



Brainstorming Sessions

Ασθενής με συνυπάρχουσα ψωρίαση και ραγοειδίτιδα

Εσκιτζής Αναστάσιος
Ιδιώτης Ρευματολόγος

Introduction

Case presentation: man with psoriatic arthritis and uveitis.

Characteristics of and treatment options for PsA-associated uveitis.

Acknowledgement to N. Κούγκας.

Disclosures: Υποστήριξη από την UCB για τη συμμετοχή στο παρόν συνέδριο.

Case presentation

50 yo male patient

- plaque **psoriasis**, moderately severe
- **psoriatic arthritis**, diagnosed 20 years ago: peripheral oligoarthritis, dactylitis

Medication History:

- **methotrexate** po, discontinued due to hepatotoxicity
- **infliximab** iv, 2ary failure with flares of arthritis
- **adalimumab** sc, 2ary failure with flares of skin psoriasis

On **secukinumab** in the past 2 yrs, with very good response (skin and joints), achieved Minimal Disease Activity.

Episode of acute unilateral anterior uveitis.



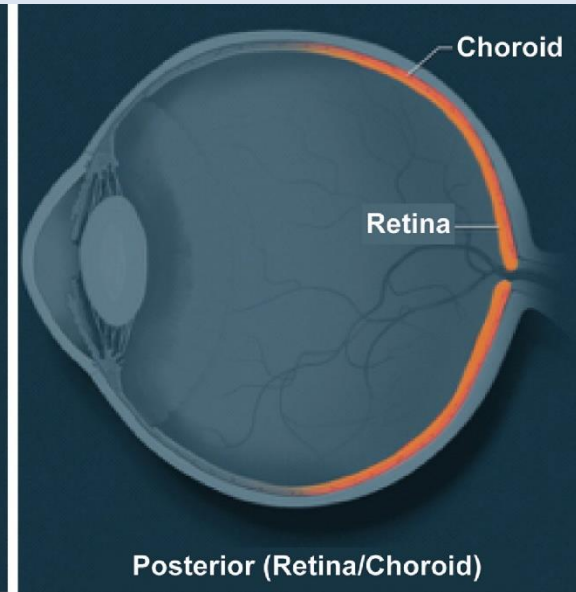
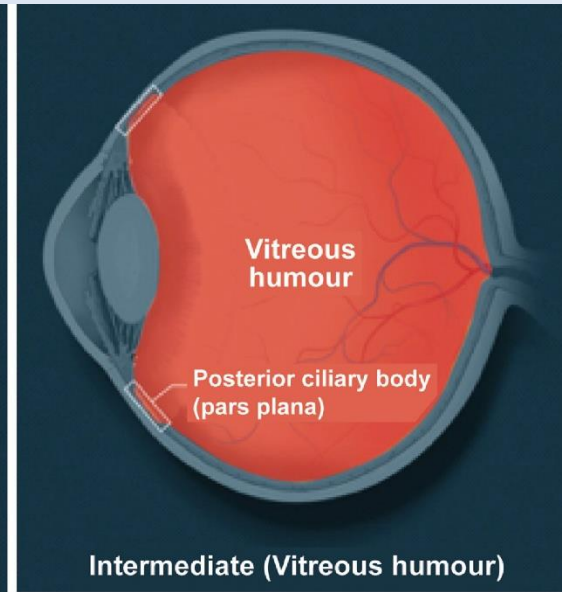
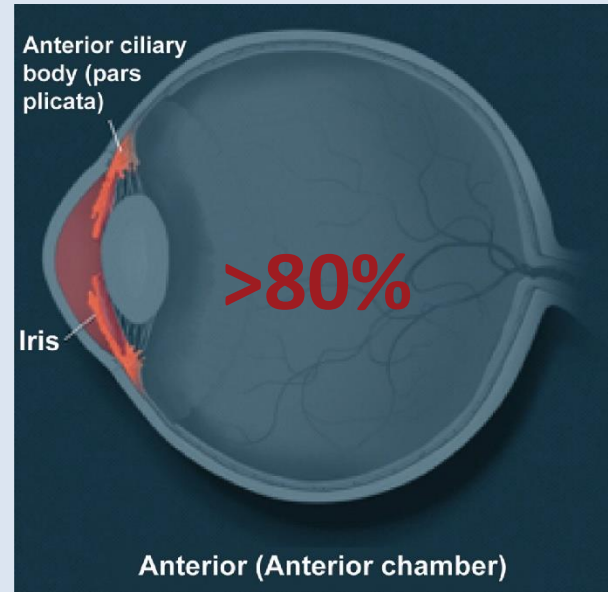
1st line treatment: **topical corticosteroids** (eye drops)

Recurrence 2 months later

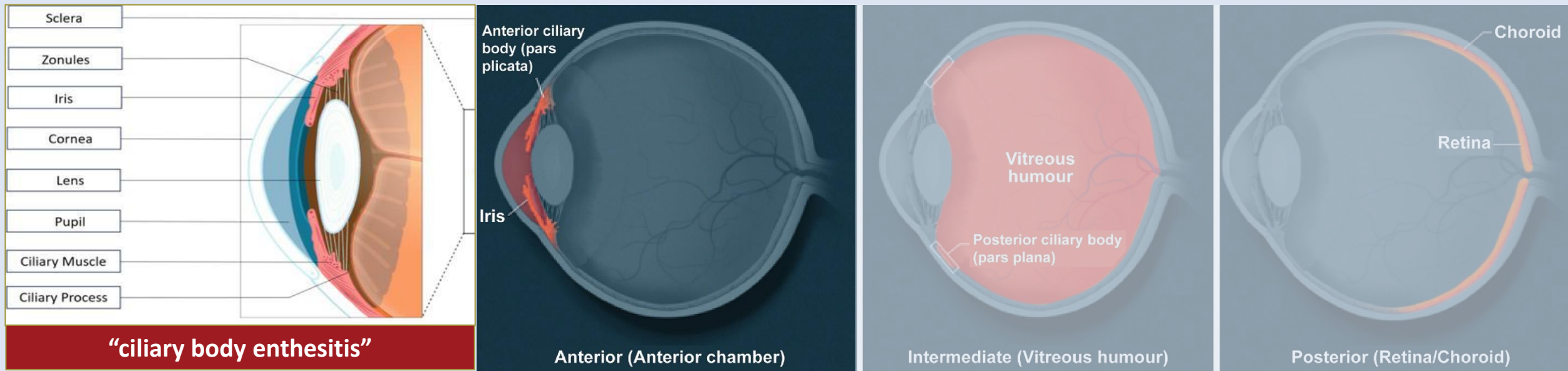


topical corticosteroids + **corticosteroids po** + ???

PsA-associated uveitis



PsA-associated uveitis



- Uveitis prevalence in PsA: 3-7% (2.7% in a large Greek study) (vs 20-40% in AS).
- Usually acute-onset.
- Can be unilateral (or alternating) or bilateral.
- Frequently recurrent (higher risk for complications).
- Associations (risk factors):

family history of SpA	HLA-B27 (+)	axial disease (or sacroiliitis)	longer disease duration	more severe disease (worse PsAID)
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Deligeorgakis, Dimitrios, et al. *Uveitis in Psoriatic Arthritis : A Comprehensive Review*. 2025, doi:10.5152/eurjrheum.2025.24078.

De Vicente Delmás, Ana, et al. “Uveitis in Psoriatic Arthritis: Study of 406 Patients in a Single University Center and Literature Review.” *RMD Open*, vol. 9, no. 1, 2023, pp. 1–8.

Kougkas, Nikolaos, et al. “Higher Frequency but Similar Recurrence Rate of Uveitis Episodes in Axial Spondylarthritis Compared to Psoriatic Arthritis. A Multicentre Retrospective Study.” *Rheumatology International*, vol. 43, no. 11, 2023, pp. 2081–88.

Treatment principles

Corticosteroids

topical corticosteroids (eye drops)

corticosteroid injections: peribulbar, sub-Tenon, intravitreal

systemic corticosteroids

Conventional synthetic DMARDs

methotrexate, cyclosporine A

sulfasalazine, MMF, azathioprine

Biologic DMARDs / targeted synthetic DMARDs

TNF inhibitors (monoclonal antibodies >> etanercept)

IL-17 inhibitors ?, IL 12/23 inhibitors ?

JAK inhibitors ??

Anterior Uveitis

- Effective therapies:
 - Strong recommendations as of July 2022:
 - TNF inhibitors (except etanercept)
 - Conditional recommendations as of July 2022:
 - Methotrexate
 - Cyclosporin
- Therapies not shown to be effective:
 - Etanercept
 - IL-17 inhibitors – studies SHIELD, ENDURE, and INSURE

Uveitis



TNFi (not ETN),
ciclosporin,
MTX

EULAR Recommendations 2023 Update

EULAR recommendations for the management of psoriatic arthritis with pharmacological therapies: 2023 update

Laure Gossec ,^{1,2} Andreas Kerschbaumer ,³ Ricardo J O Ferreira ,^{4,5} Daniel Aletaha ,³ Xenofon Baraliakos ,⁶ Heidi Bertheussen,⁷ Wolf-Henning Boehncke,⁸ Bente Appel Esbensen ,^{9,10} Iain B McInnes,¹¹ Dennis McGonagle,^{12,13} Kevin L Winthrop ,¹⁴ Andra Balanescu,¹⁵ Peter V Balint,¹⁶ Gerd R Burmester ,¹⁷ Juan D Cañete ,^{18,19} Pascal Claudepierre,^{20,21} Lihi Eder ,²² Merete Lund Hetland ,^{23,24} Annamaria Iagnocco ,²⁵ Lars Erik Kristensen,^{26,27} Rik Lories,^{28,29} Rubén Queiro ,^{30,31} Daniele Mauro ,³² Helena Marzo-Ortega ,^{12,13} Philip J Mease ,^{33,34} Peter Nash ,³⁵ Wendy Wagenaar,^{36,37} Laura Savage,³⁸ Georg Schett ,³⁹ Stephanie J W Shoop-Worrall ,⁴⁰ Yoshiya Tanaka ,⁴¹ Filip E Van den Bosch ,⁴² Annette van der Helm-van Mil,⁴³ Alen Zabotti ,⁴⁴ Désirée van der Heijde ,⁴³ Josef S Smolen³

The choice of the mode of action should reflect non-musculoskeletal manifestations related to psoriatic arthritis; with clinically relevant skin involvement, preference should be given to an IL-17A or IL-17A/F or IL-23 or IL-12/23 inhibitor; with uveitis to an anti-TNF monoclonal antibody; and with IBD to an anti-TNF monoclonal antibody or an IL-23 inhibitor or IL-12/23 inhibitor or a JAKi*.

Adalimumab

Adalimumab in Patients with Active Noninfectious Uveitis (VISUAL 1)

A trial to assess the efficacy and safety of adalimumab as a glucocorticoid sparing agent for noninfectious uveitis

STUDY DESIGN



Multinational, randomized, placebo-controlled Phase 3 Trial (2010-2014)



Inclusion Criteria

Age > 18 years
Active noninfectious uveitis
Use of prednisone (10-60 mg/day) for >2 weeks



Exclusion Criteria

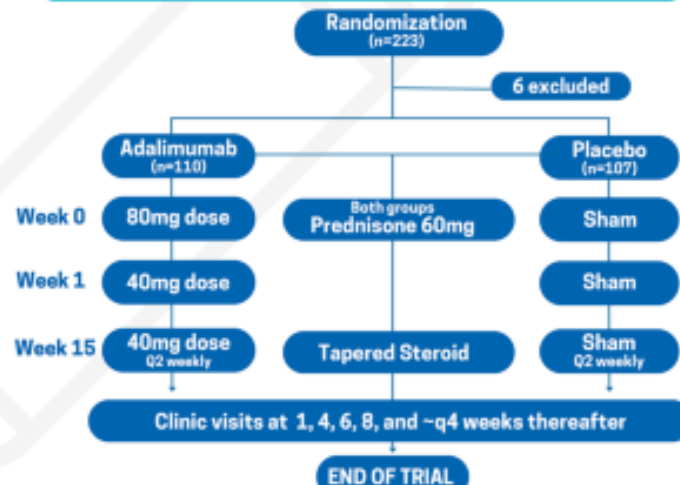
Isolated Anterior Uveitis
Inadequate response to oral corticosteroids
Confirmed/Suspected Infectious Uveitis
Presumed Ocular Histoplasmosis Syndrome



Treatment

Subcutaneous Delivery

METHODS



RESULTS

PRIMARY OUTCOME

Time to Treatment Failure

24
weeks

13
weeks

SECONDARY OUTCOMES

% cause as failure

Worse BCVA

21%

25%

Vitreous Haze

15%

36%

Anterior Chamber Cells

22%

32%

Retinal Lesions

15%

27%

ADVERSE EVENTS

(per 100 person-years)

1052.4

971.7

SERIOUS ADVERSE EVENTS

(per 100 person-years)

28.8

13.6

CONCLUSION

ADALIMUMAB
(vs. Placebo)

- Lower risk of uveitic flare or visual impairment
- More frequent adverse events

Certolizumab pegol – C-VIEW study

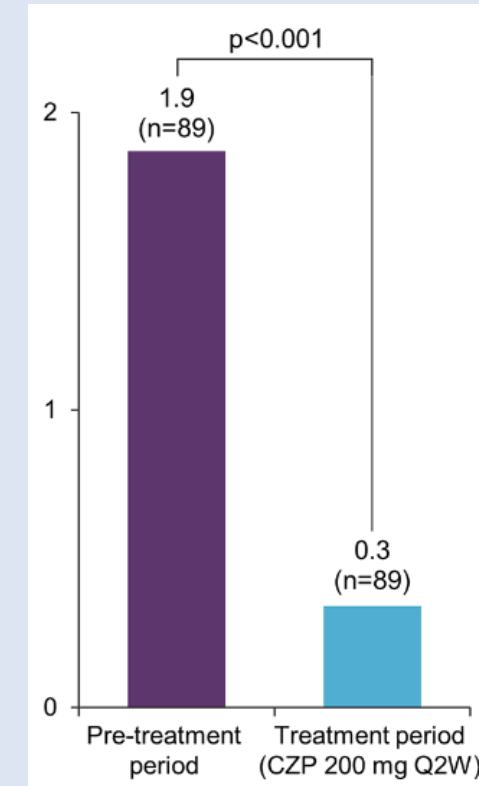
 *Therapeutic Advances in Musculoskeletal Disease*

Reduction of anterior uveitis flares in patients with axial spondyloarthritis on certolizumab pegol treatment: final 2-year results from the multicenter phase IV C-VIEW study

Irene E. van der Horst-Bruinsma , Rianne E. van Bentum, Frank D. Verbraak, Atul Deodhar, Thomas Rath, Bengt Hoepken, Oscar Irvin-Sellers, Karen Thomas, Lars Bauer and Martin Rudwaleit

Phase IV study







89 patients with axSpA, HLA-B27 (+) and history of ≥ 2 episodes of acute anterior uveitis (including 3 patients w psoriasis) received certolizumab pegol for 2 yrs



Poisson-adjusted event rate per 96 weeks

Bimekizumab

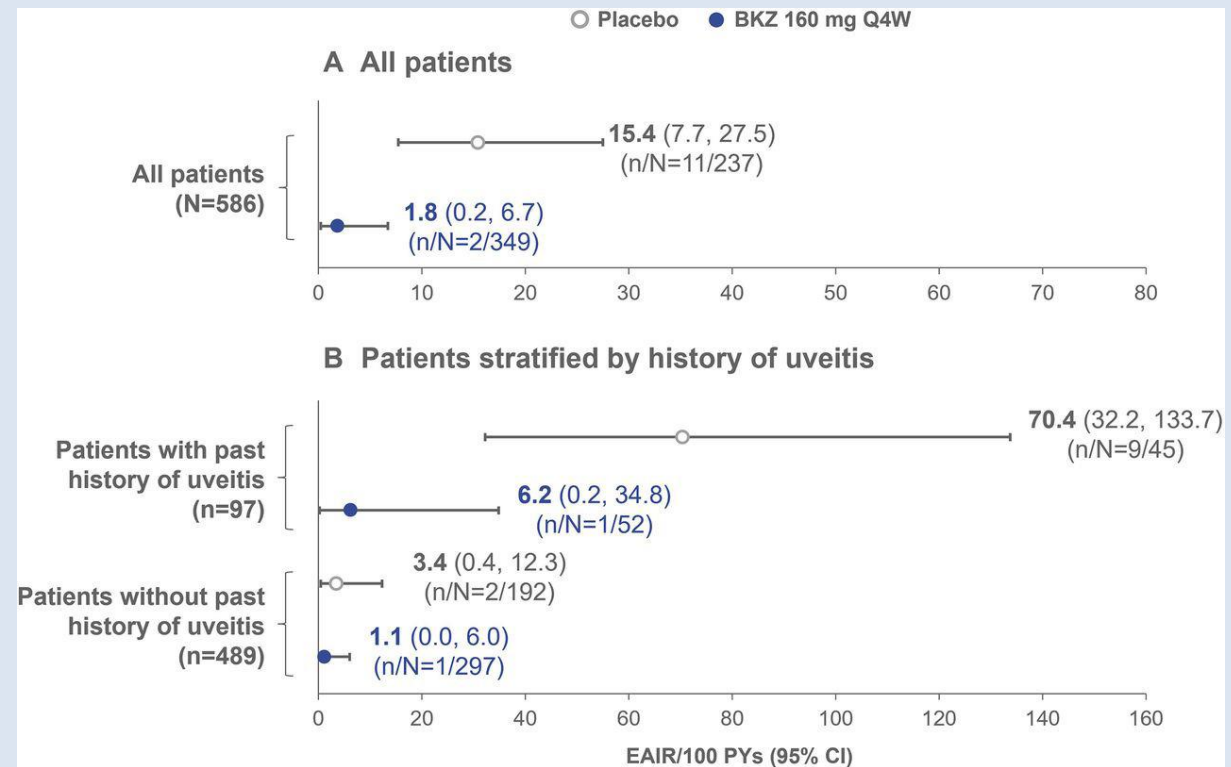
Low uveitis rates in patients with axial spondyloarthritis treated with bimekizumab: pooled results from phase 2b/3 trials

Matthew A Brown ^{1,2}, Martin Rudwaleit ³, Floris A van Gaalen ⁴,
Nigil Haroon ⁵, Lianne S Gensler ⁶, Carmen Fleurinck,⁷ Alexander Marten,⁸
Ute Massow,⁸ Natasha de Peyrecave,⁷ Thomas Vaux,⁹ Katy White,⁹ Atul Deodhar ¹⁰,
Irene van der Horst-Bruinsma¹¹

Data from BE MOBILE 1 (non-radiographic axSpA) and BE MOBILE 2 (radiographic axSpA) trials were pooled.

Exposure-adjusted incidence rates of uveitis were significantly lower with bimekizumab vs. placebo.

Separate analyses including all patients and patients with a history of uveitis.



Ustekizumab

SHORT COMMUNICATION

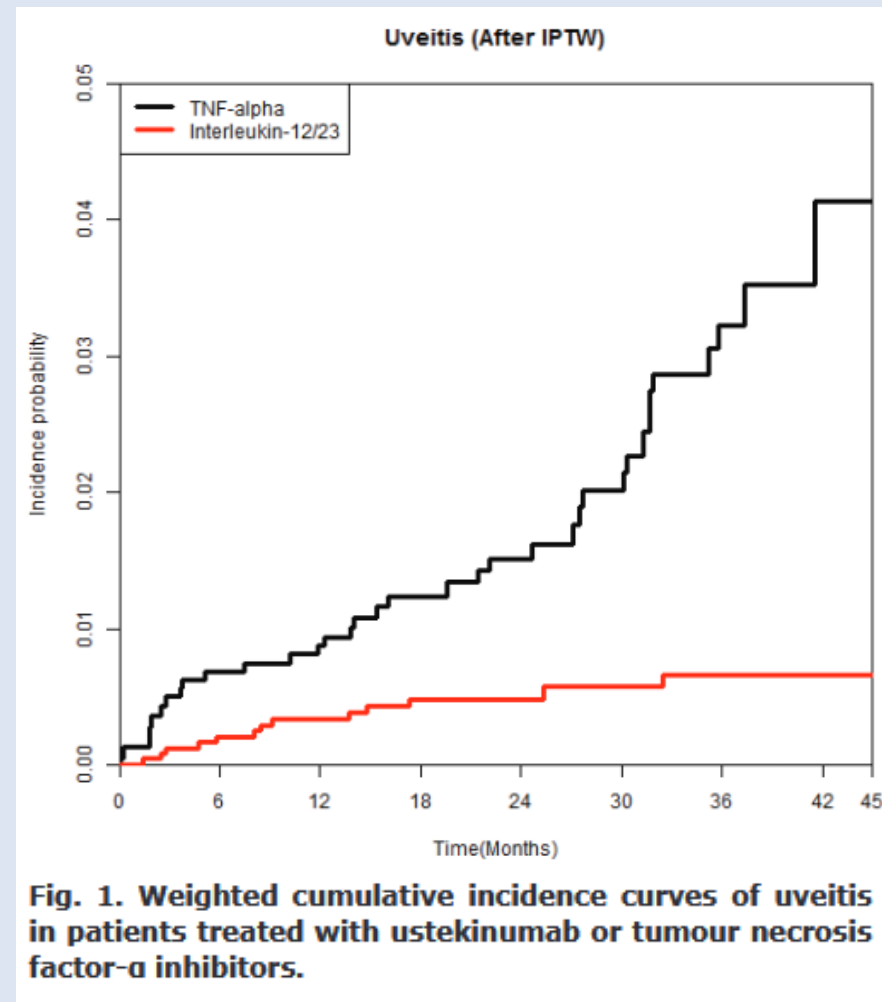
Ustekinumab Demonstrates Lower Uveitis Risk in Moderate to Severe Psoriasis Patients Compared with Tumor Necrosis Factor- α Inhibitors

Chul Hwan BANG¹, Hyun Ju OH¹, Yeong Ho KIM¹, Jin-Hyung JUNG², Ji Hyun LEE¹, Young Min PARK¹ and Ju Hee HAN^{1*}

Data from Korean health insurance database, including >4,000 patients with moderate-to-severe psoriasis receiving ustekinumab or TNFis.

Psoriatic arthritis was adjusted for.

Inverse probability of treatment weighting (IPTW) was used.



JAMA Ophthalmology

RCT: Filgotinib in Active Noninfectious Uveitis

POPULATION

29 Men, 43 Women



Adults with active noninfectious intermediate uveitis, posterior uveitis, or panuveitis

Mean age, 46 y

INTERVENTION

66 Participants analyzed



32 Filgotinib
Oral filgotinib, 200 mg,
once daily for up to 52 wk



34 Placebo
Oral placebo once
daily for up to 52 wk

SETTINGS / LOCATIONS



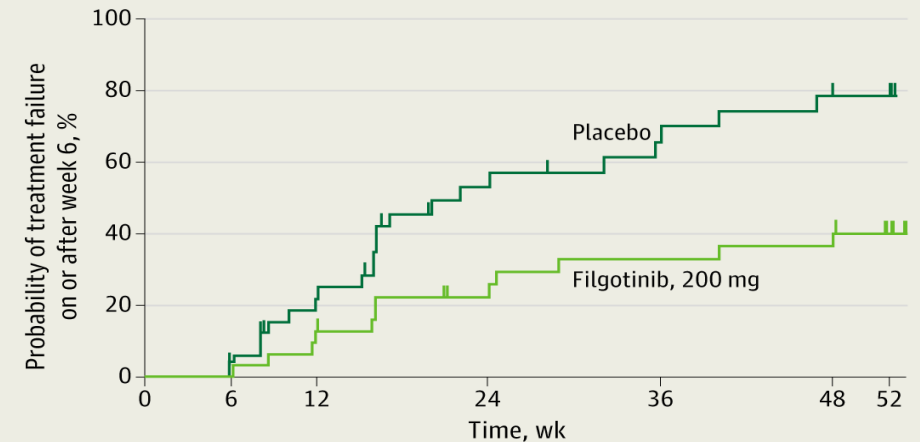
**26 Centers
in 7 countries**

PRIMARY OUTCOME

Treatment failure by week 24: composite end point represented by assessment of the presence of chorioretinal and/or retinal vascular lesions, best-corrected visual acuity, and anterior chamber cell and vitreous haze grades

FINDINGS

Despite early trial termination due to a business decision, a significantly reduced proportion of participants who received filgotinib experienced treatment failure by week 24 vs placebo



Adjusted odds ratio for treatment failure

Filgotinib vs placebo: 0.23; 95% CI, 0.08-0.69; $P = .008$

Case conclusion

Switch to certolizumab pegol.

Remission of uveitis, maintenance of minimal disease activity in skin and joints.

Concluding remarks

Uveitis is less frequent in PsA than in axSpA.

PsA arthritis patients that resemble more closely axSpA patients (axial involvement, HLA-B27) are at higher risk.

Treatment is similar to axSpA-associated uveitis and monoclonal TNF inhibitors are the drugs of choice for recurrent uveitis (but some unique targets, eg IL12-23, may also be important).

Ευχαριστώ για την προσοχή σας!