Case Report: Pseudomonas can take a toll on a patient

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Abstract

Pseudomonas aeruginosa (P. aeruginosa) is an aerobic Gram-negative bacterium that is implicated in the development of severe systemic infections among pediatric patients. It is identified in hospitalized chronically ill pediatric patients in association with genitourinary, respiratory tract, and skin or soft tissue infections as well as severe and life-threatening infection including sepsis. A variety of immunologic mechanisms play a vital role in the host defense mechanisms against invasive infections with P. aeruginosa. Rarely, specific inborn errors of immune function are implicated in deficiencies that predispose to invasive infections with P. aeruginosa. Innate immune function including germ-line encoded pattern recognition receptors such as toll-like receptors (TLRs) and their downstream signaling is vital in the host defense against P. aeruginosa through the generation of antimicrobial peptides, cytokines/chemokines, and shaping of adaptive immune responses. Herein, we describe a previously healthy two-year-old female with an invasive skin, soft tissue, and central nervous system infection secondary to P. aeruginosa. The invasive nature of this infection prompted a careful evaluation for an inborn error of immunity. Decreased cytokine response to agonists of TLRs was documented. Targeted sequencing of interleukin-1 receptor-associated kinase (IRAK)-4 documented a homozygous deletion of exons 8-13 consistent with IRAK-4 deficiency. This report provides a vital educative message in the existing scientific literature by underscoring the importance of considering inborn errors of immunity in all patients with severe P. aeruginosa infections. Functional assessments of immune function often in combination with sequencing can accurately assign a diagnosis in a timely fashion allowing for definitive treatment and the use of necessary supportive care.
Introduction

Pseudomonas aeruginosa (P. aeruginosa) is an opportunistic aerobic Gram-negative bacterial pathogen associated with a variety of genitourinary (UTI), pulmonary as well as skin and soft tissue infections (SSTI) in hospitalized pediatric patients often in association with significant morbidity. Rarely, P. aeruginosa can be associated with severe and life-threatening infections among children without previously recognized associated risk factors. In these cases, it is vital to consider the possibility of an underlying inborn error of immunity. Invasive P. aeruginosa infections have been described in the setting of inborn errors of immunity including antibody deficiencies (agammaglobulinemia) Bruton (Bruton agammaglobulinemia), combined immunodeficiency disorders (severe combined immunodeficiency, ataxia telangiectasia), defects of phagocytes (chronic granulomatous disease, leukocyte adhesion deficiency), defects in actin-polymerization (Wiskott-Aldrich syndrome, MKL1-deficiency), chronic neutropenia and innate immunity including defects in canonical NFkappaB-signaling (e.g., NEMO/NFKBIA) as well as those that impair the downstream signaling of toll-like receptors (TLRs), such as defects in interleukin-1 receptor-associated kinase (IRAK)-4 and myeloid differentiation factor 88 (MyD88). We describe the presence of an invasive soft tissue and central nervous system infection with P. aeruginosa in a previously healthy two-year-old female which prompted evaluation for an inborn error of immunity.

Case report

A previously healthy two-year-old Hispanic female was evaluated for left hip and knee pain associated with a fever and a refusal to ambulate. She had been given cephalaxin for presumed insect bites on her back and legs. Her past medical history was non-contributory. She was fully immunized. Her family history was negative for consanguinity and inborn errors of immunity. She was afebrile and her physical examination demonstrated a 3 cm circular erythematous lesion on her mid left back which was non-indurated. Her left knee was minimally swollen, warm to touch, and non-erythematous. Her hips and knees remained in flexion with demonstrated resistance to knee extension. A nodular 1.5 cm mass in her popliteal fossa was also documented with tenderness upon palpation and she was unable to walk. Her laboratory investigation demonstrated a white blood cell count of 17.2 × 10³/µL (normal range: 3.9-13.7), Hgb 10.4 g/dL (normal range:10.2-15.4), platelets 319 × 10⁹ (normal range: 150-450), 64% neutrophils (normal range: 25-72), 26% lymphocytes (normal range: 24-71), 9% monocytes (normal range: 0-14), 0.6% eosinophils (normal range: 2-10), 0.3% basophils (normal range: 0-2), erythrocyte sedimentation rate (ESR) 126 mm/hr (normal range: 0-20), and C-reactive protein (CRP) 68.8 mg/L (normal range: 0-10). A blood culture was negative. Magnetic resonance (MR) imaging of her lower extremity demonstrated myositis and fasciitis involving the soft tissues of the distal left thigh, but no abscess (Figure 1A–C). Likely pathogens included methicillin sensitive Staphylococcus aureus, methicillin resistant Staphylococcus aureus, and group A Streptococcus.

Empiric antibiotic therapy began with vancomycin (60 mg/kg/day intravenously divided every 6 hours for 3 days) and ceftriaxone (75 mg/kg/day intravenously every 24 hours for 24 hours for 3 days) which was then transitioned to cefazolin (100 mg/kg/day intravenously divided every 8 hours for 7 days), clindamycin (30 mg/kg/day intravenously divided every 8 hours for 7 days), amoxicillin-clavulanate (50 mg/kg/day orally divided every 12 hours for 4 days), and then to linezolid (30 mg/kg/day intravenously divided every 8 hours for 6 days) due to a lack of clinical and laboratory improvement. A lesion on her back was biopsied and demonstrated acute suppurative panniculitis and suppurative necrosis (Figure 2). Cultures were obtained
with growth of *P. aeruginosa* and a 3-week course of cefepime (150 mg/kg/day intravenously divided every 8 hours) for SSTI was completed. Clinical and laboratory improvement occurred as demonstrated by an ESR 57 mm/hr (normal range: 0–20) and CRP <5 mg/L (normal range: 0–10). The patient was discharged while awaiting results of a workup for an inborn error of immunity.

She then returned approximately 3 weeks later with malaise, an inability to stand upright, irritability, and pain on palpation of her back as well as a refusal to ambulate. On re-admission she was afebrile, and her laboratory investigation demonstrated an ESR 89 mm/hr (normal range: 0–20), and CRP <5 mg/L (normal range: 0–10). MR imaging of her spine demonstrated enhancement of T9-T11 with an epidural abscess (Figure 3A–B). A culture was obtained by computed tomography (CT)-guided needle aspiration and empiric therapy with meropenem (120 mg/kg/day intravenously divided every 8 hours for 2 days) was begun. Growth of *P. aeruginosa* was documented and a 6-week course of cefepime (150 mg/kg/day intravenously divided every 8 hours) was completed. Clinical improvement occurred; however, her laboratory investigation demonstrated an ESR 80 mm/hr (normal range: 0–20) and CRP <5 mg/L (normal range: 0–10) at the completion of therapy.

Secondary to the invasive nature of her *P. aeruginosa* infection, evaluation for an inborn error of immunity was completed. A normal neutrophil oxidative burst was noted. Additional laboratory assessments included a serum IgG 1331 mg/dL (normal range: 407-1009), IgA 85 mg/dL (normal range: 22-220), an elevated IgM 362 mg/dL (normal range: 43-163), and a IgE 3.8 kU/L (normal range: <97). Antibody responses documented a non-protective *Haemophilus influenza* type b antibody titer (0.30 mcg/L), a tetanus antibody titer which was protective (0.21 IU/mL), and pneumococcal titers which were protective (> 1.3 mcg/L) for nearly all serotypes covered by Prevnar 13. Lymphocyte immunophenotyping demonstrated a CD3+ count of 1340/ μL (normal range: 1484-5327), CD4+ count of 633 / μL (normal range: 733-3181), CD8+ count of 628/ μL (normal range: 370-2555), CD19+ count of 493/ μL (normal range: 370-2306), CD16/56+ count of 109/ μL (normal range: 43-526). Decreased cytokine response to TLR agonists were documented by a commercial lab (ARUP Laboratories) using peripheral blood mononuclear cells (Figure 4). LPS-induced CD62L shedding in neutrophils is an additional potential screening test for TLR defects, but was not completed in this case7. Targeted sequencing of IRAK-4 and MyD88 was performed. A homozygous deletion of exons 8-13 was documented in IRAK-4 consistent with a diagnosis of IRAK-4 deficiency. There were no siblings; however, it is important to underscore the importance of screening other family members for the same pathogenic variants.

Following the diagnosis of IRAK-4 deficiency she was start on prophylaxis with intravenous immunoglobulin (0.5 g/kg/dose intravenously every 4 weeks) as well as amoxicillin (250 mg orally each day). From her diagnosis at 2 years of age until 4 years of age she continued to experience infrequent infectious complications including a urinary tract infection (*Escherichia coli*), left knee swelling in association with a abscess (methicillin-susceptible *Staphylococcus aureus*), and a single admission for fever, cough, and post-tussive emesis. She is now 6 years of age and doing well without any recent infectious complications.

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**Figure 2.** Skin punch biopsy demonstrates acute suppurative panniculitis with suppurative necrosis (20×, Hematoxylin and Eosin).

**Figure 3.** Axial (A) and sagittal (B) fat-saturated volumetric interpolated breath-hold examination (VIBE) postcontrast magnetic resonance imagery (MRI) images of the thoracic spine demonstrate multilevel thoracic osteomyelitis and discitis with associated epidural and paravertebral abscesses.
She remains compliant with her prophylaxis therapy with intravenous immunoglobulin therapy and amoxicillin which she has tolerated without complications.

**Discussion**

Detection of lipopolysaccharide and flagellin by TLRs results in the elaboration of pro-inflammatory cytokines. TLRs possess an intracellular domain known as the Toll–IL-1R domain (TIR). Upon activation of TLRs, the recruitment of TIR-containing cytosolic adaptors such as MyD88 occurs. The adaptor MyD88 then recruits cytosolic kinases, including the IRAK complex. The IRAK complex includes two kinases including IRAK-4 and two non-catalytic subunits. This results in the activation of downstream effectors including nuclear factor κB (NF-κB) and mitogen-activated protein kinases which support the synthesis of pro-inflammatory cytokines and chemokines, such as IL-1β, -6, -8, and -12 and tumor necrosis factor alpha.

IRAK-4 deficiency is an autosomal recessive disorder which requires that affected patients have homozygous or compound heterozygous mutations in the IRAK-4 gene. IRAK-4 deficient
patients typically have normal basic immunological evaluations. Importantly, inflammatory responses are markedly blunted as demonstrated by the severe and life-threatening invasive bacterial infection with \textit{P. aeruginosa} in our patient accompanied by an absence of CRP elevation. In these patients, CRP concentrations can be strikingly misleading as IRAK-4 deficient patients demonstrate impairment in the ability to increase CRP concentrations and to mount fever responses.

Among IRAK-4 deficient patients with invasive bacterial infection, \textit{S. pneumoniae} is the most frequently (~50% of episodes) implicated organism. \textit{S. aureus} and \textit{P. aeruginosa} are less frequently (~20% of episodes) implicated organisms. Patients with IRAK-4 deficiency may also experience a variety of minor non-invasive bacterial infections such as upper respiratory tract infections (otitis, sinusitis, pharyngitis) as well as SSTI (furunculosis, folliculitis, cellulitis). Once again, the most frequently implicated organisms in these non-invasive bacterial infections among IRAK-4 deficient patients are \textit{S. pneumoniae}, \textit{S. aureus}, and \textit{P. aeruginosa}.

Careful institution of aggressive supportive care measures is necessary for IRAK-4 deficient patients. Vaccines should include conjugated and nonconjugated vaccines for \textit{S. pneumoniae} and \textit{N. meningitidis}. \textit{Haemophilus influenzae} type b conjugated vaccine should also be provided. Lifelong antibiotic prophylaxis with cotrimoxazole in combination with penicillin should be administered. Prophylaxis with intravenous or subcutaneous IgG should also be provided as a significant proportion of IRAK-4 deficient patients have impaired responses to glycanics. Empiric parenteral antibiotic therapy against \textit{S. pneumoniae}, \textit{S. aureus}, and \textit{P. aeruginosa} is critical whenever an infection is suspected or if the patient develops a fever. Inflammatory markers such as CRP should be considered unreliable. Importantly, patients may die from invasive bacterial infection despite prophylaxis even in the absence of fever or laboratory evidence of inflammation. The long-term prognosis of IRAK-4 deficient patients is positive as the risk of invasive infections tend to improve with age.

Although the occurrence of invasive \textit{P. aeruginosa} infections in IRAK-4 deficiency is not novel, there is a vital and important educative message that bears repeating. Astute clinical judgment is necessary in the evaluation of patients with a potential inborn error of immunity. Clinical findings such as severe infections with \textit{P. aeruginosa} should support the consideration of inborn errors of immunity including IRAK-4 deficiency.

**Data availability**

All data underlying the results are available as part of the article and no additional source data are required.

**Consent**

Written informed consent for publication of their clinical details and/or clinical images was obtained from the parent of the patient.

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Very well-written case report of a child with IRAK4 deficiency. I have no major comments, few minor comments include:
- Briefly discuss the role of CD62L shedding assay as another screening test.¹
- Explain briefly why there is a need for Ig replacement in a disorder that is mainly myeloid lineage being affected- it might be of interest to the readers
- A line on long term prognosis might be helpful- the risk of invasive infection tend to improve with age

References
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Is the background of the case's history and progression described in sufficient detail?
Yes

Are enough details provided of any physical examination and diagnostic tests, treatment given and outcomes?
Yes

Is sufficient discussion included of the importance of the findings and their relevance to future understanding of disease processes, diagnosis or treatment?
Yes
Is the case presented with sufficient detail to be useful for other practitioners?
Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Immunobiology of Immune dysregulation disorders; HSCT and gene therapy for immune defects

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 18 Aug 2021

David K. Buchbinder, CHOC Children's Hospital, Orange, USA

Reviewer #2:

Comment #1:
Briefly discuss the role of CD62L shedding assay as another screening test.

Response #1:
Thank you for this excellent and important suggestion. Additional discussion regarding the role of the CD62L shedding assay as a potential screening test has been added. The suggested reference has also been added.

von Bernuth H, Ku CL, Rodriguez-Gallego C, Zhang S, et al.: A fast procedure for the detection of defects in Toll-like receptor signaling. Pediatrics. 2006; 118 (6): 2498-503.

Comment #2:

Explain briefly why there is a need for Ig replacement in a disorder that is mainly myeloid lineage being affected- it might be of interest to the readers

Response #2:
Thank you for this suggestion. Additional discussion focusing on explaining why immunoglobulin prophylaxis has a role in IRAK4 deficiency has been added.

Comment #3:

A line on long term prognosis might be helpful- the risk of invasive infection tend to improve with age

Response #3:
We apologize for this omission and agree that it is important to comment on the long-term
I had a pleasure reading this case report. The authors nicely describe a rare case of IRAK-4 deficiency in a female patient with invasive pseudomonal infection.

The case report highlights several important issues. First, and most important, unusual and/or severe presentation should trigger further evaluation for possible underlying inborn errors of immunity (IEI). In this case, invasive infection and failure to respond to antibiotics raised this concern.

Next, the case report highlights additional important laboratory findings, such as normal CRP levels in the presence of severe infection. In that regard, this case report shows that 'normal' CRP should not always be considered a reassuring sign, but rather the opposite, and in some cases could suggest a failure to mount a required inflammatory response.

Finally, the case report describes a very elegant stepwise work-up approach, starting with identifying 'clinical clues' (such as normal CRP), followed by ordering the correct diagnostic assay, and finally performing targeted sequencing of a single candidate gene (sadly, not that common anymore).

There are several minor points I would like to suggest:

1. Decreased cytokine response to TLR agonists: Was that performed by a commercial lab? If so, I think I would mention it so readers would know that it can be ordered. Also, which cells were used (fibroblasts/PBMCs)?

2. Were the authors able to test for LPS-induced CD62L shedding in neutrophils?

3. I don't know if the patient has siblings, but perhaps the Discussion should include a recommendation to screen other family members for the same pathogenic variants.

Is the background of the case's history and progression described in sufficient detail?
Yes

Are enough details provided of any physical examination and diagnostic tests, treatment given and outcomes?
Yes

Is sufficient discussion included of the importance of the findings and their relevance to future understanding of disease processes, diagnosis or treatment?
Yes

Is the case presented with sufficient detail to be useful for other practitioners?
Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Inborn Errors of Immunity/Primary Immunodeficiency Disorders

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 18 Aug 2021

David K. Buchbinder, CHOC Children's Hospital, Orange, USA

Reviewer #1:

Comment #1:
Decreased cytokine response to TLR agonists: Was that performed by a commercial lab? If so, I think I would mention it so readers would know that it can be ordered. Also, which cells were used (fibroblasts/PBMCs)?

Response #1:
Thank you for this comment. The cytokine response to TLR agonists was performed by a commercial lab. We agree that this is important to point out for readers. Moreover, peripheral blood mononuclear cells were utilized for testing. This was also pointed out to help other readers/clinicians that may be facing similar clinical situations.

Comment #2:
Were the authors able to test for LPS-induced CD62L shedding in neutrophils?

Response #2:
We did not test for LPS-induced CD62L shedding in neutrophils. As suggested by Reviewer #2, additional discussion regarding the role of the CD62L shedding assay as a potential
screening test has been added.

Comment #3:

I don't know if the patient has siblings, but perhaps the Discussion should include a recommendation to screen other family members for the same pathogenic variants.

Response #3:

The patient does not have any siblings, but we agree that this is an important point to make from a clinical perspective. Additional discussion regarding the importance of screening other family members for the same pathogenic variants has been added.

**Competing Interests:** I have no competing interests.