

Impact of Inititing Biologics in Patients With Severe Asthma on Longterm Oral Corticosteroids or Frequent Rescue Steroids (GLITTER): Data From the International Severe Asthma Registry

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Aim and Methods



Aim

To examine the effectiveness of initiating biologics in a large, real-world cohort of adult patients with severe asthma and high oral corticosteroid exposure (HOCS)*.

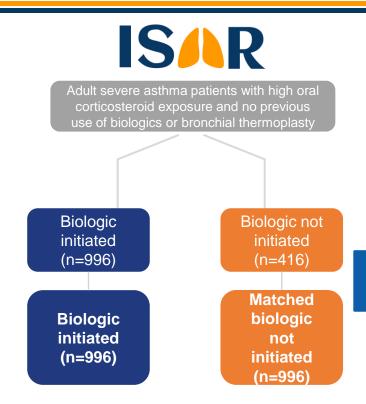
Outcomes

Primary outcome: reduced rate of asthma exacerbations

Secondary outcomes

- · improvement in asthma control
- reduction in OCS dose
- reduced number of asthma-related emergency department visits and asthma-related hospital admissions

Outcomes were estimated over a 12-month follow-up period.

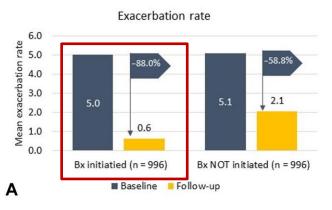


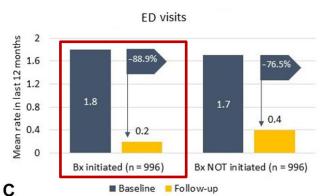
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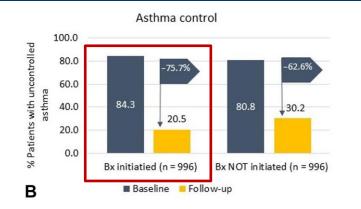


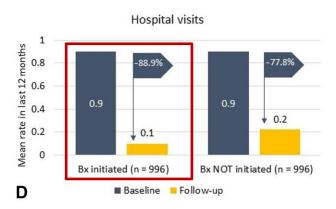
Change from baseline in exacerbation rates, asthma control, emergency department visits, and hospital visits











Patients who initiated a biologic experienced an 88.0% reduction in exacerbation rates, an 89% reduction in emergency department visits and hospital visits, and a 76% reduction in asthma control in the 12-month follow-up period.



Effectiveness of biologic initiation vs non-initiation on OCS reduction



Outcome	Biologic not initiated	Biologic initiated	Marginal difference in % probability (95% CI)	Relative risk (95% CI)
Total OCSs				
Increased dose (%)	27.6	16.0	-11.6 (-29.8 to 6.7)	0.51 (0.17 to 1.51)
Low reduction (%)	63.6	54.4	-9.2 (-24.8 to 6.4)	0.87 (0.61 to 1.24)
Moderate reduction (%)	5.5	16.2	10.7 (4.2 to 17.3)	3.82 (1.58 to 9.25)
Optimal reduction (%)	3.3	13.4	10.0 (-0.6 to 20.7)	7.73 (0.71 to 84.27)
Long-term OCSs				
Increased dose (%)	14.3	8.6	-5.7 (-18.0 to 6.5)	0.51 (0.12 to 2.17)
Low reduction (%)	73.6	68.5	-5.1 (-22.5 to 12.3)	0.94 (0.69 to 1.28)
Moderate reduction (%)	4.2	8.9	4.8 (-1.7 to 11.2)	2.55 (0.78 to 8.37)
Optimal reduction (%)	7.9	14.0	6.1 (-7.7 to 19.9)	4.16 (0.21 to 82.18)

Patients who initiated a biologic were **2.48 times more likely** to achieve a daily total OCS dose of <5 mg compared with those who did not (estimated risk probability of 38.0% vs 15.3%; P = .011) and **2.20 times more likely** to achieve a daily long-term OCS dose (i.e., maintenance dose only) of <5 mg (risk probability, 49.6% vs 22.5%; P = .002).

Compared with those who did not initiate a biologic, those who initiated a biologic were **7.73 times more likely** to have an optimal (>75%) total OCS reduction.



Effectiveness of biologic initiation vs non-initiation on healthcare resource utilisation



Outcome	Biologic not initiated	Biologic initiated	Marginal difference in % probability (95% CI)	Relative risk (95% CI)
ED Visits				
Risk of ED visit (%)	14	6	-9 (-14, -3)	0.35 (0.21, 0.58)
Hospitalisation				
Risk of hospitalization (%)	12	5	-7 (-10, -3)	0.31 (0.18, 0.52)

Compared with patients who did not, patients who initiated a biologic had approximately **one-third the risk and frequency** of asthma-related emergency department visits and hospitalizations (i.e., serious exacerbations).



Conclusions





In a real-world setting, keeping severe asthma patients on HOCS or initiating biologics can both result in improvements in severe asthma.



However, HOCS patients who received biologics experienced the combined benefit of improvements in health outcomes (including exacerbation rates, and healthcare resource utilization) whilst being able to reduce high levels of both short- and long-term oral steroid exposure.

