Infection after knee arthroplasty in an immunosuppressive patient caused by Mycobacterium senegalense: a case report and literature review

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Abstract

Background The infection caused by nontuberculous Mycobacterium (NTM) is gradually increasing in both hospital-acquired infection and community-acquired infection. The most common NTM mainly includes Mycobacterium fortuitum group, the M. chelonae/abscessus group and the M. smegmatis group. The infection caused by the third biological variety in M. fortuitum group is rarely reported worldwide.

Case presentation A 33-year-old female patient with 8-year history of rheumatoid arthritis underwent a right knee arthroplasty and developed postoperative infection. In hospital A, debridement was carried out to preserve the prosthesis joint. M. fortunatum was cultured in biopsy tissue taken out during operation and puncture fluid. Unfortunately, it is impossible to carry out the drug sensitivity test of Mycobacterium in hospital A. Local doctors used amikacin and levofloxacin to treat the patient empirically. In order to treat the infection accurately, the patient went to hospital B (a specialized hospital of tuberculosis) for drug sensitivity test. According to the results of drug sensitivity, cefoxitin, clarithromycin, amikacin and moxifloxacin were used in combination treatment. After two weeks of treatment, there was no significant remission, so the patient went to hospital C (a large teaching hospital) for joint prosthesis removal and bone cement frame transfer operation, and anti-infection treatment was carried out according to the drug sensitivity test results. One month later, the patient's condition improved significantly and the wound healed.

Conclusions NTM should be considered as main pathogens in immunosuppressive patients when the wound did not heal due to infection. Simple antibiotic treatment is not good for deep abscess, but treatment combined with surgical debridement and appropriate antibiotics is obviously effective.

Background

Nontuberculous Mycobacterium (NTM) is mainly distributed in soil, dust, river and tap water. According to the different growth speed, it can be divided into rapid growing mycobacterium (RGM) and slow growing mycobacterium (SGM). RGM refers to the colony that can be seen by naked eyes within 7 days when grow on solid medium. The mycobacteria with colonies more than 7 days old can be seen are SGM. For skin and soft tissue infection, RGM is the main pathogen usually. In clinic,
Mycobacterium fortunatum, Mycobacterium cheloni and Mycobacterium abscess are the most common.

In taxonomy, M. fortuitum group includes M. fortuitum, M. peregrinum, and the unnamed third biovariant complex [1]. Traditionally, according to the result of sorbitol test, the third biological variety of M. fortuitum group can be divided into two groups: negative and positive of sorbitol test. With the development of molecular biology and whole genome sequencing technology, the classification of the third biological variety of M. fortuitum group is becoming more and more precise. At present, the third biological variety of M. fortuitum group included M. senegalense, M. mageritense, M. septicum, M. porcinum, M. neworleanense, M. brisbanense, M. houstonense and M. bonickei [2]. The gene sequences of these strains are very close, and the difference of 16sDNA sequence is 15 bp or less [1].

There are very limited reports of infection caused by the third biological variant of M. fortuitum. Here we report a case of infection after arthroplasty caused by M. senegalense. From this case, we summarize the diagnosis and treatment process of patients, hoping to provide some reference for the future clinical treatment.

Case Presentation

A 33-year-old female patient developed incision pain accompanied by exudation of yellow purulent secretion after right knee replacement (fig 1). The patient had a history of rheumatoid arthritis for more than 8 years. She took methotrexate and leflunomide for a long time. The patient first went to hospital A and found that the right knee joint was swollen and the skin temperature was slightly high. The incision of the operation was interrupted, which could be seen to pass through the superficial subcutaneous sinus about the size of mung bean all the time, and the yellow brown purulent secretion could be seen to flow out (fig 1). The main laboratory test positive indexes were C-reactive protein 32.97mg/l, rheumatoid factor 78.7u/ml, ESR 120mm / h, hemoglobin 85g / L, albumin 27g / L, blood potassium 3.36mmol/l, blood calcium 1.72mmol/l. The swollen right knee joint was punctured, and a large number of leukocytes were found in the puncture fluid and M. fortunatum was detected in the puncture fluid culture. The surgeons performed surgical debridement while preserving the prosthesis
joint. *M. fortuitum* was detected in the tissues taken during operation and the culture of puncture fluid. Because in hospital A the drug sensitivity test of *M. fortuitum* is unable to be carried out, amikacin 0.4g per day, levofloxacin 0.6g per day intravenous drip were used to treat the patients. In order to accurately use antibiotics for treatment, the patient went to hospital B (a special hospital of tuberculosis) to carry out the drug sensitivity test of *M. fortuitum*. According to the results of drug sensitivity, the doctors adjusted the treatment drug, and changed it into intravenous drip of cefoxitin 4.0 g twice a day, intravenous drip of amikacin 0.5 g once a day, intravenous drip of moxifloxacin 0.4 g once a day, and oral administration of clarithromycin 0.5 g twice a day. However, after two weeks of treatment, the patient's condition did not improve significantly. In order to relieve the pain as soon as possible, the patient went to hospital C (a large teaching hospital). In the hospital C, doctors carried out a systematic test on the patient. The detection results of rheumatoid factor and autoantibody of patient were detailed in Table 1. The analysis of the immune function of the patients, including the lymphocyte classification count and its function analysis, were detailed in Table 2. Lymphocyte technology and function analysis were performed by flow cytometry (FACSC CANTO, BD, USA). The results showed that the number of lymphocytes was normal, but the function of T and B lymphocytes and NK cells decreased, demonstrating the patient was in the state of immunosuppression. Surgeons removed the prosthesis and moved the bone cement frame into the patient. The preoperative and postoperative joint imaging data are shown in Figure 2 and figure 3, respectively. *RGM* was detected by bacteriological culture of the tissue samples taken out during the operation. It was identified as *M. senegalense* by IVD-MALDIBIOTYPER (Antu, zhengzhou, China) (fig 4). The strain was re identified as *M. senegalense* by sequencing technology. First generation sequencer (Life Technology 2500 DX, ABI, Japan) was used for sequencing operation. PCR amplification of 16S rDNA and sequencing of PCR products were performed using primers described previously [3]. Compared with those available in the National Center for Biotechnology Information GenBank database, the expansion products of the strain is 99.83% similar to that of M. Senegal (GenBank accession no. Lc082328.1). The drug sensitivity test used the *RGM* drug sensitivity test strip provided by thermo company, and the test principle was micro broth dilution method. The results were interpreted according to CLSI M24-A2 [4],
as shown in Table 3. According to the drug sensitivity results, doctors found that the antibiotics selected in hospital B did not contradict the drug sensitivity results, so they continued to treat according to this plan. After one month of treatment, the wound healed and the patient's condition improved significantly.

**Literature review**

We searched PubMed for cases of *M. fortuitum* group infection and found 10 case reports in addition to ours (Table 4). We found that the previous case reports showed that the *M. fortuitum* group mainly caused wound infection and skin and soft tissue infection. In most cases, the immune function of the patients was not evaluated but drug sensitivity tests were carried out for all of them. The antibiotics used in the treatment were mainly fluoroquinolones such as ciprofloxacin, levofloxacin, moxifloxacin, ofloxacin, aminoglycosides such as amikacin and gentamycin, carbapenems such as imipenem and meropenem, doxycycline and trimethoprim-sulfamethoxazole. When the patient had deep abscess formation, the combination of surgical debridement and appropriate antibiotics had a very significant therapeutic effect. All the patients were cured of the infection caused by *M. fortuitum* group.

**Discussion And Conclusions**

NTM is a common pathogen and a life-threatening infection for people with low immune function, especially for organ transplant patients or stem cell transplant patients [14]. In this case, the patient had a history of rheumatoid arthritis for eight years, and took methotrexate and leflunomide for a long time. The lymphocyte classification of the patients showed that the number of total T-lymphocytes, total B-lymphocytes, helper / inducer T-lymphocytes and suppressor / cytotoxic T-lymphocytes were normal. However, the analysis of lymphocyte function showed that the proportion of IFN - γ positive cells in total CD4 positive T cells, total CD8 positive T cells and NK cells decreased after induction. It showed that the lymphocyte function of the patients was decreased. The results were consistent with the history of rheumatoid arthritis and long-term use of immunosuppressive drugs. For people whose immune function was inhibited, NTM should be considered as a common pathogen.

The pathogenic bacteria in this case is *M. senegalense*, which was identified by IVD-MALDIBIOTYPER
and sequencing in hospital C. However, the identification results of both hospital A and hospital B were M. fortuitum. In fact, these results were not contradictory. M. senegalense belongs to the third biological variety of M. fortuitous group. The commonly used laboratory identification methods include IVD-MALDIBIOTYPER and sequencing. Adela ALCOLEA Medina et al. showed that the accuracy of IVD-MALDIBIOTYPER in identifying Mycobacterium was not 100% compared with sequencing method. The results showed that the accuracy of identification of RGM and SGM were 75% and 85%, respectively compared with sequencing technology [15]. It can be seen that the identification results of NTM by IVD-MALDIBIOTYPER are not 100% accurate. At present, the level of NTM identification to species must depend on the means of sequencing.

M. senegalense was first found in Africa and was mainly prevalent in Africa. In taxonomy, it belongs to the third biological variety of M. fortuitum. The third biological variation of M. fortuitum mainly caused catheter-related infection in immunosuppressed population and surgical wound infections, osteomyelitis following open fractures and localized post-traumatic wound infections in normal immune function population [1]. As NTM mainly come from the soil, dust and water flow in the natural environment, the outbreak of nosocomial infection was mainly seen in the pollution of medical water, and community acquired infection was mainly seen in post-traumatic infection, such as foot on nail or motor vehicle accident trauma [1]. Based on the review of previous reports, we found that M. fortuitous group mainly caused wound and skin soft tissue infection.

Through the summary of this case, we know that for the treatment of deep abscess, only relying on antibiotics treatment, the effect is not good. Surgical debridement combined with appropriate antibiotics is very effective. If it is treated by surgery alone, the infection will recur easily 4-6 weeks after operation [1]. Early studies by Wallace et al. showed that 17% (13 / 76) of the M. fortunatum group and 6% (3 / 47) of the M. cheloni-abscess group could be successfully cured only by surgical means [16].

Conclusions

For the infection of immunosuppressive patients, NTM is a very important pathogen. The treatment of deep abscess only depends on antibiotics or surgical debridement alone, and the therapeutic effect is
not good enough. Only surgical debridement combined with appropriate antibiotics can achieve satisfactory therapeutic effect.

Abbreviations
NTM: nontuberculous Mycobacterium, RGM: rapidly growing mycobacterium, SGM: slow growing mycobacterium

Declarations

Ethics approval and consent to participate
The study protocol was approved by the Tongji Hospital ethics committee for research in health.
Informed written consent was obtained from the patient.

Consent to publish
The patient provided written consent for the case details to be published.

Availability of data and materials
All data is contained within the manuscript. Clinical isolates will be made available upon requests from Dr. Ziyong Sun.

Competing interests
The authors declare that they have no competing interest.

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Authors’ contributions
LT drafted the manuscript. FW analyzed the immune function of patients by flow cytometry. YZ was in charge of the patient’s treatment process. XW, LG and HZ completed the sequencing operation. ZC and ZS were in charge of the laboratory work. All authors read and approved the final manuscript.

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Tables

**Table 1. Detection of rheumatoid index**

| Testing items                         | Results         | Reference range | Unit       |
|---------------------------------------|-----------------|-----------------|------------|
| Rheumatoid factor                     | 118↑T positive  | 0-30            | IU/ml      |
| Anti-keratin antibody                 | positive        | negative        |            |
| Anti-cyclic citrullinated peptide antibody | >400↑           | 0-20            | RU/ml      |
| Antinuclear antibody                  | 1:100 Nuclear particle type | <1:100 negative |            |
|                                       | 1:100 homogeneous type |            |            |

**Table 2. Lymphocyte classification and function analysis**

| Testing items                         | Results         | Reference range | Unit       |
|---------------------------------------|-----------------|-----------------|------------|
| Total T lymphocytes (CD3+CD19-) (#)   | 1202            | 955-2860        | Quantity/ul |
| (CD3+CD19-) (%)                       | 77.44           | 50-84           | %          |
| Total B lymphocytes (CD3-CD19+) (#)   | 324             | 90-560          | Quantity/ul |
| (CD3-CD19+) (%)                       | 20.91↑          | 5-18            | %          |
| Helper / inducible T lymphocytes (CD3+CD4+) (#) | 807            | 550-1440        | Quantity/ul |
| (CD3+CD4+) (%)                        | 52.04↑          | 27-51           | %          |
| Suppressor / cytotoxic T lymphocytes (CD3+CD8+) (#) | 369            | 320-1250        | Quantity/ul |
| (CD3+CD8+) (%)                        | 23.79           | 15-44           | %          |
| NK cell (CD3-/CD16+CD56+) (#)         | 19↓             | 150-1100        | Quantity/ul |
| (CD3-/CD16+CD56+) (%)                 | 1.20↓           | 7-40            | %          |
| Percentage of IFN-γ positive cells after induction in total CD4 + T cells | 6.80↓          | 14.54-36.96     | %          |
| Percentage of IFN-γ positive cells after induction in total CD8 + T cells | 16.72↓          | 34.93-87.95     | %          |
| Percentage of IFN-γ positive cells after induction in total NK cells | 24.14↓          | 61.2-92.65      | %          |
### Table 3. Antimicrobial sensitivity of *Mycobacterium porcinum*

| Antibacterial drugs | MIC (µg/ml) | Susceptible | Breakpoints (µg/mL) | Resistant | Results |
|---------------------|-------------|-------------|---------------------|-----------|---------|
| Trimethoprim/sulfamethoxazole | 8/152 | ≤2/38 | - | ≥4/76 | resistant |
| Ciprofloxacin | ≥4 | ≤1 | 2 | ≥4 | resistant |
| Moxifloxacin | 2.0 | ≤1 | 2 | ≥4 | Intermediate |
| Cefoxitin | 32 | ≤16 | 32-64 | ≥128 | Intermediate |
| Amikacin | ≤1 | ≤16 | 32 | ≥64 | sensitive |
| Doxycycline | 0.5 | ≤1 | 2-4 | ≥8 | sensitive |
| Tegafycline | 0.5 | - | - | - | - |
| Linezolid | 4 | ≤8 | 16 | ≥32 | sensitive |
| Imipenem | 16 | ≤8 | 8-16 | ≥32 | Intermediate |
| Cefepime | ≥32 | - | - | - | - |
| Amoxicillin-clavulanate | ≥64/32 | - | - | - | - |
| Cefatriaxone | ≥64 | - | - | - | - |
| Minocycline | ≤1 | - | - | - | - |
| Clarithromycin | ≤0.06 | ≤2 | 4 | ≥8 | Intermediate |
| Note: - No Breakpoint in CLSI M24-A2 |

### Table 4. Published cases of *Mycobacterium fortuitum* group infection

| Case No | Author Country Year | Sex/Age | Cause of infection | NTM | Infection location | Underlying diseases |
|---------|---------------------|---------|-------------------|-----|-------------------|---------------------|
| 1 | Lei tian/China/2019[5] | Male/68-year-old | M. houstonense | Surgical wound infection | Open humeral fracture |
| 2 | Shih-Chen Tsai/China/2016[6] | Male/40-year-old | M. fortuitum | Skin and soft tissue infection | Sebaceous cyst in the abdominal wall |
| 3 | Bharti Chogtu/India/2017[7] | Male/51-year-old | M. fortuitum | Wound infection | Umbilical hernia |
| 4 | Carlos A. Torres-Duque/ Colombia/2010[8] | Female/seventeen-year-old | M. peregrinum | Pulmonary infection | Prosthetic aortic valve endocarditis |
| 5 | T. Hod/ Israel/2008[9] | Male/65-year-old | M. fortuitum | Peritonitis | Diabetic nephropathy |
| 6 | T. Hod/ Israel/2008[9] | Male/59-year-old | M. fortuitum | Catheter tunnel abscess | Hypertensive nephrosclerosis |
| 7 | Miki Nagao/Japan/2008[10] | Female/58-year-old | M. peregrinum | Surgical site infection | Lipoma |
| 8 | Ian Cheung/ Australia/2007[11] | Female/58-year-old | M. peregrinum | Wound infection | Knee arthroplasty |
| 9 | Corrado Serra/ Italy/2007[12] | Female/32-year-old | M. fortuitum | Lumbar pain | Ureteral colic and lithiasis |
| 10 | MICHAEL SACCENTE/USA/2005[13] | Female/44-year-old | M. fortuitum | Wound infection | Bilateral total knee arthroplasty |

**Figures**
Figure 1
Continuous exudation of tawny secretion from the wound after operation

Figure 2

Imaging showed swelling of the implanted prosthesis joint and surrounding tissue.
Figure 3

Imaging showed that the prosthesis tissue had been removed and implanted into the cement frame for fixation.
Figure 4

The result of IVD-MALDIOTYPHER showed that the strain was Mycobacterium senegalense.