Immunodeficiency Screening and Management for Patients with Recurrent Sinusitis

Christine Reger, DNP, CRNP

Katherine Schmitt, BSN, RN



Importance of understanding immunodeficiencies for rhinology patients

Introduction



Common presentations of immunodeficiencies in rhinology practice

The Immune System Basics

Innate, humoral, and cellular immunity

Importance of immune system balance for health

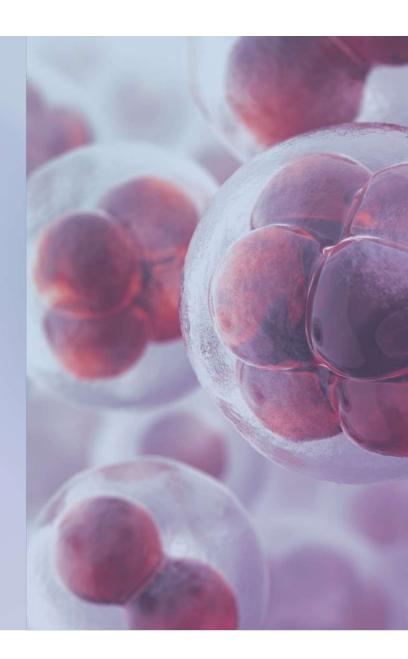
Clinical Presentations

Symptoms indicating potential immunodeficiency

Role of rhinology team in early recognition and referral

Types of Immunodeficiencies

- •Primary Immunodeficiency (PIDs)
 - •Definition and genetic basis
 - •Common types: Specific Antibody Deficiency (SAD), Common Variable Immunodeficiency (CVID)
- •Secondary Immunodeficiency (SIDs)
 - •Causes: infections, malignancy, medications



Evaluation Algorithm



Steps for evaluating adult patients with suspected immunodeficiency



Importance of assessing vaccine responses

Step 1: Has the patient been previously diagnosed or evaluated for an immunodeficiency?

Step 2: Is there a history of recurrent infections as an adult?

Step 3: Was the patient born preterm or full term? Any complications surrounding the delivery, including neonatal intensive care unit (NICU) stay, illnesses, antibiotic use, ventilator support?

Step 4: Was the patient primarily healthy during childhood, excluding the expected viral infections that are known to occur? Or was the childhood marked by recurrent infections and antibiotic use and frequent absences from school?

Step 5: If there is a history of recurrent infections in adulthood and/or childhood, obtain additional history regarding the types and frequency of infections, duration of symptoms, and treatments rendered. Refer to Step 6 and 7.

Step 6: Assess if there is a history of the following:

- Recurrent bacterial infections, such as acute otitis media, sinusitis, and pneumonia?
- Recurrent respiratory viral infections or prolonged duration of these illnesses?
- *Systemic viral infections, such as Epstein-Barr virus (EBV), Cytomegalovirus (CMV), Herpes Simplex Virus (HSV)?
- Bacteremia, meningitis, abscesses, skin or soft tissue infections?
 Fungal infections, such as thrush or fungal sinus or lung infections?
- Recalcitrant warts or eczema?
- •Recurrent or chronic diarrhea or GI disorders?
- · Autoimmune disorders?
- · Hospitalization for illness?
- Travel?
- Did they receive the standard vaccinations for their age, including childhood vaccinations?(17)
- Family history of recurrent infections, clinically diagnosed immunodeficiencies or autoimmune disorders?
 Does the patient have infections and/or physical features of a syndromic disorder?

Has the patient ever undergone clinical genetic testing?

Step 7: Assess for possible etiologies of secondary immunodeficiencies.

Step 8: Based on the aforementioned history, if there are frequent cases of bacterial infections then consider evaluation for a humoral immunodeficiency. If there are frequent viral or fungal infections then consider evaluation for a cellular immunodeficiency.

Step 9: Initial immunology laboratory assessment:

- •Quantitative immunoglobulins (IgA, IgG, IgM)
- Complete blood count (CBC) with differential. Assess Absolute Lymphocyte Count (ALC), Absolute Neutrophil Count (ANC) and Absolute Eosinophil Count (AEC).
- Streptococcus pneumoniae titers. This may test evaluate either 14 or 23 serotypes depending on the lab. Determine if and when pneumococcal vaccine(s) have been received in the patient's lifetime. As noted previously, pneumococcal conjugate vaccine is now part of the childhood vaccine series.
- •Tetanus titer. Determine when tetanus vaccine had been most recently administered in the patient's lifetime. This is

•If any of these labs have been previously checked, compare the trend of the results over time.

Step 10: If any of the aforementioned labs are abnormal, consider a referral to a Clinical Immunologist. If the aforementioned labs are normal but the history is concerning for an immunodeficiency, consider a referral to a Clinical Immunologist.

Specific Antibody Deficiency (SAD)



Characteristics and clinical implications



Relationship with chronic rhinosinusitis (CRS)



Management strategies: antibiotic use, immunoglobulin replacement

Common Variable Immunodeficiency (CVID)



Epidemiology and diagnostic criteria



Treatment options such as immunoglobulin replacement therapy and prophylactic antibiotics

Management Options



Antibiotic prophylaxis



Immunoglobulin replacement therapy administration methods



Considerations for long-term care

Collaborative Care

Importance of multidisciplinary care (rhinology team, clinical allergy/immunologists)

Enhancing patient outcomes through collaborative management

Nursing Implications

Key Responsibilities:

Patient Education

- Explain the importance of follow-up appointments and adherence to prescribed therapies.
- Teach patients and caregivers about recognizing signs of infection and when to seek medical attention.

Support in Treatment Administration

- Assist in administration of immunoglobulin replacement therapy (IV or SC) as per physician's orders.
- Monitor for infusion reactions and provide supportive care as needed.

Documentation and Reporting

- Document patient assessments, treatments, and responses accurately.
- Communicate effectively with the healthcare team to ensure continuity of care.

Role of Nurses in Immunodeficiency Care

Nursing Implications

Educating patients about signs and symptoms of immunodeficiency.

Providing instructions on medication adherence and infection prevention.

Monitoring patients for treatment responses and adverse effects.

Collaborating with rhinologists and clinical immunologists in patient care



Recap of key points discussed

Conclusion



Future directions in managing immunodeficiencies in rhinology practice

References

- 1.Sikora AG, Lee KC. Otolaryngologic manifestations of immunodeficiency. Otolaryngol Clin North Am 2003; 36:647–672
- .2.Wasserman RL, Manning SC. Diagnosis and treatment of primary immuno-deficiency disease: the role of the otolaryngologist. Am J Otolaryngol 2011;32:329–337.
- 3. & Abbas AK, Lichtman AH, Pillai S, Baker DL. Cellular and molecular immunol-ogy. Tenth edition. Philadelphia, Pennsylvania: Elsevier; 2022. 587 p. This reference provides education on the various components of immune system.
- 4. § Samargandy S, Grose E, Yip J, Lee JM. Endoscopic sinus surgery outcomes of endoscopic sinus surgery inpatients with an immunodeficiency.
- 5. &Otani IM, Lehman HK, Jongco AM,etal.Practical guidance for the diagnosisand management of secondary hypogammaglobulinemia: a Work GroupReport of the AAAAI Primary Immunodeficiency and Altered Immune Re-sponse Committees. J Allergy Clin Immunol 2022; 149:1525–1560. This report reviews causes of secondary hypogammaglobulinemia.
- 6.s.Mustafa SS. Steroid-induced secondary immune deficiency. Ann AllergyAsthma Immunol 2023; 130:713-717. This review discusses the immunosuppressive effects of oral corticosteroids.
- 7. EBousfiha A, Moundir A, Tangye SG, etal. The 2022 update of IUIS pheno-typical classification for human inborn errors of immunity. J Clin Immunol2022; 42:1508–1520. This article details the clinical and laboratory phenotypes of PIDs.
- 8. a Tangye SG, Al-Herz W, Bousfiha A, et al. Human inborn errors of immunity: 2022 update on the classification from the International Union of Immuno-logical Societies Expert Committee. J Clin Immunol 2022; 42:1473–1507. This article details the classification of inborn errors of immunity.
- 9. &Meyts I, Bousfiha A, Duff C,etal.Primary immunodeficiencies: a decade ofprogress and a promising future. Front Immunol 2021; 11:625753. This article describes the increase in understanding of PIDs and details concernsthat will need to be addressed in the future.
- 10.Bousfiha AA, Jeddane L, Ailal F,etal.Primary immunodeficiency diseasesworldwide: more common than generally thought. J Clin Immunol 2013;33:1–7.Adult immunodeficiency evaluationBuckeyandBosso1068-9508 Copyright © 2023 Wolters Kluwer Health, Inc. All rights reserved.www.co-otolaryngology.com53Copyright © 2023 Wolters Kluwer Health, Inc. All rights reserved.
- 11.Keswani A, Dunn NM, Manzur A,etal.The clinical significance of specificantibody deficiency (SAD) severity in chronic rhinosinusitis (CRS). J AllergyClin Immunol Pract 2017; 5:1105–1111.
- 12.Tam JS, Routes JM. Common variable immunodeficiency. Am J Rhinol Allergy2013; 27:260–265.13.Odnoletkova I, Kindle G, Quinti I,etal.The burden of common variableimmunodeficiency disorders: a retrospective analysis of the European Societyfor Immunodeficiency (ESID) registry data. Orphanet J Rare Dis 2018;
- 13:201.14.s.Common variable immune deficiency (CVID)jimmune Deficiency Foundation[Internet]. [cited 15 October 2023]. https://primaryimmune.org/understand-ing-primary-immunodeficiency/types-of-pi/common-variable-immune-defi-ciency-cvid. [Accessed 15 October 2023]This reference details common variable immune deficiency.
- 15.McCusker C, Upton J, Warrington R. Primary immunodeficiency. AllergyAsthma Clin Immunol 2018; 14:61.
- 16.Cooney TR, Huissoon AP, Powell RJ, Jones NS. Investigation for immuno-deficiency in patients with recurrent ENT infections. Clin Otolaryngol AlliedSci 2001; 26:184–188.
- 17. Orange JS, Ballow M, Stiehm ER, etal. Use and interpretation of diagnostic vaccination in primary immunodeficiency: a working group report of the Basicand Clinical Immunology Interest Section of the American Academy of Allergy, Asthma & Immunology. J Allergy Clin Immunol 2012; 130:S1–24.
- 18.Tay L, Leon F, Vratsanos G,etal. Vaccination response to tetanus toxoid and 23-valent pneumococcal vaccines following administration of a single dose of abatacept: a randomized, open-label, parallel group study in healthy subjects. Arthritis Res Ther 2007; 9:R38.
- 19. «Vaccines for PneumococcaljCDC [Internet]. 2023 [cited 11 October 2023].https://www.cdc.gov/vaccines/vpd/pneumo/index.html.This reference provides information on pneumococcal vaccinations
- .20.aCDC. Centers for Disease Control and Prevention. 2023 [cited 12 October2023]. Immunization schedules for 18 & younger. https://www.cdc.gov/vaccines/schedules/hcp/imz/child-adolescent.html.This reference provides the immunization schedule for pediatric patients.
- 21.Agarwal S, Cunningham-Rundles C. Treatment of hypogammaglobulinemia inadults: a scoring system to guide decisions on immunoglobulin replacement. J Allergy Clin Immunol 2013; 131:1699–1701.
- 22.s.Immunoglobulin replacement therapyjImmune Deficiency Foundation [Inter-net]. [cited 12 October 2023]. https://primaryimmune.org/understanding-primary-immunodeficiency/treatment/immunoglobulin-replacement-therapy. This reference provides information on history, administration, and management ofimmunoglobulin replacement therapy