Baseline characteristics of severe asthma patients initiating biologic treatment worldwide

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Figure 1: BEAM study population

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Introduction

To identify responders to biologic treatment, patients' pre-therapy characteristics must be understood.

Aim

To describe the characteristics of biologic users at initiation.

Methods

Study Design and Population

 In this analysis, we described and compared the baseline characteristics of ISAR patients who initiated anti-IgE or anti-IL5/5R

Patient Inclusion

- Observational, cohort study using patient (≥18 years old, GINA Steps 5 or uncontrolled GINA 4) data from the International Severe Asthma Registry (ISAR; http://isaregistries.org/)
- Patients treated with anti-IL5/5R or anti-IgE during study period (May 2015 to July 2021)
- Available registry data from biologic initiation date until a follow-up visit that is closest to a 24-weeks period
- Had at least one year of pre-therapy information

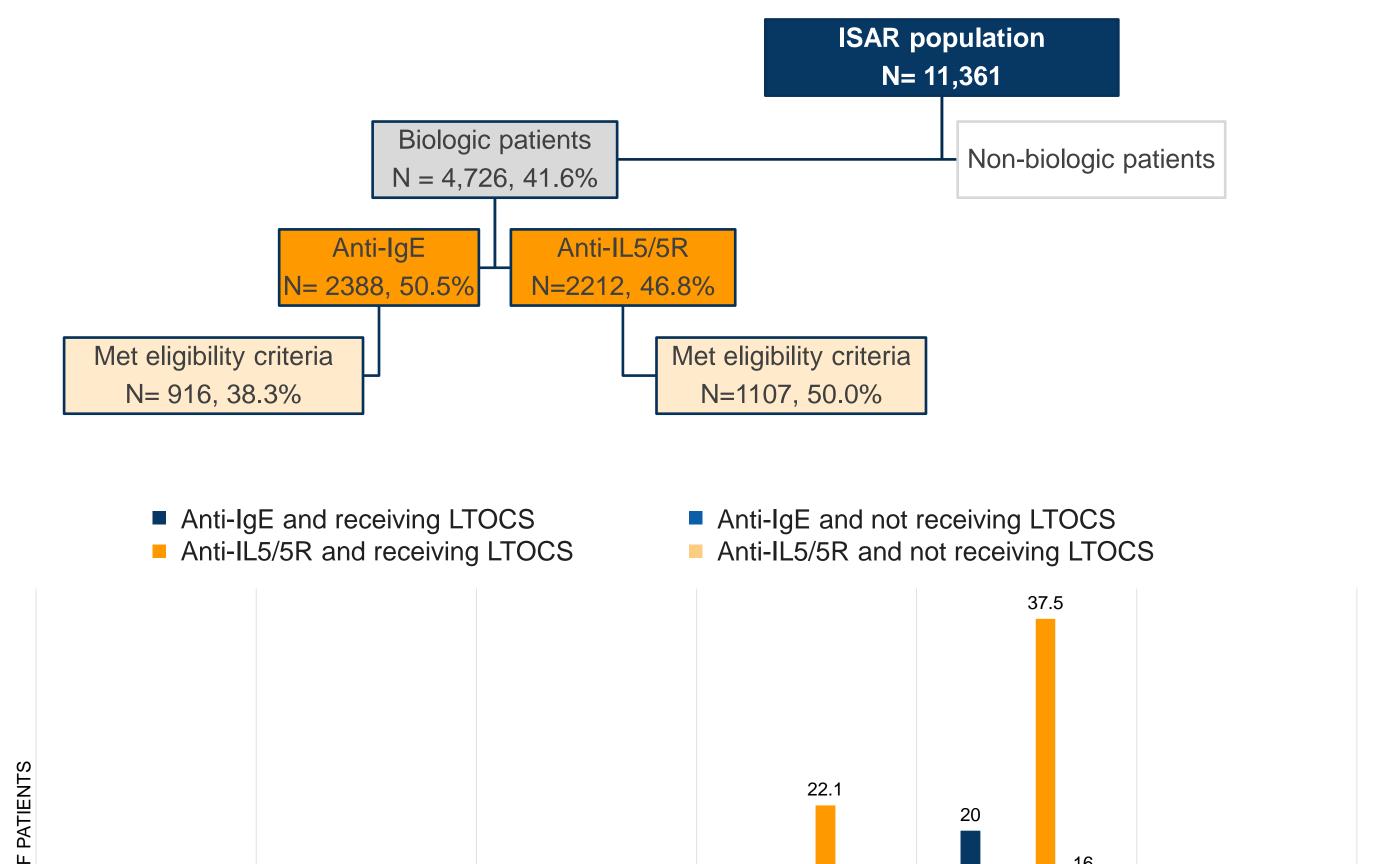


Figure 2: OCS related comorbidities among anti-IgE and anti-IL5/5R patients

(N=394)

Results

Table 1: Baseline characteristics among anti-IgE and anti-IL5/5R patients in the BEAM study

patients in the BEAW Study			
Characteristics	Anti-IgE	Anti-IL5/5R	P-value
Characteristics	(N = 916)	(N = 1107)	r-value
Age, N (%)	916 (100%)	1107 (100%)	
18 - 34, N (%)	159 (17.4%)	98 (8.9%)	< 0.001
35 - 54, N (%)	351 (38.3%)	419 (37.9%)	
55 - 79, N (%)	382 (41.7%)	567 (51.2%)	
80+, N (%)	24 (2.6%)	23 (2.1%)	
Gender, N (%)	916 (100%)	1107 (100%)	
Female	593 (64.7%)	642 (58%)	< 0.001
Age at onset, N (%)	429 (46.8%)	780 (70.5%)	
mean (SD)	24.3 (17.2)	31.35 (18.6)	< 0.001
BMI, N (%)	862 (94.1%)	989 (89.3%)	
mean (SD)	33.57 (11.3)	29.97 (8.4)	< 0.001
Smoking status, N (%)	843 (92%)	981 (88.6%)	
Ex, N (%)	205 (24.3%)	311 (31.7%)	
Current, N (%)	33 (3.9%)	16 (1.6%)	< 0.001
Never, N (%)	605 (71.8%)	654 (66.6%)	
Asthma control, N,(%)	527 (57.5%)	689 (62.2%)	
Well controlled	153 (29%)	102 (14.8%)	
Partially controlled	44 (8.3%)	108 (15.7%)	< 0.001
Uncontrolled	330 (62.7%)	479 (69.5%)	
Receiving LTOCS	426 (46.5%)	667 (60.3%)	< 0.001
LTOCS dose (mg), N (%)	224 (52.6%)	454 (68.1%)	
mean (SD)	13.53 (10.4)	13.1 (10.2)	0.608
≤5 mg	67 (29.9%)	124 (27.3%)	0.049
Percent of predicted FEV ₁ , N (%)	835 (91.2%)	967 (87.4%)	
mean (SD)	81.93 (23.0)	77.2 (23.5)	< 0.001
<80%	397 (47.5%)	531 (54.9%)	0.003
FEV₁/FVC Ratio, N (%)	864 (94.3%)	1045 (94.4%)	
mean (SD)	0.71 (0.1)	0.67 (0.2)	< 0.001
<0.7	375 (43.4%)	573 (54.8%)	0.01
	Annual even	ts	
Exacerbation, N (%)	442 (48.3%)	902 (81.5%)	
2+	353 (79.9%)	690 (76.5%)	0.288
Hospital admission, N (%)	897 (98%)	1068 (96.5%)	
mean (SD)	0.29 (1.2)	0.53 (1.6)	< 0.001
Emergency room admission, N (%)	897 (98%)	1068 (96.5%)	
mean (SD)	1.26 (4.1)	0.87 (2.8)	0.002
Invasive ventilation, N (%)	897 (98%)	1068 (96.5%)	
mean (SD)	0.03 (0.3)	0.09 (0.5)	0.005
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Anti-IL5/5R patients at or before biologic initiation compared with anti-IgE patients:

- Were older
- Had a greater age of asthma onset
- Had lower proportion of female patients
- Had lower BMI
- Had worse lung function
- Had more uncontrolled asthma
- Received long-term oral corticosteroids (LTOCS)
- Reported more potential steroid-related comorbidities (Figure 2)
- Were more likely to report invasive ventilations and hospital admissions (Table 1)

There were no significant differences between annual events such as exacerbations. However, anti-IgE patients were more likely to report emergency room visits than anti-IL5/5R patients at biologic initiation.

Conclusions

At biologic initiation, anti-IL5/5R patients seemed to have more severe asthma characteristics.

Clinical responses such as exacerbation, asthma control, LTOCS dosages, spirometry and healthcare resource utilization at post biologic periods will inform the effect of therapy on these patients.

Abbreviations

Acknowledgments

(N=780)*

Disclosures

DEPRESSION (N=



DIABETES (N=399)* PNEUMONIA (N=395)

ANXIETY (N=481)*



disclosures



of the poster



Summary







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