Supplemental Antibiotic Injections into the Disc Eradicate Lumbar Pyogenic Spondylodiscitis and Reduce Residual Lumbago

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Abstract

The standard treatment for lumbar pyogenic spondylodiscitis is an intravenous antibiotic. If conservative treatment fails, surgery is indicated. However, many patients suffer from residual lumbago after prolonged conservative treatment, and invasive surgery is problematic in poorly conditioned patients. We developed a new treatment in which intravenous antibiotics are supplemented by multiple injections of antibiotic directly into the infected disc. Here we report our experience with twenty adult patients. Shortly after infection was diagnosed, we performed the needle biopsy that we reported followed by direct antibiotic injection into the infected disc. Antibiotic was injected twice weekly until inflammation subsided. The average number of injections was 6.8. Infection was eradicated in all cases. Surgery was needed in two cases (10%): one because of acute paralysis and one because of residual sciatica. We observed no adverse effects. Excluding two cases who died of cancer, we obtained excellent (n=12) or good (n=3) results (83%) in a total of 18 patients evaluated according to Macnab’s lumbago criteria at an average of 27.5 months follow-up. The disc height was retained in ten cases during the initial two months of treatment: nine of these patients showed excellent results on Macnab’s lumbago scale. We conclude that supplementing standard systemic antibiotic therapy with multiple injections of antibiotic directly into the infected disc provides a safe and effective method of eradicating lumbar pyogenic spondylodiscitis. This treatment also provides an excellent chance of maintaining the disc height, which leads to less residual lumbago.

Keywords: Lumbar pyogenic spondylodiscitis; Antibiotic; Lumbago; Injection; Disc

Introduction

Lumbar pyogenic spondylodiscitis is a serious condition. Many patients are elderly or immuno-suppressed, and associated medical illnesses are frequent. The standard treatment is intravenous (IV) antibiotic with external immobilization of the spine. If conservative treatment fails, surgical intervention is indicated. The standard surgical technique combines anterior debridement with bone grafting and poses great risks for debilitated patients. Prolonged conservative treatment is also associated with a high incidence of residual lumbago. We have therefore introduced a new treatment: early biopsy with a 21G needle [1] followed by supplemental direct injections of antibiotic into the infected disc. We repeat the direct injections multiple times, usually twice weekly, and also administer the same antibiotic intravenously. Here we report 20 cases in which we have applied this treatment. The purpose of this study is to evaluate the efficacy of direct injection therapy for lumbar pyogenic spondylodiscitis in (1) eradicating infection, (2) retaining disc height and (3) reducing residual lumbago.

Illustrative Case

A 60-year-old man developed severe back pain and high grade fever. He presented to our hospital five days after onset of symptoms. His white blood cell count was 13,000 and CRP was 30.9. He was under insulin therapy for type 2 diabetes. Radiographs and MRI revealed disc infection at L4/5 disc and abscess in the right psoas muscle (Figures 1 and 2).

At the third day after presentation, in addition to systematic antibiotic therapy, a 21-G needle was inserted into the L4/5 disc as for discography (Figure 3).

Fluid was taken from the disc for culture. After pus aspiration and irrigation with saline antibiotic was injected into the disc. MSSA was positive for both blood culture and needle biopsy.
We applied irrigation and supplemental antibiotic injection into the L4/5 disc eight times in 1-month period. The abscess in the psoas muscle was also irrigated through a 21-G needle and antibiotic injected. IV antibiotic was administered for 33 days. Infection eradicated, and his back pain disappeared. The result for low back pain was excellent according to Macnab’s grading at a 24-month follow-up. Radiograph showed that the height of L4/5 disc was maintained (Figure 4).

Patients and Methods

Between March 2004 and December 2014, we encountered and treated 20 cases of lumbar pyogenic spondylodiscitis. There were fourteen men and six women with a mean age of 68.0 (23 to 82). Diagnosis was based on history, elevation of C-reactive protein (CRP), white blood cell counts (WBC) and image studies including X-ray and MRI.

Eight patients had been referred from other practices or departments, and eight had already been treated with IV antibiotic. The time between onset of symptoms and presentation to our department was 19.0 (3 to 64) days. The general condition of the patients was evaluated by the grading system of the American Society of Anesthesiologists (ASA). Two patients had cancer surgery within two weeks of symptom onset. There was no septic patient at presentation. Table 1 lists the medical comorbidities and other information about the patients. All patients but two, who died of cancer, were followed for at least six months (Ave. 27.5 months, 4 to 50).

The final outcome for residual lumbago was evaluated by Macnab’s pain grading [2]. The disc height was measured on plain lateral X-ray or sagittal MR image. The levels of infection were L1/2 (N=2), L2/3 (N=3), L3/4 (N=3), L4/5 (N=10) and L5/S1 (N=2). Four cases sustained multi-level disc lesions; the most severely damaged disc was chosen for disc height measurement. There were three cases with endplate destruction on X-ray. Two cases showed segmental kyphosis, which may not be related to infection. The height was classified into A: normal height, B: half normal height or greater, C: less than half normal height, D: partial collision of the vertebrae and E: disappearance. The X-ray was taken at initial presentation, and again at two and six months, and at one, two, and three years later, as possible.

Treatment

Under fluoroscopy the patients were placed in the lateral or prone position, as for discography. At the first treatment a needle biopsy with a 21G needle was performed, as we reported [1], to isolate the causative microorganism. This needle biopsy technique consists of three parts. (A) Insert a 21G needle as for discography, aspirate pus or fluid. (B) If step A fails, inject saline and collect fluid as reflux. (C) If step B fails, insert another needle into the disc, inject saline and collect reflux from another needle.
### Table 1: The medical comorbidities and other information about the patients.

| Patient | Age | Sex | Onset to Diagnosis (days) | Level | Associated medical illness | ASA | WBC (x1000) | CRP (mg/dL) | Needle biopsy | Blood culture | Organism isolated | Abscess |
|---------|-----|-----|----------------------------|-------|---------------------------|-----|-------------|-------------|---------------|--------------|----------------|-----------------|---------|
| 1       | 79  | F   | 28                         | L4/5  | Pyelonephritis            | 3   | 15.7        | 11.5        | N/A           | N/A          | Escherichia coli | Psoas         |
| 2       | 68  | M   | 46                         | L4/5  | Previous lumbar surgery   | 1   | 12.2        | 6.9         | N/A           | N/A          | Unknown        | none            |
| 3       | 60  | M   | 3                          | L4/5  | Diabetes Mellitus, Femoral head necrosis | 2   | 13.0        | 34.5        | P             | P            | MSSA           | Psoas          |
| 4       | 73  | M   | 46                         | L3/4  | Acute myocardial Infarction, Colon Cancer | 4   | 13.5        | 16.1        | P             | N            | MSSA           | Psoas          |
| 5       | 61  | F   | 64                         | L2/3  | Poor nutrition, Epilepsy  | 2   | 12.8        | 17.6        | P             | N            | Hafnia         | none            |
| 6       | 78  | M   | 7                          | L4/5  | Diabetes Mellitus         | 2   | 7.9         | 14.3        | P             | N            | Corynebacterium | Psoas          |
| 7       | 82  | F   | 9                          | L3/4/5|                            | 2   | 22.2        | 32.0        | P             | N            | S. epidermidis | Psoas          |
| 8       | 83  | F   | 30                         | L2/3  | Bladder Cancer            | 4   | 14.5        | 16.9        | P             | P            | MRSA           | Epidural       |
| 9       | 66  | M   | 6                          | L4/5  | Liver cirrhosis           | 3   | 15.8        | 9.8         | P             | P            | Klebsiella pneumoniae | none        |
| 10      | 23  | M   | 14                         | L5/S  | Obesity (BMI=39), Atopic dermatitis | 1   | 13.0        | 5.0         | P             | P            | MSSA           | Epidural       |
| 11      | 67  | M   | 10                         | L5/S, L3/4 | Hepatocellular carcinoma | 4   | 12.3        | 12.3        | P             | N            | S. oralis      | none            |
| 12      | 53  | M   | 3                          | L5/S  | Acute myocardial infarction | 4   | 6.8         | 5.6         | P             | P            | MRSA           | none            |
| 13      | 62  | M   | 3                          | L1/2  |                            | 1   | 12.6        | 1.8         | P             | N            | Preptostreptococcus | none        |
| 14      | 82  | M   | 4                          | L2/3  |                            | 2   | 6.7         | 1.8         | N             | N            | Unknown        | none            |
| 15      | 79  | F   | 14                         | L3/4  | Colon Cancer, Pyelonephritis | 3   | 17.7        | 33.8        | N             | P            | Streptococcus B | Psoas          |
| 16      | 55  | M   | 7                          | L1/2/3| Obesity (BMI=42), Diabetes Mellitus | 2   | 16.8        | 34.8        | N/A           | N/A          | Streptococcus B | Epidural       |
| 17      | 87  | M   | 5                          | L2/3/4| Old myocardial infarction  | 3   | 6.7         | 2.4         | N             | N            | Unknown        | none            |
| 18      | 75  | F   | 20                         | L3/4  |                            | 2   | 10.4        | 16.5        | P             | N            | Salmonella     | Psoas          |
| 19      | 61  | M   | 40                         | L5/S  | Diabetes Mellitus, decubitus | 4   | 14.6        | 22.8        | P             | P            | MSSA           | none            |
| 20      | 66  | M   | 21                         | L4/5  | Diabetes Mellitus         | 1   | 6.3         | 3.9         | N             | N            | Unknown        | none            |
| Average | 68  | M   | 19                         |       |                           |     | 12.1        | 15.0        |               |               |                |                 |

ASA: Grading System of the American Society of Anesthesiologists, BMI: Body Mass Index, MSSA: Methicillin-Susceptible Staphylococcus aureus, S. epidermidis: Staphylococcus epidermidis, MRSA: Methicillin-resistant Staphylococcus aureus, S. oralis: Streptococcus oralis, P: Positive, N: Negative, N/A: Not Applicable

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Co. Ltd. Osaka, Japan) was used for cases with Methicillin-resistant Staphylococcus aureus (MRSA). Patients were not required to remain at bed rest and were allowed to walk with bracing when pain diminished. The treatment continued until inflammation subsided, as determined by relief of spine pain and decrease in levels of CRP to less than 0.5 mg/dl.

| Patient | Age | Sex | Diagnosis to direct Injection (days) | Number of Injection | Duration of direct injection (days) | Duration of IV treatment | Follow up (months) | Residual lumbago | Disc height change presentation→2M→last follow-up | Other information |
|---------|-----|-----|-------------------------------------|--------------------|------------------------------------|-------------------------|-------------------|---------------|---------------------------------------------|------------------|
| 1       | 79  | F   | 3                                   | 3                  | 20                                 | 41                      | 49                | Excellent     | A→A→B                                      |                  |
| 2       | 68  | M   | 0                                   | 3                  | 10                                 | 10                      | 44                | Excellent     | B→B→B                                      |                  |
| 3       | 60  | M   | 4                                   | 8                  | 24                                 | 33                      | 38                | Excellent     | A→A→A                                      |                  |
| 4       | 73  | M   | 3                                   | 2                  | 11                                 | 22                      | 28                | Excellent     | B→B→B                                      |                  |
| 5       | 61  | F   | 6                                   | 5+5                | 22+20                              | 31+24                   | 23                | Good          | D→E→E                                      | Relapsed two months later |
| 6       | 78  | M   | 3                                   | 8                  | 30                                 | 55                      | 18                | Poor          | A→D→D                                      | Surgery for residual lumbago |
| 7       | 82  | F   | 4                                   | 12                 | 41                                 | 51                      | 9                 | Excellent     | A→A→A                                      |                  |
| 8       | 83  | F   | 3                                   | 4                  | 9                                  | 23                      | 9                 | N/A           |                                            | Died of cancer. Infection subsided. |
| 9       | 66  | M   | 4                                   | 21                 | 35                                 | 8                       | Excellent       | A→D→D                                      |                  |
| 10      | 23  | M   | 0                                   | 12                 | 36                                 | 36                      | 12                | Good          | A→B→B                                      |                  |
| 11      | 67  | M   | 1                                   | 10+6               | 37+22                              | 48+25                   | 4                 | N/A           |                                            | Relapsed at another disc. Died of cancer. Infection subsided. |
| 12      | 53  | M   | 0                                   | 12                 | 46                                 | 51                      | 22                | Good          | B→C→C                                      |                  |
| 13      | 62  | M   | 0                                   | 5                  | 18                                 | 18                      | 21                | Fair          | C→C→C                                      |                  |
| 14      | 82  | M   | 3                                   | 7                  | 38                                 | 52                      | 21                | Excellent     | A→A→A                                      |                  |
| 15      | 79  | F   | 0                                   | 10                 | 10                                 | 40                      | 12                | Excellent     | B→B→B                                      |                  |
| 16      | 55  | M   | 6                                   | 3                  | 18                                 | 24                      | 24                | Excellent     | A→A→A                                      | Paralysis due to epidural abscess, posterior surgery |
| 17      | 67  | M   | 0                                   | 5                  | 15                                 | 25                      | 15                | Excellent     | A→B→B                                      |                  |
| 18      | 75  | F   | 0                                   | 1                  | 1                                  | 23                      | 12                | Fair          | A→C→C                                      |                  |
| 19      | 61  | M   | 1                                   | 7                  | 27                                 | 36                      | 6                 | Excellent     | B→C→C                                      |                  |
| 20      | 66  | M   | 6                                   | 4                  | 20                                 | 17                      | 12                | Excellent     | C→C→C                                      |                  |
| Average |     |     | 2.15                                | 6.8                | 24.8                               | 36.8                    | 18.4             |               |                                            |                  |

Table 2: Summary of treatment.

There were abscesses in the psoas muscle of eight patients. Five of these patients also received injections of antibiotic into the psoas muscle under fluoroscopy in a prone position. Psoas abscesses resolved after initiation of the disc treatment and control of the infection.

There were three patients with epidural abscess. One individual needed emergent posterior decompression because of paralysis; surgery was followed by direct injection of antibiotic into the disc in addition to IV antibiotic. In the other two cases disc injection therapy was performed without surgical treatment. Table 2 demonstrates summary of treatment.

Results

According to the ASA scale, four cases each were grade 1 and 3, five were grade 4, and seven were grade 2. The average white blood cell count was 12600/mm³ (6300 to 22200) and the mean value of CRP was 15.0 (1.8 to 32.0) mg/L. The average time from onset to presentation to our department was 19.0 (3 to 64) days.

Direct injection started an average of 2.2 (0 to 6) days after presentation. The average duration of direct antibiotic injections into
the disc was 24.8 days and the average number of injections was 6.8 (1 to 12). The average duration of IV treatment was 36.0 (14-63) days.

In eight of the 17 patients in whom it was performed (47%), the causative microorganism was identified by blood culture, and in 12 of 17 patients in whom it was performed (70.6%) an organism was isolated from the samples taken during needle aspiration biopsy. In total, 16 patients had a positive culture. The most commonly identified organism was Staphylococcus aureus (4 cases). MRSA and Streptococcus Group B were each identified in two cases. Mycobacterium tuberculosis was not found.

An abscess was found in the psoas muscle of eight patients and an epidural abscess was found in three. The epidural abscess caused severe neurological deficits in one patient (Frankel A, patient 16); this patient underwent posterior decompression surgery on the next day after presentation and then received disc injection treatment. His neurological deficits recovered fully. Except for this case, the combination of multiple disc injections of antibiotic and IV antibiotic eradicated the infection in all cases, and all abscesses disappeared without major surgery. No patients showed neurological deterioration or other adverse effects.

In one patient who received five local injections and one who received ten local injections, infection recurred within two months after the first course of treatment. The second biopsy was positive for the same microorganism in one patient and negative in the other. A second course of five and six local injections in combination with IV antibiotic eradicated the recurrent infections. One of these patients died of cancer within three months; the other showed no signs of relapse in a one-year follow-up period.

As for drug related adverse effect, two cases (Patient 1 and 3) showed elevation of liver enzymes after over one-month’s treatment. The antibiotic treatment was ceased, however, their infection eradicated shortly after halt of the treatment.

Excluding the two cases who died of cancer within four months after the onset of infection, the outcome for lumbago in the other 18 patients was excellent in 12 (66.7%), good in three (16.7%), fair in two (11.1%) and poor in one (5.6%). The poor result (patient 6) was caused by spinal deformity and spinal canal stenosis. This patient underwent posterior decompression surgery four months later.

Figure 5 shows the overall disc height change for patients with fair and poor results on the lumbago scale.

The disc height decreased in ten cases; eight of these showed a decrease in the first two months. All patients but one whose disc height was maintained or decreased slightly (one-class down) showed excellent or good outcomes for lumbago.

Figure 6 demonstrates the relationship between residual lumbago and disc height change for 18 cases during the initial two months. In this period, the disc height was maintained in 10 patients: nine of these (90%) showed excellent results and one showed a fair result for lumbago.

![Disc height](image)

**Figure 5:** Overall disc height change. A: normal height, B: half or higher, C: less than half, D: partial collision of the vertebrae, E: disappearance. Only fair and poor results are shown in the figure. Results in the other cases were excellent or good.

In five cases the disc height moved one-class down (A to B, B to C or D to E); two of these showed excellent results for lumbago, and results were good for three. In one patient whose disc height moved down two classes (A to C), the result for lumbago was fair. Two patients showed a three-class decrease in disc height (A to D); in one of these the result for lumbago was excellent and in the other, the outcome was poor.

**Discussion**

The diagnosis and treatment of lumbar pyogenic vertebral osteomyelitis remain difficult, in part because increasing numbers of individuals are elderly or immuno-compromised. Standard treatment consists of strict immobilization and IV antibiotics, but invasive surgery is often necessary.

The widely accepted surgical indications [3,4] are neurological deficit, significant disc and adjacent vertebral body destruction, spinal instability or developing kyphosis, psoas or epidural abscesses, intractable pain, and failed medical management. The timing of surgery is controversial. Surgery is usually selected after conservative management for one or two months has failed. The standard procedure, anterior debridement and bone grafting, is highly invasive and should not be used for a patient in poor general condition. In the current study, nine out of twenty patients (45%) were at high risk, ASA grade 3 or 4, because of medical comorbidities or advanced age.
Another problem in treating lumbar pyogenic spondylodiscitis is that, even if infection subsides, prolonged conservative treatment is often followed by residual low back pain.

Variable rates for surgery are reported in the literature. Legrand et al. [5] stated that surgery is rarely needed, whereas Hadjipavlou et al. [6] and Sundararaj et al. [7] reported rates greater than 50%.

Legrand et al. [5] insisted that conservative treatment alone eradicated infection in almost all cases, but the rate of residual lumbago reached 40% and functional impairment 15%. Hadjipavlou et al. [6] recommended surgical treatment in part because it resulted in less residual pain (26%) than prolonged conservative treatment (64%).

Many authors [8–10], however, suggested that surgery is needed in approximately 25 to 30% of cases of lumbar pyogenic spondylodiscitis. In the current study, we succeeded in avoiding surgery in all but two cases (10%), and the final outcome was extremely favorable: in 15 (83%) out of 18 patients results were excellent or good, and we observed no functional impairment. Even in the two cases which required surgery, we were able to choose the less invasive posterior decompression.

One possible reason for the successful outcome of our strategy is that early direct treatment reduced destruction of the involved disc compared to IV treatment alone, which improved stabilization of the spine.

Nearly all patients (90%) who maintained disc height during the initial two months showed excellent outcomes for lumbago (Figure 6). We therefore consider that preventing disc destruction in the early stage is the most important factor in reducing residual lumbago and that, if loss of disc height in the initial two months cannot be entirely prevented, keeping it to the smallest amount possible will have the best chance of reducing lumbago.

There have been several attempts to treat pyogenic spinal infections with minimally invasive surgical techniques. Since 1991, Yu et al. [11] and other authors [12–14] used percutaneous discectomy to treat pyogenic spondylodiscitis. A technique for continuous percutaneous suction drainage [15] was also developed and later modified [16]. More recently, endoscopic debridement following continuous irrigation was reported [17]. All of these methods eradicated infection in 90 to 100% of cases. The advantages of our approach are that no special instrument or technique is needed and that the procedure is safer and considerably less invasive than previously described methods.

The key element of our therapy is that the treatment should start soon after pyogenic spondylodiscitis has been diagnosed. In the previously reported minimally invasive techniques, treatment did not begin until conservative management for three weeks to two months had failed. We believe that the main focus of infection is in the disc and that early initiation of treatment of the disc is critical, because it increases the chances for isolating the causative organism [2] and preventing infection from spreading. The minimal invasiveness and absence of adverse effects justify beginning this treatment at an early stage and using multiple injections. The average number of direct antibiotic injections into the disc was 6.8 and the average duration was 24.8 days. Most patients recovered clinically within this period, but on average IV treatment continued for 10.5 days after the cessation of direct injections. The main reason for prolonged IV treatment was to provide an extra margin of safety and because CRP did not return to normal in some cases even after clinical symptoms had disappeared.

An important element of the technique is that the injection should cover the entire area of the disc. We believe that a small focus of infection in the disc is isolated from blood flow and that injecting the loculated area may open the barrier and restore the blood supply. The combination of revascularization, drainage due to needle insertion and pharmacological effects of the antibiotic helps to eradicate the disc infection.

In general, systemically administered antibiotic penetrates very poorly into the disc. For example, after intravenous Cephradine (Cephalosporine) or Flucloxacillin (Penicillin), no antibiotic was detected in discs removed during scoliosis surgery [18]. We consider that direct injection enhanced the pharmacological effect of the antibiotic and that this strategy runs little risk of creating antibiotic-resistant microorganisms because the same agent is used for IV and local injection.

During the procedure, we made sure to maneuver the needle tip so as to administer antibiotic to the whole disc area and to apply the injection alternately from right or left.

There is a concern that direct injection may spread infection to the epidural space or the psoas muscle, but we encountered no case in which direct injections caused epidural or psoas abscesses. We first used this technique for a patient with psoas abscess, but are now comfortable applying it to patients without psoas abscess.

Our experience includes three cases of epidural abscess. One case with severe paralysis (Frankel A) needed emergent posterior decompression surgery on the day following presentation to our hospital. We added direct antibiotic injection to IV treatment on the sixth day. His neurological deficit recovered fully after three direct injections. The result for low back pain was excellent according to Macnab's grading at the 12-month follow-up. His disc height was maintained (A to A).

Because the other two cases with epidural abscess showed no paralysis, we started direct injection treatment along with IV administration. Infection was eradicated in both cases. The low back pain of one of these patients recovered to "good" and his disc height changed from A to B. The other case died of cancer within four months and was excluded from the lumbago study.

Our series included eight cases of psoas abscess. In five of these patients we injected antibiotic into the psoas muscle (in the prone position) in addition to injecting it directly into the disc. The technique was as follows: a 21G needle was placed on the transverse process, as for selective radiculography, and then advanced gradually according to the distance measured in CT image or MRI. Antibiotic injection followed aspiration of pus and saline irrigation. Infection was eradicated in all eight cases. We think that the epidural space and psoas muscle are more resistant to microorganisms than the disc because their circulation is richer, and we are confident that disc injection therapy barely spread infection to the epidural space or the psoas muscle.

Although our results show that most of the patients recovered successfully with this treatment, the study has several limitations. The number of patients was relatively small, for example, and we did not encounter cases with tuberculosis or fungus infections. Direct injection is a painful procedure for patients and an additional work for surgeons. The overall usefulness of the technique therefore remains to be established by studying more patients and different types of infection. Comparison of the present results with those produced by conservative
treatment is important but difficult, because of large differences in patients’ general condition and severity of infection. We think, however, that, in this series, our outcome was favorable both in eradicating infection and reducing residual lumbago.

Conclusion

In conclusion, repeated antibiotic injection into the infected disc provides a simple and effective treatment for eradicating lumbar pyogenic spondylodiscitis that also assists in maintaining disc height and reducing residual lumbago. It deserves broader application, especially in debilitated patients.

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