Autoantibodies: The mystery revealed

Connective Tissue Diseases

Ansar MEFTEH

&

Fatma ZAABI

Faculty of Medicine of Monastir

Faculty of Pharmacy of Monastir

- •Firstly reported in 1978 in a Japanese woman diagnosed with systemic lupus erythematosus (SLE)
- Initially designated as 'Ne' then as blastogenic nuclear antigen (high expression of the target antigen in mitogen-stimulated or proliferating lymphocytes) and ultimately identified as PCNA (elevated Antigen expression in rapidly Proliferating Cell Nuclei)
- Formerly considered as highly specific for SLE, but this specificity is actually debated.
- ●Low prevalence <5% in SLE
- High titers of anti-PCNA antibodies may be specifically associated with SLE

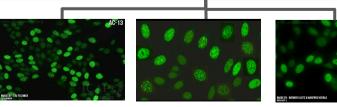
- Family: Antinuclear antibodies
- Target antigen: <u>34 kDa</u> auxiliary protein of DNA polymerase δ , involved in DNA replication and repair.

Anti-PCNA antibodies

- Besides SLE, can be present in;
- Rheumatoid arthritis/ Systemic sclerosis/ Sjogren syndrome
- Autoimmune thyroiditis
- -Hepatitis B and C virus infection
- Even in healthy individuals
- Associated with malignancies (lymphoma)

Screening technique: IIF (Indirect immunofluorescence) on HEp-2 cells (IIF prevalence of 0.07%)

IIF pattern on Hep-2 cells: speckled pleomorphic fluorescence (30-60% of cells) grains of variable size and distribution, no marking of cells in mitosis



Confirmation techniques:

- Double immunodiffusion according to Ouchterlony, line/dot blot, ELISA, ALBIA, NALIA, LIA...
- •At cut-off values resulting in 100% specificity, 52.5% (ELISA), 42.5% (ALBIA) and 35% (LIA) of samples with a proliferating cell nuclear antigen-like IIF staining pattern were positive

In case of pleomorphic staining not confirmed on confirmation techniques:

- consider pseudo-PCNA antibody testing
- rule out first a malignancy, rather than SLE.



