

Prevalence of Comorbidities in Adults with Severe Asthma: Results from the International Severe Asthma Registry (ISAR)

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Introduction

- Comorbidities complicate asthma management and have implications for treatment.
- Comorbidities may worsen asthma outcomes.
- Published data are scarce in severe asthma patients.
- Three main categories:
 - Potentially T2-related comorbidities
 - Potentially OCS-related comorbidities
 - Comorbidities mimicking/exacerbating asthma

Aim

- To understand the pattern of comorbidities in adults with severe asthma and investigate their association with asthma-related outcomes.

Methods

- Study population:** Adult patients from 22 countries enrolled in ISAR.
- 30 individual comorbidities**
- Assessment of comorbidity history:**
 - Case report forms (most contributing countries): specific categorical fields + free-text fields
 - Electronic medical record data mining (USA)
 - Information collected from all available visits (period prevalence)



Table 1. Study sample characteristics (N=11,821).

Characteristics	N	(%)
Sex: Female	7,352	(62.2)
Age:	Median = 56 (IQR: 45, 66)	
Selected therapeutics:		
Long-term OCS	2,792	(23.8)
Biologics	5,428	(45.9)
Age at asthma onset:	Median = 30 (IQR: 15, 45)	
<12 years old	1,194	(20.7)
≥12 years old	4,584	(79.3)
Duration between enrolment and latest visits (years):	Median = 0.71 (IQR: 0, 2.1)	

Results

Table 2. Prevalence of individual comorbid conditions in patients enrolled in the ISAR (24/01/2022 database).

Comorbidities	Prevalence	Sample size*	N countries contributing
Potentially T2-related			
Allergic rhinitis	49%	11,281	22
Chronic rhinosinusitis ¹	38%	11,177	21
Nasal polyposis	21%	11,613	22
Eczema/atopic dermatitis	10%	11,600	22
Urticaria	3.5%	6,849	4
Food allergy	3.3%	6,977	5
Aspirin sensitivity	1.6%	7,498	7
Eosinophilic esophagitis	0.52%	6,149	3
Potentially OCS-related			
Obesity	42%	11,583	22
Hypertension	23%	9,252	12
Sleep apnea	22%	10,094	21
Dyslipidemia	16%	6,849	4
Anxiety/depression ²	14%	11,019	21
Osteoporosis	13%	10,742	21
Diabetes	12%	11,422	22
Coronary heart disease	8.9%	11,039	22
Pneumonia	8.5%	10,300	20
Other significant infections	8.1%	6,918	20
Peptic ulcer	2.6%	10,323	20
Pulmonary embolism/VTE	2.5%	9,972	20
Cataract	2.4%	10,923	21
Renal failure	1.5%	11,032	21
Adrenal insufficiency	1.3%	6,149	3
Glaucoma	1.3%	10,888	21
Cerebrovascular accident	0.63%	9,968	20
Mimicking/exacerbating asthma			
Gastro-esophageal reflux disease ³	44%	7,400	7
Chronic obstructive pulmonary disease	14%	7,508	7
Bronchiectasis	11%	7,509	7
VCD/laryngeal spasms	11%	7,199	5
Dysfunctional breathing	3.2%	7,389	6

*Variations in sample size are due to missing values for individual patients and/or at the country level.
 1. With or without nasal polyposis; with or without allergic rhinitis.
 2. Can also mimic/exacerbate asthma.
 3. Can also be OCS-related.

Figure 1. Co-occurrence of comorbidities across three categories in patients with complete data (N=7,561 patients; 7 countries).

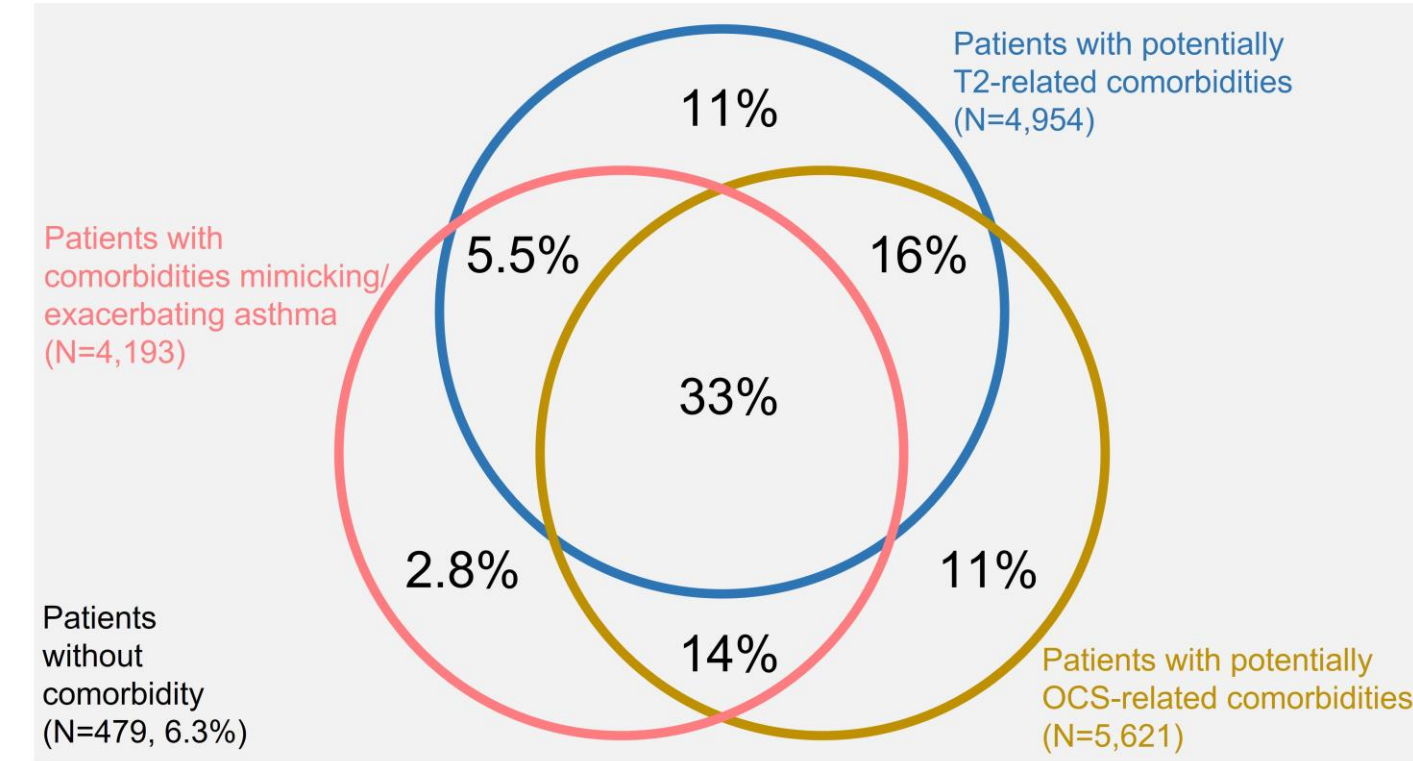


Figure 2. For those with data on at least 3 comorbidities of any type, number of comorbidities reported in ISAR patients overall and by categories.

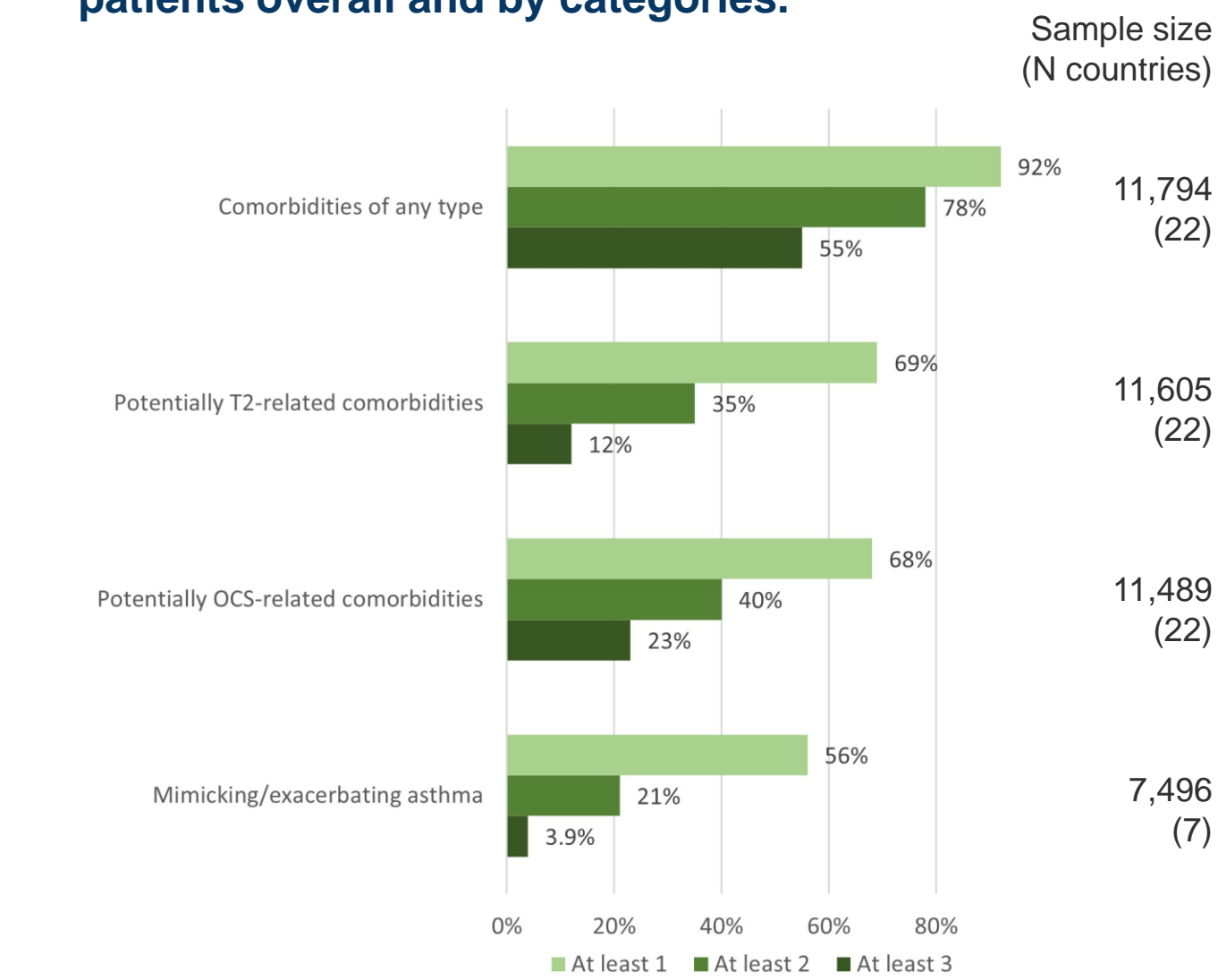
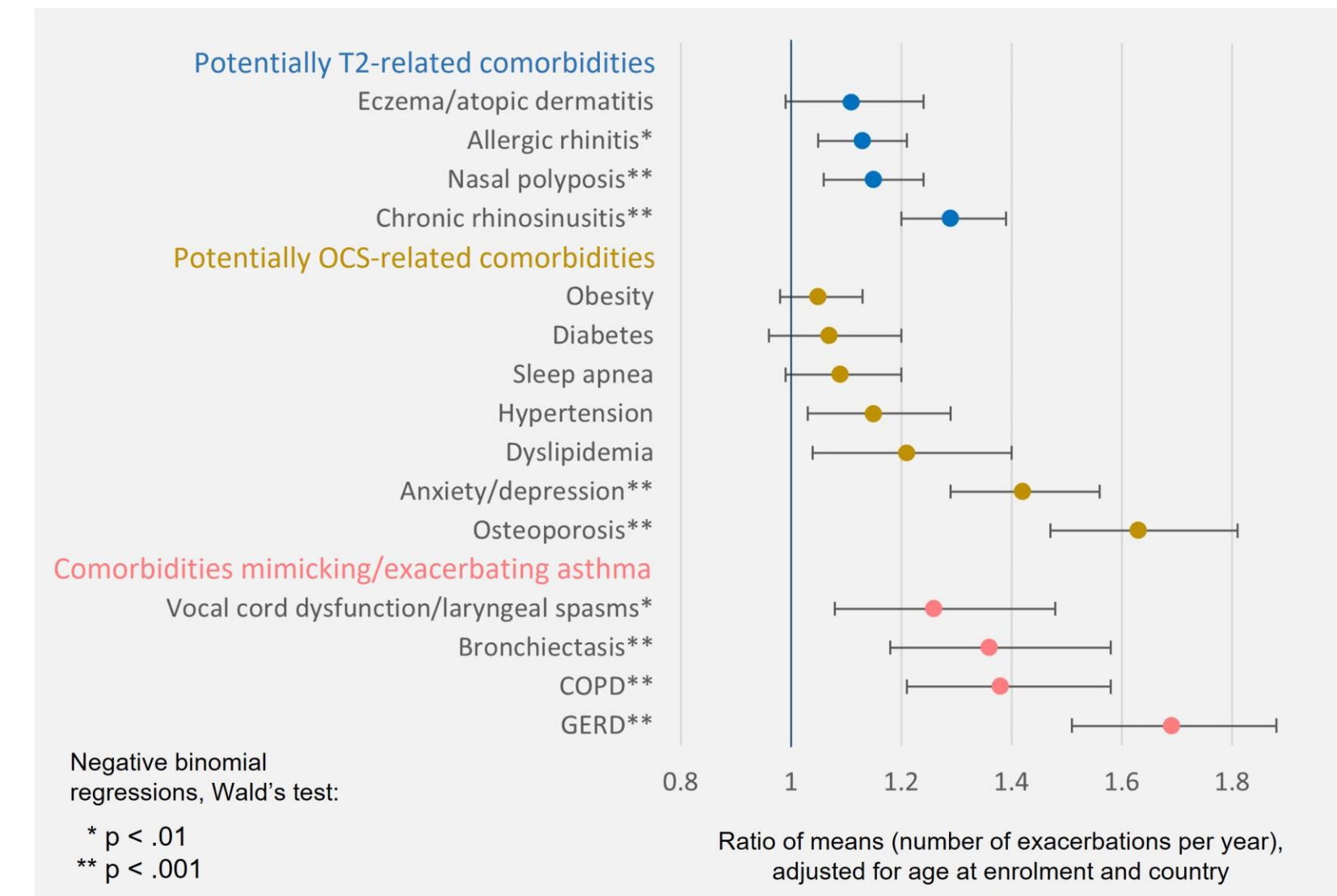


Figure 3. Association between comorbidities and exacerbation rates at enrolment.



Note: Pre-biologic exacerbation rates for patients undergoing biologic therapy.

Conclusions

- High prevalence of comorbidities (T2- and non T2-related) in real-world severe asthma patients, underscoring the importance of systematic evaluation for comorbidities and multidisciplinary approach.
- Co-occurrence of multiple comorbidities within patients:
 - 55% of patients with at least 3 comorbidities
 - 33% of patients with comorbidities spanning the 3 categories
- Comorbidities are associated with higher exacerbation rates.
- Future work: investigate the impact of T2-related comorbidities on response to biologics.

Abbreviations

COPD: Chronic obstructive pulmonary disease; GERD: Gastro-esophageal reflux disease; IQR: Interquartile range; ISAR: International Severe Asthma Registry; OCS: Oral corticosteroids; T2: Type 2 inflammation.

Acknowledgements

Writing, editorial support, and/or formatting assistance in the development of this poster was provided by Joash Tan, BSc (Hons), of the Observational and Pragmatic Research Institute, Singapore, and was funded by AstraZeneca.

Disclosures

This study was conducted by the Observational and Pragmatic Research Institute (OPRI) Pte Ltd and was partially funded by Optimum Patient Care Global and AstraZeneca Ltd. No funding was received by the Observational & Pragmatic Research Institute Pte Ltd (OPRI) for its contribution. **Presenter's conflict of interest disclosure:** David Price full COI disclosures can be found in "COI disclosures" QR

