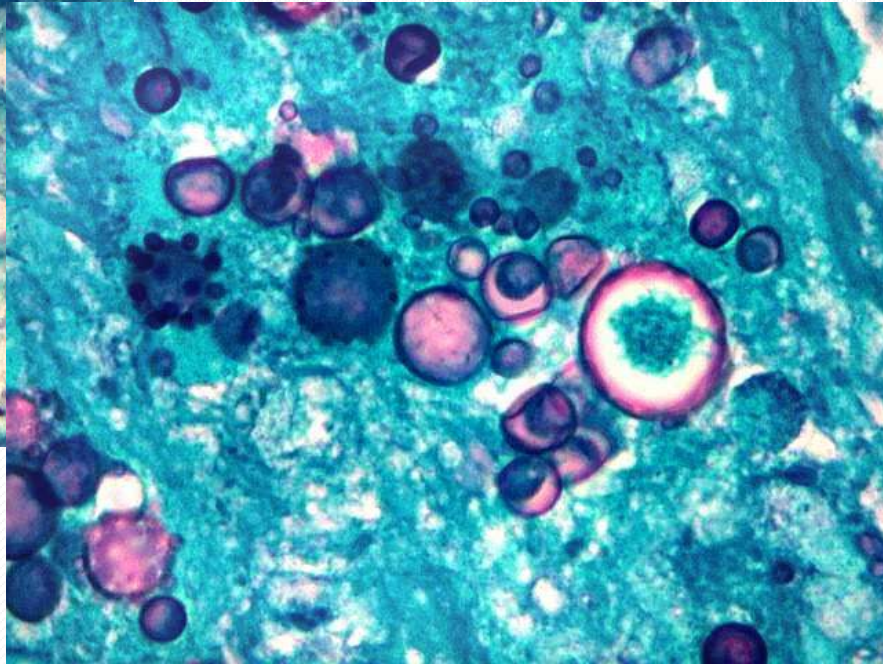
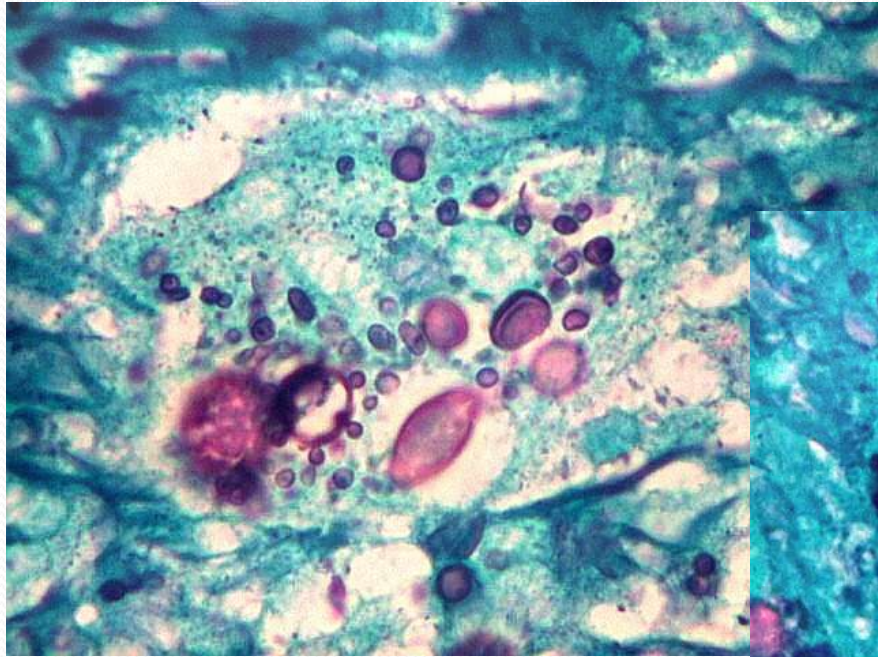
GROCOTT+VERDE LUZ(PARACOCO)



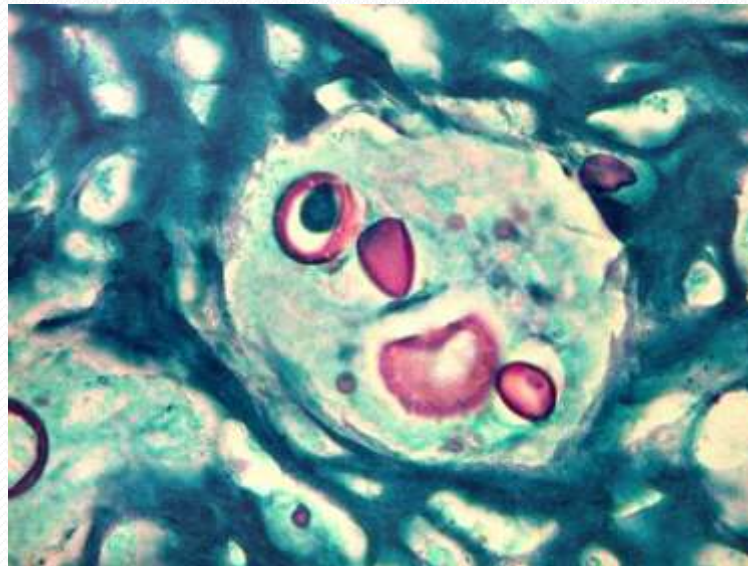
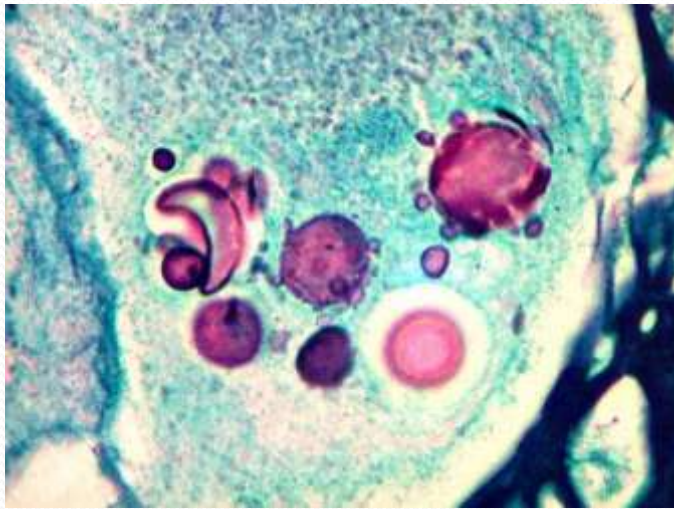
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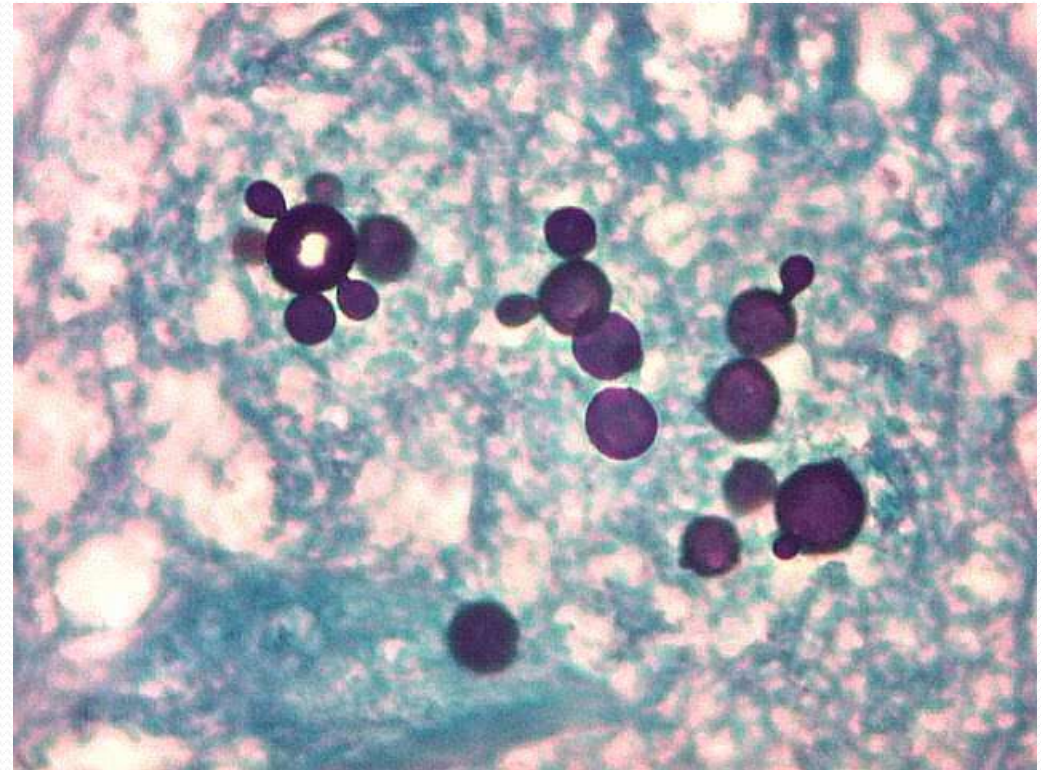
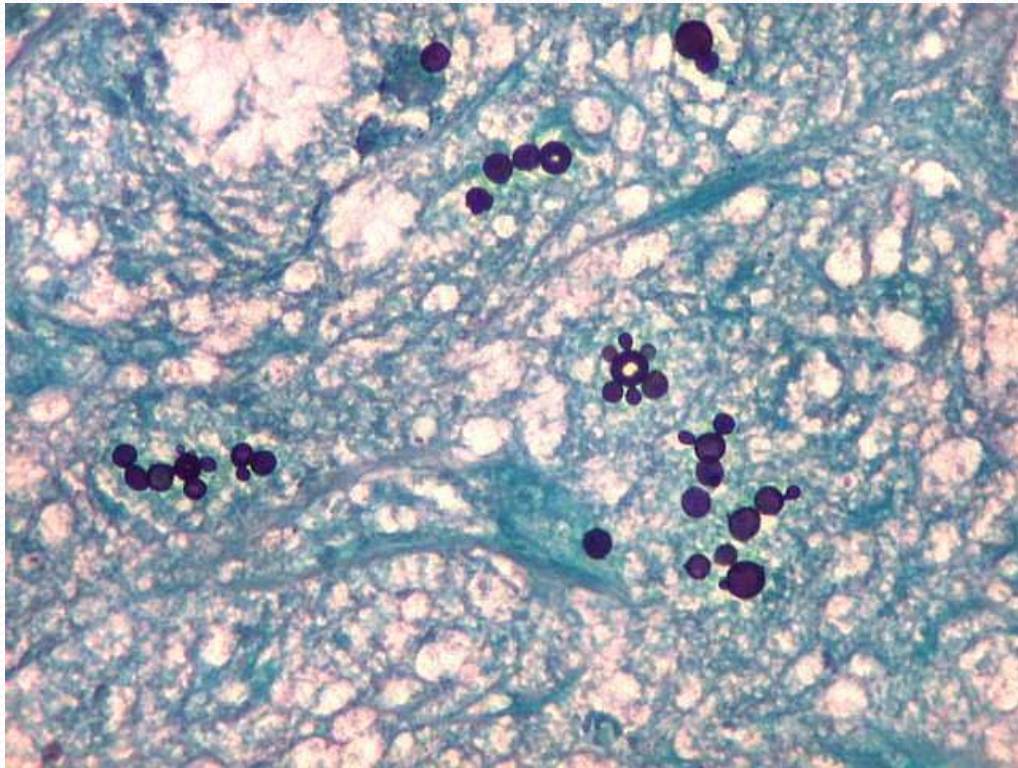
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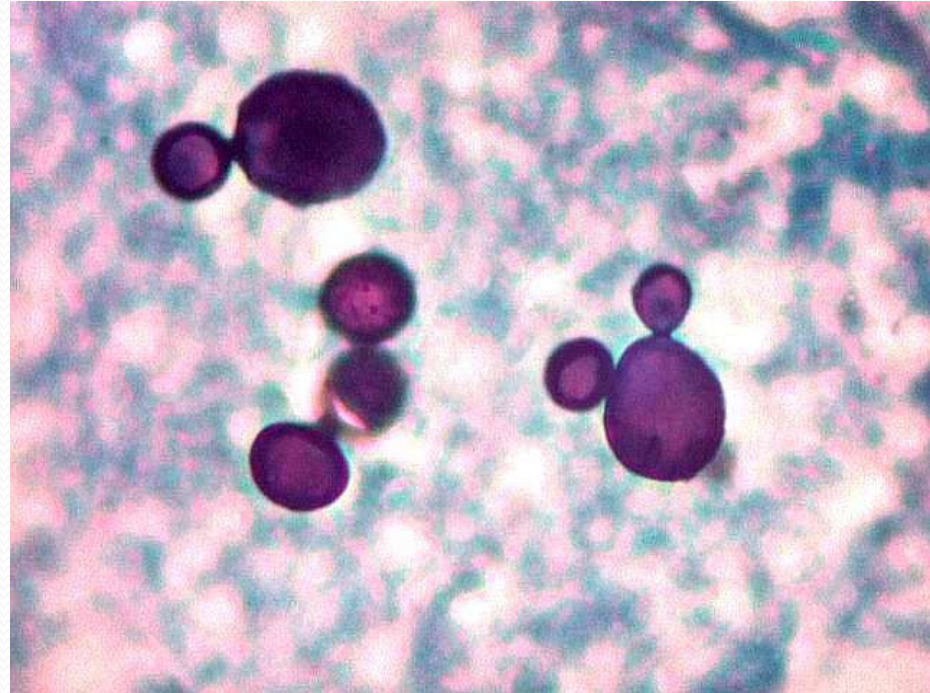
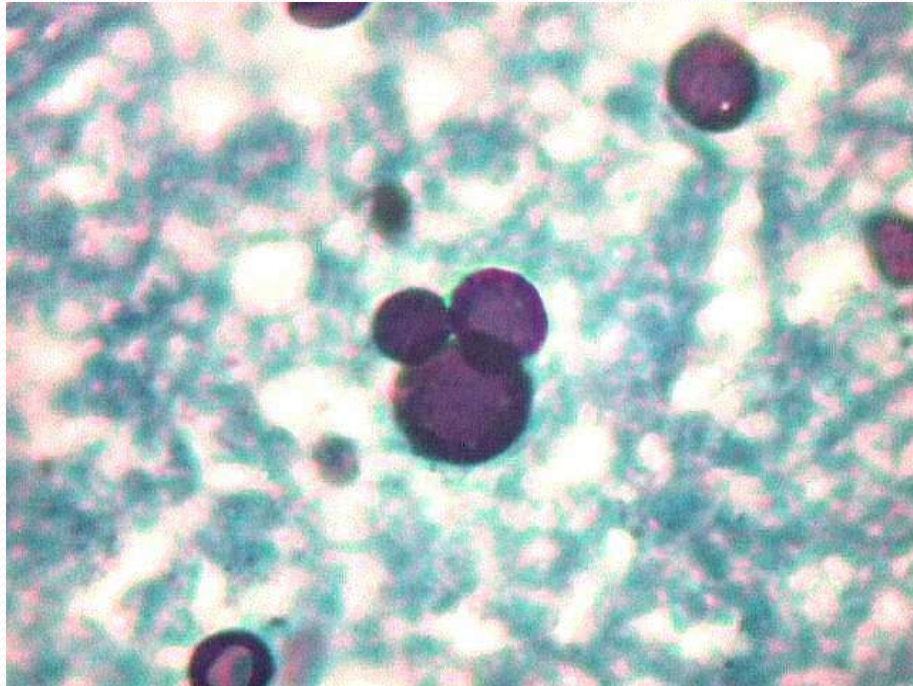
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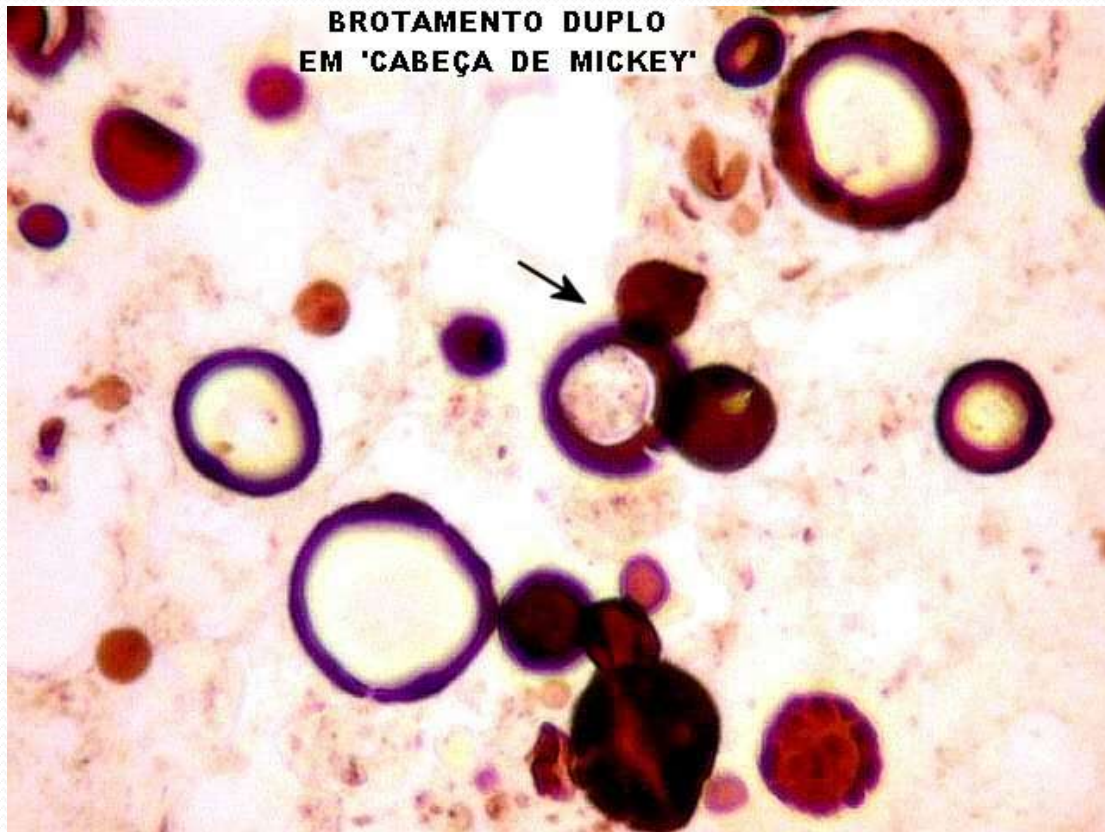
GROCOTT+VERDE LUZ(PARACOCO)



GROCOTT+VERDE LUZ(PARACOCO)



GROCOTT SEM COLORAÇÃO DE FUNDO



BAAR - BACILO ÁLCOOL ÁCIDO RESISTENTE

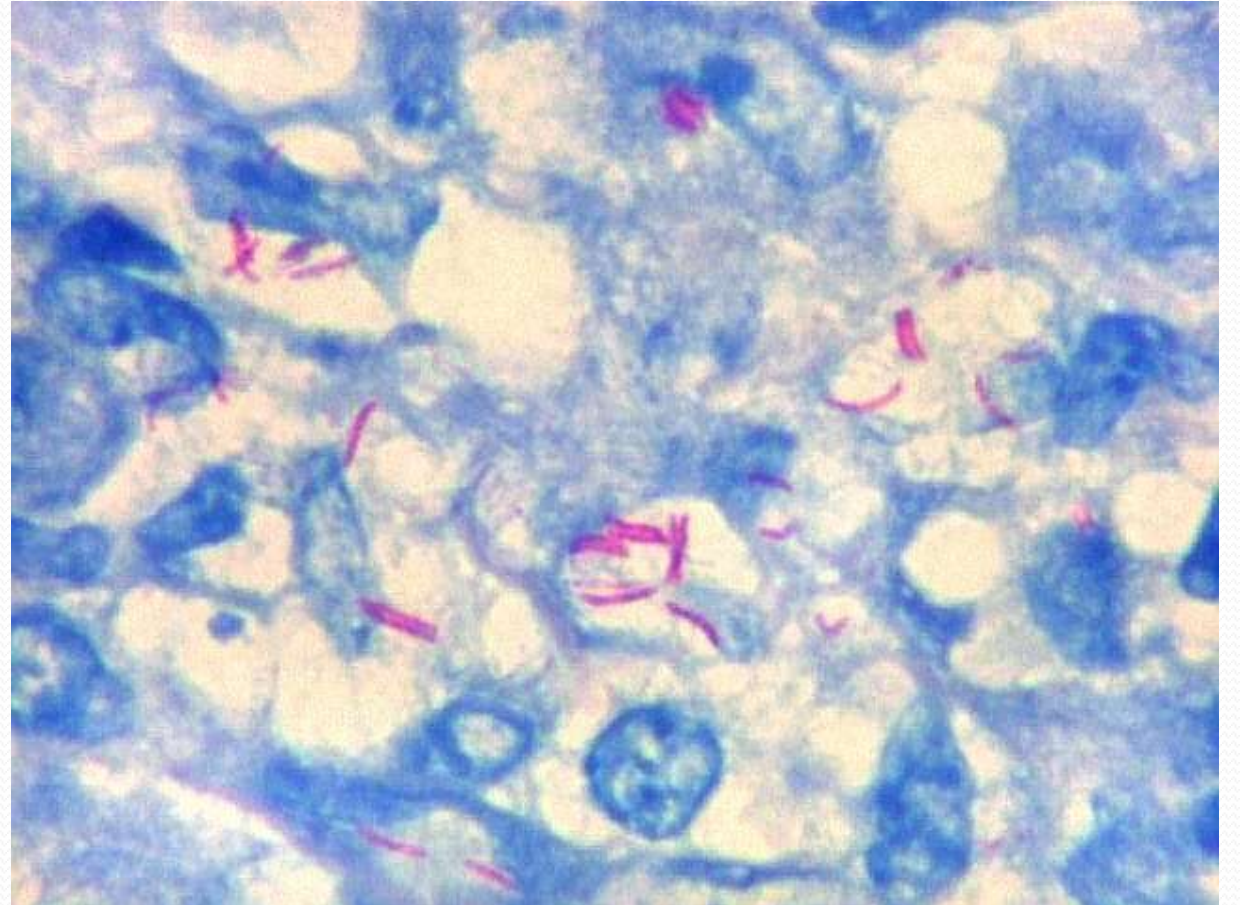
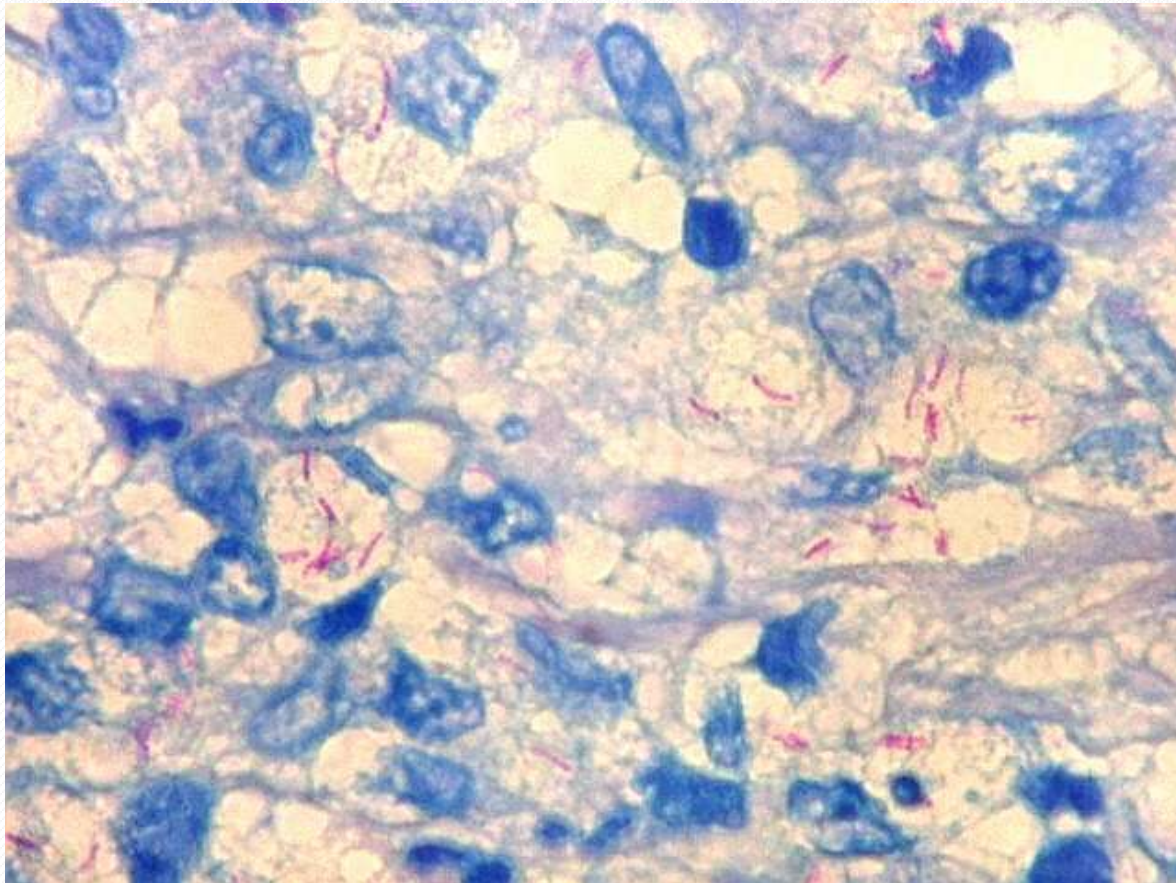
Resistência ácida- É uma característica de micobactérias, devido a composição das paredes das células de micobactérias ricas em lipídeos.

Bactérias patogênicas:

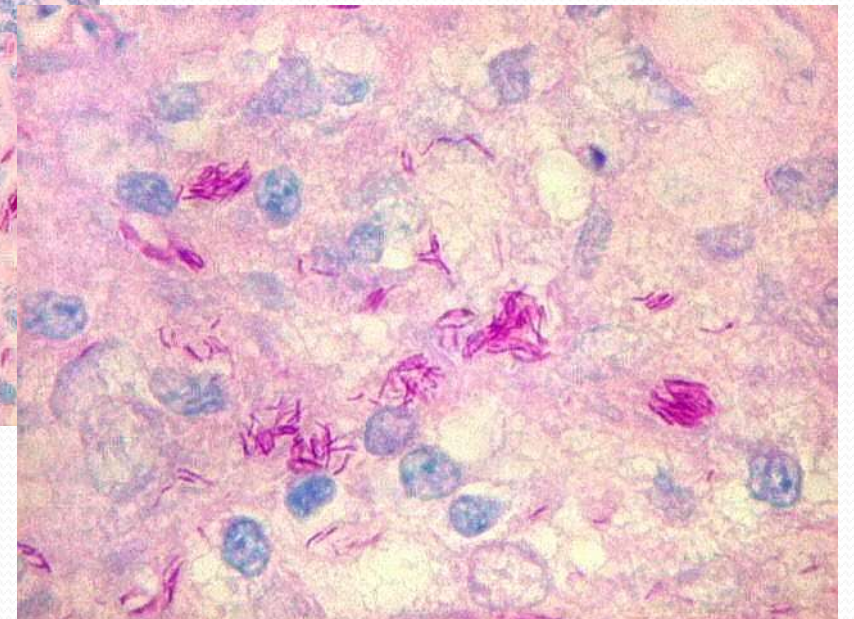
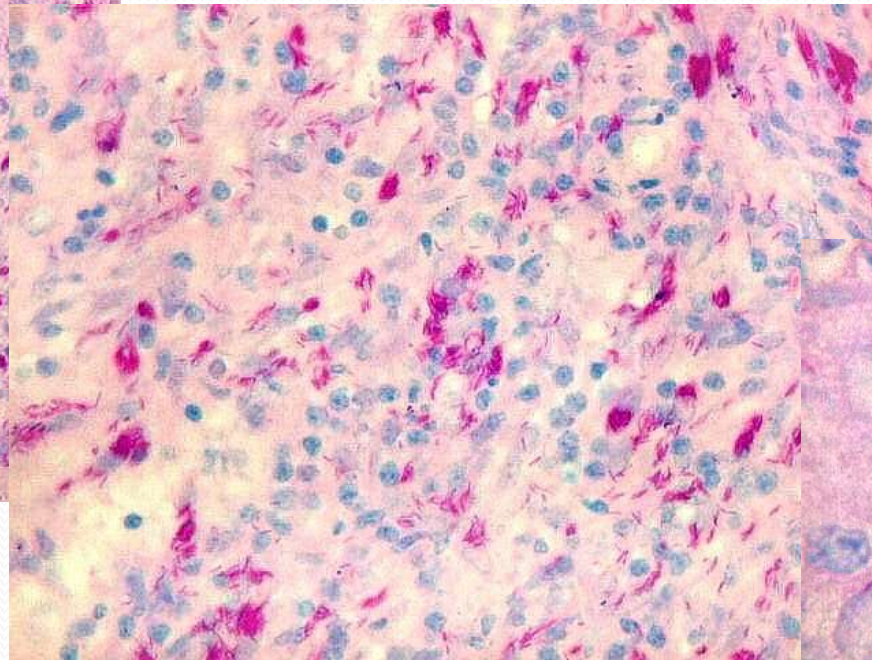
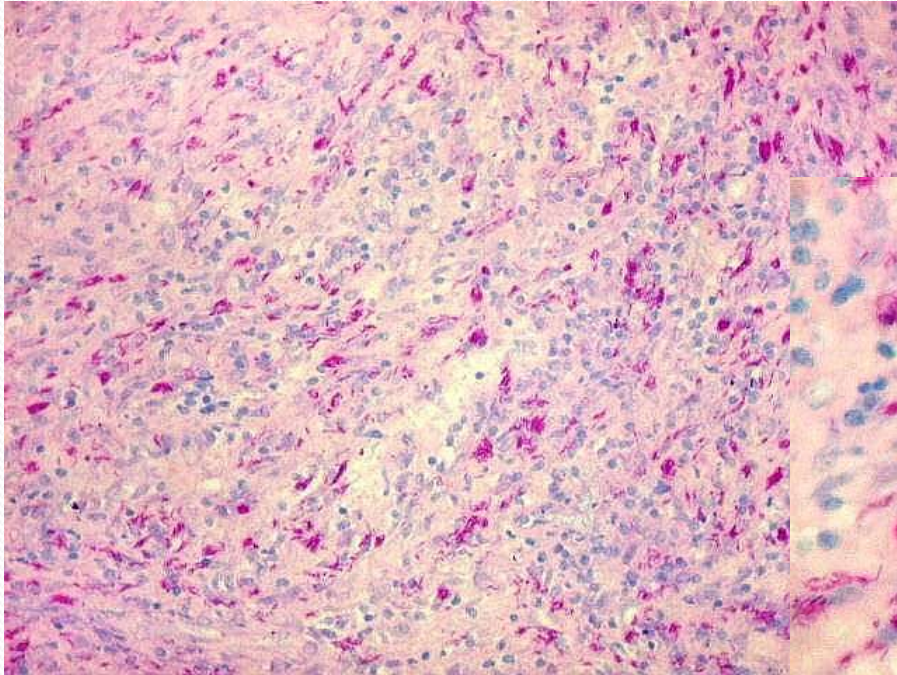
Bacilo de Koch – causador da tuberculose.

Bacilo de Hansen- causador da hanseníase ou lepra.

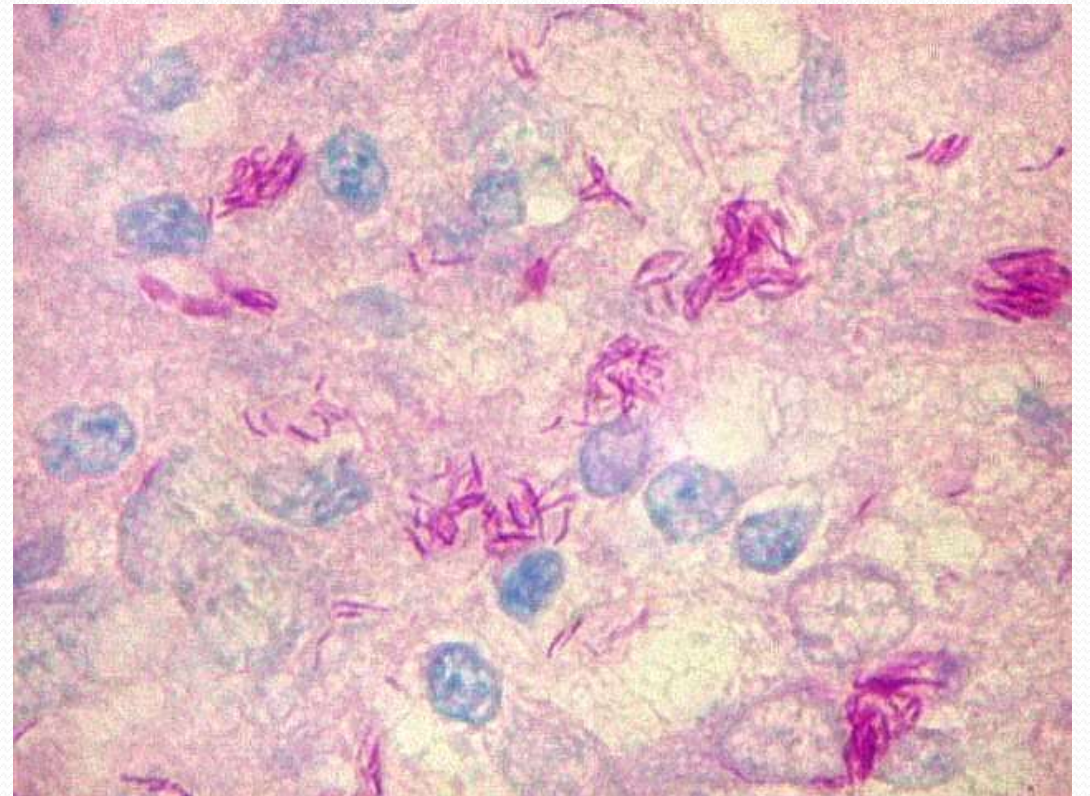
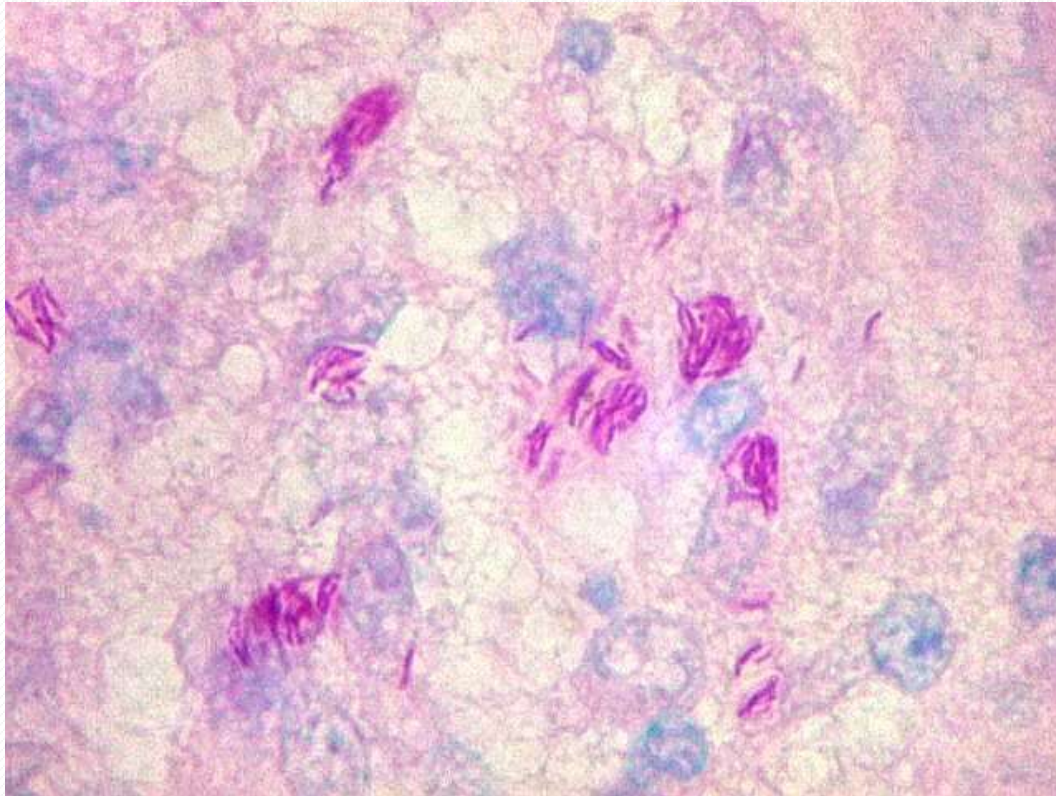
BACILOS DE KOCK



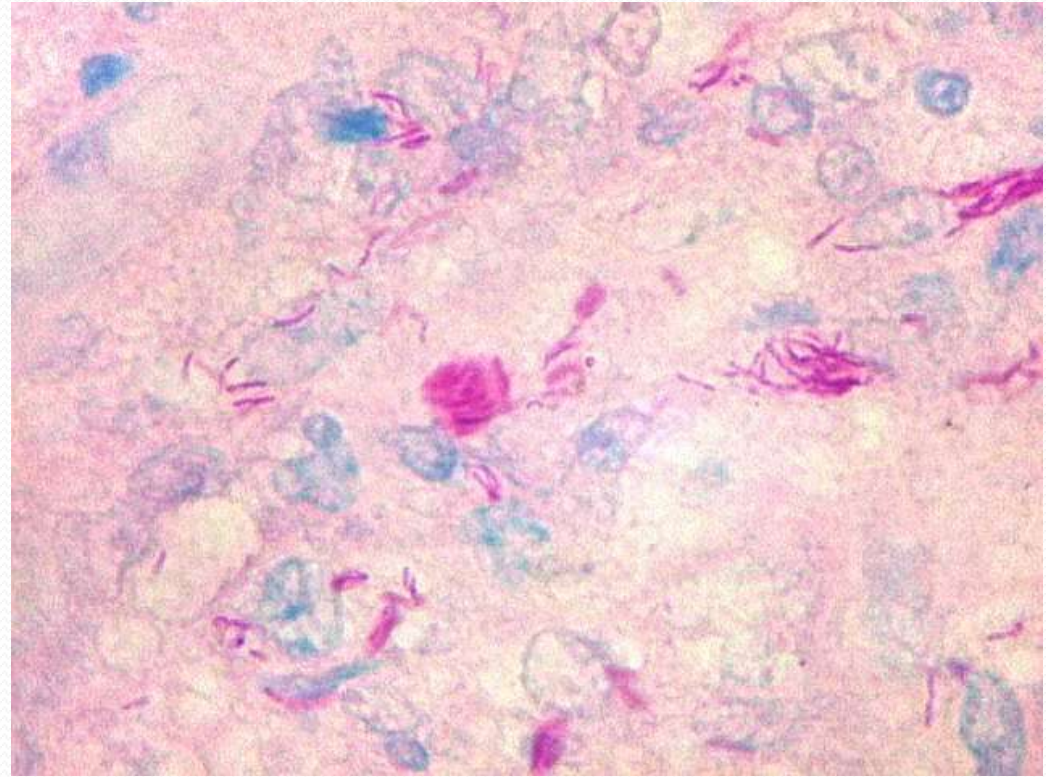
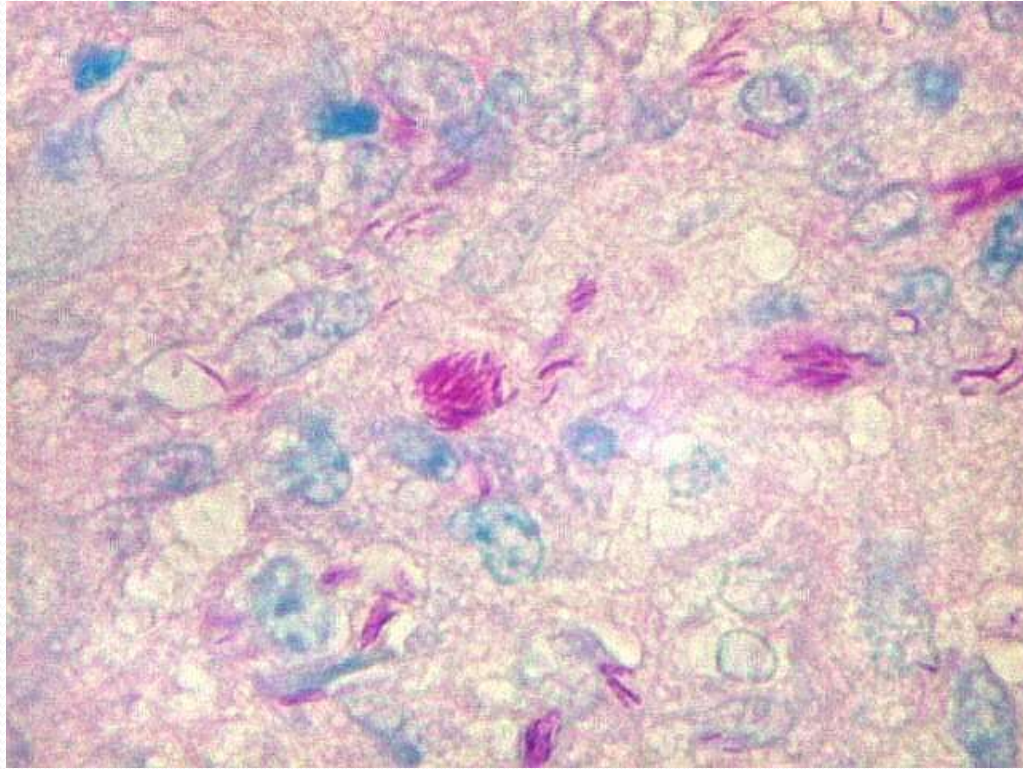
BACILOS DE HANSEN



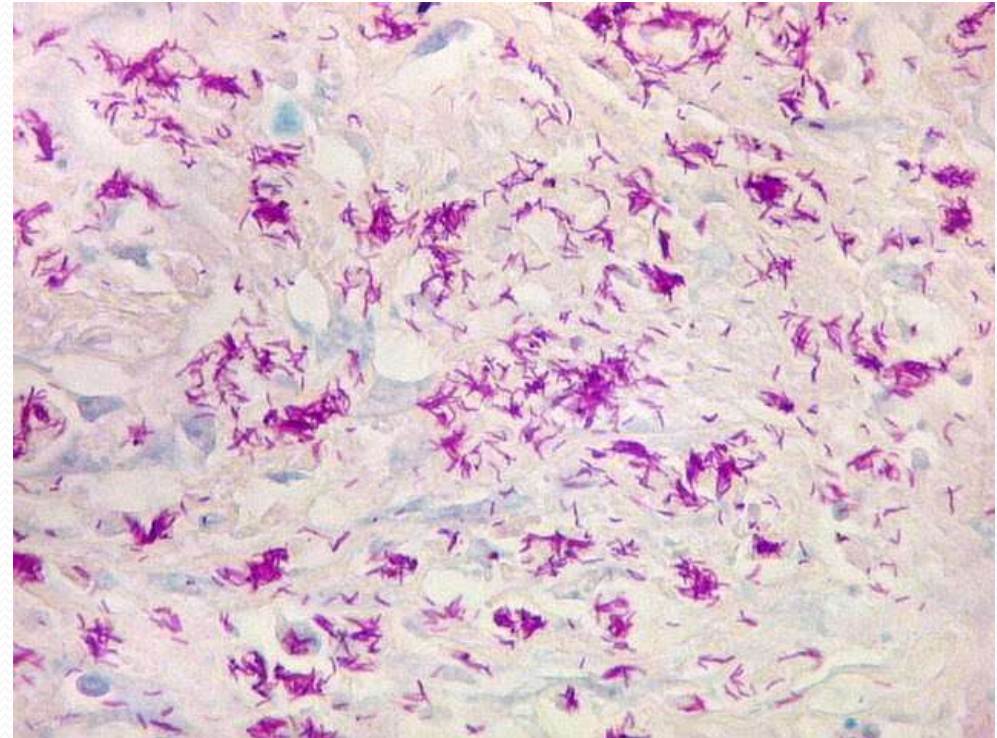
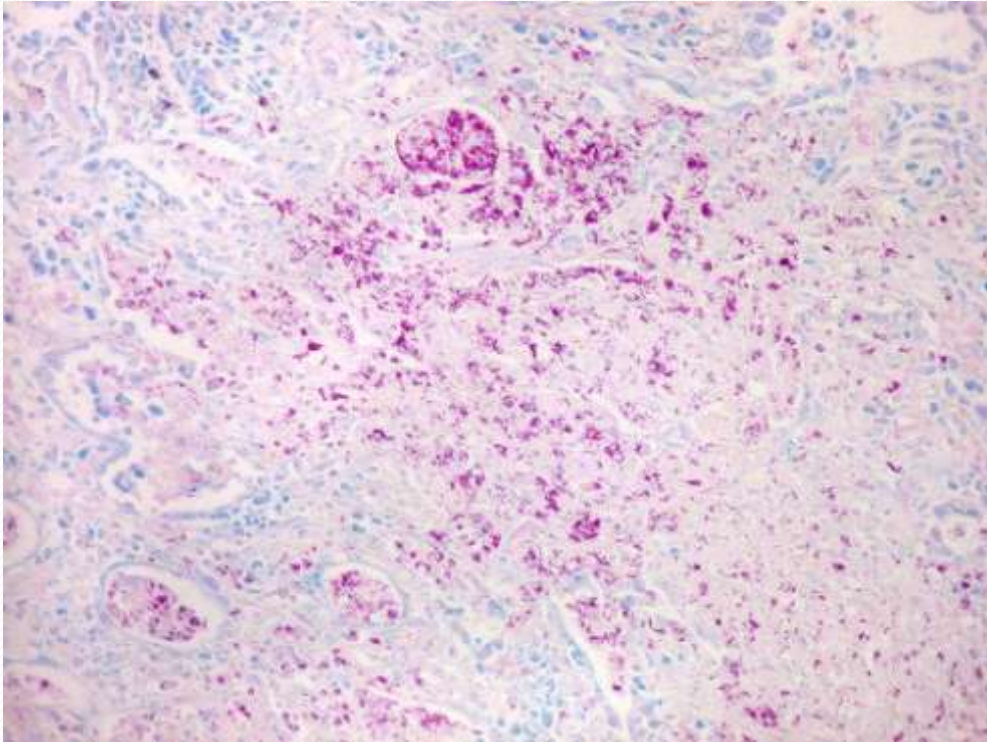
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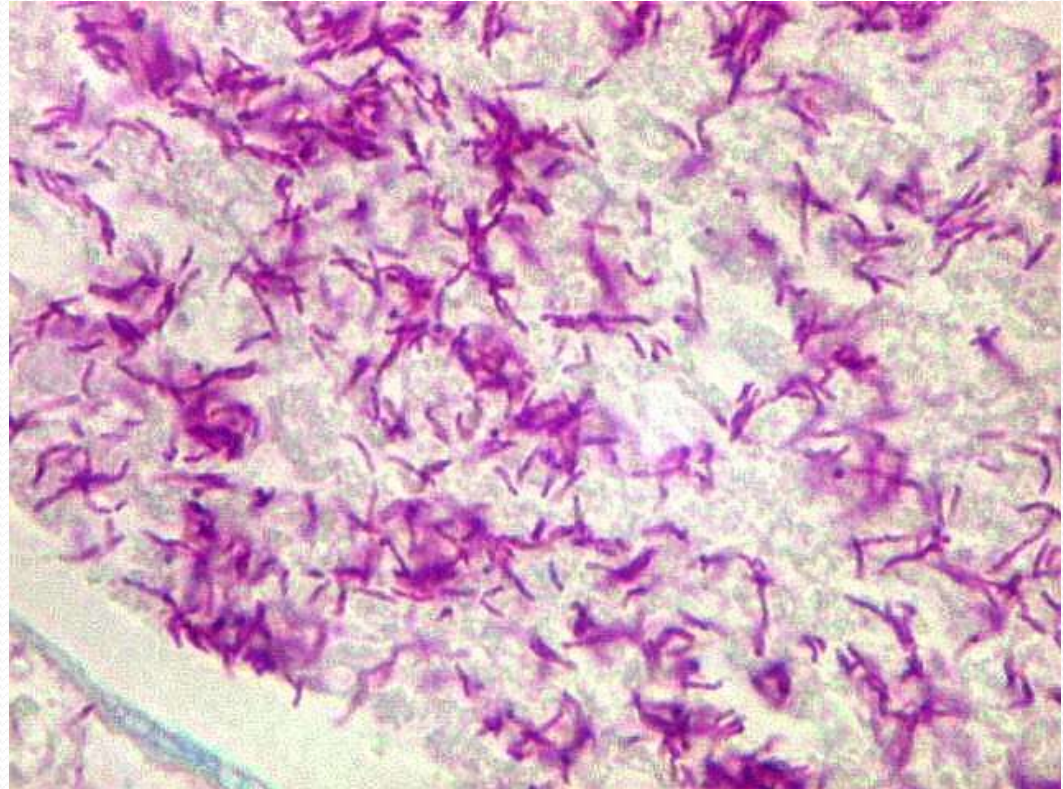
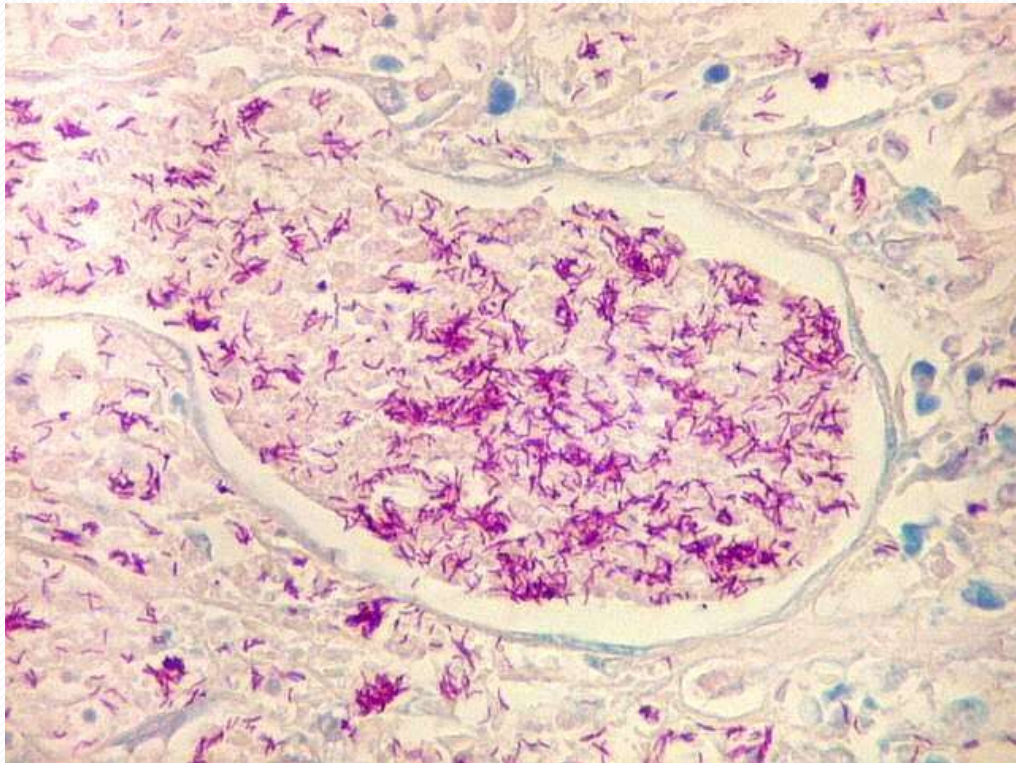
BACILOS DE HANSEN



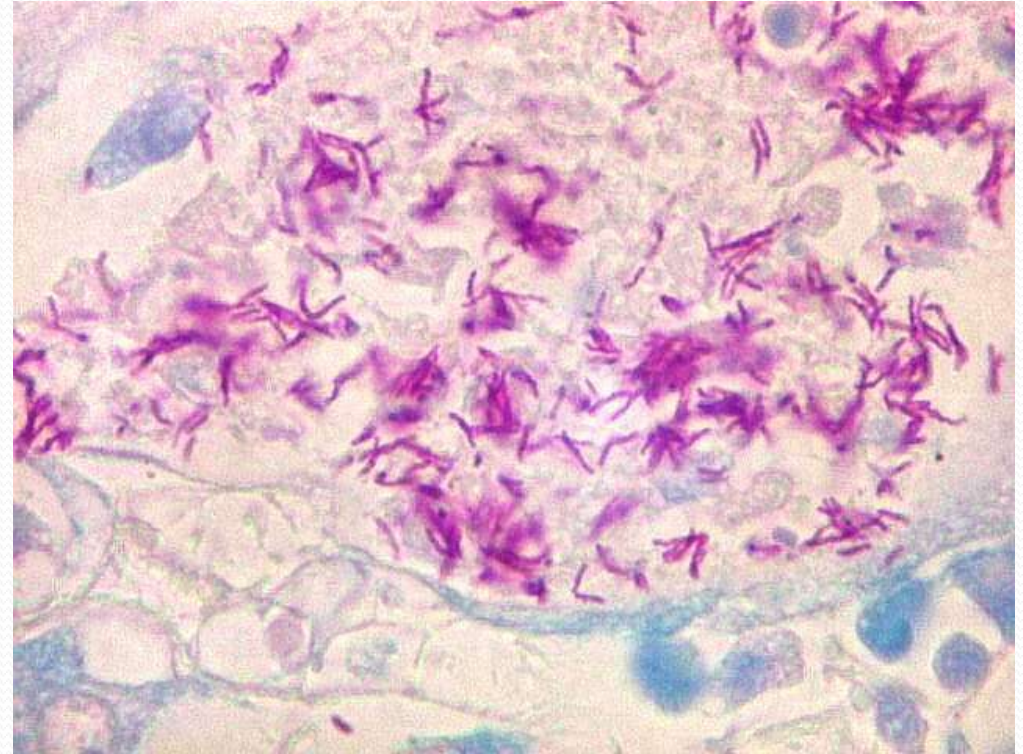
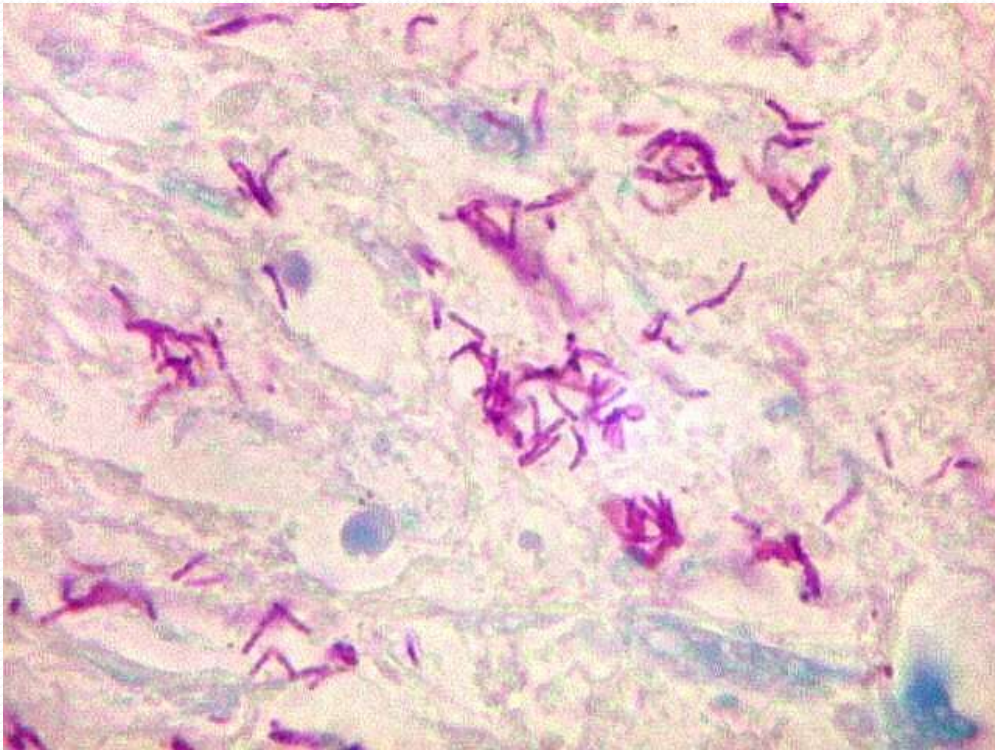
ZIEHL NEELSEN (MICOBACTERIOSE)



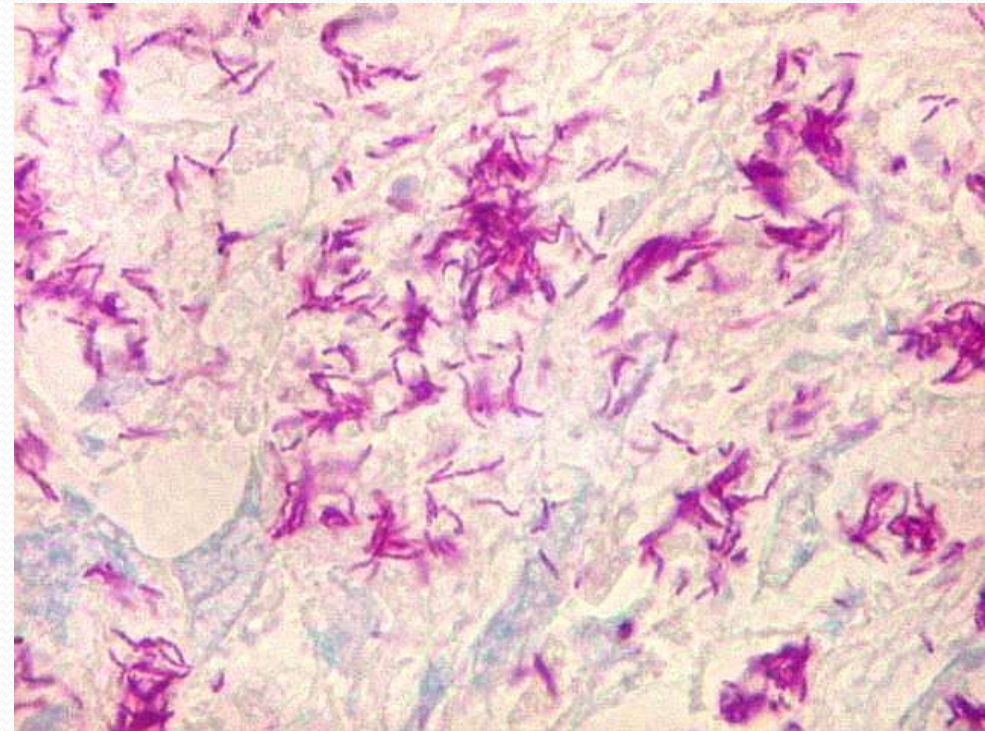
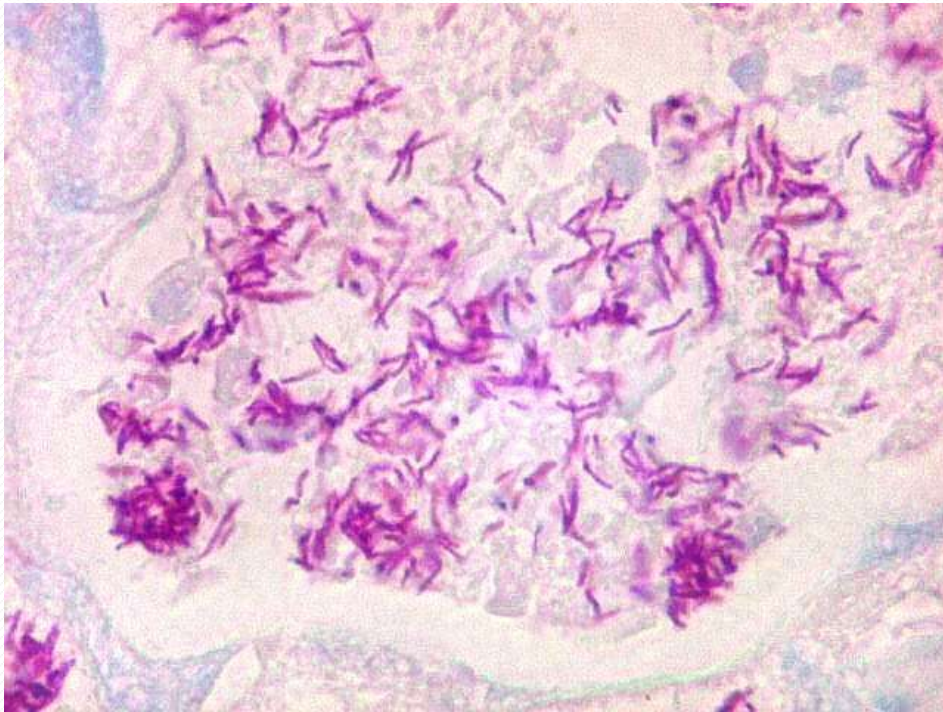
ZIEHL NEELSEN (MICOBACTERIOSE)



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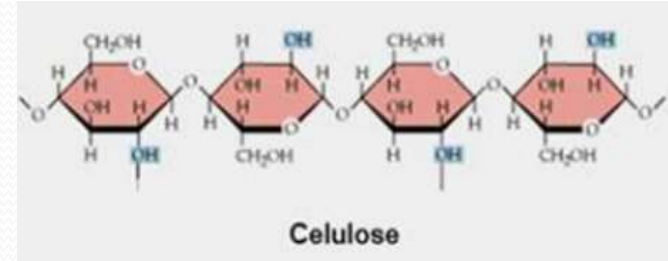


ZIEHL NEELSEN (MICOBACTERIOSE)



MUCOPOLISSACARÍDEOS

São longas cadeias de moléculas de açúcares usadas na construção dos ossos, cartilagens, pele e tendões.



O fluído que lubrifica as suas articulações contém mucopolissacarídeos.

Todos os tecidos tem um pouco desta substância como parte normal de sua estrutura.

ENZIMAS: proteínas especiais responsáveis pelas reações que ocorrem em nosso corpo.

MÉTODO DE FERRO - COLOIDAL DE MOWRY

1-SOLUÇÃO ESTOQUE DE CLORETO FÉRRICO 29%

- Água destilada 100 mL
- Cloreto férrico 29 g

2-SOLUÇÃO MATRIZ DE FERRO COLOIDAL DE MULLER

- Água destilada 250 mL

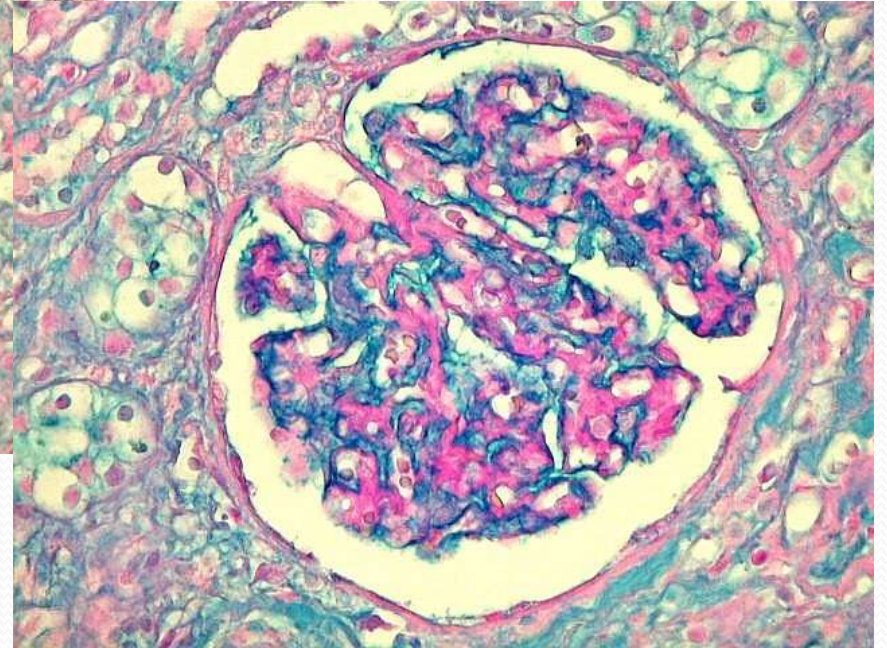
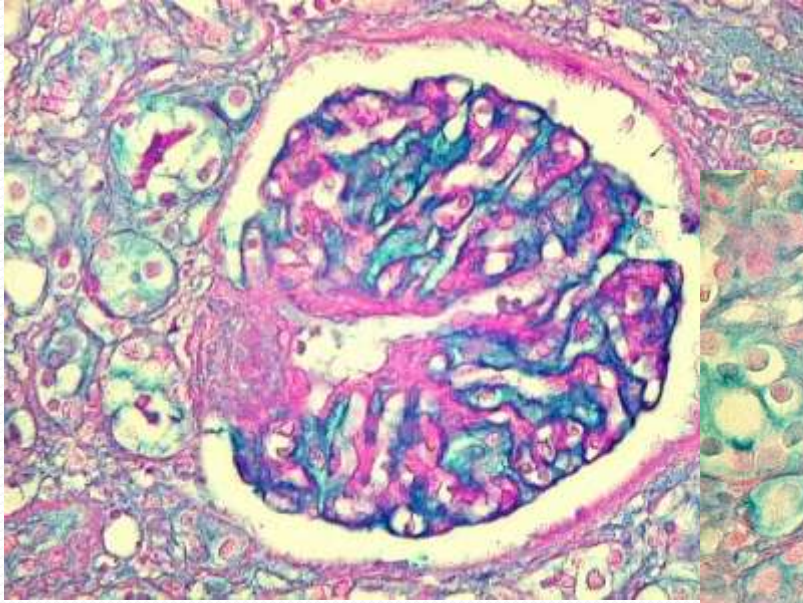
***Ferver até ebulição**

- Adicionar cloreto férrico 29% 4 mL

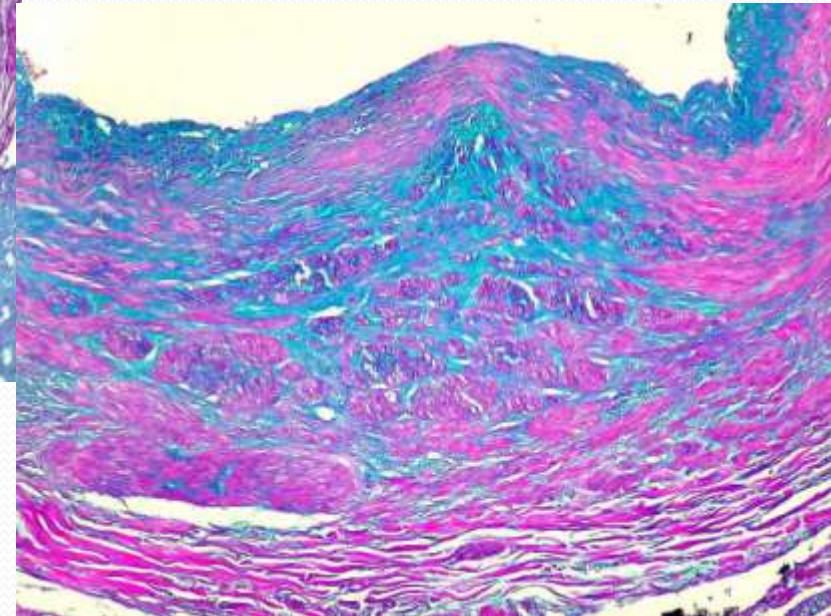
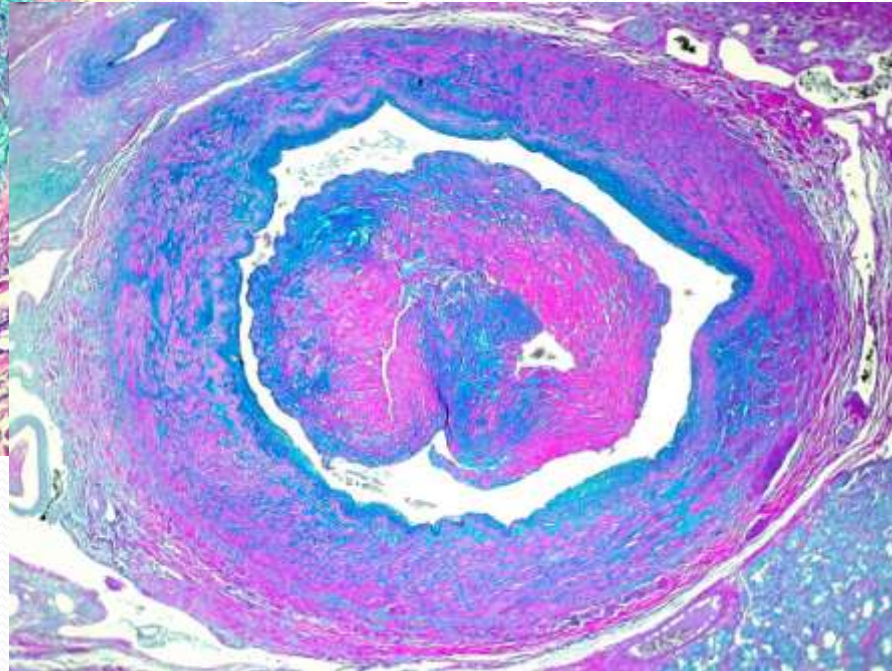
3- SOLUÇÃO USO DIÁRIO DE FERRO COLOIDAL DE MULLER

- Solução matriz de ferro coloidal de Muller 20 mL
- Água destilada 15 mL
- Ácido acético glacial 5 mL

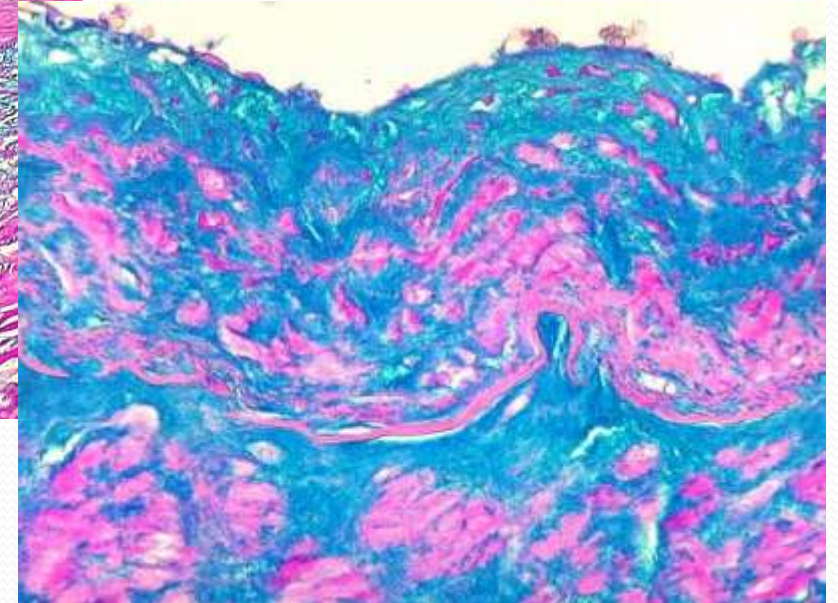
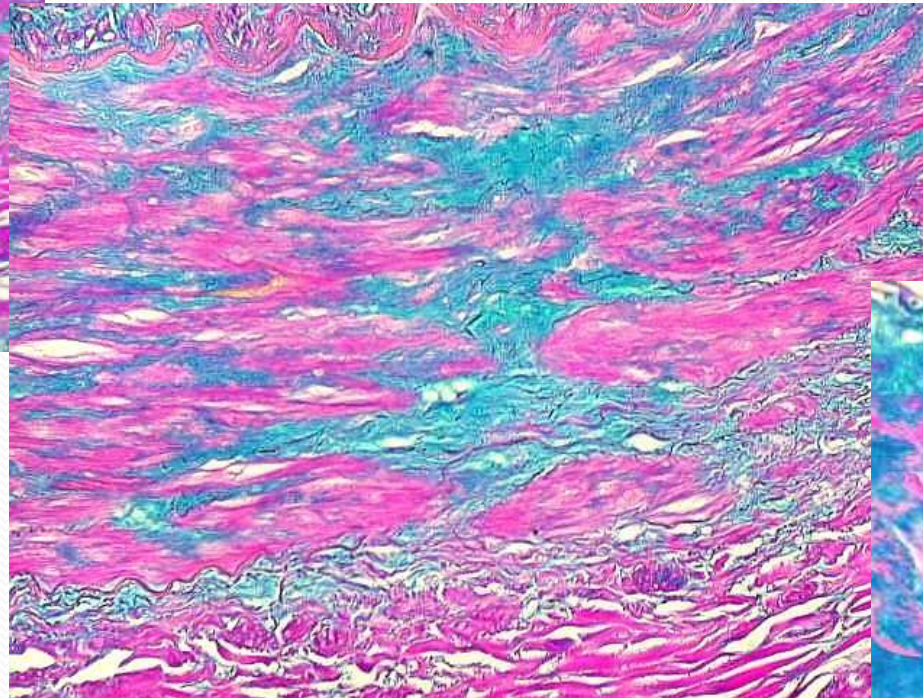
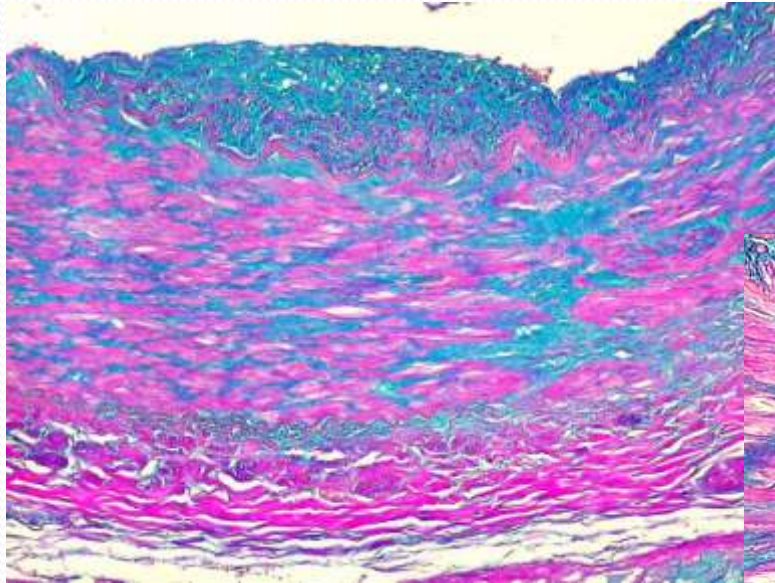
FERRO COLOIDAL



FERRO COLOIDAL



FERRO COLOIDAL



BIBLIOGRAFIA

- Técnica histológica em anatomia patológica – prof. Jorge Michalany;
- Métodos histológicos (Instituto de Patologia das Forças Armadas dos Estados Unidos da América (AFIP);
- Introdução as mucopolissacaridoses;
- McManus JFA, MowryRW;
- Staining methods histologic and histochemical, New York, NY Hoeber, 1960:133

AGRADECIMENTOS

- Edna Tereza Fidelix (minha esposa)
- Irineu Antônio Mantovanelli Neto – técnico de laboratório
- Isabella Domingues Guize – técnica de laboratório
- Dra. Dulcelena Ledo de Oliveira- patologista do laboratório
- Dr. Luciano de Souza Queiroz – patologista Unicamp (Imagens)

