

# Intravenous immunoglobulin for the treatment of quetiapin-induced bullous pemphigoid in an HIV-infected patient

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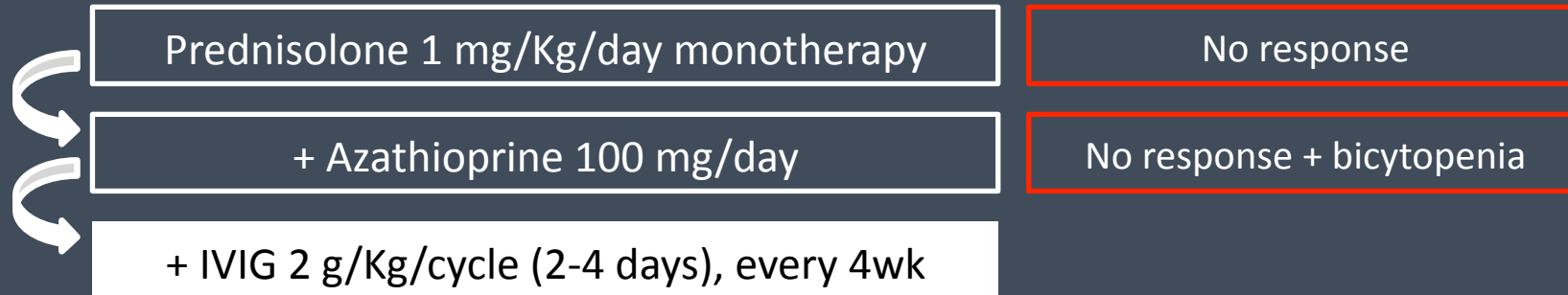
## Case Presentation

- 48-year-old woman
- HIV infection (CD4 319/ $\mu$ L, viral load <50 c/mL), dementia, dyslipidemia
- Polymedicated (HAART, neuroleptics, statin)
- **Bullous pemphigoid**
  - Light microscopy, DIF
  - Persistent eosinophilia
  - ↑ PCR
  - Excluded autoimmune and paraneoplastic association



# Case Presentation

## Treatment:



- ✓ Temporary disease control
- ✓ Well tolerated, no severe side-effects

but:

- ➔ Refractory disease with corticosteroid tapering
- ➔ Persistently elevated eosinophilia, and rising...

Persistent  
trigger? DRUG?

# Case Presentation

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## Reviewing drug history:

- **Quetiapine:** the last introduced drug; 1200 mg/day (above the maximum recommended daily dose)
  - Quetiapine was discontinued
    - Complete clinical resolution
    - Resolution of eosinophilia
  - Inadvertent rechallenge with olanzapine
    - New flare and eosinophilia
  - Complete sustained response with quetiapine/related drug withdrawal
    - No need for immunosuppressors or immunomodulators





# Discussion

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## 1. Bullous pemphigoid in HIV patient

- HIV and autoimmunity: a disease continuum?
- Autoimmune diseases are common, but bullous pemphigoid is rare (4 cases)

## 2. Quetiapine-induced bullous pemphigoid

- Neuroleptics are associated with bullous pemphigoid
- Quetiapine: 1 previous report

## 3. IVIG in HIV patient

- An alternative to immunosuppression
- Widely used in other diseases in immunosuppressed patients

## Conclusions:

- This rare case of drug-induced bullous pemphigoid in an HIV patient suggests quetiapine as a drug trigger for this condition
- IGIV appears to be a safe and effective alternative to immunosuppressors in immunodeficient patients with severe bullous pemphigoid
  - Safety, efficacy, and minimum effective dose studies in this subpopulation are warranted

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