

# **Leukaemia Section**

**Short Communication** 

# **Cutaneous T-cell lymphomas**

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# Clinics and pathology

### Phenotype/cell stem origin

These are peripheral T-cell tumours, usually expressing CD4 along with other T-cell markers. The CD30 molecule is expressed in a histologically distinct subset (vide infra).

Classification: The WHO identifies some well-defined clinicopathological entities with distinct clinical behaviour. Indolent forms include mycosis fungoides, primary cutaneous CD30 (Ki1) positive anaplastic large cell lymphomas and its benign variant lymphomatous papulosis; aggressive forms include Sezary's syndrome and peripheral CD30-negative T-cell lymphoma.

#### **Epidemiology**

Primary cutaneous non Hodgkin's lymphomas (NHL) has an approximate 0,4/100.000 incidence in

the U.S. The majority of patients are between 40 and 60 years old.

### **Clinics**

Details on clinics, pathology, cytogenetics and molecular cytogenetics can be found in disease-specific cards (i.e. mycosis fungoides / Sezary's syndrome, primary cutaneous CD30 positive anaplastic large cell lymphoma).

## References

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