

Geographic Variability and Pathogen-Specific Considerations in the Diagnosis and Management of Chronic Granulomatous Disease

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Abstract: Chronic granulomatous disease (CGD) is a rare but serious primary immunodeficiency with varying prevalence and rates of X-linked and autosomal recessive disease worldwide. Functional defects in the phagocyte nicotinamide adenine dinucleotide phosphate oxidase complex predispose patients to a relatively narrow spectrum of bacterial and fungal infections that are sometimes fastidious and often difficult to identify. When evaluating and treating patients with CGD, it is important to consider their native country of birth, climate, and living situation, which may predispose them to types of infections that are atypical to your routine practice. In addition to recurrent and often severe infections, patients with CGD and X-linked female carriers are also susceptible to developing many non-infectious complications including tissue granuloma formation and autoimmunity. The DHR-123 oxidation assay is the gold standard for making the diagnosis and it along with genetic testing can help predict the severity and prognosis in patients with CGD. Disease management focuses on prophylaxis with antibacterial, antifungal, and immunomodulatory medications, prompt identification and treatment of acute infections, and prevention of secondary granulomatous complications. While hematopoietic stem-cell transplantation is the only widely available curative treatment for patients with CGD, recent advances in gene therapy may provide a safer, more direct alternative.

Keywords: CGD, primary immunodeficiency diseases, nicotinamide adenine dinucleotide phosphate oxidase complex, DHR-123 oxidation assay

Introduction

Chronic granulomatous disease (CGD) is a rare primary immunodeficiency that is caused by defects in the nicotinamide adenine dinucleotide phosphate (NADPH) oxidase complex, which is critical for superoxide production. The disease was initially described in 1957 and appropriately named "fatal granulomatous disease of childhood" as all 4 patients died before 6 years of age.¹ Although it was initially thought to solely be an X-linked disease, the first females were diagnosed in 1968, leading to the identification of autosomal recessive (AR) forms.² Affected individuals develop recurrent and severe infections from a narrow range of characteristic bacteria and fungi and suffer from granulomatous inflammation that has become pathognomonic for the disease. Since its original description, improvements in antibacterial, antifungal, and immunomodulatory prophylaxis as well as hematopoietic stem-cell transplantation (HSCT) have allowed patients to live well into adulthood. Herein, we will review

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important concepts in the diagnosis and management of CGD as well as the genetic and geographic variability of the disease.

Epidemiology and Pathophysiology

CGD has a minimum estimated prevalence of 1 in 200,000–250,000 live births in the United States (US).^{3,4} Around the world, the prevalence is somewhat variable. While Japan appears to have a similar rate to the US, the prevalence of CGD has been reported to be as high as 3.43 in 100,000 live births in Jeju Island, Korea, due in part because of a common ancestor. In Israel, the prevalence of CGD is estimated to be 1.17 in 100,000 live births and is predominantly autosomal recessive.⁵ Even within Europe, however, the prevalence ranges from 0.1 in 100,000 in Italy to 0.85 in 100,000 live births in the United Kingdom and Ireland.^{6,7}

The NADPH oxidase complex is comprised of 5 protein subunits and is responsible for generating the reactive oxygen species necessary for phagocyte killing of bacteria and fungi both directly and by the activation of intraphagosomal proteases.^{8,9} Pathogenic variants in any of the 5 genes that encode the 2 membrane-bound subunits (gp91^{phox} encoded by *CYBB* and p22^{phox} encoded by *CYBA*) or the 3 cytosolic subunits (p47^{phox} encoded by *NCF1*, p67^{phox} encoded by *NCF2*, and p40^{phox} encoded by *NCF4*) of the complex lead to the clinical presentations that are CGD.^{10,11} Recently, biallelic, loss-of-function mutations in *CYBC1* have also been found to cause decreased NADPH function resulting in CGD.^{12,13} While *CYBC1* is not directly involved in the production and release of reactive oxygen species, it appears to be essential for gp91^{phox} and p22^{phox} dimerization and formation of the NADPH complex.¹²

Pathogenic variants in the *CYBB* gene result in the X-linked form of the disease. This is the most common form of CGD worldwide with most cases being hemizygous males.¹⁰ X-linked, female carriers of *CYBB* mutations are usually protected from typical CGD infections since they have two separate neutrophil populations secondary to random X chromosome lyonization – one with normal and the other with aberrant NADPH oxidase function.^{14,15} Some carriers, however, demonstrate skewed X chromosome lyonization, in which less than 20% of their phagocytes express the wild type *CYBB* gene. These patients can present with CGD-like manifestations often later in life, with carriers that have less than 10% normal phagocytes having the highest risk for severe infection.^{15–20}

Biallelic pathogenic variants in *NCF1*, the most common AR cause of the disease, tend to be associated with a better

prognosis compared to *CYBB* and *NCF2*, and account for approximately 25% of CGD cases in North America and Europe.^{3,11} Together, *CYBB* and *NCF1* variants account for 75–90% of cases of CGD in North American, European, and Japanese populations.^{3,4,6,21} In contrast, areas with higher rates of consanguinity have predominantly AR disease.^{4,5,22,23} Cohorts have shown that in populations with >50% rates of consanguinity, 62–100% cases of CGD are secondary to autosomal recessive forms of the disease.^{5,22–24}

Phenotype-Genotype Considerations

Phenotypic differences have been described between X-linked and AR disease. In X-linked disease, the mean age of symptom onset is between 4 and 14 months of age, and the mean age of diagnosis is 2.7–3.1 years.^{4,6,7,24} In comparison, AR disease often presents later, with a mean symptom onset at 30–36 months of age and mean age of diagnosis at 5.8 years–7.8 years.^{4,6} The difference in the age at diagnosis between X-linked and AR CGD persists even when controlling for gender.⁴ Patients with X-linked disease also have a more frequent and severe infections leading to a higher mortality compared to AR disease.⁴

In addition to phenotypic differences seen between X-linked and AR disease, there can be variability in presentation resulting from how the specific mutations affect gene function. Studies have shown the level of phagocyte-derived reactive oxygen intermediate (ROI) production is inversely proportional to survival and is the most important factor in predicting disease severity regardless of the gene affected.²⁵ Generally, patients with nonsense, frameshift, and splice-site variants, or deletions are more likely to have decreased residual ROI production and have worse survival than patients with missense variants. The specific gene that is defective can also have a variable effect on ROI production. Individuals with defects in *NCF4* and *CYBC1* typically have more residual superoxide production and present with more mild infections compared to those with defects in the 4 other known genetic causes of CGD.²⁶ Information regarding a patient's ROI and specific genetic variant can be beneficial in guiding clinical decision-making and providing individualized patient care. These values can often be derived directly from the sequence data.

Clinical Presentation

Infection continues to be the most common clinical feature of the disease and the leading cause of death in patients with

CGD.^{4,27} The majority of patients present with characteristic infections of the skin, lymph nodes, lung, liver, and bone before 5 years of age. Depending on the genotype and residual ROI production, however, diagnosis may be delayed well until the third decade of life.⁴

Skin, soft tissue infections, and lymphadenitis are frequently the initial presenting sign of CGD.^{7,28} In Middle Eastern countries, lymphadenitis is the most common infection overall.^{4,29} While *Staphylococcus aureus*, *Serratia marcescens*, and *Klebsiella* spp are common causes of skin and lymph node infections, occasionally no organism is identified. In these cases, careful consideration must be taken to identify more unusual and fastidious organisms (see pathogen-specific considerations below). Pulmonary infections are generally the most common type of infection in CGD patients overall, primarily presenting as pneumonia, although lung abscesses and empyemas have also been reported.²⁷ Recurrent pulmonary infections can lead to chronic complications such as fibrosis, pleural thickening, pulmonary hypertension, and bronchiectasis. Mucous pneumonia, a severe inflammatory response to inhaled fungal elements in decayed organic matter, can also be a presenting sign of CGD.^{30,31} *Aspergillus* spp. are the most common causes of pneumonia in CGD, but other organisms, including *Staphylococcus* spp. and *Burkholderia cepacia* complex, may be identified. Liver abscesses, primarily due to *S. aureus*, are an important cause of morbidity in CGD, and often present with non-specific symptoms of fever and an elevated erythrocyte sedimentation rate.²⁸ Similar to skin and lymph node infections, development of a liver abscess may lead to the diagnosis of CGD.

In addition to recurrent infections, patients with CGD have aberrant inflammation leading to the formation of granulomata and can rarely present with associated macrophage activation syndrome.^{32,33} While granuloma formation can occur in a variety of organs, it most commonly affects the gastrointestinal and genitourinary tracts.^{34,35} Granulomatous colitis is commonly seen in children with CGD and nearly half of the patients with the disease eventually develop inflammatory bowel disease (IBD).³⁴ Gastrointestinal inflammation can present with broad symptoms of fever, abdominal pain, diarrhea, rectal bleeding, and weight loss or in some cases bowel obstruction.³⁶ Within the genitourinary tract, granulomatous inflammation can lead to outlet obstruction and in rare cases eosinophilic cystitis.^{35,37} It is important to consider CGD in any patient presenting with granuloma formation as it can be an early clue to establishing a diagnosis of CGD.⁴

Patients with CGD and X-linked female carriers are both at increased risk for developing a wide range of autoimmune conditions. These manifestations do not seem to have a specific age or geographic predisposition with 5–15% of CGD patients being affected and up to two-thirds of X-linked carriers reporting symptoms in some studies.^{36,38} Commonly reported autoimmune conditions include the following: discoid and systemic lupus erythematosus, rheumatoid arthritis, immunoglobulin A (IgA) nephropathy, chorioretinitis, idiopathic thrombocytopenia (ITP), autoimmune hepatitis, Raynaud phenomenon, and stomatitis.³⁶ It is important to note that ROI production does not seem to be related to the risk of autoimmune disease.¹⁵

Pathogen Variability Within the United States

Recent comprehensive data on the pathogens causing infection in patients with CGD in the US primarily come from two single-center cohorts.^{27,39} There are also data prior to 2000 that came from a national registry of patients with CGD.⁴ It is notable that both of the more recent single-center cohorts included patients who were diagnosed either because of affected siblings or IBD but had never had a serious infection (3–11%), which was not reported in the older registry data.

The pathogen profile in patients with CGD is small, but those few pathogens cause a high percentage of infections. Most bacteria causing infection in CGD are catalase positive, as are most pathogenic bacteria in general. Catalase negative bacteria can also cause infection however, and catalase is not necessary for pathogenicity in CGD model systems.⁴⁰ Filamentous molds are the most common fungal pathogens in patients with CGD, but patients do not show an increased susceptibility to the dimorphic molds like coccidioidomycosis, histoplasmosis, or blastomycosis that are endemic in the United States.³¹

Aspergillus fumigatus is the most common fungal pathogen. Other *Aspergillus* species including *A. nidulans* and *A. tanneri* have also been isolated and have been associated with increased azole resistance.⁴¹ With the advent of fungal prophylaxis, several species of non-*Aspergillus* fungi have been found to cause serious infection in patients with CGD and also have increased azole resistance, including *Phellinus* spp., *Paecilomyces variotti*, *Paecilomyces lilacinus*, and *Neosartorya udagawae*.^{31,42} Fungal pathogens are associated with higher mortality than bacterial infections in patients with CGD. In the larger cohort study in the US,

55% of non-stem cell transplant-related deaths were caused by fungal infections.²⁷

In CGD, the most common bacterial pathogens in the United States are *S. aureus*, *Serratia marcescens*, *Burkholderia cepacia* complex, *Nocardia* spp., and *Klebsiella* spp.^{27,39} Both cohort studies revealed a higher fatality rate with *Burkholderia* than other bacterial infections. Other bacteria found in soil and brackish water that uncommonly cause disease in healthy individuals but are more common pathogens in CGD are *Granulibacter bethesdensis*, *Chromobacterium violaceum*, and *Francisella philomiragia*.^{43–45} While most of the cases of *Chromobacterium violaceum* within the US have been primarily described in the southeastern United States, this facultatively anaerobic, Gram-negative bacillus can also be found in tropical and sub-tropical soil worldwide.^{44,46} *Granulibacter bethesdensis*, a fastidious Gram-negative organism that causes indolent lymphadenitis and splenitis in CGD occurs in the US and Europe, but its natural reservoir is unknown.⁴⁵

Pathogen Variability Outside of the United States

The pathogen profile of CGD outside the United States varies greatly depending on vaccination history, antibiotic prophylaxis, country of origin, and likely other incompletely understood factors. When evaluating infectious symptoms in patients with CGD, it is important to consider their country of birth, climate, and living situation, which may predispose them to types of infections that may be atypical to your routine practice. While there does not appear to be an ethnic predisposition, it is also important to consider cultural practices as incidence rates may be higher in cultures where consanguinity is more common.^{4,5,22,23}

Similar to patients with CGD who are born in the United States, *Aspergillus* spp. and/or *Staphylococcus* spp. were the most common microorganisms cultured in cohorts from Europe, Northern India, Israel, Iran, Latin America, and Turkey.^{7,22,23,47–51} Unlike the United States., *Salmonella* spp. had higher incidences in Europe and Israel and were found to be the most frequent causes of septicemia in both of these studies.^{47,49} Higher rates of *Pseudomonas* spp. were seen in CGD patients from Northern India and Israel.^{48,49} Also, dissimilar to CGD patients within the United States was the low rate of *Burkholderia cepacia* in the European CGD cohorts.^{7,47} The biggest difference in presentation of pathogens,

however, is in *Bacillus Calmette-Guerin* (BCG)-related disease and tuberculosis, which is more frequently found outside of the United States.

Tuberculosis continues to be one of the top ten causes of death and the leading single infectious cause of death among all individuals worldwide.⁵² Incidence rates are low in the United States and new cases are more frequently seen in patients who are non-U.S.-born.⁵³ The BCG vaccine is a live attenuated strain of *M. bovis* that was first used in humans in 1921 to help combat tuberculosis in children, and it continues to be the only vaccine currently used to protect against this disease.⁵⁴ The World Health Organization recommends vaccination of infants at birth or as soon thereafter to protect against meningeal and miliary tuberculosis in endemic areas.⁵⁵ Patients with CGD and other immune deficiencies are susceptible to adverse outcomes from the vaccine, ranging from regional disease like BCG lymphadenitis to disseminated BCGosis.⁵⁶ Many European countries no longer give the BCG vaccine except to high-risk patients. Among reports from countries who still give BCG, the incidence of BCG adenitis ranged widely from 16.6% to 59.2% (Table 1).^{22,23,50,51,57,58} The BCG World Atlas is a helpful database to help determine BCG status of patients from other countries.⁵⁹

CGD patients living in areas endemic for tuberculosis have seen rates of infection that exceeds what is considered usual for those populations.^{51,60,61} In this population, pulmonary tuberculosis was the most common manifestation followed by peripheral lymph node involvement; however, a subset of patients with CGD can present with more disseminated, recurrent, and severe disease.^{51,61} In some of these endemic areas that are resource poor, the inability to distinguish BCG from tuberculosis has led to

Table 1 CGD Patients with High Rates of Reactions to BCG in Countries with a National Vaccination Policy

Country/Region	Adverse Reactions to BCG	Reference
China	59.2%	Zhou et al ⁵⁷
Iran	55.9%	Fattahi et al ²³
Latin America	30%	De Oliveira-Junior et al ⁵⁰
Mexico	58%	Blancas-Galicia et al ⁵¹
Morocco	16.6%	Baba et al ⁵⁸
Turkey	22.5%	Koker et al ²²

a possibly higher rate of mortality in CGD patients compared to what is seen in other countries.⁶²

Pathogen Specific Considerations

Many of the common CGD pathogens discussed in preceding sections (*Aspergillus* spp., *Staphylococcus aureus*, *Burkholderia cepacia*, *Serratia marcescens*, *Nocardia* spp., *Klebsiella* spp.) can present with varied clinical manifestations with substantial overlap in disease manifestations across these organisms.^{4,7,27,39} Nevertheless, certain organisms are more commonly associated with particular disease manifestations. For example, *Aspergillus* most frequently causes pulmonary disease, ranging from asymptomatic imaging findings to locally invasive disease involving the chest wall.⁶³ It is also among the top causes of osteomyelitis and brain abscesses.^{4,64} In contrast, *Staphylococcus aureus* and *Serratia marcescens* are more frequently isolated from skin abscesses, lymph nodes and liver abscesses.^{4,27} In addition to well-known pathogens, new pathogens are being identified with the advent of molecular diagnostics. For example, a syndrome of fever and lymphadenitis refractory to typical broad-spectrum antibacterials and prone to relapse was associated with *Granulibacter bethesdensis* infection.^{43,65}

While some patients with CGD develop fungemia or bacteremia as a result of their infections, the majority of these infections are localized and require invasive sampling of involved tissues (eg, bronchoalveolar lavage, lung and bone biopsies, lumbar puncture for cerebrospinal fluid, and abscess aspirates) to establish a microbiological diagnosis and guide antimicrobial therapy. The most frequent pathogens associated with CGD can be grown on aerobic bacterial media and in fungal cultures. Acid-fast smears and cultures are also important as CGD patients may have infections with both tuberculous and non-tuberculous mycobacteria. However, despite invasive testing, a microbiological diagnosis is often elusive in half or more of cases.^{7,39} This is likely multifactorial, including infection due to fastidious organisms that are difficult to culture and/or inhibition of organism growth due to antimicrobial exposures prior to obtaining diagnostic specimens. With some organisms, it can also be difficult to discern whether they are truly pathogenic or a colonizer. For example, *Actinomyces* is a common colonizer of the gastrointestinal tract and *Paecilomyces* is a common environmental saprophyte, but both have also been described as pathogenic in CGD.^{40,66} Cytology and histopathology provide important clues to the presence of fastidious or

non-viable organisms. For example, *Granulibacter bethesdensis* was initially detected by special Warthin-Starry stain and subsequently cultured using buffered charcoal yeast extract agar.⁴³

More recently, additional culture-independent techniques have been utilized to increase the rates of microbiological diagnosis. For common pathogens, targeted testing for specific microbial products can be a helpful adjunct to traditional culture. For *Aspergillus*, detection of galactomannan antigen and *Aspergillus* by PCR in blood and bronchoalveolar lavage have high sensitivity and specificity for the diagnosis of invasive aspergillosis in BAL specimens.⁶⁷

For detection of rare and novel pathogens, additional approaches are needed and becoming more readily available. One such technique involves amplification and sequencing of a conserved region of the bacterial 16S or fungal 28S ribosomal RNA. This can lead to the identification of a causative organism either from culture or directly from specimens.^{43,68} More recently, metagenomic sequencing of whole blood and bronchoalveolar lavage fluid has been used for the detection of pathogens with the potential to detect a broader range of pathogens and mixed pathogens.^{69,70} While these techniques hold promise for diagnosing challenging infections, they are also at risk of detecting colonizing and contaminating organisms and data on their practical clinical utility is still emerging.

CGD Diagnosis

It is important to have a high index of suspicion of CGD in patients with recurrent or severe infections with any of the characteristic organisms described above, as prompt diagnosis and treatment are crucial to a better prognosis. The primary diagnostic tests used in CGD functionally assess the NADPH complex in stimulated neutrophils. The nitro-blue tetrazolium (NBT) reduction test was historically used; however, it has been largely superseded by the dihydrorhodamine (DHR) assay as the gold standard for diagnosing CGD. While positive results in either of these tests are generally diagnostic, genetic sequencing confirms the diagnosis.⁷¹

DHR-123 Oxidation Assay

The DHR-123 oxidation assay is the gold-standard method to diagnose CGD. In this test, stimulated phagocytes normally oxidize DHR-123 to a fluorescent green signal, rhodamine, by the NADPH complex. A flow cytometer is then used to measure the oxidative burst activity.

Phagocytes with a defective NADPH complex are unable to effectively oxidize DHR-123 and a change in fluorescence is not seen. The DHR-123 assay is widely commercially available, and it does not rely on operator subjective interpretation. Based on quantitative results, it can distinguish between X-linked and recessive forms of CGD, as well as carriers of pathogenic variants in gp91phox. This assay is highly sensitive and combined with the capacity to assess residual superoxide production, it can predict the severity and prognosis in patients with CGD.²⁵ There are limitations to its use, however as this test can be abnormal in ehrlichiosis infection, myeloperoxidase deficiency, and the syndrome of synovitis, acne, pustulosis, hyperostosis, and osteitis (SAPHO).^{72,73}

NBT Reduction Test

This is one of the oldest qualitative assays used to test neutrophil function. In this test, a drop of blood is placed on a slide and incubated with an activating agent and NBT. Normal granulocytes reduce the yellow NBT dye to blue formazan, a dark blue precipitate. Although it does not require sophisticated equipment, its use has been limited. While it can be used to identify gp91phox carriers, the interpretation of this test is subjective leading to increase false-negative results.⁷⁴ It is also unable to differentiate between X-linked and autosomal recessive CGD.⁷⁵

Genetic Sequencing

An abnormal neutrophil function test should be followed by confirmatory genetic sequencing. Knowledge about how a specific variant effects protein function as well as residual superoxide production have been predictive of disease severity and mortality risk.²⁵ Both single-gene and multi-gene panels are available to sequence DNA from peripheral blood leukocytes. In patients suspected to have AR disease, it is also important to note that NCF1 is flanked by pseudogenes, which may complicate sequencing and interpretation.⁷⁶

Management of Infectious and Granulomatous Complications

Given the risk for recurrent and/or severe infections in patients with CGD and the characteristic organisms that are known to commonly cause disease, the mainstay of management of this disease is targeted prophylaxis. Optimal prophylactic regimens should include a combination of antibacterial agents, antifungal agents, and immunomodulation with interferon-gamma (IFN- γ) therapy. In addition, the prompt

evaluation and treatment of acute infections along with adjunctive anti-inflammatory medications are important to minimize infectious and granulomatous complications.

Daily oral trimethoprim-sulfamethoxazole (5mg/kg of the trimethoprim component with a maximum daily dose of 1 double-strength tablet) is recommended for antibacterial prophylaxis.⁷⁷ Other antibacterial agents used include trimethoprim alone, 2nd or 3rd generation cephalosporins, and quinolones.⁷⁸ Antifungal prophylaxis with itraconazole has been shown to significantly reduce the incidence of invasive fungal disease in patients with CGD.^{4,27,47} First-line antifungal prophylaxis is itraconazole (100mg for <13 years old or <50kg and 200mg daily for >13 years old or >50kg), but voriconazole and posaconazole are being increasingly used.^{77,79} Transaminitis has been associated with the azoles, so liver function tests should be periodically monitored. Due to variability in azole absorption, some clinicians assess drug levels, especially in those CGD patients with gastrointestinal granulomas. Voriconazole can lead to significant phototoxicity and, over long periods of time, there is a concern for fluoride toxicity.

The use of interferon gamma in patients with CGD allows for a more targeted therapeutic immunomodulatory approach. Interferon gamma has been shown to stimulate superoxide release and improve phagocyte-mediated bacterial killing in vitro.⁸⁰⁻⁸² Studies have also shown a significant reduction in infection frequency, duration and hospitalization rates in patients with CGD.^{83,84} In the United States, recombinant interferon gamma is FDA approved and commonly used for infection prophylaxis in CGD. In Europe, interferon gamma therapy is often reserved for patients with persistent, recurrent infections despite appropriate antibacterial and antifungal prophylaxis.⁶ It may also be beneficial in CGD populations living in tuberculosis endemic areas. Interferon gamma is started at a dose of 50 $\mu\text{g}/\text{m}^2$ (or 1.5 $\mu\text{g}/\text{kg}$ if BSA is <0.5 m^2) and administered subcutaneously three times weekly. Common side effects include fevers, malaise, chills, fatigue and location injection site erythema or tenderness.

If and when there is a concern for acute infection, patients with CGD warrant a prompt and thorough physical and laboratory evaluation as there can be great variability in clinical presentation. Some patients, especially the younger children, with CGD may not present with classic infectious signs or symptoms of fever, fatigue, malaise, leukocytosis, or neutrophilia. In patients with CGD, an erythrocyte sedimentation rate and C-reactive protein can be very helpful in both assessing for possible infection and

monitoring for response to treatment. There should be a low threshold for imaging the chest and/or abdomen (eg, liver) given the high risk for indolent bacterial and fungal infections. Early diagnosis of infections not only may keep the patient from being hospitalized, but also will reduce the risk for further organ damage and inflammation (eg, microgranuloma formation) at sites of infection.

Granulomatous complications in CGD can be quite profound and at times debilitating to patients. When treating acute infections in patients with CGD, an anti-inflammatory agent may be necessary in addition to antimicrobial therapy since granuloma formation can be seen at infection sites. Systemic corticosteroids have been used as adjunctive therapy to antibiotics in the treatment of persistent infections, such as staphylococcal liver abscesses. Staphylococcal liver abscesses can be very difficult to treat in CGD and may even require surgical intervention. Steroids are typically dosed at 1mg/kg/day for 2–3 weeks, followed by a taper over several months (on average 5 months).^{85,86} Corticosteroids have also been used to treat obstructive granulomatous lesions in CGD, such as bladder granulomas causing bladder outlet obstruction or gastrointestinal lesions causing obstruction.⁸⁷ The use of systemic immunosuppressive medications, such as ustekinumab or vedolizumab, continues to be explored in patients with CGD colitis and has had varying effects to date.^{88–90} TNF-inhibitors should be avoided in patients with CGD, as they have been associated with serious bacterial and fungal infections and increased mortality.⁹¹

Long-Term Management of CGD

Allogeneic HSCT is the only widely available curative treatment for CGD with the potential for resolution of both infectious and inflammatory complications. Outcomes have improved significantly over the last few decades with reported overall survival rates now consistently >90% in pediatric patients less than 14 years of age regardless of donor source.^{92–99} Furthermore, pediatric patients who undergo HSCT have fewer infections, improved growth parameters, and higher quality of life measures compared to those treated conventionally.^{100,101}

Patients with intractable infection or autoinflammation at the time of HSCT and adolescents or young adults have traditionally been difficult to transplant with historically high rates of graft-versus-host disease (GVHD) and transplant-related mortality.^{92,93,102,103} However, in 2014, Güngör et al published a large, multi-center HSCT trial that included 13 adult patients with impressive overall and event-free survival rates,¹⁰⁴ and

several institutions have subsequently reported excellent transplant outcomes in adolescent and young adult patients, including those with severe infection and significant autoinflammation at time of transplantation.^{105–107} Another multi-center study also recently demonstrated the presence of CGD-related colitis does not appear to negatively affect survival or significantly increase the risk of GVHD.¹⁰⁸ The role of autoinflammation and organ dysfunction on transplant outcomes remains unclear; however, these recent publications indicate that HSCT for definitive cure may be considered for high-risk patients with reasonable outcomes.

Gene therapy is an appealing alternative to HSCT and a number of small trials have been conducted to treat gp91phox deficiency using γ -retroviral vectors and reduced intensity conditioning.^{109–112} All trials demonstrated initial engraftment of transduced neutrophils at 10% to 30% of circulating neutrophils, and gene therapy resulted in full or partial resolution of infection in most patients. However, cell engraftment progressively decreased with time, and several patients developed myelodysplastic syndrome (MDS) due to insertional activation of proto-oncogenes.¹¹² In response to the high incidence of MDS seen with γ -retroviral vectors, gene therapy trials are currently underway using self-inactivating (SIN) lentiviral vectors. Encouragingly, studies using similar vectors for the treatment of Severe Combined Immunodeficiency and Wiskott Aldrich have showed good success.¹¹³ Long-term outcomes with gene therapy are unknown, and as with HSCT, it is unclear what level of oxidase-positive neutrophils is necessary for resolution of autoinflammation and sustained correction of the CGD phenotype.

Currently, there are no standard HSCT guidelines and no consensus surrounding transplant protocols for CGD patients. Large, multi-center studies are currently underway to further investigate long-term transplant outcomes, identify optimal conditioning regimens, and determine the level of donor chimerism necessary for resolution of autoinflammation and sustained protection against new-onset inflammatory or autoimmune disease manifestations. Biomarkers predictive of disease course and transplant outcomes are also needed to help guide treatment decisions. With the new SIN lentiviral vectors and optimization of conditioning regimens, gene therapy may also become a viable alternative to HSCT in the future.

Conclusion

CGD is a rare but serious primary immunodeficiency with varying prevalence and rates of X-linked and AR disease worldwide. Functional defects in the phagocyte NADPH oxidase complex predispose patients to a relatively narrow spectrum of

bacterial and fungal infections that are sometimes fastidious and often difficult to identify. When evaluating and treating patients with CGD, it is important to consider their native country of birth, climate, and living situation, which may predispose them to types of infections that are atypical to your routine practice. In addition to recurrent and often severe infections, patients with CGD and X-linked female carriers are also susceptible to developing many non-infectious complications including tissue granuloma formation and autoimmunity. The DHR-123 oxidation assay is the gold standard for making the diagnosis and it along with genetic testing can help predict the severity and prognosis in patients with CGD. Disease management focuses on prophylaxis with antibacterial, antifungal, and immunomodulatory medications, prompt identification and treatment of acute infections, and prevention of secondary granulomatous complications. While HSCT is the only widely available curative treatment for patients with CGD, recent advances in gene therapy may provide a safer, more direct alternative.

Abbreviations

CGD, chronic granulomatous disease; NADPH, nicotinamide adenine dinucleotide phosphate; AR, autosomal recessive; BCG, Bacillus Calmette-Guerin.

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