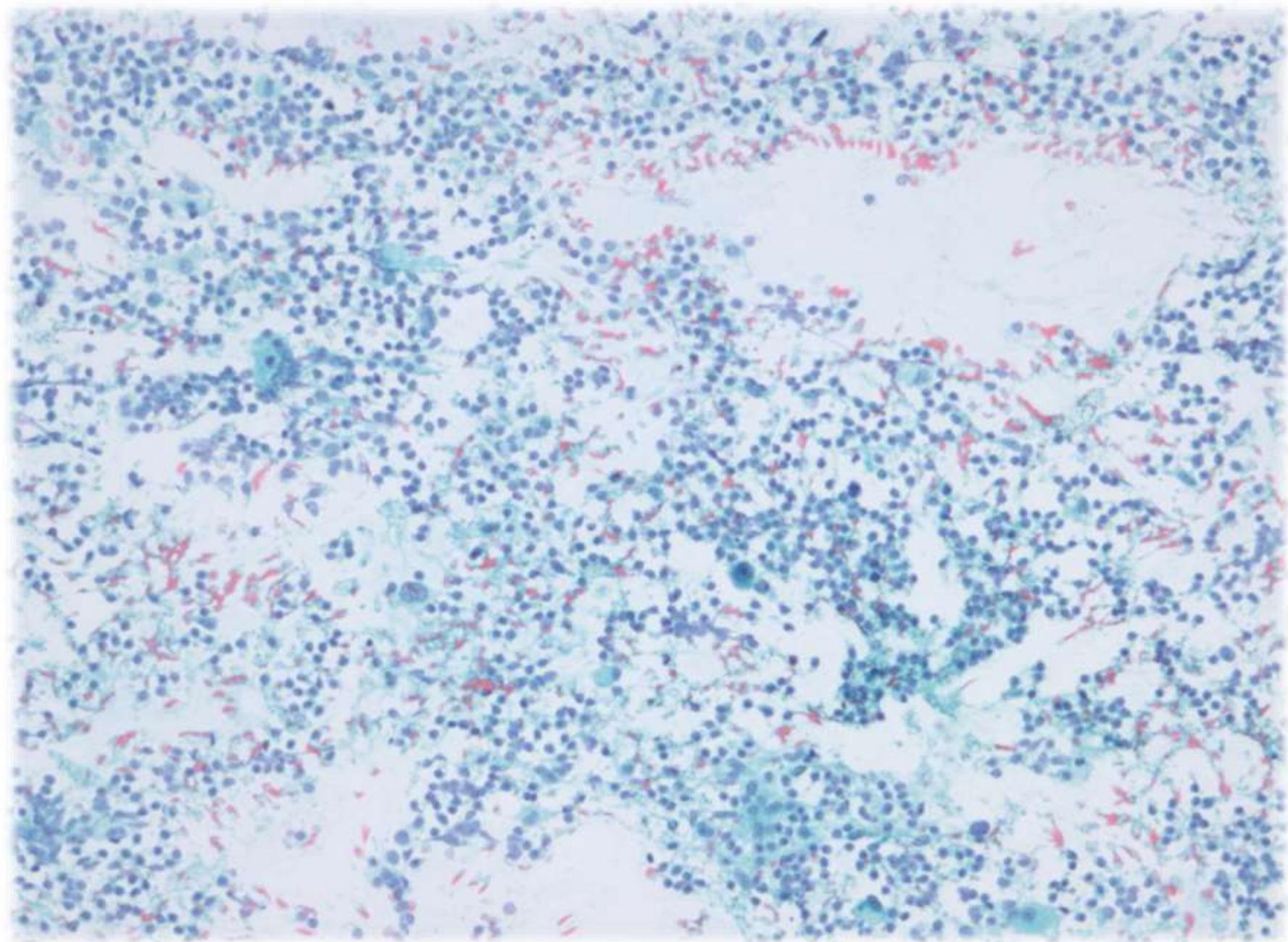


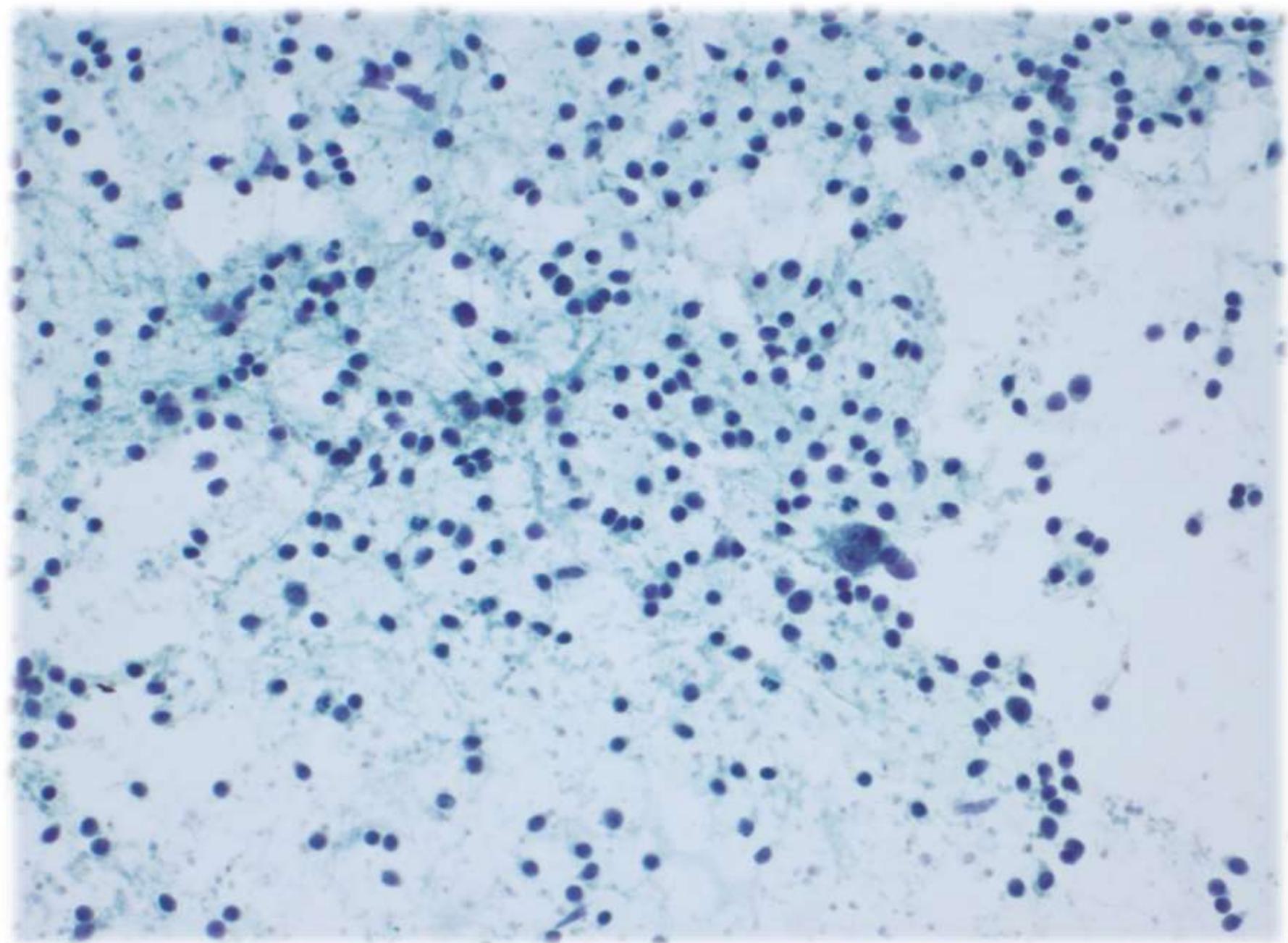
39-jähriger Mann

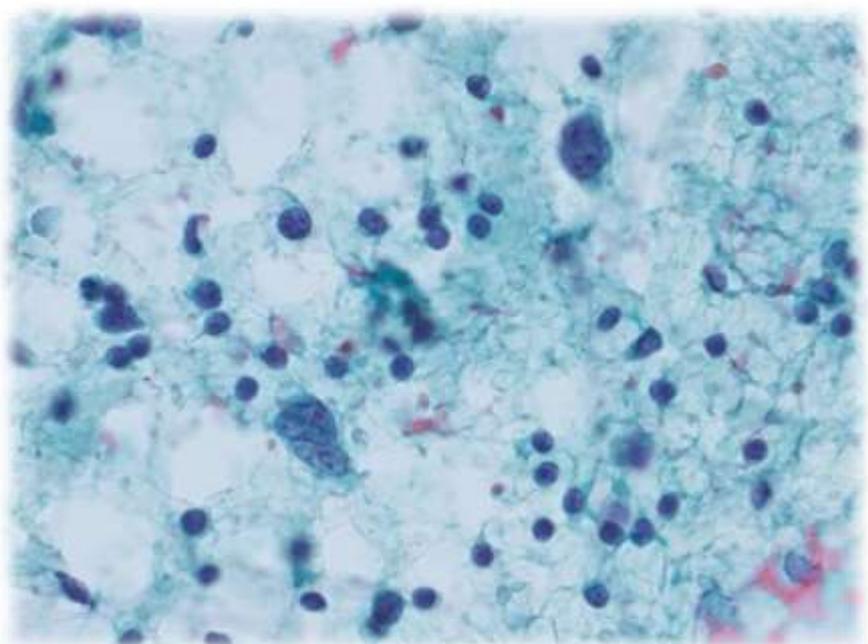
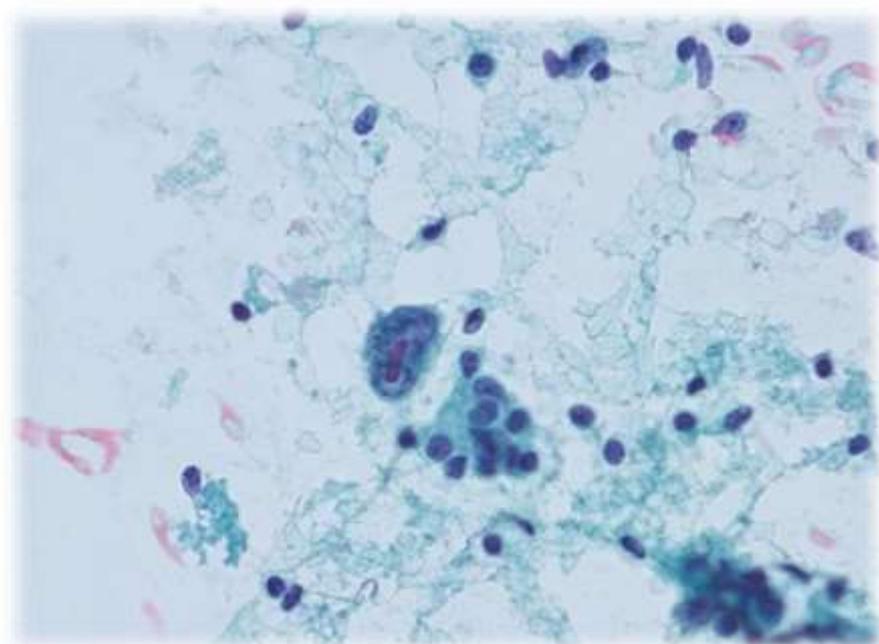
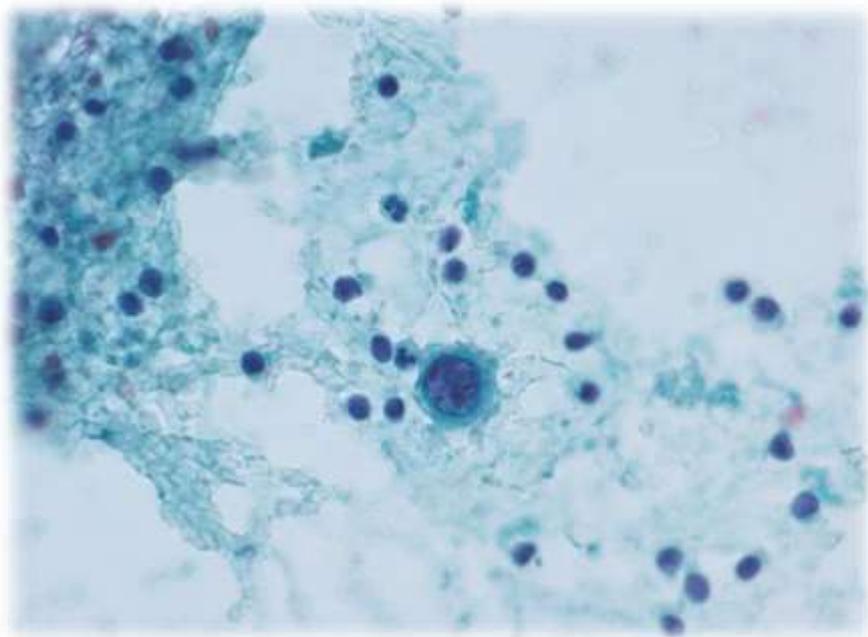
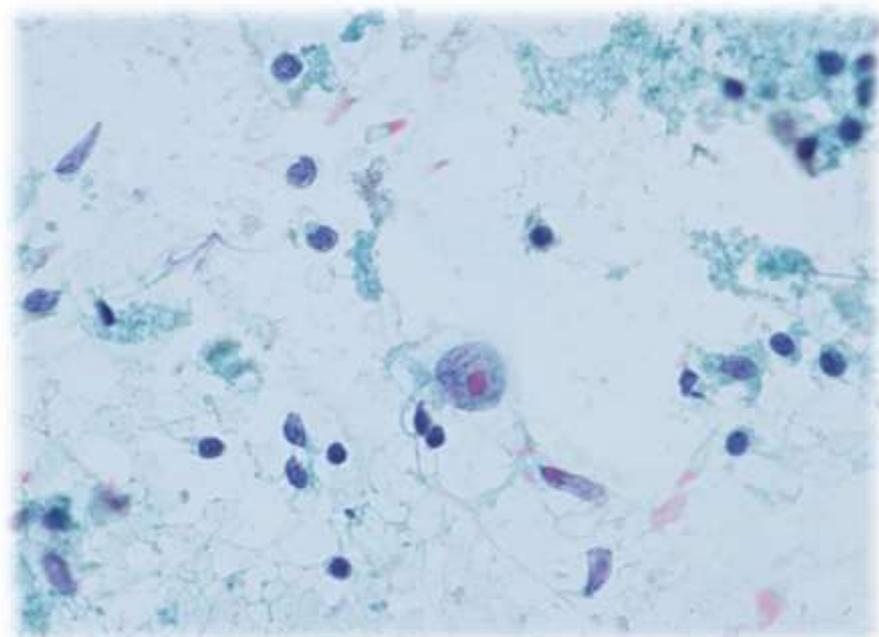
schmerzlose Schwellung zervikal seit drei Wochen

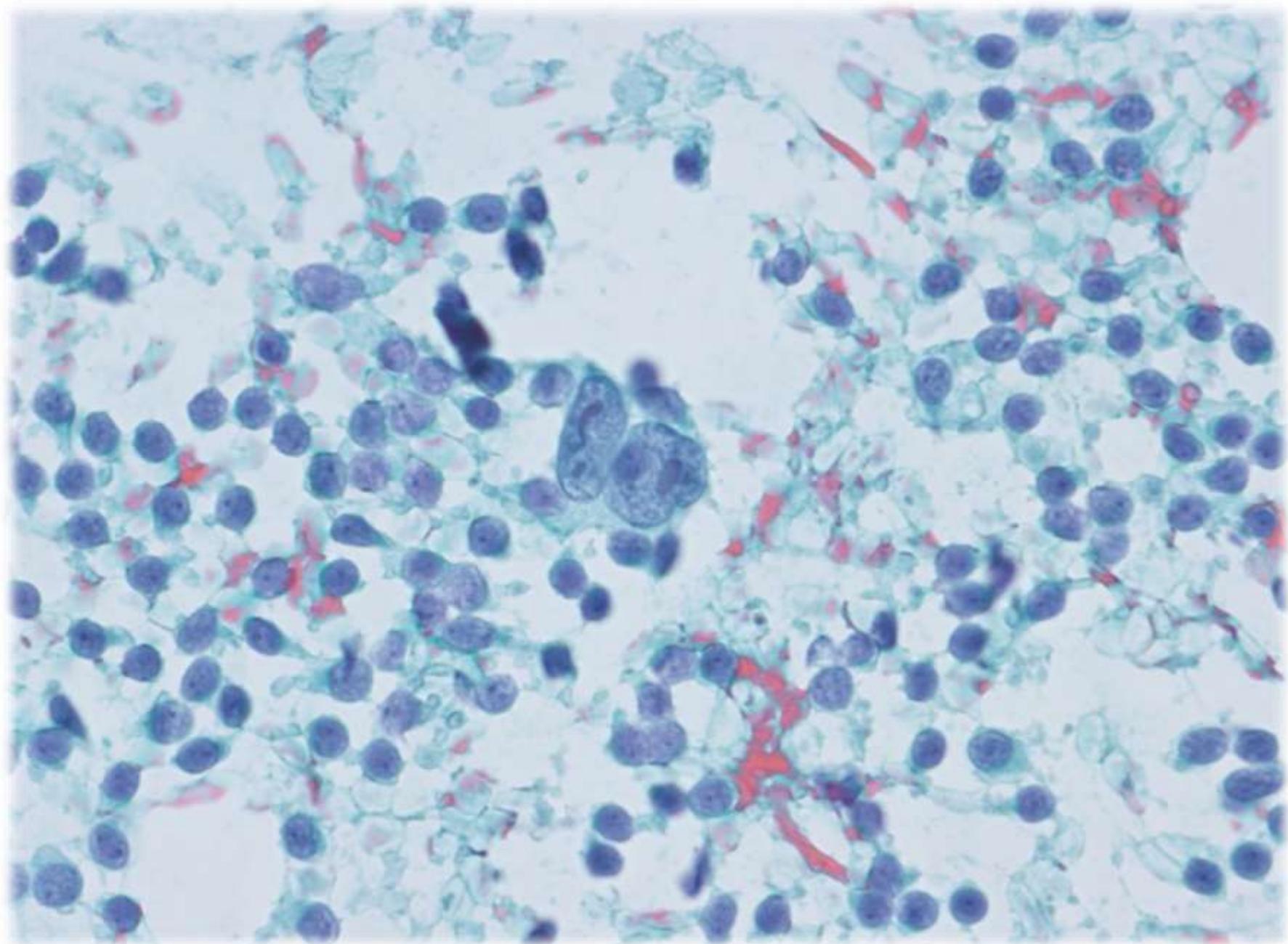


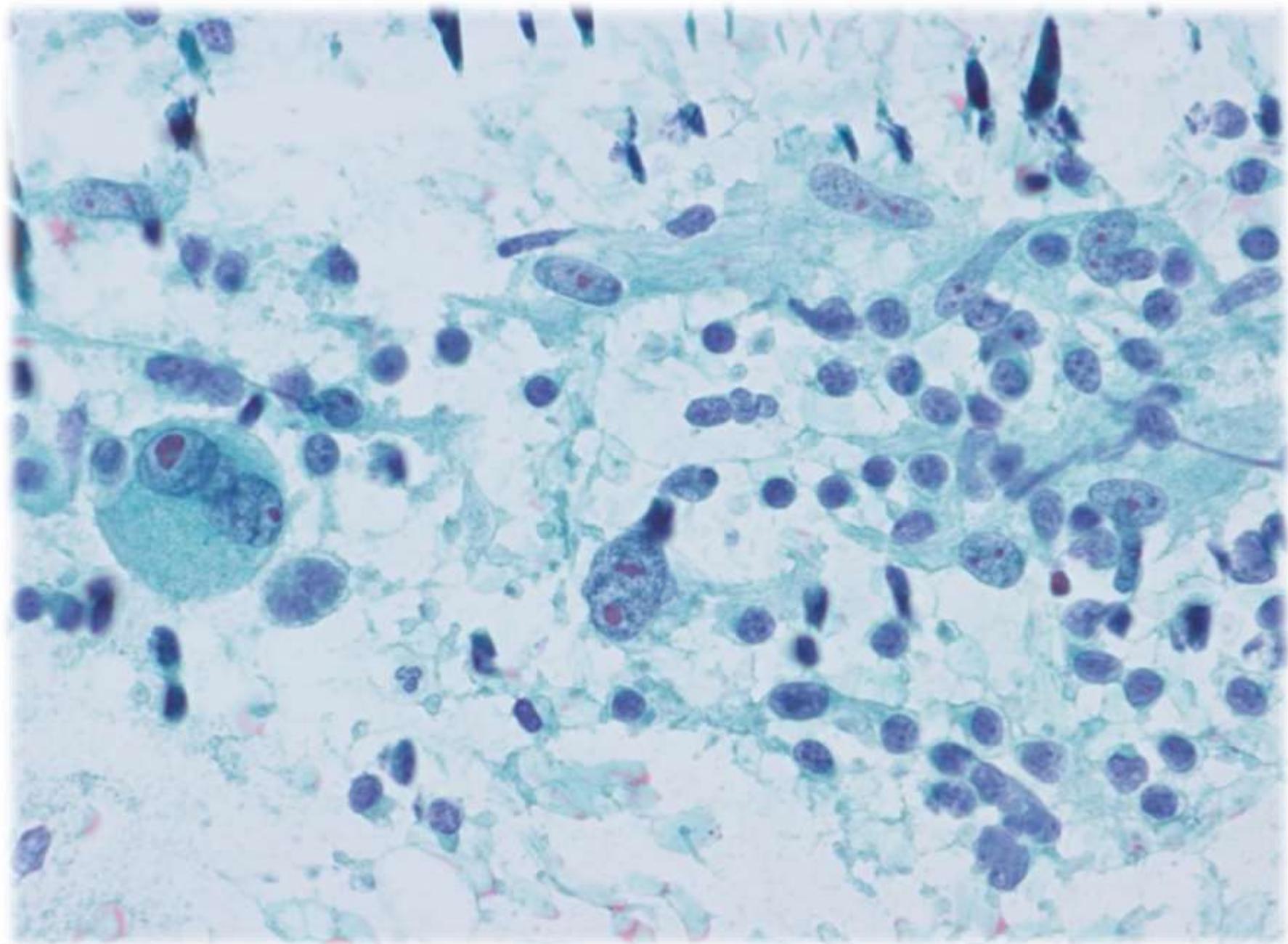
FNP

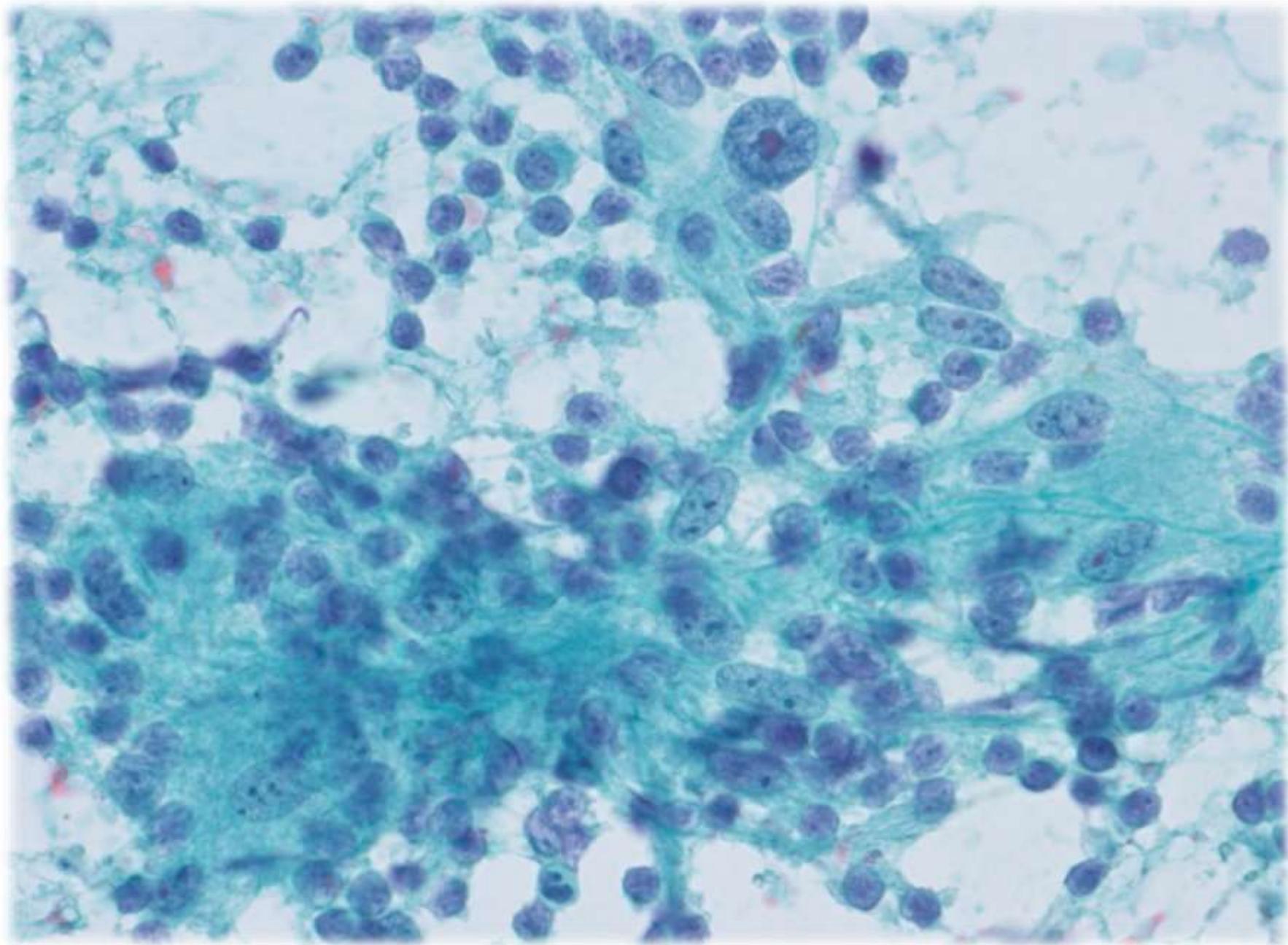






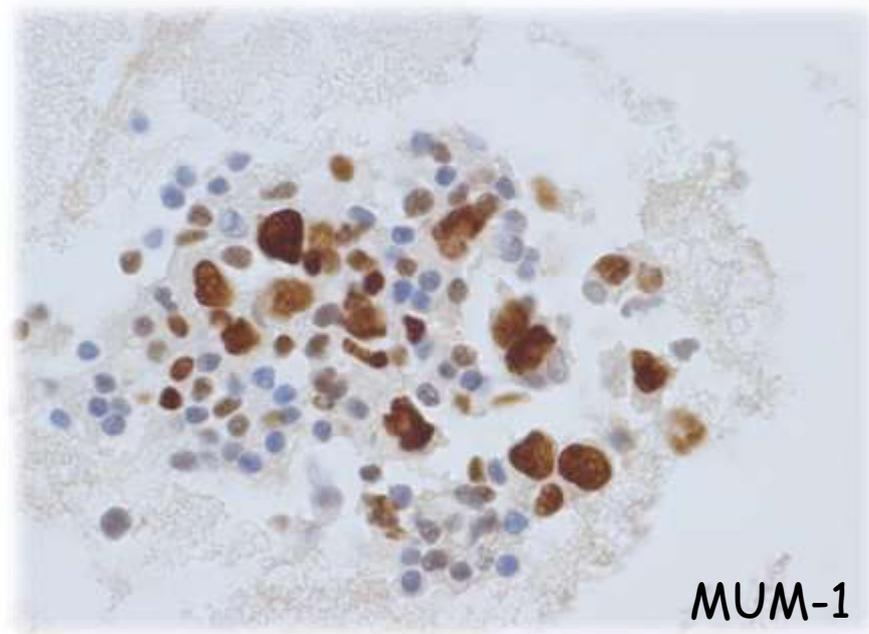
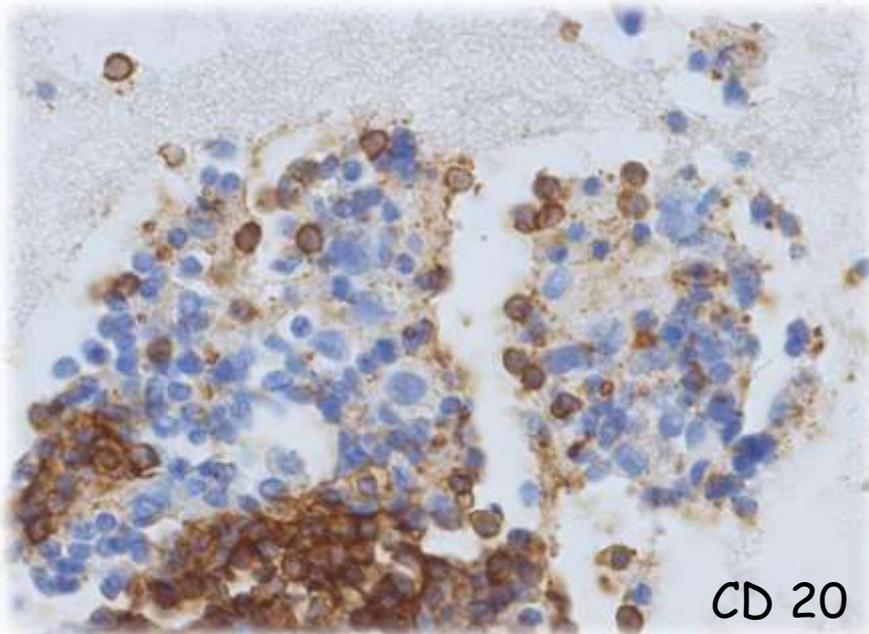
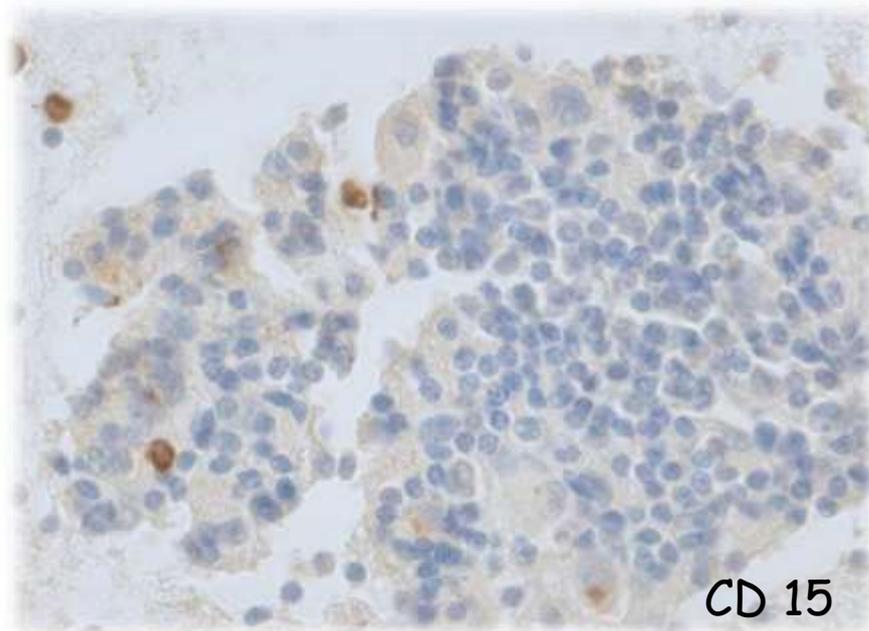
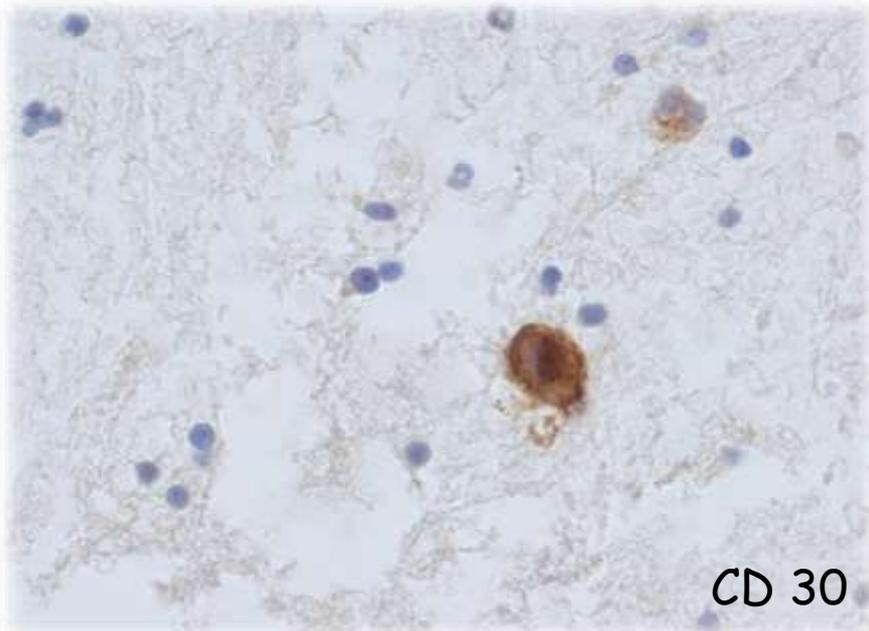






Differentialdiagnose

- T-Zell- und Histiocyten-reiches B-NHL
- granulomatöse Lymphadenitis
- Hodgkin Lymphom
- Melanommetastase
- anaplastisches NHL
- Karzinommetastase

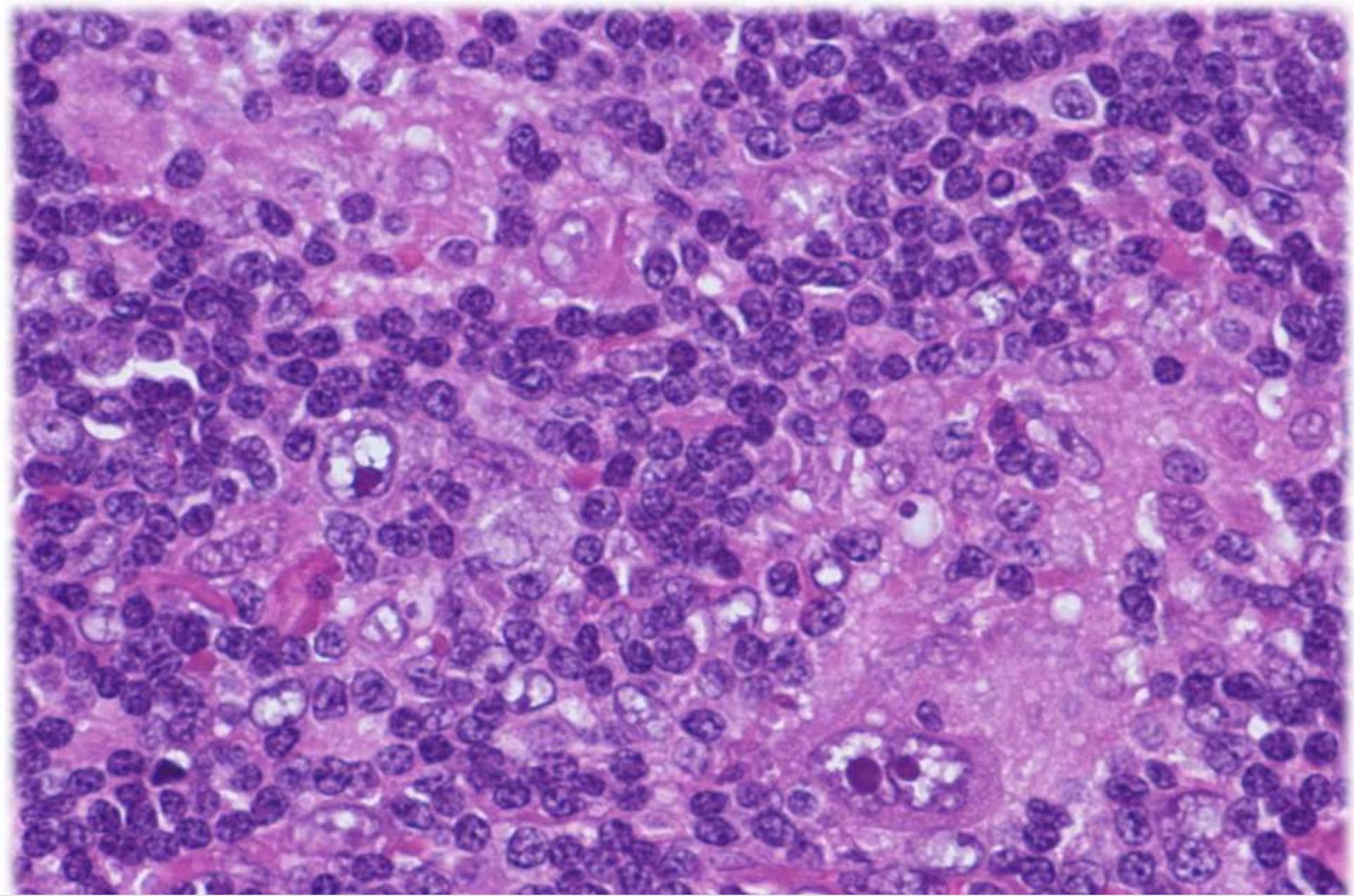


Hodgkin Lymphom

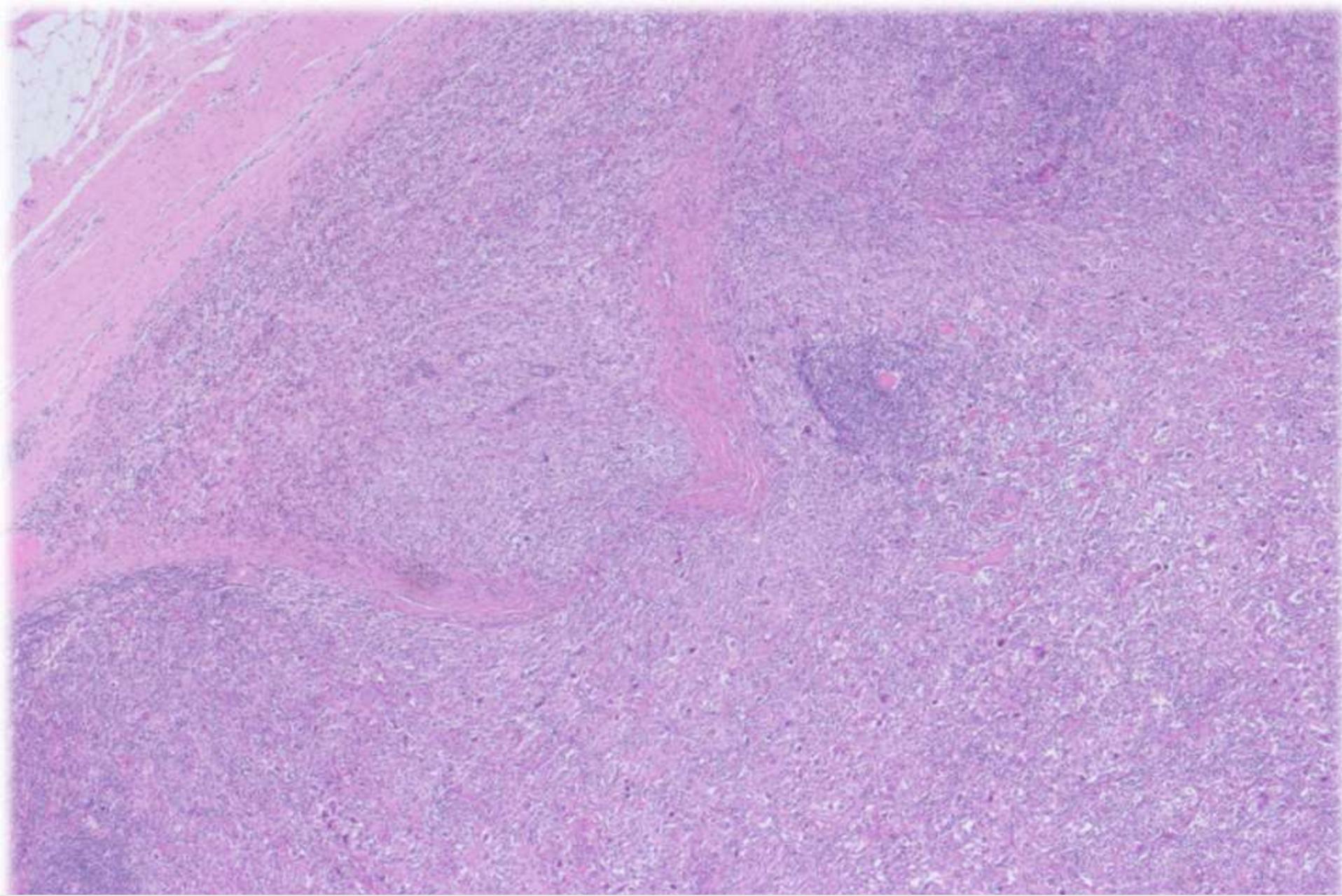
- junge Erwachsene; zweiter Gipfel im Senium
- bevorzugter Befall von zervikalen und mediastinalen Lymphknoten, oftmals Lymphknotenkonglomerate
- primär extranodaler Befall insgesamt selten, wenngleich die Milz öfters befallen ist

Hodgkin Lymphom

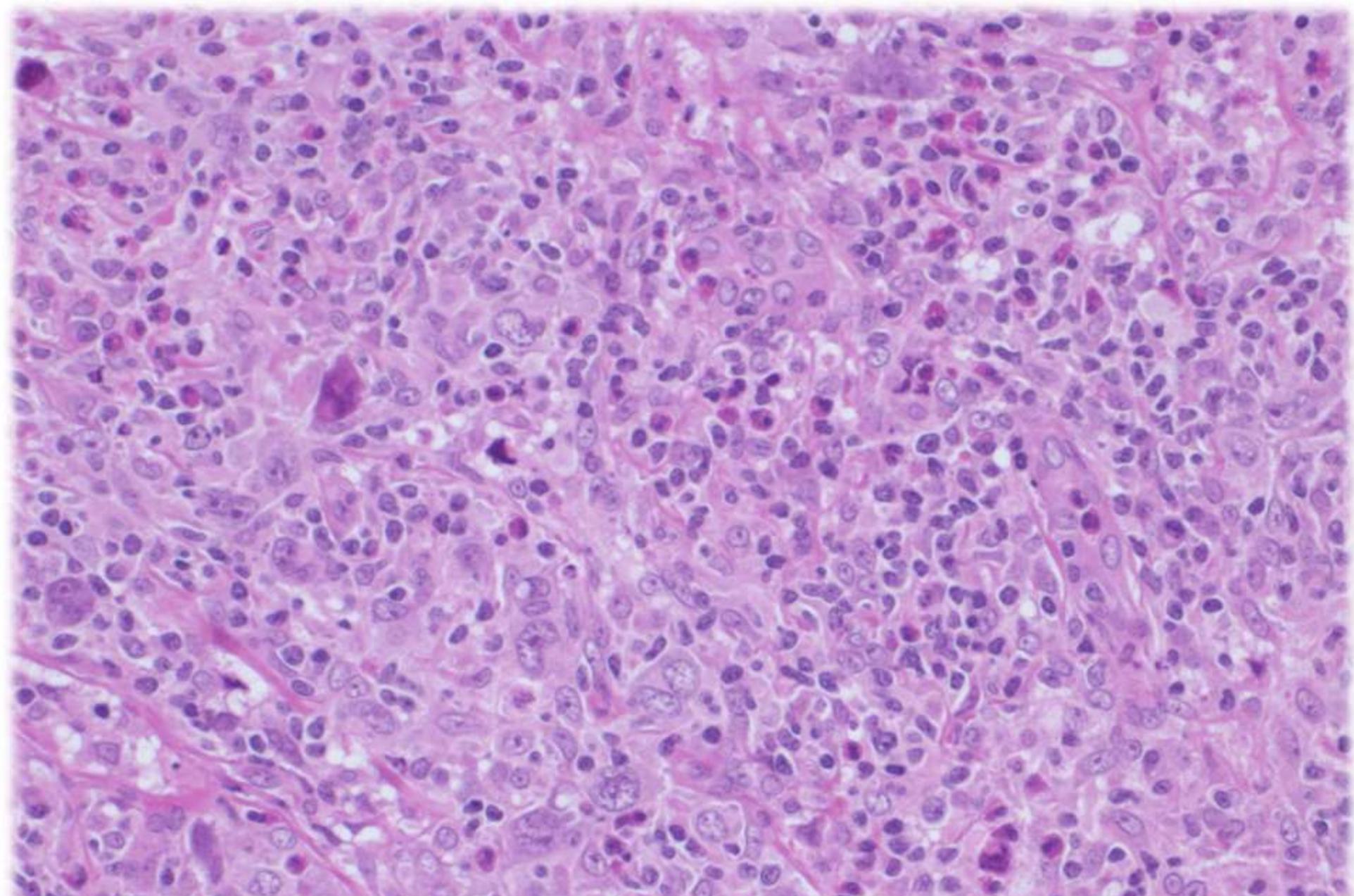
- lymphozytenreiches klassisches HL
- Klassisches HL, noduläre Sklerose
- gemischtzelliges klassisches HL
- lymphozytenarmes klassisches HL
- noduläres lymphozytenprädominantes HL
(früher Paragranulom)



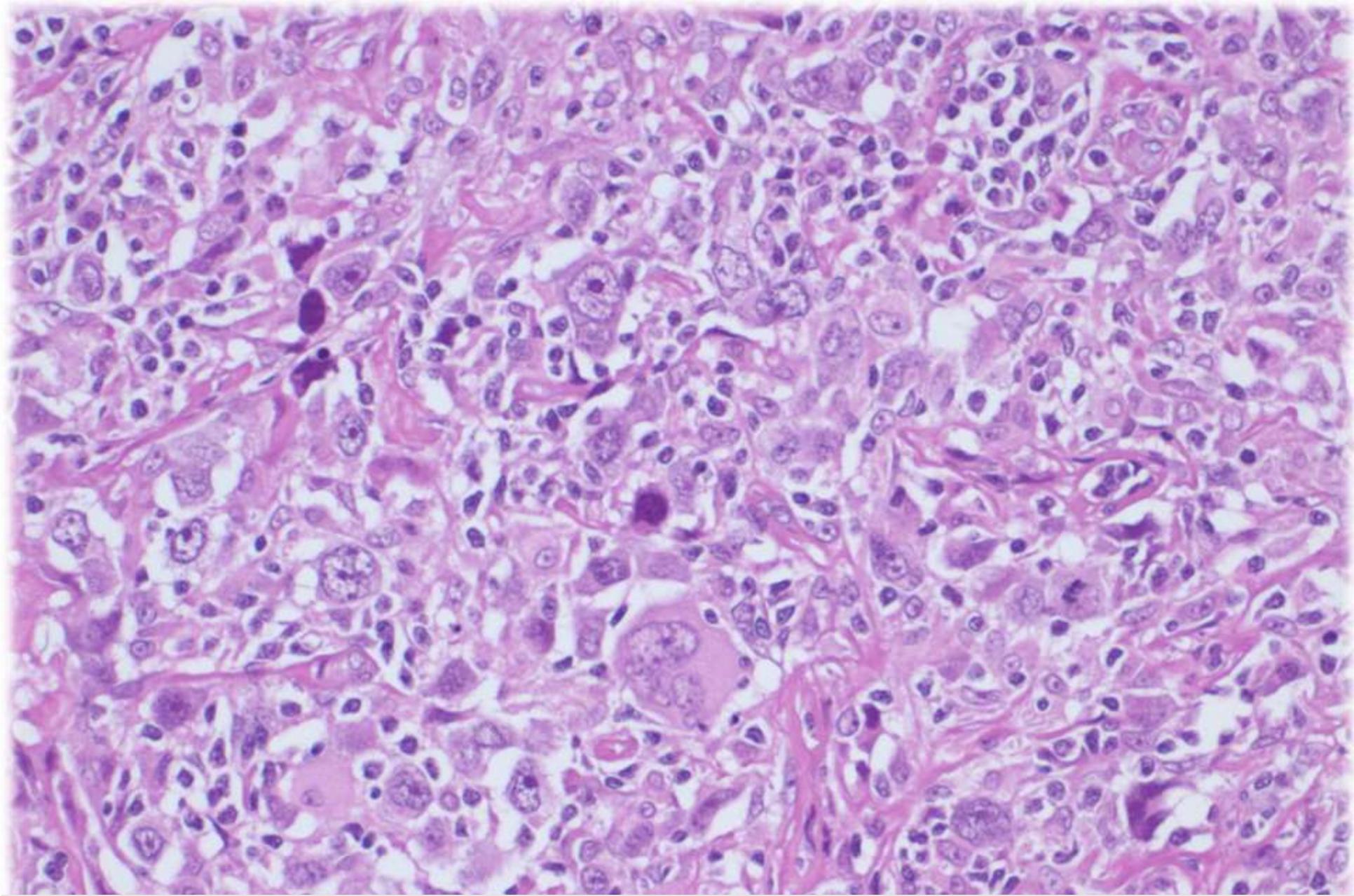
lymphozytenreiches klassisches HL



klassisches HL, noduläre Sklerose

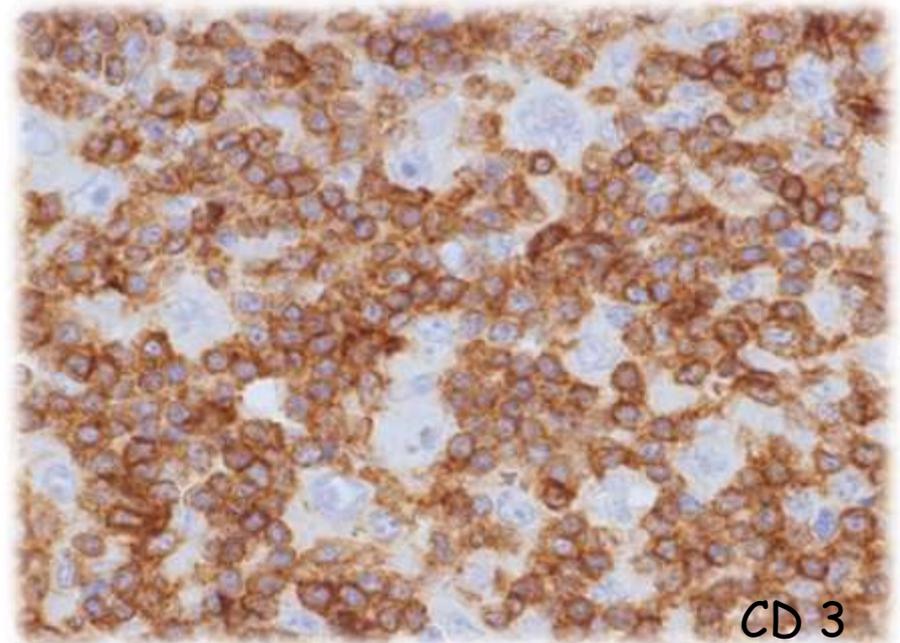
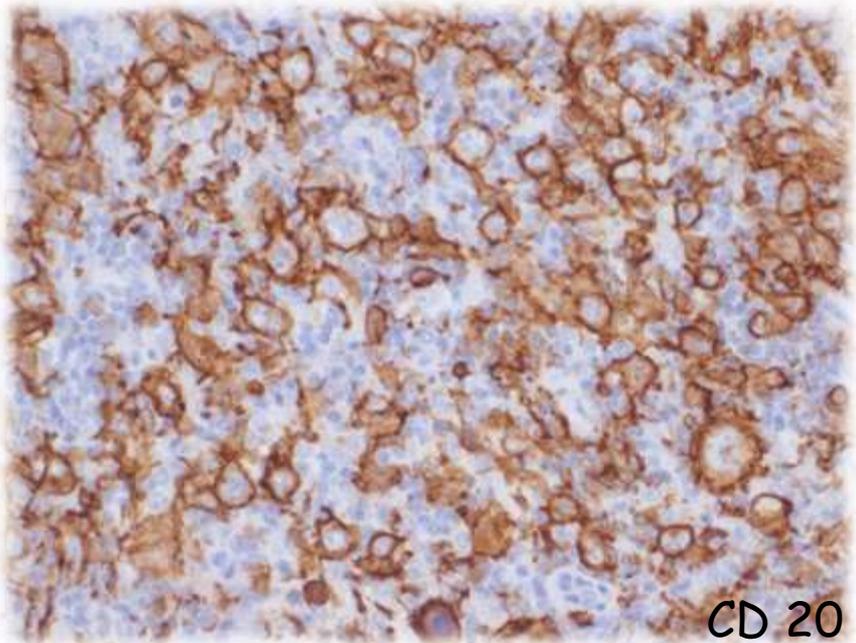
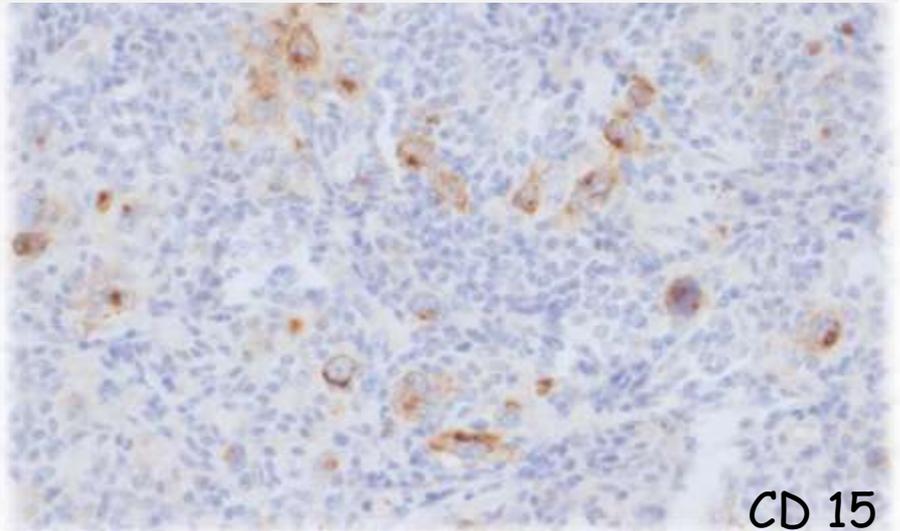
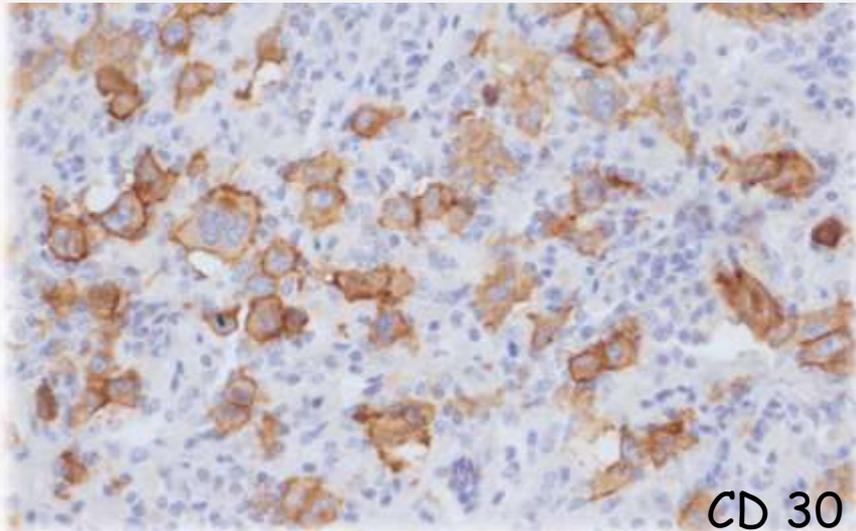


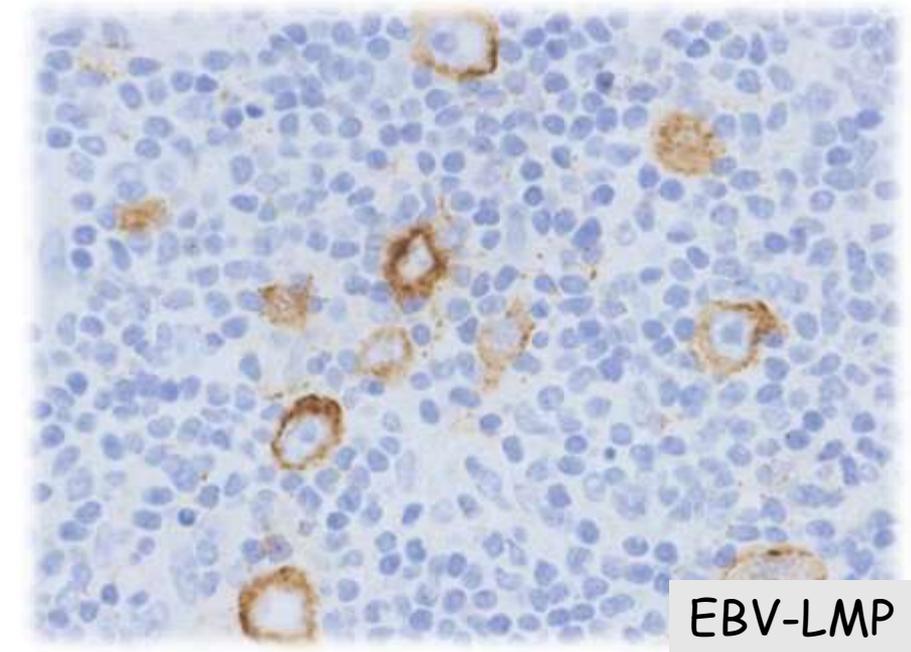
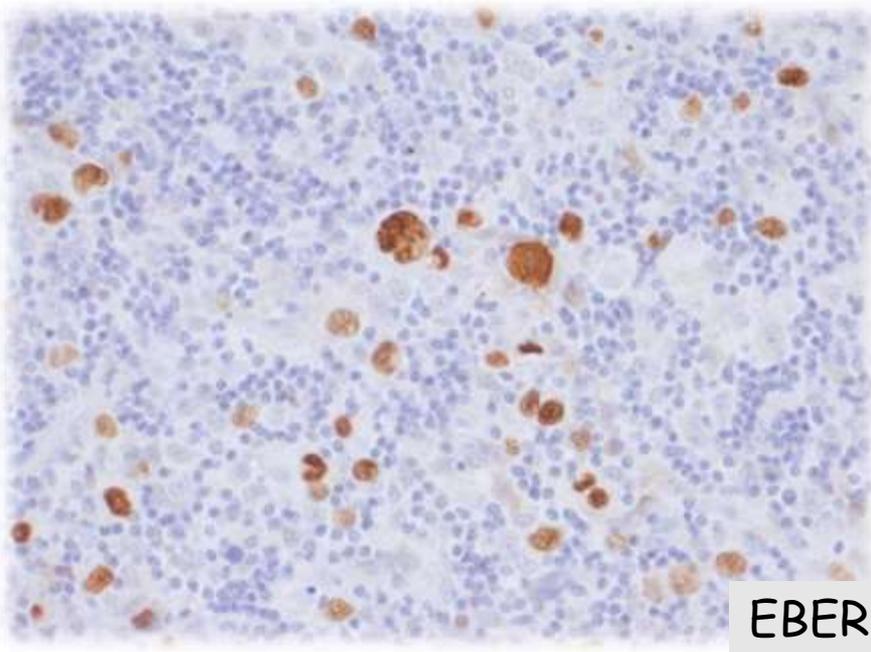
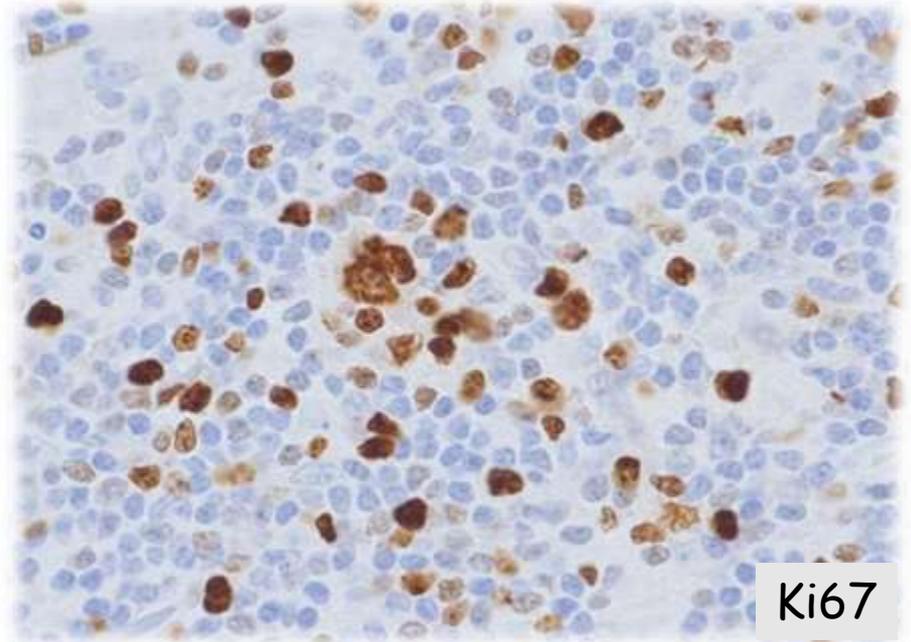
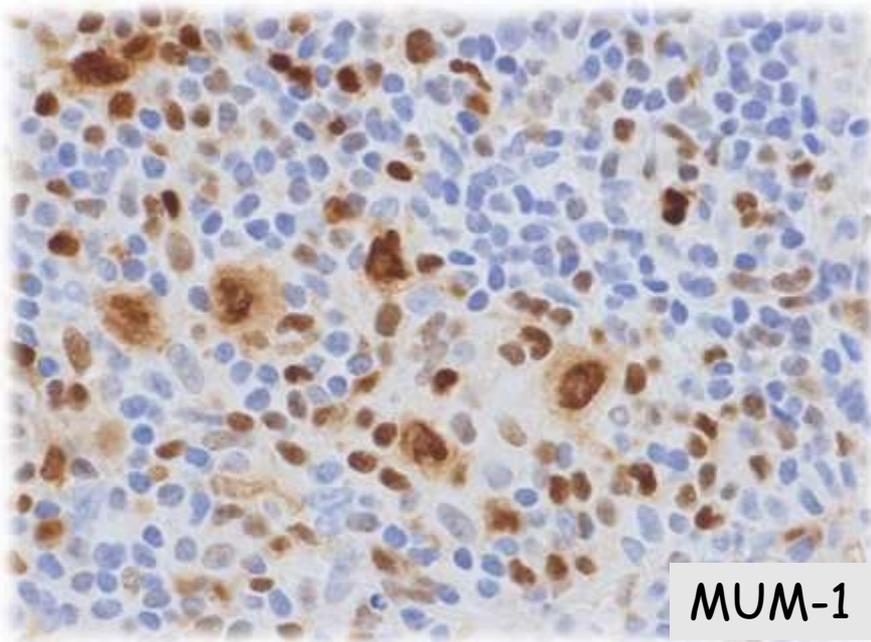
gemischtzelliges klassisches HL



lymphozytenarmes klassisches HL

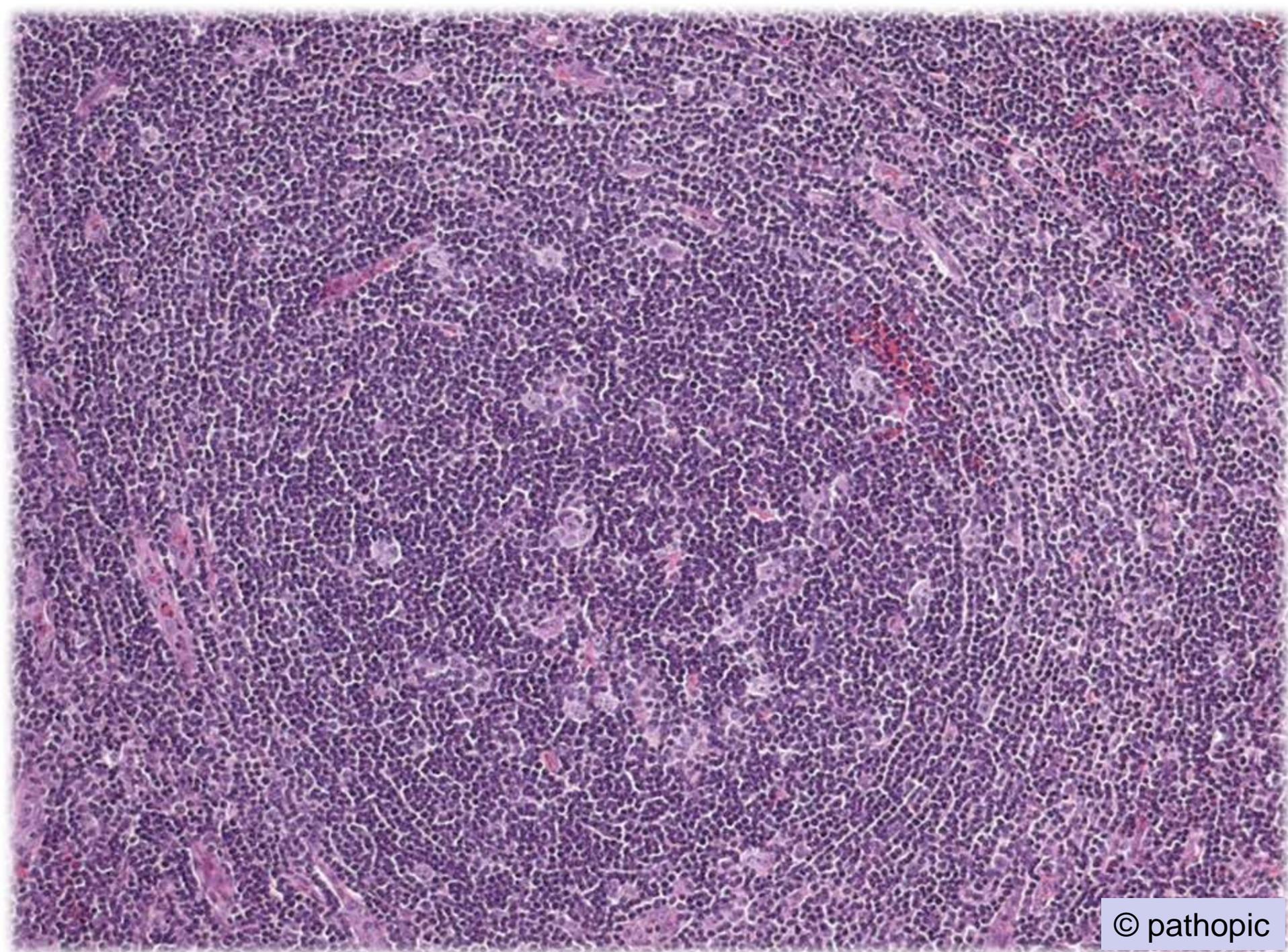
klassischer Immunophänotyp



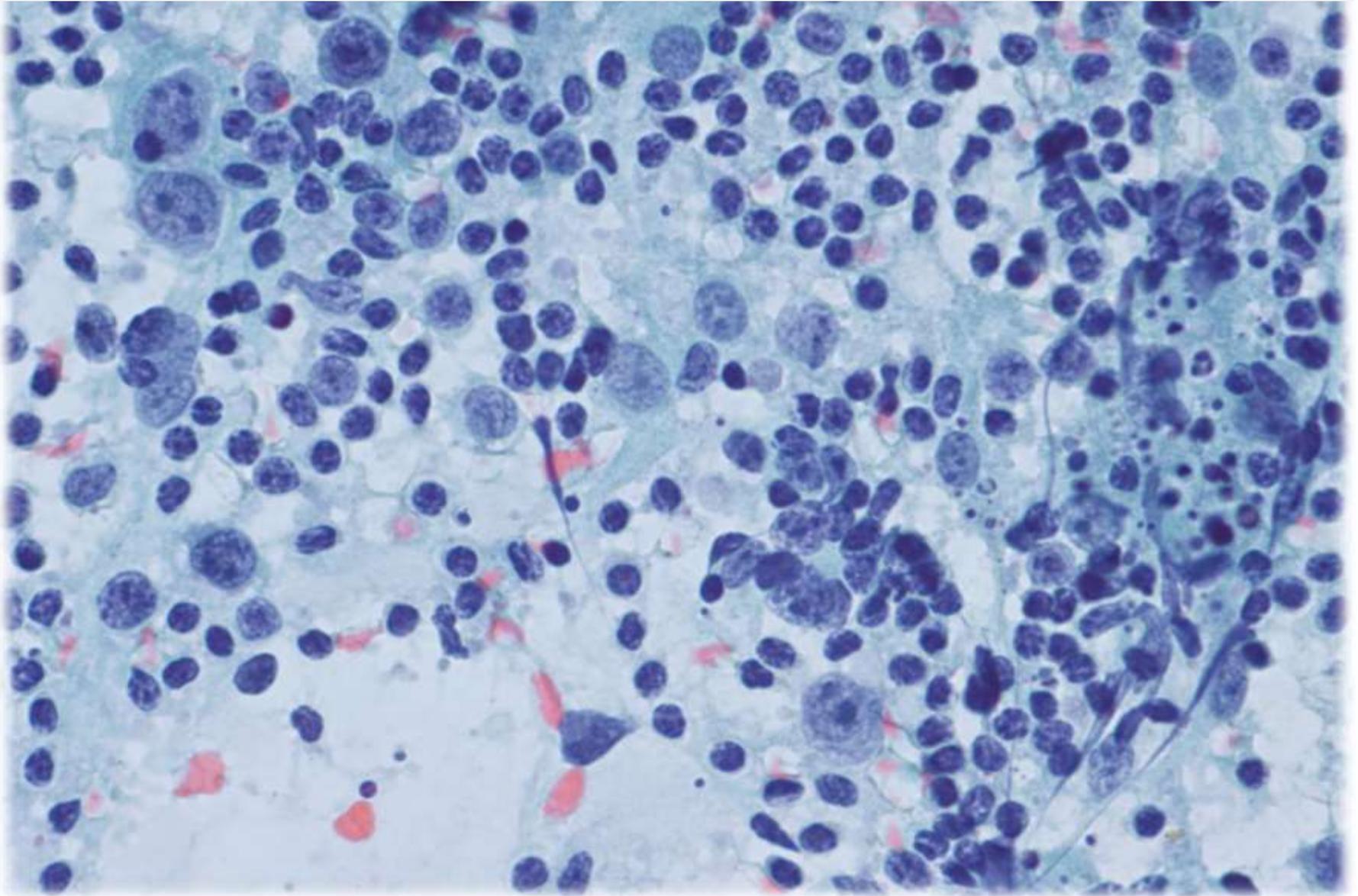


noduläres lymphozytenprädominantes Hodgkin Lymphom

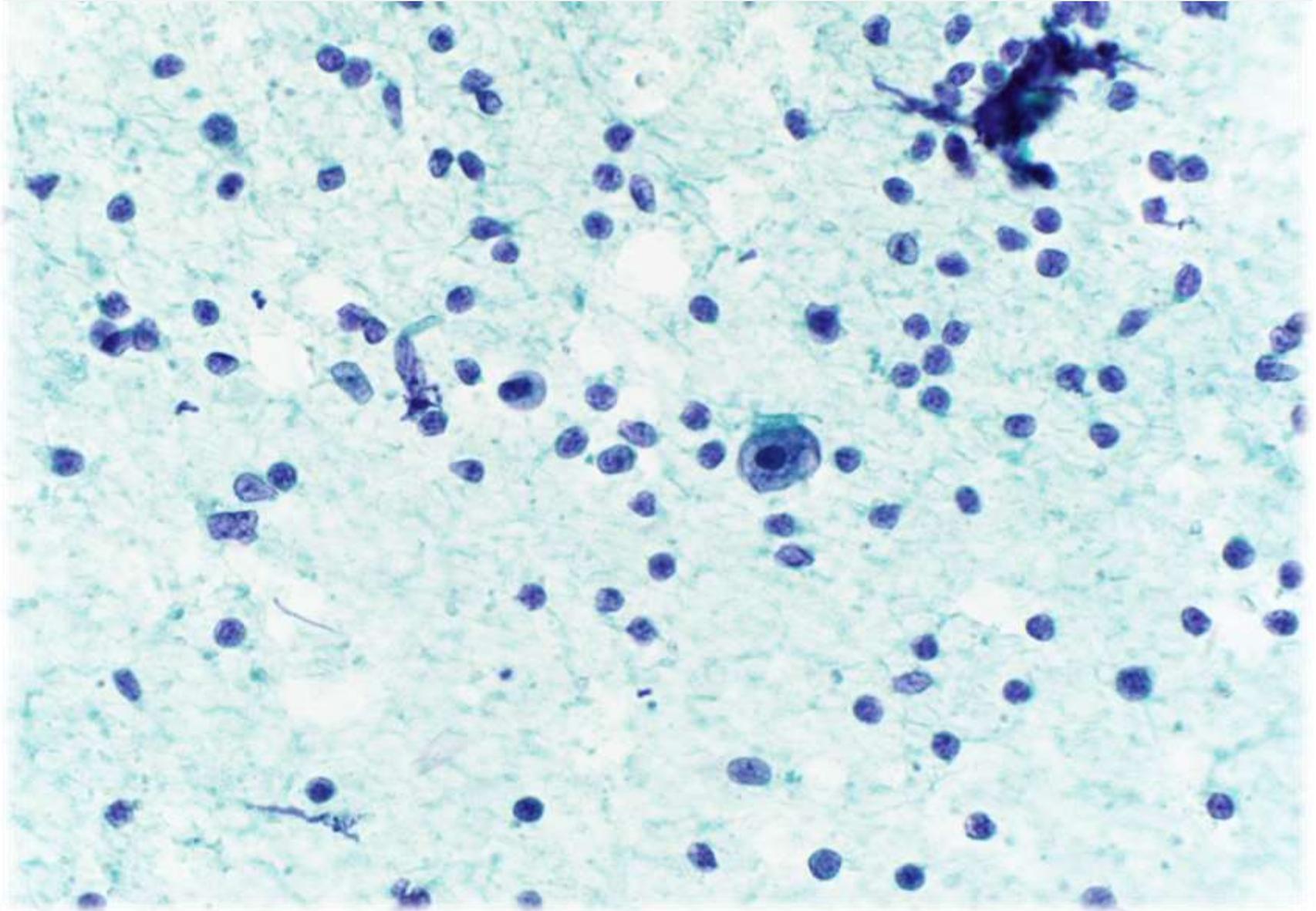
- vereinzelte LP-Zellen (ehemals L&H-/Popkornzellen) auf B-Zell-reichem Hintergrund
- CD20+; J-chain+; EMA+; CD30-/+; CD15-; EBV-



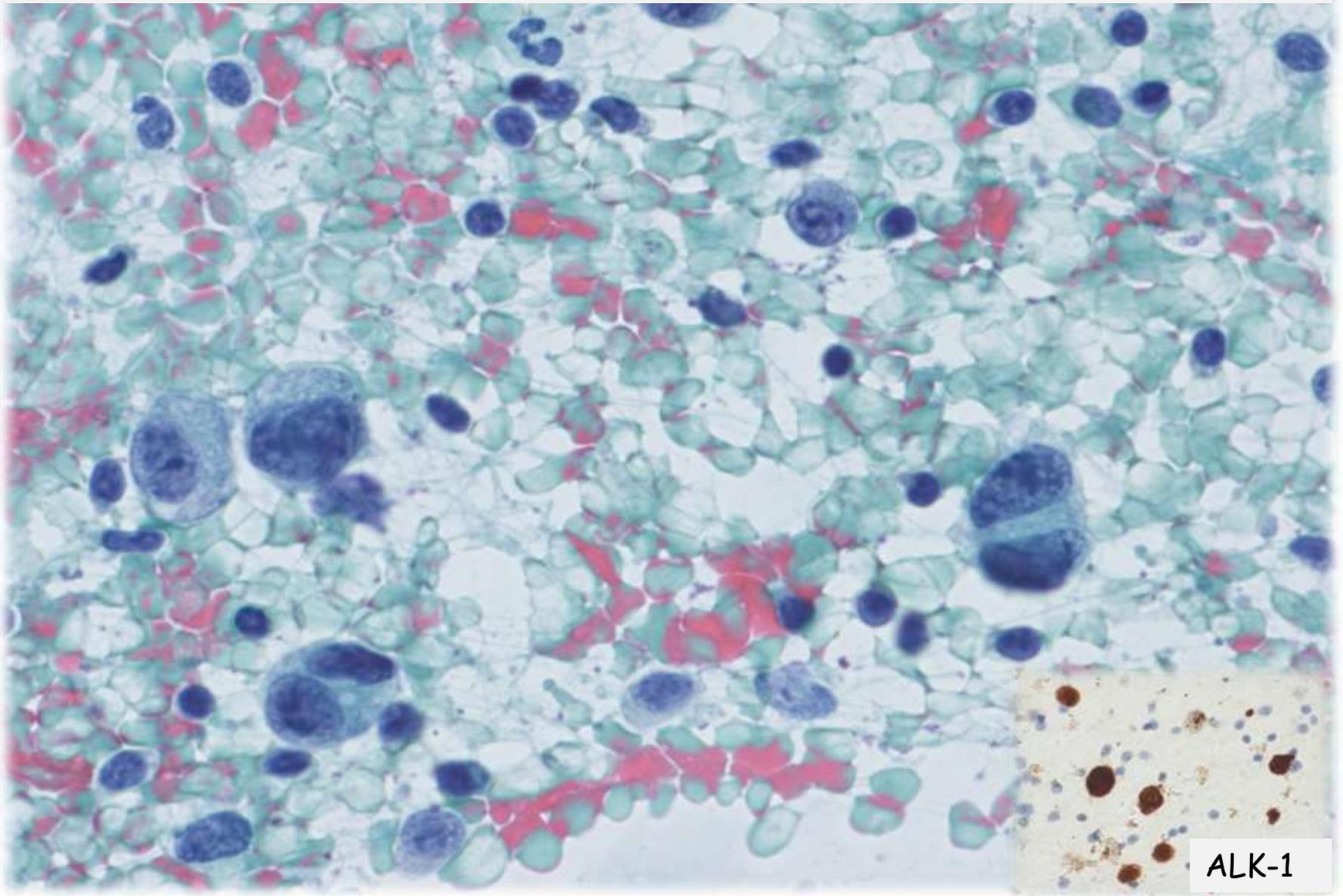
reaktive Lymphadenopathie



T-Zell und histiozytenreiches B-NHL

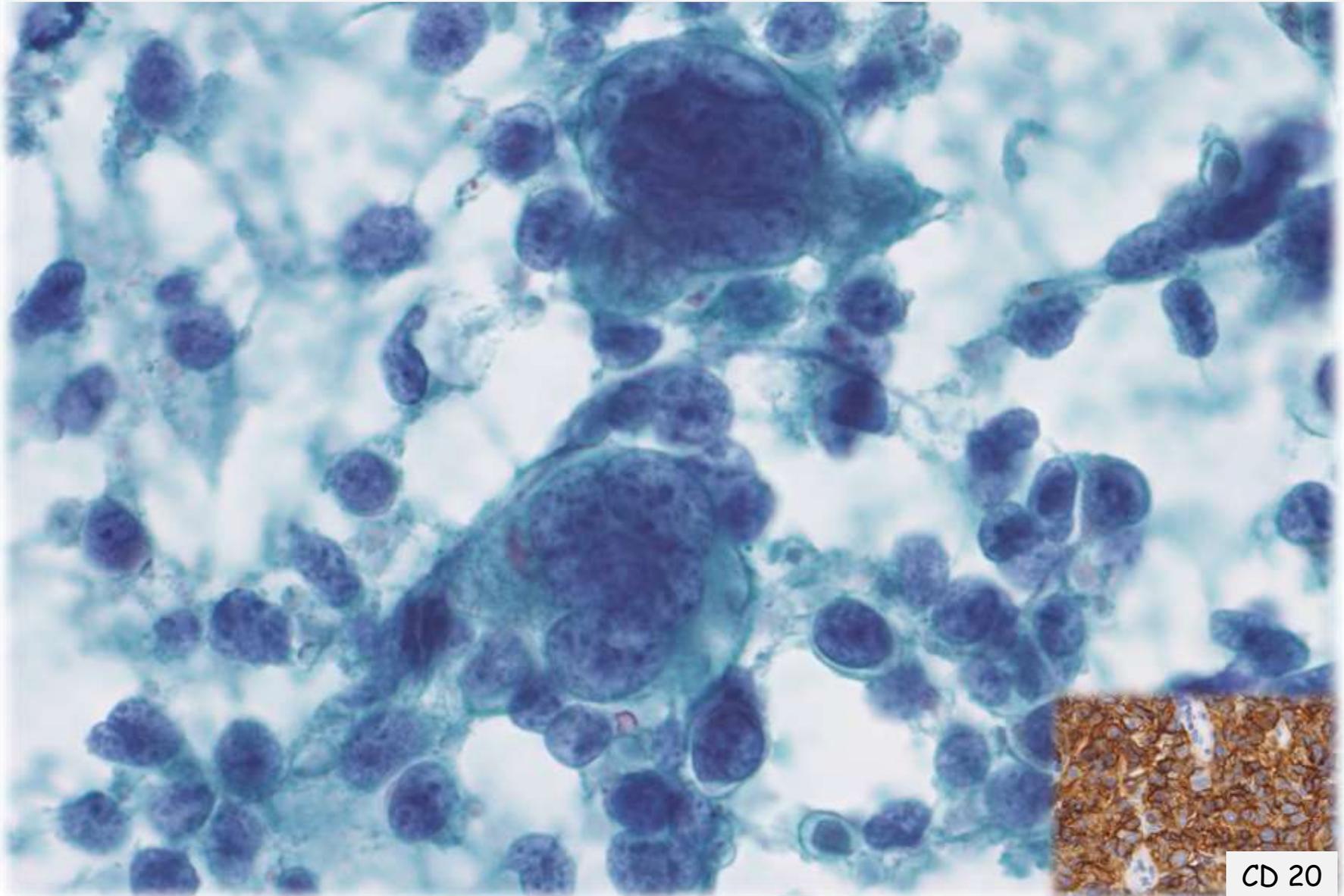


grosszellig-anaplastisches Lymphom (ALK-1+)



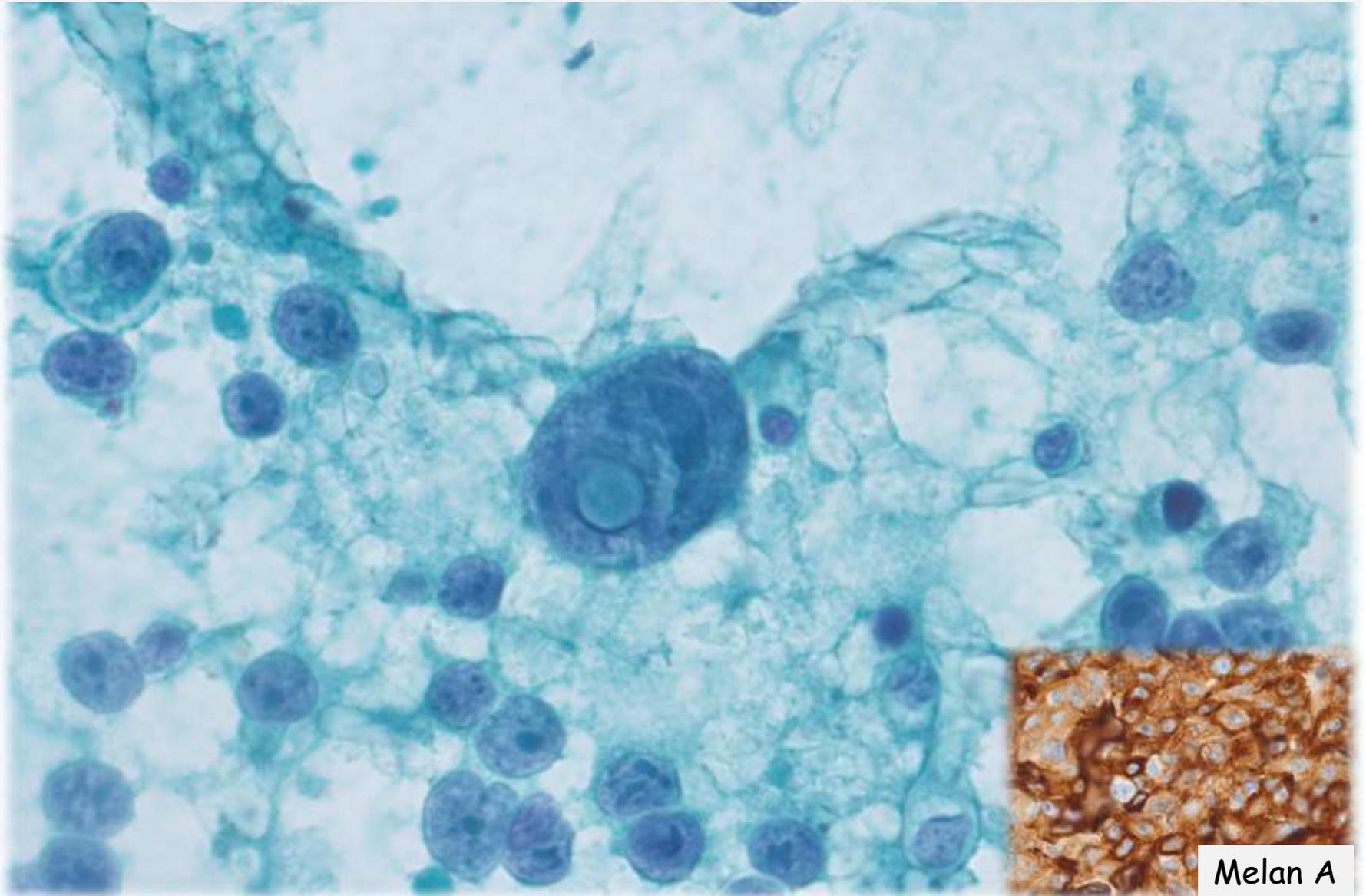
ALK-1

DLBCL, anaplastische Variante



CD 20

Melanom



Melan A