

Patient-Reported Outcome Measure: CRS-PRO

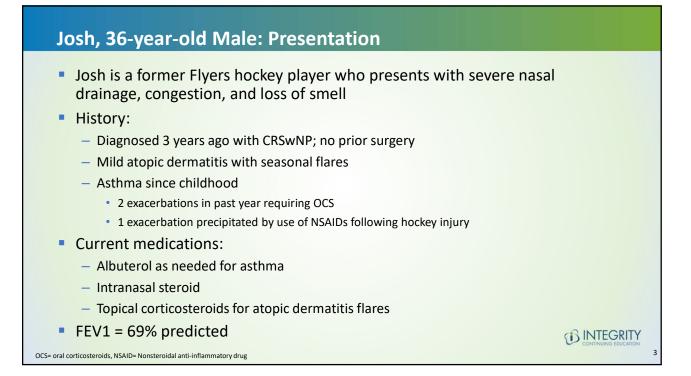
- 12-item patient-completed questionnaire assessing impact of CRS in previous 7 days
- Concise, valid, and reliable measure of CRS patient impact
- Developed with extensive input from patients with CRS
- Correlated highly with SNOT-22 in validation study

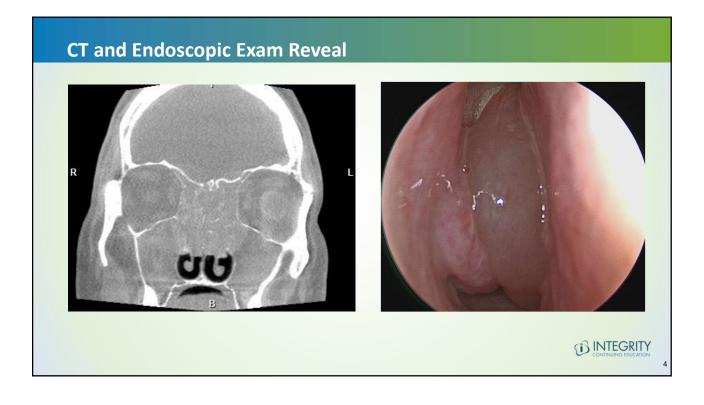
Physical Symptoms	Not at All	A Little Bit	Somewhat	Quite a Bit	Very Much
1. I had difficulty breathing through my nose	0	1	2	3	4
2. I felt pressure in my face	0	1	2	3	4
3. My face hurt	0	1	2	3	4
4. I had to blow my nose	0	1	2	3	4
5. I have been coughing	0	1	2	3	4
6. I had mucus in my throat	0	1	2	3	4
7. I had mucus in my nose	0	1	2	3	4
Sensory Impairment	Not at All	A Little Bit	Somewhat	Quite a Bit	Very Much
8. I had problems with my sense of smell	0	1	2	3	4
Psychosocial Effects	Not at All	A Little Bit	Somewhat	Quite a Bit	Very Much
9. My symptoms kept me awake at night	0	1	2	3	4
10. I felt fatigued	0	1	2	3	4
11. I worried that my condition will get worse	0	1	2	3	4
12. I was frustrated by my condition	0	1	2	3	4

CRS-PRO, Chronic Rhinosinusitis Patient-Reported Outcome; SNOT-22, Sino-nasal Outcome Test.

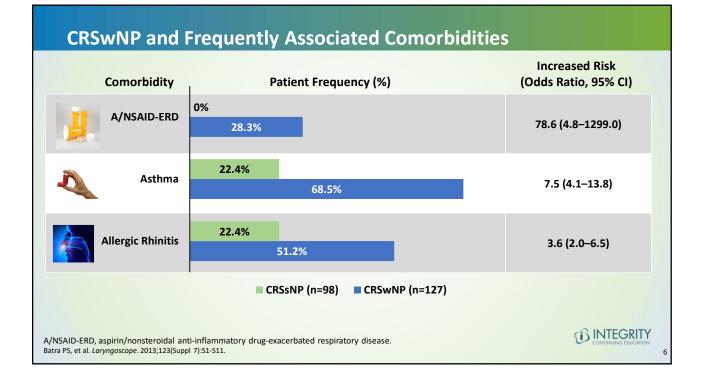
Ghadersohi S, et al. J Allergy Clin Immunol Pract. 2020;8(7):2341-2350.

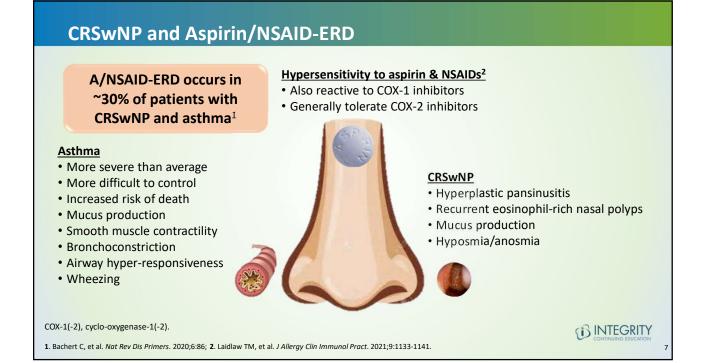
CONTINUING EDUCATION





<section-header> Josh, 36-year-old Male: Care Plan and Follow-up Josh was prescribed: Short course OCS Low-dose ICS for astma Do Fu Advised to avoid NSAIDS or ASA On follow-up 2 months later: Josh reports compliance with therapies and described temporary relief following previous visit After OCS, symptoms returned and he is experiencing a loss of smell and inability to sleep Had asthma exacerbation 10 days ago requiring acute treatment with repeated bursts of albuterol and reports he had increase in daily asthma symptoms in week preceding exacerbation Apspital pulmonologist prescribed a medium dose ICS + LABA for maintenance therapy and to be used as a reliever

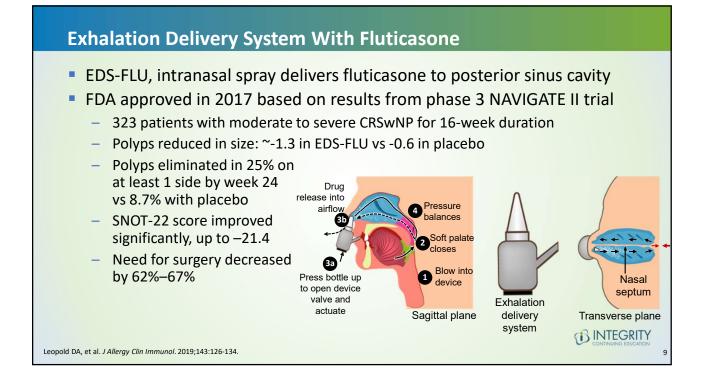


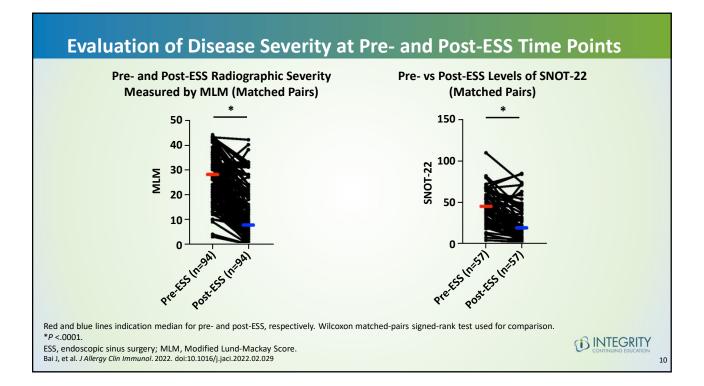


Standard-of-Care Medical Management

- Saline irrigation
 - Some benefit (compared with placebo) from daily, large-volume (150 mL) irrigation with hypertonic solution
 - No benefit from low-volume (5 mL) nebulized saline spray
- Intranasal corticosteroids
 - Nasal sprays, nasal installations/drops, EDS-FLU
- Oral corticosteroids
 - Acute relief for severe blockage and loss of smell
- Antibiotics
 - Prior RCT demonstrating small but significant benefit using doxycycline, possibly due to contribution of *S. aureus* to CRSwNP pathogenesis

Blaiss MS. Allergy Asthma Proc. 2020;41:413-419; Chong LY, et al. Cochrane Database Syst Rev. 2016;4:CD011995; Naclerio R, et al. J Allergy Clin Immunol Pract. 2020;8:1532-1549; Kern RC, et al. Int Forum Allergy Rhino. 2018;8:471-481.







Mean SNOT-22 Score at Each Time Point*

	ESS Plus Medical Therapy Group (n=118)	Medical Therapy Only Group (n=116)
Lund-Mackay score, points		
Mean	18.4 (4.3)	18.5 (4.9)
0–4	1/112 (1%)	0/111 (0%)
5–9	1/112 (1%)	4/111(4%)
10-14	20/112 (18%)	27/111 (24%)
15–24	90/112 (80%)	80/111 (72%)
Aeroallergen sensitization	64 (54%)	62 (53%)
SNOT-22 score		
Mean	51.9 (20.4)	50.5 (19.7)
0 to <20	10 (8%)	4 (3%)
20 to <40	24 (20%)	33 (28%)
40 to <60	39 (33%)	42 (36%)
60 to <80	34 (29%)	28 (24%)
≥80	11 (9%)	9 (8%)
EQ-5D-5L utility score	0.8 (0.2)	0.8 (0.2)
EQ-5D-5L VAS, mm	70.9 (17.02)	70.0 (17.2)

*Error bars indicate SDs. In the medical therapy group, 116 patients were assessable at baseline, 113 at 3 months, 107 at 6 months, and 103 at 12 months. In the ESS plus medical therapy group, 118 at baseline, 106 at 3 months, 107 at 6 months, and 103 at 12 months. *The minimal clinically important difference of SNOT-22 is 9 points. Adjusted mean differences at 3, 6, and 12 months, were -15.2, -8.3, and -4.9, respectively. EQ-5D-5L, EuroQuol Five Dimension, Five Level scale; SD, standard deviation; VAS, visual analogue scale. Lourijsen E, et al. *The Lancet*. 2022;10(4):337-346.

Martha, 41-year-old Female: Presentation

 Martha is a 41-year-old female with asthma and allergic rhinitis (AR). She presents to your clinic with nasal congestion and a decrease in the sense of smell. On exam you notice bilateral nasal polyps. Her asthma is not bothering her.

PMHx:

- Asthma
- AR (skin test + dust mites, molds)

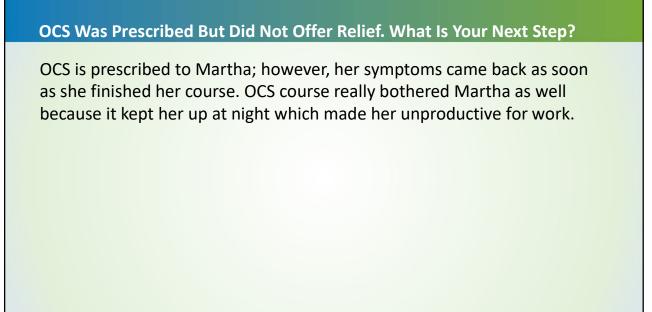
Clinical Values:

FEV₁: 69% predicted

12

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FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity; PMHx, prior medical history.



OCS, oral corticosteroids.

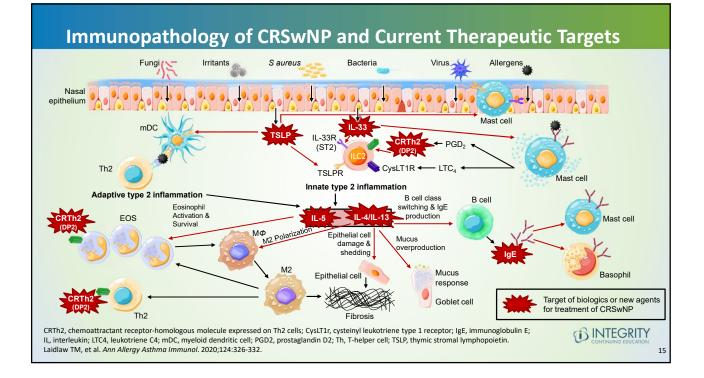
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Treatment Options



ENT evaluation for surgery and/or biologic

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FDA-Approved Biologics for CRSwNP

Biologic	Target	FDA Indication	Phase 3 Trials	Number of Patients
Dupilumab	IL-4 IL-13	Add-on maintenance treatment in adult patients with inadequately controlled CRSwNP	SINUS-24 ¹ SINUS-52 ¹	276 448
Omalizumab	lgE	Nasal polyps in adult patients 18 years of age and older with inadequate response to nasal corticosteroids, as add-on maintenance treatment.	POLYP-1 ² POLYP-2 ²	138 127
Mepolizumab	IL-5	Add-on maintenance treatment of adult patients 18 years and older with CRSwNP.	SYNAPSE ^{3,4}	407

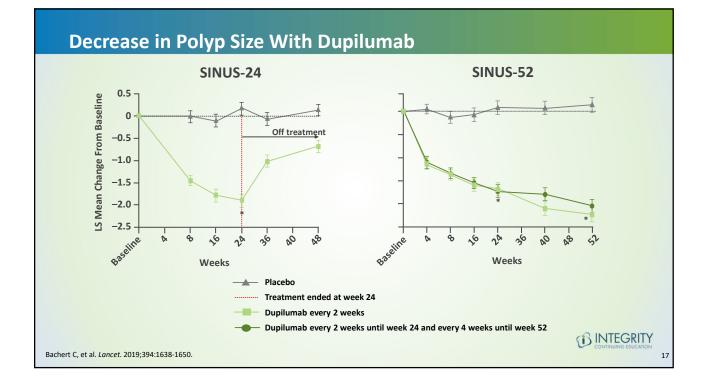
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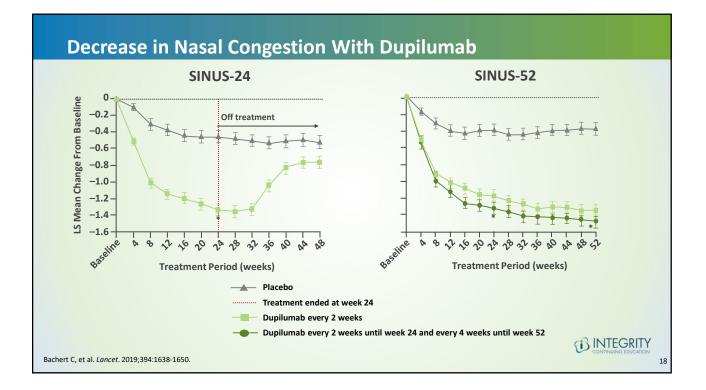
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1. Bachert C, et al. Lancet. 2019;394:1638-1650. 2. Geveart P, et al. J Allergy Clin Immunol. 2020;146:595-605. 3. Han JK, et al. Lancet Respir Med. 2021;9:1141-1153. 4. Hopkins C, et al. Eur Respir J. 2020;56:4616.

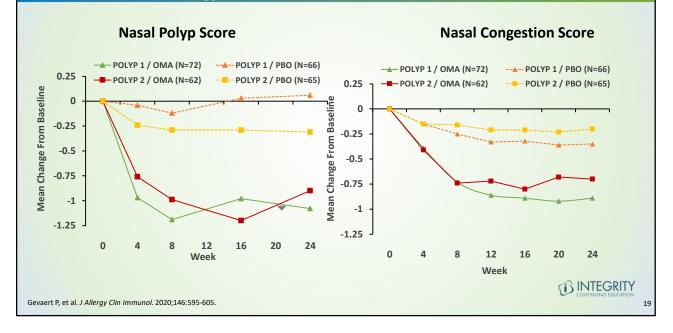
Hopkins C, et al. Eur Respir J. 2020;50:4010.
 Dupixent (dupilumab) [package insert]. Revised 2022. Accessed 2022. https://www.regeneron.com/downloads/dupixent_fpi.pdf
 Xolair (omalizumab) [package insert]. Revised 2021. Accessed 2022. https://www.gene.com/download/pdf/xolair_prescribing.pdf

Nucala (mepolizumab) [package insert]. Revised 2021. Accessed 2022. https://gskpro.com/content/dam/global/hcpportal/en_US/Prescribing_Information/Nucala/pdf/NUCALA-PI-PIL-IFU-COMBINED.PDF.





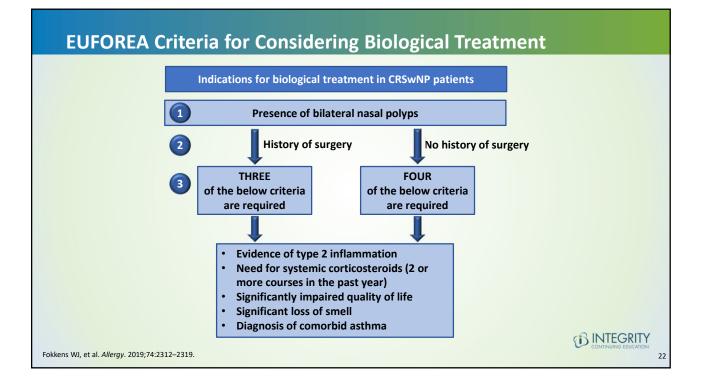
Omalizumab: Change from Baseline to Week 24 in Nasal Congestion Score and Nasal Polyp Score



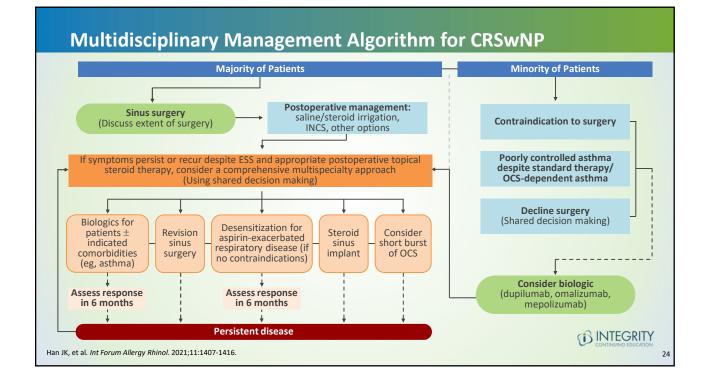
Mepolizumab: SYNAPSE Results Nasal Obstruction VAS Score (Weeks 49-52) Total Endoscopic NPS (Week 52) Median difference (95% Cl): -0.73 (-1.11, -0.34); P <.001 Median difference (95% Cl): -3.14 (-4.09, -2.18); P <.001 1.0-LS Mean Change From Baseline (95% Cl) in Total Endoscopic NPS LS Mean Change From Baseline (95% CI) in Nasal Obstruction VAS Score 0.0 0.0 -1.0 -2.0--0.5 -3.0-4.0 -1.0 5.0

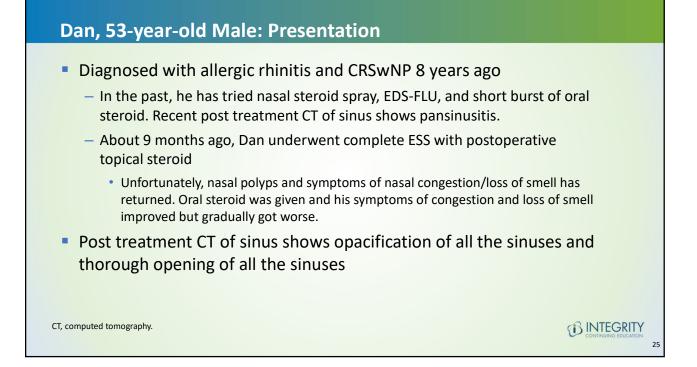


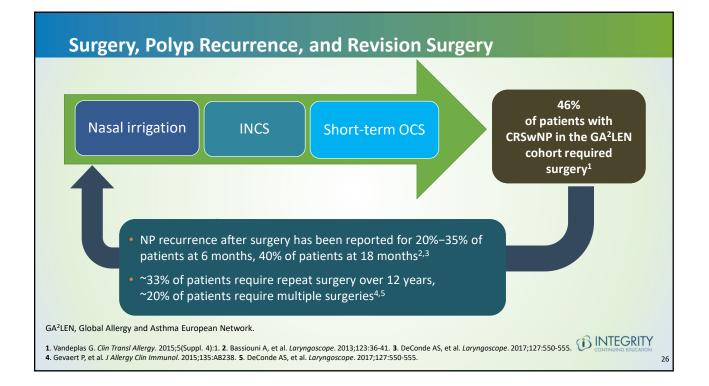
		Patient-important Outcomes						Surrogate Outcomes		
	HRQol SNOT-22 (0-110)	Symptoms VAS (0-10 cm)	Smell UPSIT (0-40)	Rescue OCS	Rescue polyp surgery	Advers	e events	Nasal polyp size (0-8)	CT score LMK (0-24)	
tandard care	50.11	6.84	14.04	31.96%	21.05%	73.78%		5.94	18.35	
Dupilumab	-19.91 (-22.50, -17.32)	-3.25 (-4.31, -2.18)	10.96 (9.75, 12.17)	- 21.73 (-24.61, -18.22) RR 0.32 (0.23, 0.43)	-16.35 (-18.13, -13.48) RR 0.22 (0.14, 0.36)	0.13 (-8.12, 9.88) RR 1.00 (0.83, 1.33)		-2.04 (-2.73, -1.35)	- 7.51 (-10.13, -4.89)	
Dmalizumab	-16.09 (-19.88, -12.30)	-2.09 (-3.15, -1.03)	3.75 (2.14, 5.35)	-12.46 (-23.65, 12.78) RR 0.61 (0.26, 1.40)	-7.40 (-11.04, -2.43) RR 0.65 (0.48, 0.68)	-2.60 (-15.58, 13.28) RR 0.96 (0.79, 1.18)		-1.09 (-1.70, -0.49)	-2.66 (-5.70, 0.37)	
Mepolizumab	- 12.89 (-16.58, -9.19)	-1.82 (-3.13, -0.50)	6.13 (4.07, 8.91)	- 10.23 (-15,98, -2.88) RR 0.68 (0.50, 0.91)	-12.33 (-15.56, -7.22) RR 0.41 (0.26, 0.66)	- 3.07 (-13.44, 9.07) RR 0.96 (0.82, 1.12)		-1.06 (-1.79, -0.34)		
Benralizumab	- 7.68 (-12.09, -3.27)	-1.15 (-2.47, 0.17)	2.95 (1.02, 4.88)	- 9.91 (-16.30, -0.96) RR 0.69 (0.49, 0.97)	-2.53 (-9.05, 7.16) RR 0.88 (0.57, 1.34)	-1.48 (-13.28, 12.54) RR 0.98 (0.82, 1.11)		-0.64 (-1.39, 0.12)	- 1.00 (-3.83, 1.83)	
ASA Desensitization	-10.61 (-14.51, -6.71)	-2.74 (-3.92, -1.57)	2.72 (-1.17, 6.61)		- 16.00 (-19.79, 0.21) RR 0.24 (0.06, 1.01)	(8.30, RR	9.21 901.87) 3.84 13.22)	-0.95 (-2.44, 0.55)	-0.31 (-3.50, 2.88)	
Classification of	intervention cold	or						Certainty	(shading)	
Among most beneficial Among most harmful		Among interme	ediate beneficial	clearly different from		clearly different from		No data	High/moderate (solid)	
		Among interme	ediate harmful			(blank)	Low/very low (shaded)			

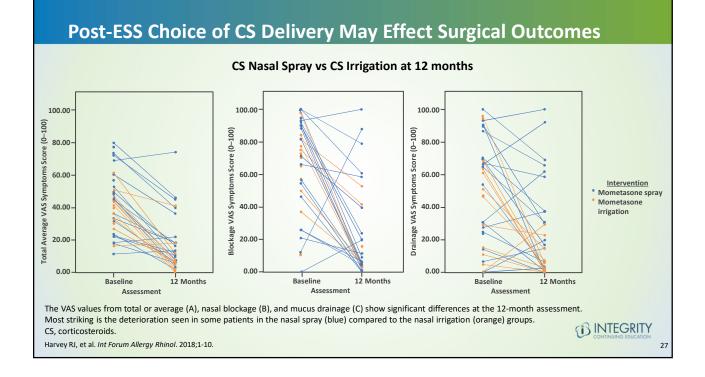


Indications for biological treatment in CRSwNP				
Presence of bilat	eral polyps in a patient who had ESS*			
	THREE criteria are required			
Criteria	Cut-off Points			
Evidence of type 2 inflammation	• Tissue eos ≥10/hpf, OR blood eos ≥250 u/L, OR total IgE ≥100 IU/mL			
Need for systemic corticosteroids or contraindication to systemic steroids	 • ≥2 courses per year, OR long term (>3 months) low dose steroids 			
Significantly impaired quality of life	• SNOT-22 ≥40			
Significant loss of smell	Anosmic on smell test (score depending on test)			
Diagnosis of comorbid asthma	Asthma needing regular inhaled corticosteroids			









Median Time to Revision Surgery or Polyp Recurrence in Patients With CRSwNP Alone or With Comorbidities

Patient Category	Median Years to Revision Surgery	Median Years to Polyp Recurrence
CRSwNP alone	20	20
CRSwNP with asthma	11	4
CRSwNP with asthma and NSAID-ERD	7	0.66

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Bachert C, et al. J Asthma Allergy. 2021;14:127-134; Leung RM, et al. Int Forum Allergy Rhinology. 2014;4(11):871-876.

