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Table 1. MOG-AR risk scores

# Introduction

- Myelin oligodendrocyte glycoprotein antibody-associated disease (MOGAD) can present with relapsing or monophasic disease courses.
- Monophasic patients may not require immunosuppressants,

Risk factor	Points
Onset age (45y or older)	2
Female sex	2
Attack phenotype	
Cortical encephalitis	4
ADEM	3
Optic neuritis	2
Cerebral monofocal	1
Brainstem or cerebelar	1
Myelitis	0
No use of immunosupressive therpay	5
Oral steroids (< 3 months)	3

## highlighting the need to identify this group.

- Previous studies proposed monitoring anti-MOG antibody seroreversion or utilizing clinical scores to pinpoint patients with low risk of relapse.
- However, these methods lack reproducibility across cohorts and utilize only select predictors.

Adapted from Yun, 2024. Risk grade: grade 1 = 0-4; grade 2 = 5-8; grade 3 = 9-12; grade 4 = 13-16.

#### Figure 1. Time to relapse in MOGAD patients



# Objective

 This study aims to assess the association of seroreversion and MOGAD-AR scores<sup>1</sup> with monophasic MOGAD.

#### Methods

• We conducted a retrospective observational study involving MOGAD

adult patients from the Demyelinating Diseases Outpatient Clinic at HCFMUSP, São Paulo, Brazil, in 2024.

- Patients were identified through a REDCap database, with inclusion based on the 2023 international diagnostic criteria.
- Exclusion criteria included those without follow-up appointments or with initially negative anti-MOG antibodies.
- Data on demographics, disease characteristics, treatment, and seroreversion were collected.
- Cox regression was used to evaluate hazard ratios (HR) for relapse based on MOG-AR risk grades (1-2 versus 3-4) (Table 1), seroreversion and a combination of both (multivariate analysis)

## Results

• The study included 46 patients (37% male, median age at first attack

## Discussion

- Our study was characterized by a high relapse rate in MOGAD.
  Monophasic disease was linked to seroreversion, but not MOG-AR scores. Further validation of the MOG-AR score is needed.
- Our study may be limited by selection bias due to the nature of a referral tertiary outpatient clinic for adult demyelinating diseases.

31 (IQR 24 - 40) years).

- The most common initial manifestation was optic neuritis (76%).
  Relapses occurred in 76% of patients, with a median time to relapse of 17.5 (IQR 6.5, 57.25) months (Figure 1).
- No significant association was found between higher MOG-AR risk grades and relapse (univariate HR 0.55, 95% CI 0.28 - 1.10, p = 0.09; multivariate HR 0.75, CI 0.26-2.12, p = 0.59).
- Seroreversion occurred in 67% of those tested, acting as a protective factor against relapse (univariate HR 0.29, 95% CI 0.09 0.88, p = 0.03; multivariate HR 0.27, 95% CI 0.09 0.86, p = 0.03).

## Conclusion

Seroreversion of anti-MOG antibodies significantly predicted monophasic disease, while MOG-AR risk grades, isolated or combined with anti-MOG seroreversion, did not effectively indicate relapse risk in our study.

# Disclosure

• Authors have nothing to disclose.

1. Xu, Yun, et al. "A simple score (MOG-AR) to identify individuals at high risk of relapse after MOGAD attack." *Neurology: Neuroimmunology* & *Neuroinflammation* 11.6 (2024): e200309.

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