Case Report,

Cat-Scratch Disease Pneumonia: An Atypical Presentation

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Abstract:

Purpose: Cat-scratch disease is caused by Bartonella henselae infection and it commonly presents in children with a mild cutaneous lesion and adjacent lymphadenitis. It has rarely been observed that it can present as a disseminated infection, such as encephalitis or pneumonia. We present a case report of a 35-year old, immunocompetent, patient who suffered pneumonia and encephalitis, resulting from Bartonella henselae infection. We also present a review of the literature published regarding this rare complication.

Methods: We’ve searched the literature in the databases of Pubmed and Google Scholar to find articles that report pulmonary presentations of cat-scratch disease.

Results: We’ve found only 16 other reported similar cases with pulmonary involvement in cat-scratch disease.

Conclusions: Pulmonary presentation in cat-scratch disease is an uncommon presentation of a common disease, and it should be ruled out, even in cases it seems unlikely.

Key words: Bartonella, Cat, Pneumonia, Encephalitis, Lymphadenitis

Abbreviations: Computed Tomography (CT), Cerebrospinal Fluid (CSF), Indirect Fluorescence Assay (IFA), Enzyme Immunosorbent Assay (EIA), Polymerase Chain Reaction (PCR)

Introduction:

Cat-scratch disease was first described by Debré [1] in 1954 and has since been examined in various reviews, as methods of diagnosis have evolved. The most common causative agent is Bartonella henselae (formerly Rochalimaea henselae), although other similar organisms have been reported to be causative, less common, such as Afipia felis [2] and Bartonella claridgeiae [3]. It is transmitted to human via contact of broken skin areas or mucosal surfaces with cat saliva or after scratch or bite from an infected cat, cat fleas, while there are reports of cases with exposure to dogs [4]. The most common clinical feature is a cutaneous lesion at the site of inoculation, often manifested with a vesicular, erythematosus, and papular phase, accompanied by regional tender lymphadenopathy and fever. Atypical manifestations have been described, in up to 33% of patients older than 60 years, compared to 14% in younger patients [5] and include encephalitis and endocarditis, resulting from disseminated infection. Other atypical manifestations are Parinaud's ocuoglandular syndrome, neuroretinitis, erythema nodosum, and granulomatous hepato-splenitis, myalgias, arthralgias and osteomyelitis. Bacillary angiomatosis and peliosis hepatis are complications only found in immunocompromised patients. Pertaining respiratory complications, only rare cases have been reported, including pneumonia, pleural effusion or pleural thickening, with an unknown prevalence. In the past, skin testing was used for diagnosis, but it has now been replaced by serology tests, including Indirect Fluorescence Assay (IFA) and Enzyme Immunosorbent Assay (EIA). Polymerase Chain Reaction (PCR) amplification testing for Bartonella DNA in the affected tissues can confirm
the diagnosis, while PCR in the serum has low sensitivity [6]. Identification by culture is very difficult because it requires specific laboratory conditions. The lymph node histopathology usually reveals lymphoid hyperplasia, microabscess formation and stellate granulomas with acellular and necrotic centers and peripheral accumulation of lymphocytes and histiocytes.

Florin et al [4] revised the criteria for diagnosis in 2008, requiring three of the following four:

1. History of contact with a cat with or without primary skin inoculation
2. Exclusion of other causes of lymphadenopathy
3. Positive serology
4. Lymph node biopsy showing microabscess or granulomas and positive Warthin-Starry silver stain.

**Table 1: Laboratory Workup**

| Parameter | Value                  |
|-----------|------------------------|
| WBC       | 11640 x10^9/μL         |
| NE        | 8260 x10^9/μL          |
| LY        | 2390 x10^9/μL          |
| MONO      | 910 x10^9/μL           |
| BASO      | 40 x10^9/μL            |
| EO        | 40 x10^9/μL            |
| HCT       | 41%                    |
| HB        | 13.7 g/dl              |
| PLT       | 462000 x10^9/μL        |
| GIU       | 148 mg/dl              |
| URE       | 31 mg/dl               |
| CRE       | 1 mg/dl                |
| NA        | 135 mg/dl              |
| K         | 4.6 mg/dl              |
| TP        | 7.3 mg/dl              |
| AST       | 21 mg/dl               |
| ALT       | 25 mg/dl               |
| ALP       | 79 mg/dl               |
| GGT       | 140 mg/dl              |
| TBIL      | 0.42 mg/dl             |
| LDH       | 305 mg/dl              |
| AMY       | 45 mg/dl               |
| CRP       | 17.4 mg/dl (normal values 0-0.4 mg/dl) |
| CSF cells | 19 (Poly:15, Lympho:4)  |
| CSF Glu   | 68 mg/dl               |
| CSF TP    | 36.1 mg/dl             |
| Pleural fluid cells | 1100 (Poly: 12%, Lympho: 80%, Mono: 8%) |
| Pleural fluid TP | 4.1 mg/dl             |
| Pleural fluid LDH | 171 mg/dl             |

**Abbreviations:** WBC: White Blood Cell Count, NE: Neutrophil Count, LY: Lymphocyte Count, MONO: Monocyte Count, BASO: Basophile Count, EO: Eosinophil Count, HCT: Hematocrit, HB: Hemoglobin, PLT: Platelet Count, GLU: Glucose, URE: Urea, CRE: Creatinine, NA: Sodium, K: Potassium, TP: Total Protein, AST: Aspartic Aminotransferase, ALT: Alanine Aminotransferase, ALP: Alkaline Phosphatase, GGT: Gamma-glutamyltransferase, TBIL: Total Bilirubin, LDH: Lactic Dehydrogenase, AMY: Amylase, CRP: C-Reactive Protein, CSF: Cerebrospinal fluid

**Case report:**
A 35-year-old male patient came to the emergency department with loss of consciousness and first onset generalized epileptic seizures. He didn’t have any known health problems and he is considered immunocompetent. His relatives reported that for 20 days he had been suffering from high grade fever and night sweats. He also complained about painful swelling in the axilla and elbow of the left arm. He had seen a doctor for that who prescribed common antibiotics and had visited the emergency department once more, 5 days ago, but then decided to leave against medical advice. Upon his arrival at the emergency department he developed generalized tonic seizures, which were followed by interchanges of consciousness between coma, lethargy and a confusional, disoriented state. He also had tachypnea and low arterial oxygen saturation. His brain CT scan was normal, and a lumbar puncture was conducted. The results from the cerebrospinal fluid (CSF) indicated towards encephalitis and they are shown on table 1. A CT scan of the lungs revealed bilateral pleural effusion and a bilateral nodular pattern of opacities in the upper pulmonary lobes (Figure 1), while the CT scan of the abdomen only revealed a mild enlargement of the spleen. He was administered anti-seizure treatment with Levetiracetam, antimicrobial coverage for common central nervous system pathogens and required low flow oxygen administration. Pleural fluid was sent for analysis after thoracentesis, which revealed...
lymphocytic exudate. An ultrasound of the left elbow revealed enlargement of epitrochial lymph nodes, and a biopsy revealed abscess formation and inflammation without specific characteristics. The abscess in the elbow and axillary areas were both drained subcutaneously and were sent for microbiologic analysis.

During the second day of inpatient care, he showed signs of recovery of consciousness, with alertness and orientation, with full recovery of consciousness on the fourth day. He developed worsening crackles at auscultation of the lungs and required oxygen administration for at least ten more days. A second CT of the chest revealed improvement of opacities in the lung parenchyma with persistent pleural effusions. From the laboratory workup, increased inflammatory markers were observed, with increased neutrophil count and C-reactive protein. A thorough laboratory workup for viral and bacterial pathogens’ antibodies was negative, and the cultures and direct stains of blood, CSF, pleural fluid and abscess drainage fluid were negative.

The patient remembered that he had petted a stray cat and kittens about a month before the onset of the symptoms and that the cat scratched him on his left hand. The IgG antibody tests for Bartonella henselae came out positive at a titer of 1:512 which was doubled six days later, and at that time the IgM antibodies were found positive as-well. His antibiotic regimen was changed to Doxycyclin and Rifampicin and the fever and the rest of clinical signs were attenuated gradually. The patient was discharged thirty days later feeling well.

The purpose of this article is to review the pulmonary manifestations of cat-scratch disease in the literature. Pulmonary manifestations can include pneumonia, appearing on CT scans as opacities, pulmonary nodules and reticulonodular or interstitial pulmonary infiltrates. Pleural effusions and pleural thickening are often described, and when available, Bartonella DNA is detected with PCR after thoracentesis. 16 patients are described in this review with pulmonary manifestations of Bartonella henselae disease, 3 females and 13 males. 7 of them are children aged 3-12 years old and the rest are adults. 5 of them were immunocompromised and 1 of them suffered from type 2 diabetes mellitus, which can be regarded as a risk factor for infections. The rest did not have any significant medical history and were considered immunocompetent. Patient characteristics and clinical signs and symptoms are presented in table 2. Only 4 patients had developed both central nervous system and pulmonary involvement, like the patient in this case report. Two cases were complicated with septic shock and one of them died; surprisingly, only suffering from type 2 diabetes as an underlying disease.

Discussion:
We present an uncommon manifestation of a rather common disease. To our knowledge, only 16 similar cases of pulmonary involvement in cat-scratch disease were reported in the past. Positive antibody titers, along with the exclusion of other causes via a thorough microbiological workout, placed the diagnosis in our patient. Unfortunately, neither PCR testing nor Warthly-Stain silver stain were available in our hospital. As we were able to witness it in our patient, disseminated infection of Bartonella henselae, follows a rather slow course of response to antibiotic treatment. The axillary and elbow abscesses required subsequent drainage and

**Literature review on Cat-scratch disease associated pneumonia:**

**Figure 1:** Chest CT scan of a patient with pneumonia of Bartonella Henselae. Nodular pulmonary infiltrates and pleural effusion can be seen.
he was treated with oral antibiotics for a total of 6 months to completely recover. This case was presented to us amidst the 2020 pandemic of SARS-COVID-19 [24]. Although most signs in our patient didn’t fit the clinical profile of COVID-19 disease, the persistent respiratory disease, along with high grade fever, placed a high suspicion for COVID-19. Thus, the patient was transferred, at first, to the COVID-19 treatment unit, until two PCR tests from nasopharyngeal swab came out negative. We’d like to report this as a take-home message, and as a recommendation towards clinicians, during this pandemic; even though COVID-19 might be a prevalent disease that could fit a certain clinical profile, clinicians should be alert for other diseases as-well, whose diagnosis and treatment must not be delayed.

Declarations:
- **Funding:** This study received no funding.
- **Conflict of Interest:** Authors Fotis Konstantinou, Ioanna Skrapari, Asimoula Megkou and Evangelos Kokkinakis declare that they have no conflict of interest.

- **Availability of data and material:** All data generated or analyzed during this study are included in this published article.

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Table 2: A review of the literature on pulmonary manifestations in cat-scratch disease

| Patient characteristics | Year | Clinical signs and symptoms | Atypical manifestations | Diagnosis |
|-------------------------|------|-----------------------------|------------------------|-----------|
| Female, 46, housewife   | 1957 [7] | Fever, Headache, Dyspnea, Enlarged epitrocheal LN | Consolidation of right lower lobe, pleural reaction | Cat-scratch antigen skin test (+) |
| Male, 20, termite technician | 1986 [8] | Fever, Enlarged epitrocheal LN | Mental confusion, increased sleepiness | Lymph node biopsy: Warthin-Starry stain (+) |
| Male, 24 | 1987 [9] | Fever, Pleuritic chest pain, Enlarged left axillary LN | Left apical pleural thickening, Mediastinal mass in the right posterior side | Cat-scratch antigen skin test (+) Mediastinal mass biopsy: necrotizing granulomas, Warthin-Starry stain (-) |
| Male, 34, Renal allograft recipient | 1989 [10] | Fever, Dry cough, Headache, Enlarged left epitocheial and axillary LN | Seizure, Bilateral infiltrates, Septic shock | Lymph node biopsy: Warthin-Starry silver stain (+) |
| Male, 27, HIV (+) | 1990 [11] | Fever, Diarrhea, Enlarged left epitocheial and axillary LN | Pleural effusion, Papilledema, Retinitis, Liver abscess | Lymph node biopsy: Warthin-Starry stain (+) |
| Male, 4 | 1994 [12] | Fever, Vomiting, Right side abdominal pain, Enlarged left axillary LN | Left side pleural effusion and pneumonia | PCR of pleuritic fluid (+) Serology antibody testing (+) |
| Female, 19, renal allograft recipient | 1995 [13] | Fever, Diarrhea, Vomiting | Bilateral pulmonary nodules in the lower lobes, retroperitoneal adenopathy, Low-density lesion in the spleen | PCR on lung nodules biopsy specimens (+) Serology antibody testing (+) |
| Male, 8, | 1995 [14] | Fever, Vomiting, Enlarged left axillary LN | Seizures, Diffuse bilateral interstitial infiltrates | Indirect fluorescent antibody titers 1:512 |
| Male, 11 | 1995 [22] | Fever, Tachypnea, Enlarged inguinal LN | Seizures, bilateral pleural effusions | Lymph node biopsy: Stellate abscess Cat-scratch antigen skin test (-) |
| Male, 6 | 2001 [15] | Fever, Submaxillary LN | Multiple hypodense hepatosplenic lesions, diffuse bilateral reticulonodular pulmonary infiltrates | Positive serology testing |
| Male, 3 | 2007 [16] | Fever | Pneumonia, pleural effusion, and pericarditis | Serology antibody testing (+) |
| Male, 21, HIV (+) | 2010 [17] | Fever, vomiting, blurred vision | Neuroretinitis, Peripheral pulmonary nodules with mediastinal lymphadenopathy | Lung biopsy: granuloma with necrotic center PCR (+) from biopsy sample Serology antibody testing (+) |
| Male, 12 | 2013 [18] | Fever | Bilateral reticulonodular pulmonary infiltrates, Multifocal hepatosplenic granulomas | Serology antibody testing (+) |
| Female, 3 | 2013 [19] | Fever, Cough, Diarrhea | Scattered 2–3mm nodular densities throughout the lungs Focal hypodensities throughout the liver and left kidney | Pulmonary biopsy: necrotizing granulomas PCR (+) from biopsy sample Serology antibody testing (+) |
| Male, 51, selective IgA deficiency | 2016 [20] | Right sided chest pain | Multiple pulmonary nodules, Hypoenhancing hepatosplenic lesions | Lung biopsy: giant cell reaction, caseation necrosis, Liver biopsy: granulomatous hepatitis with scattered non-necrotizing granulomas Serology antibody testing (+) |
| Male, 44, type 2 | 2019 [21] | Hypoxia, Cough | Bilateral opacities, Left | Mediastinal lymph node biopsy: |
| Diabetes | Upper lobe pulmonary nodule, Prominent mesenteric, mediastinal, and hilar lymph nodes | Warth Starry stain (+) Serology antibody testing (+) |
|----------|---------------------------------------------------------------------|-------------------------------------------------|
| Male, 35 | 2020 current case report, Hypoxia, Left epitrochial and axillar LN | Epileptic seizures, Bilateral opacities, nodules and pleural effusions Serology antibody testing (+) |