

Selected Topics in Clinical Immunology - Biologics-

Biomedical Sciences March 1, 2018 S. Adler



today's menu

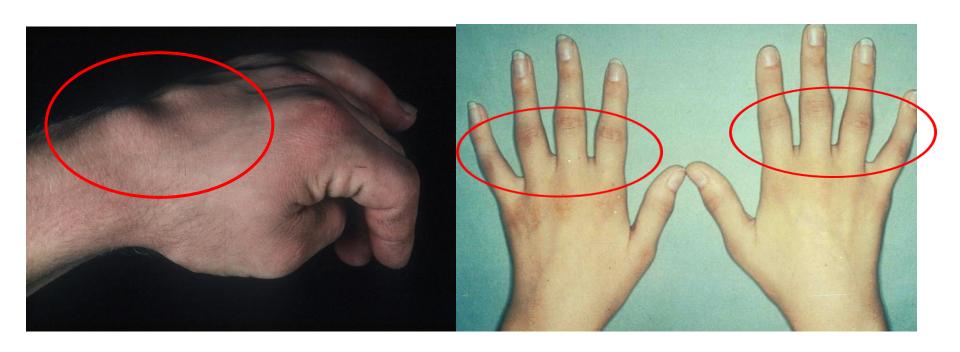
- lessons regarding pathogenesis
- cytokines
- cytokine inhibitors / cytokine antagonists
- therapeutic antibodies beyond anti-cytokines
- immunoreconstitution



what are we talking about

pathogenesis of inflammatory diseases

rheumatoid arthritis, early

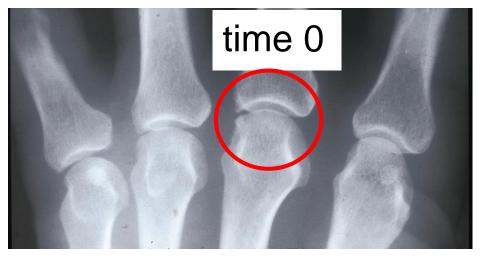




what are we talking about

pathogenesis of inflammatory diseases

- rheumatoid arthritis
- joint space narrowing







what are we talking about

pathogenesis of inflammatory diseases

rheumatoid arthritis, late







Clinical example

Rheumatoid Arthritis (RA)

joint inflammation

tendons and bursae

systemic inflammation

ESR, CrP

anemia, thrombocytosis

rheumatoid factor

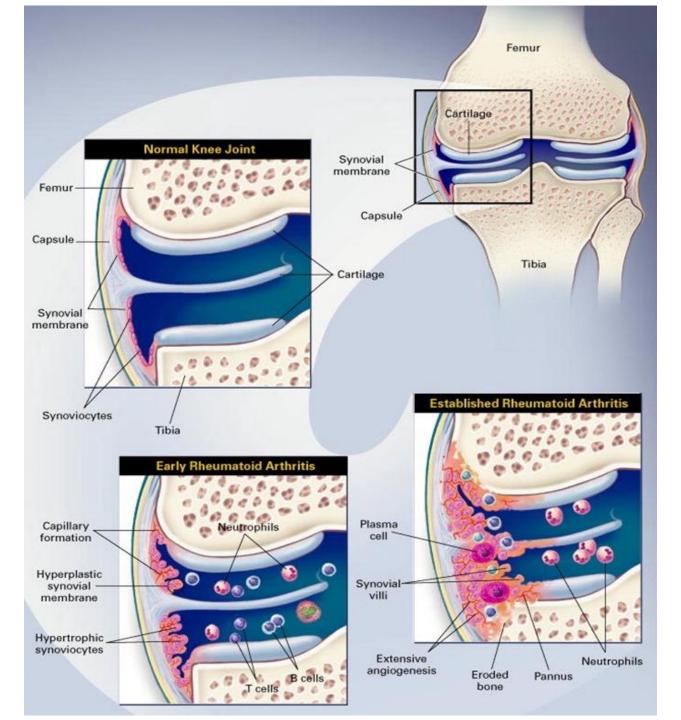
CCP-antibodies



pathogenesis of inflammation

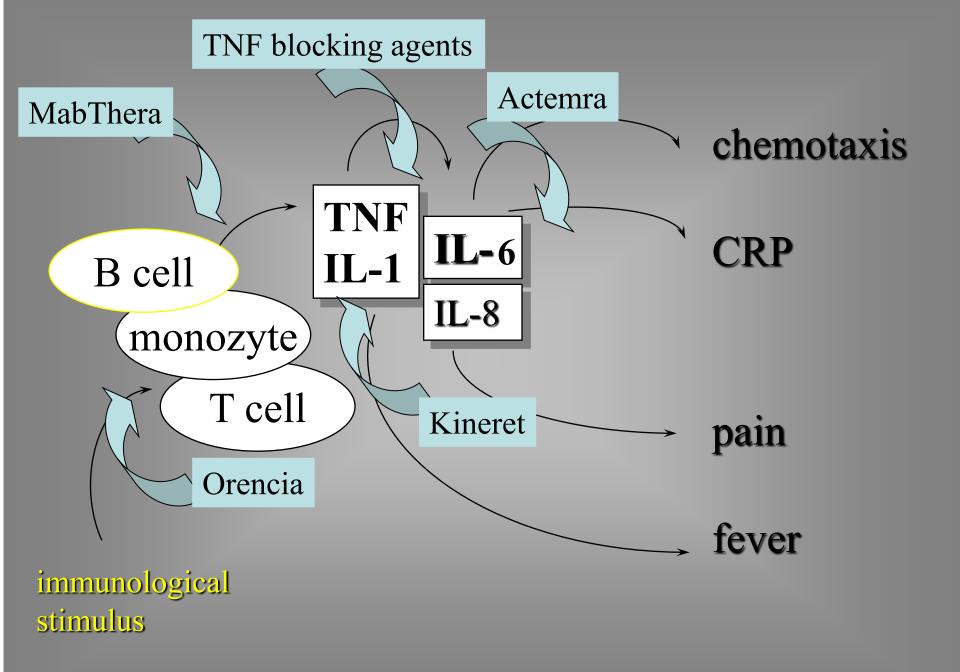
molecular mechanisms rheumatoid arthritis







Choy, 2001, NEJM



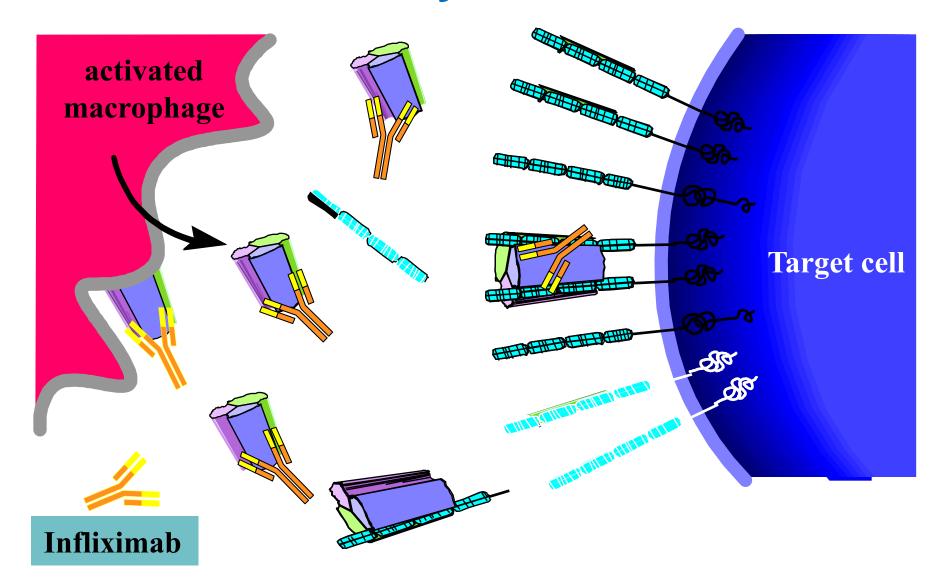


anti-cytokine mechanisms

Tumor necrosis factor (TNF)



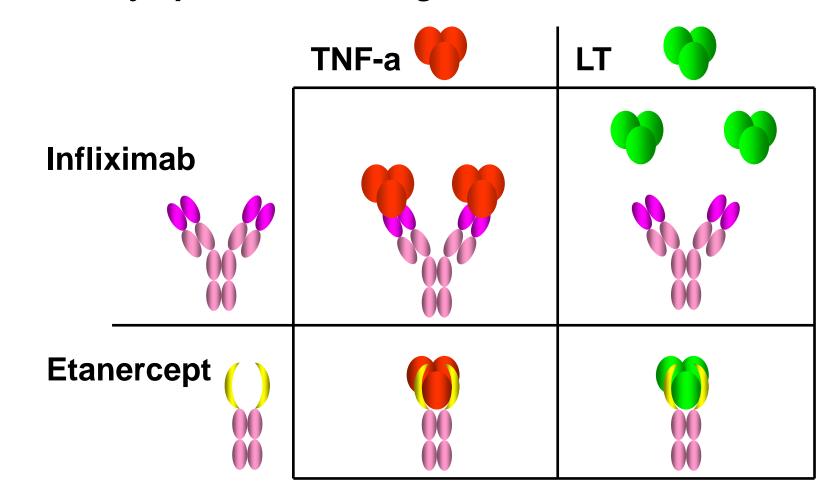
TNF-blockade by Infliximab





specificity of monoclonal antibody versus fusion protein

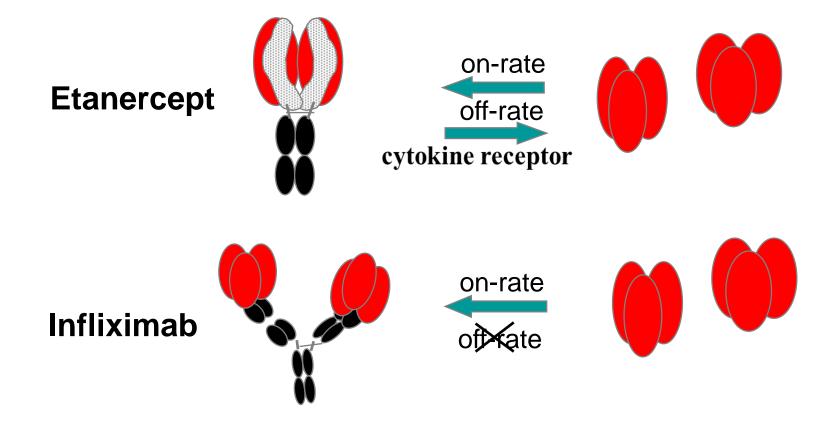
TNFa and lymphotoxin binding





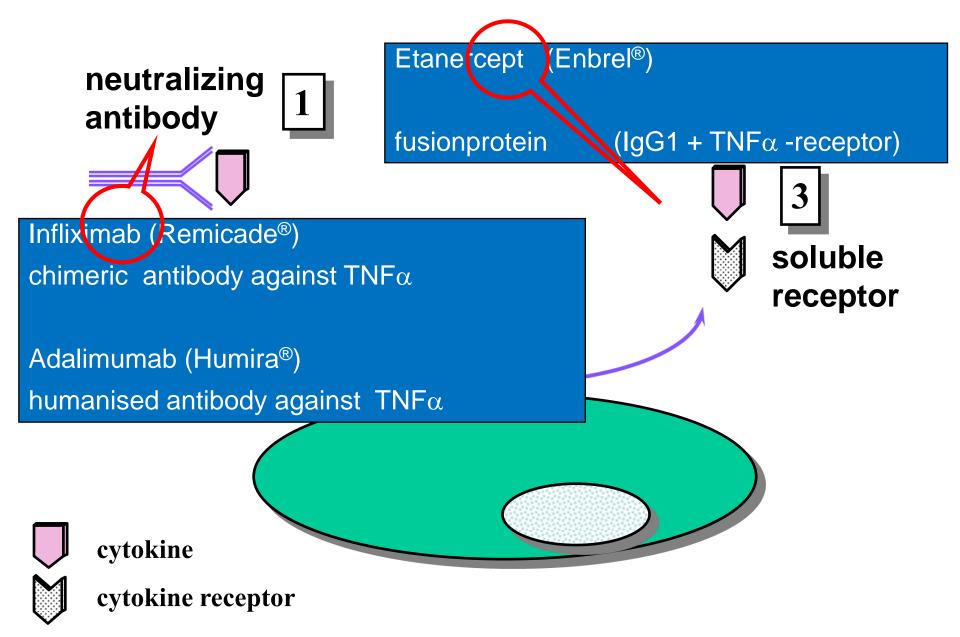
avidity

reversibility of binding



mechanisms of inhibition







treatment goals

- reduction of disease activity
- prevention of joint destruction
- long-lasting remission
- minimalized side effects



treatment goals

how to measure...

reduction of disease activity («DiseaseActivityScore»)
 number of tender joints
 number of swollen joints
 ESR/ CrP

 prevention of joint destruction / disease damage ultrasound
 X-rays

MRI (magnetic resonance imaging)



summary anti-cytokine strategies

neutralization of

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TNF
IL-1
IL-6
IL-17
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competitive blocking of receptor

IL-1 receptor antagonist (IL-1ra)

use of soluble receptor

TNF R abatacept



beyond anti-cytokine antibodies



pathogenesis of RA

B-cells as important factors

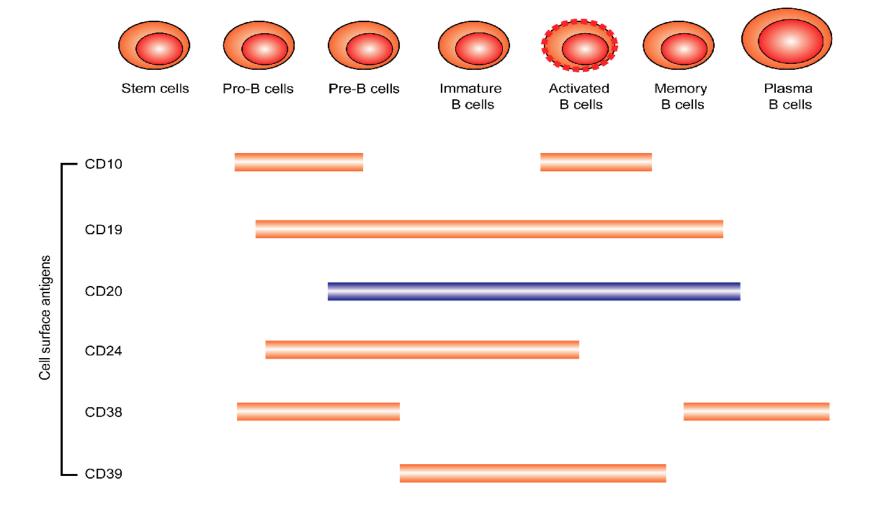
 abundance of B-cells in the synovium of affected joints organized into lymphoid structures

3 critical roles of B-cells

 antigen presentation and T-cell activation autoantibody production
 cytokine production



Steps in the maturation of B cells

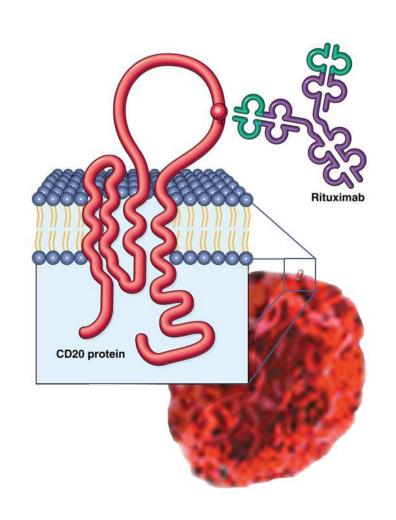




Rituximab (MabThera®/Rituxan®)

Rituximab

- novel
- genetically engineered
- anti-CD20 therapeutic monoclonal antibody
- selective depletion of CD20+ B-cells

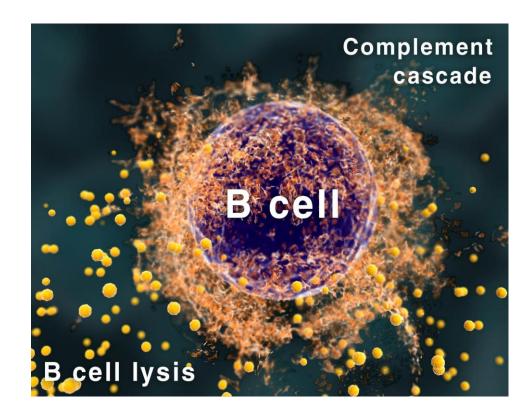




complement-dependent cytotoxicity

Rituximab bound to CD20

- interacts with C1q
- triggers activation of the complement system
- leads to B cell lysis via formation of pores in the membrane

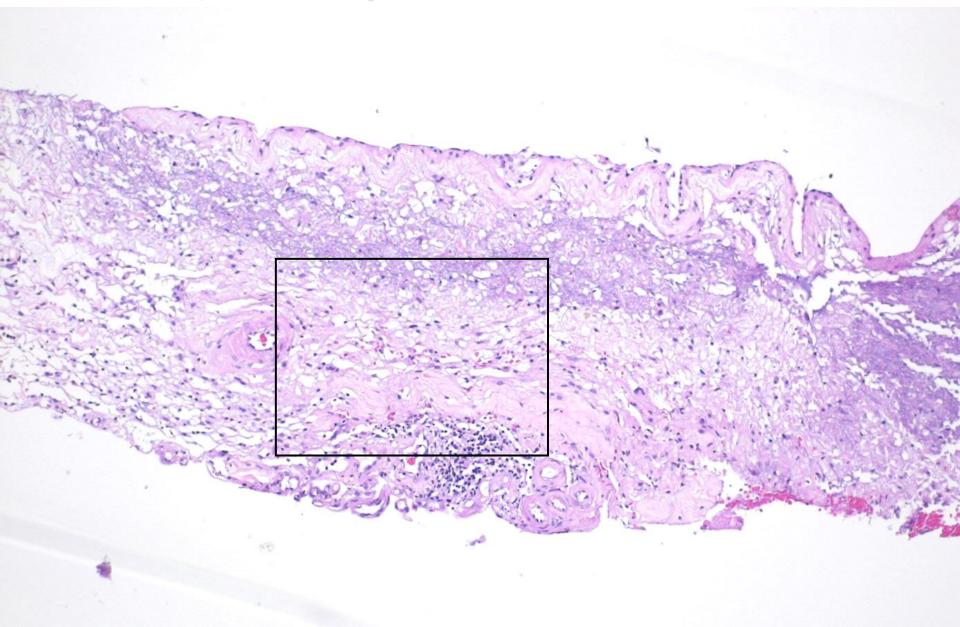


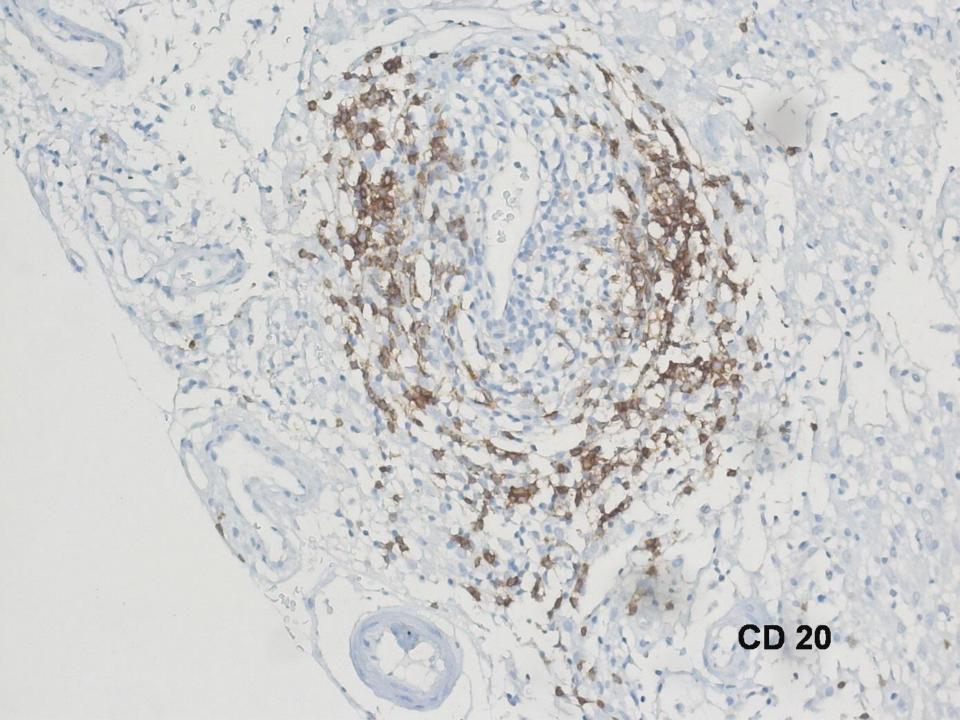


pachymeninigitis in a patient with RA



pachymeninigitis in a patient with RA

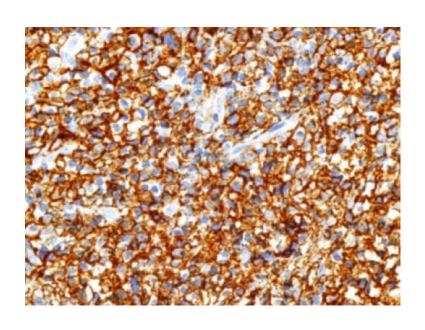


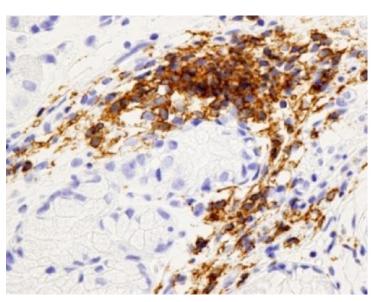




pre-post rituximab therapy

CD20 stain example: gastric MALT- lymphoma







summary cell-targeted strategies

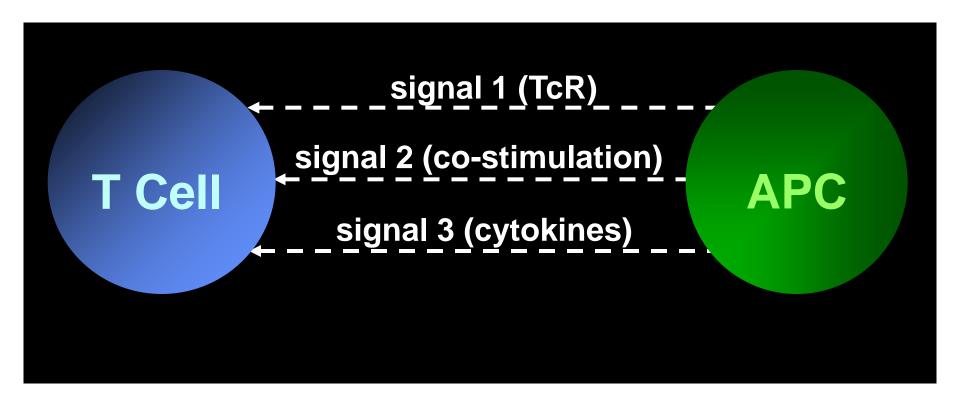
- depletion of B lymphocytes
- depletion of CD+ lymphocytes
- classical immunosuppressive agents ciclosporine
 - => inhibits function / activity of T-lymphocytes
 - => used in organ transplantation



mechanisms of co-stimulation



how are T cells recruited?





Anti-TNF antibodies and the risk of malignancies *JAMA 2006, 295 (19): 2275*

- systematic review and meta-analysis in randomized controlled trials
- 144 trials, 9 suitable for analysis
- Etanercept excluded (why?)
- RA patients only
- 3493 patients, 1512 controls
- Infliximab up to 10mg/kg, every 4 wk
- Adalimumab up to 40mg per wk
- Duration of therapy until diagnosis of malignancy: 2 -114 weeks (!!)



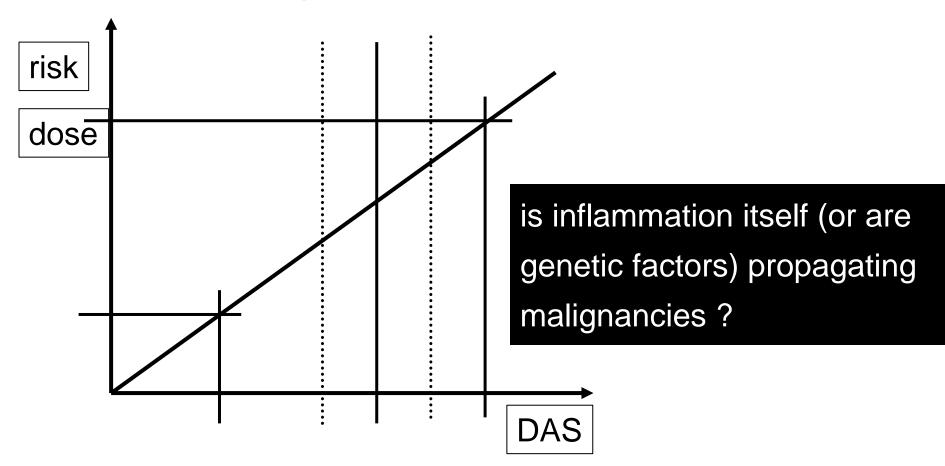
Anti-TNF antibodies and the risk of malignancies *JAMA 2006, 295 (19): 2275*

results

- 29 malignancies in verum, 3 in placebo
- OR 3.3 (1.2 9.1)
- however:
 - Low dose: OR 1.4 (0.3 5.7)
 - High dose: OR 4.3 (1.6 11.8)
- number needed to harm (NNH): 154 (91 500)



relation: risk of malignoma / disease activity



case control study showing an increased lymphoma risk of up to 25 (BMJ 1988)



after review of the existing literature and thorough discussion:

 screening for Tbc and latent Tbc infection should be performed in all patients prior to any anti- TNF-a therapy

- screening should be based on history, chest X-ray and an IGRA test.
 - history: detailed history of exposure to or prior treatment for Tbc, considering the risk associated with birthplace or country of origin
 - chest X-ray: for detecting past or present Tbc
 - IGRA test