EUVAS update June 5th 2012

Marinka Twilt

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

Submitted by Jennette et al

Reported for EUVAS meeting by N. Rasmussen

- Large Vessel Vasculitis (LVV)
 - Takayasu Arteritis (TAK)
 - Giant Cell Arteritis (GCA)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
 - Polyarteritis Nodosa (PAN)
 - Kawasaki Disease (KD)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
 - > AAV
 - Microscopic Polyangiitis (MPA)
 - Granulomatosis with Polyangiitis (GPA)
 - Eosinophilic Granulomatosis with Polyangiitis (EGPA)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
 - Immune complex SVV
 - Anti-GBM disease
 - Cryoglobulinemic Vasculitis
 - IgA Vasculitis (HSP)
 - Hypocomplementemic Urticarial Vasculitis
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
 - Behçet Disease (BD)
 - Cogan's Syndrome
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
 - Cutaneous Leukocytoclastic Angiitis
 - Cutaneous Arteritis
 - Primary Angiitis of the CNS (PACNS)
 - Isolated Aortitis
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
 - Lupus vasculitis
 - Rheumatoid vasculitis
 - Sarcoid vasculitis
- Vasculitis with Probable Etiology

- Large Vessel Vasculitis (LVV)
- Medium Vessel Vasculitis (MVV)
- Small Vessel Vasculitis (SVV)
- Variable Vessel Vasculitis (VVV)
- Single Organ Vasculitis (SOV)
- Vasculitis Associated with Systemic Disease
- Vasculitis with Probable Etiology
 - HCV-associated cryoglobulinemic vasculitis
 - Drug-associated immune complex vasculitis
 - Drug-associated ANCA-associated Vasculitis
 - Cancer associated vasculitis

DCVAS

Diagnostic and classification criteria in Vasculitis

- Prospective study to design and validate classification and diagnostic criteria in vasculitis (total sites 108!)
- still recruiting. N=1096, aiming for 4000
- data: clinical features, serology, pathology, radiology, baseline data + 6 months f-up
- Online database <u>www.dcvas.org</u>
- DCVAS is partner in CIHR pediatric vasculitis grant

R. Luqmani

5-year follow-up study

535 pts from first 4 RCTs
281 MPA, 254 GPA

▶ 87% filled in questionnaire

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Kerstin Westman

Vascular function in AAV (study1)

Single centre study vascular effects

Rituximab

- Patients: I5 AAV (rituximab), I5 AAV (CYC), 30 healthy controls (matched)
- Primary outcome: aortic pulse wave velocity
- Secondary outcomes: FMD, lipids

Alina Casian

Vascular function in AAV (study 2)

Part of Pexivas

- Immunomodulatory/ vasculoprotective effects of PLEX
- Based on removal anti-oxLDL antibodies and proinflammatory HDL, oesteopoetin, ADMA, prothrombotic anti-plasminogen antibodies

Renal histology

> 2 different studies

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- Anti-plasminogen antibodies at diagnosis and relation to renal histology
- Histological determinants for long-term renal outcome

Anti-plasminogen

- anti plasminogen antibodies at diagnosis
- relation to renal biopsy
- anti-plasminogen antibodies during follow-up
- presence anti-plasminogen and renal outcome
- "new" patients (clear diagnosis according ACR criteria)
 - N=80 biopsies and sera
- "old" patients (Dx according ACR criteria not known)
 - N=70 biopsies and sera

MYCYC trial

- MMF versus CYC
- recruitment finished July 2011
- ▶ n=140 (70 MMF, 70 CYC)
- Results to be presented soon (ANCA Workshop):

PEXIVAS

Plasma exchange and glucocorticoid dosing in treatment of ANCA-associated vasculitis

Goal inclusion 500 pt

> 250 PLEX

I25 standard steroids / I25 reduced dose steroids

> 250 no PLEX

I25 standard steroids /I25 reduced dose steroids

Start June 2010

- ▶ GFR < 50 ml or pulmonary hemorrhage
- I51 pt included

59 centers active

RITAZAREM

Superiority of fixed interval repeat rituximab against AZA for prevention relapse in AAV

- Rituximab preferred to CYC for relapses
- Relapse rate 71% in non-repeat Rituximab group (@ 24 months)
- Relapse 24% in repeat group (rituximab every 6 months)(@ 24 months)

Smith et al in press (A&R)

CORTAGE

Corticosterod-based treatment for AAV pt > 65 years of age

- FVSG study
- Reduce treatment related morbidity (death and SAE's) by 30% at 3 years (68 → 38%)
- Treatment based on Five Factor Score
 - FFS = $0 \rightarrow cs$ alone
 - FFS > I \rightarrow cs + iv CYC 500 mg/m²
- Experimental treatment
 - CS reduced doses
 - CYC 500 mg per pulse for all

Expected results oct 2012

MAINRITSAN

- GPA & MPA patients (FFS > I)
- Newly diagnosed or recent relapse
- Treatment cs + CYC followed by
 - Rituximab
 - Azathioprine
- Hypothesis: absolute reduction by 25% of relapse in rituximab maintenance group
- N = 118 (last inclusion june 2010)
- Expected results oct 2012

CLEAR

C5A receptor-inhibitor on leucocytes exploratory ANCAassociated Renal vasculitis trial

- Industry trial
- CCX168 = specific human C5a-receptor antagonist
- Orally administered
- C5A = powerful neutrophil chemo-attractant and leads to pro-inflammatory cytokines
- ▶ In mice 30 mg BID \rightarrow effectively blocked
- Phase I \rightarrow well tolerated in healthy volunteers

CLEAR

- Phase 2
- Multinational randomized double-blind, placebo controlled phase 2 clinical trial in 39 study centers in Europe
- Primary outcome: efficacy and tolerability CCX168
- Age 18-75
- Positive ANCA/PR3/MPO
- Inclusion completed
- Step I of phase 2 completed (no SAE's, no rescue ivMP necessary)

BREVAS

- Belimumab in AAV
- Industry trial
- Goal n = 400
- Induction rituximab + steroids or CYC + steroids
- N = 200 belimumab + AZA
- N= 200 placebo + AZA
- Primary outcome: relapse
- Stratified by ANCA type, initial dx
- Inclusion: GPA/MPA

mepolizumab

- mepolizumab in EGPA (CSS)
- anti-II-5 antibodies
- specific for human II-5 (eosinophils decrease)
- 2 open label studies in EGPA
 - showed reduction steroids requirements
 - reduction exacerbations during treatment
 - well tolerated
- double blind, randomized, placebo controlled study
- treatment duration I year
- start 2013
- goal n = 140 (1:1 randomized)

Pediatric studies

- Brainworks and ARChiVe
- PVAS

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MYPAN

PVAS

- pediatric modification of BVAS
- 22 modified + 6 new items
- training of users
- manuscript submitted

MYPAN

- open label randomized control trial
- non-inferiority MMF to CYC for induction
- primary outcome: remission @ 6 mos
- goal n=40 (I:I randomization)
- non-inferiority margin 15%
- bayesian approach

2nd annual CanVasc meeting

ANDIAN VASCULITIS NETTA

Montréal, QC November 22nd, 2012

Registration and information on http://www.canvasc.ca



Scientific committee : Pr. Loïc Guillevin (president)

Organisation :

Maud Placines-Charier Nex & Com Medical Events 159 rue de Silly 92100 Boulogne Billancourt Tel : +33 1 46 43 33 23 Fax : +33 1 46 43 33 24 Email : m.placines@nex-com.com

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