Debridement and Corpectomy Via Single Posterior Approach to Treat Pyogenic Spondylitis after Vertebral Augmentation

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Research article

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Abstract

**Background** Infection after VA often limits the daily activities of patients and even threatens their life. Operation has become an indispensable choice for such patients. sPVRIF may be one of the effective treatments if the patient suffers from intolerable severe pain, neurological deficits, and damage to spinal stability. This study aimed to investigate the clinical efficacy of treatment of PSVA with sPVRIF.

**Methods** The study was performed on 19 patients with PSVA who underwent VA at 4 hospitals in the region between January 2010 and July 2020. Nineteen patients were included. Among them, 16 patients underwent sPVRIF to treat the PSVA.

**Results** A total of 2267 patients underwent VA at 4 hospitals in the region. Of the 19 patients with postoperative PSVA, suppurative spondylitis was misdiagnosed as an osteoporotic vertebral fracture in 4 patients and they underwent VA. Besides osteoporosis, 18 patients had other comorbidities. The average interval between the first surgery and the diagnosis of PSVA was 96.4 days. Of the 19 patients, 16 received surgical treatment. Among the patients undergoing surgery, one died of refractory septic shock after the surgery, and one died of prostate cancer. The surgical time was 175.0±16.8min, and the intraoperative blood loss was 465.6±166.0 mL. Fourteen patients recovered from the infection. Pathogenic microorganisms were cultured in 12 patients.

**Conclusion** PSVA is an extremely serious complication that can even threaten the life of the patients. sPVRIF may be one of the effective treatments if the patient suffers from intolerable severe pain, neurological deficits, and damage to spinal stability.

**Background**

Vertebral augmentation (VA) has the advantages of immediate pain relief, minimal surgical trauma, and easy surgical techniques. Since its first clinical application by Galibert in 1984[1], VA has gradually become one of the main methods for treating osteoporotic vertebral fractures (OVFs). Despite the extensive development of VA, preventing multiple complications related to surgery is still a key issue that cannot be ignored in improving clinical efficacy. Previous studies reported that the incidence of pyogenic spondylitis after vertebral augmentation (PSVA) fluctuated at 0%–1.6%, which was low, but the consequences were extremely serious and even threatened the lives of patients[2-4]. At present, most of the relevant studies on PSVA are case reports. These studies focused on the high-risk factors of postoperative infection, clinical manifestations after infection, and clinical diagnosis[5-16]. However, no in-depth research has been conducted on the surgical treatment of such patients. Antibiotics cannot form an effective antibacterial film around the infected lesion, and conventional anti-infective treatment measures are not sufficient because bone cement is a nonbioactive artificial prosthesis. In addition, once such patients develop PSVA, they often experience unbearable severe pain, spinal deformity and loss of stability, and neurological deficits. Surgery inevitably becomes one of the main treatment methods for such patients. Park JW et al. [17] reported 11 patients with PSVA, 10 of which underwent surgery. The surgical procedure was thorough debridement of infected tissue and material, including polymethylmethacrylate (PMMA) following anterior column reconstruction via the anterior approach and percutaneous pedicle screw fixation. However, patients with OVF are mostly older people with multiple comorbidities; they cannot bear the trauma caused by one-stage anterior and posterior surgery. Most spinal diseases can be treated via a single surgical approach with the familiarity of spinal anatomy and the highly developed surgical instruments related to spinal surgery. This study was novel in using single posterior debridement, vertebral body resection, and intervertebral bone graft fusion and internal fixation (sPVRIF) to treat PSVA so as to shorten the surgical time, reduce the surgical trauma, and avoid interference with the abdominal organs of the patient. The relevant surgical techniques, surgical experience, and preliminary results of the surgery are discussed in the following sections.

**Methods**

All patients with postoperative spinal infections between January 2010 and July 2020, who primarily underwent VA, were examined after obtaining approval from the institutional review board of the hospital. Postoperative infection occurred in 9 patients (out of 1286 patients operated, with an infection rate of 0.7%), and 10 were referred from the other 3 institutions for the treatment of infection (out of 981 patients operated, with an infection rate of 1.0%). Finally, 19 patients were included in this study (Table 1).
PMMA was used for all patients. Kyphoplasty was performed in 12 and vertebroplasty in 7 patients. The baseline data of the patients, such as sex, age, comorbidities, C-reactive protein (CRP) level, erythrocyte sedimentation rate (ESR), and visual analog scale (VAS) scores, were recorded before the first and the second surgeries. After the patient was diagnosed with PSVA, urine culture, sputum culture, and blood culture were routinely performed. Blood for blood culture should be drawn blood before using antibiotics besides drawing blood during chills. In addition, three blood samples from both sides of the human body are required for blood culture to increase the detection rate of pathogenic microorganisms. One of the 19 patients in this study gave up treatment, 2 chose conservative treatment, and 16 chose surgery, based on whether the patient had severe pain, neurological deficits, and loss of spinal stability, combined with the patient's own desire for treatment.

The brief procedure of the surgery was as follows. The patient's information was checked. Tracheal intubation was performed under general anesthesia, and the patient was placed in a prone position, with a suspended abdomen. Before the surgery, the C-arm was used to confirm the surgical site, and the surgical area was sterilized with 5% iodophor solution three times. The skin and subcutaneous tissue were incised, and the bilateral paravertebral muscles were separated and exposed. The Wiltse approach was used to implant the pedicle screw of the intended fixed segment (the decision to implant a shorter pedicle screw into the destroyed vertebral body was made based on the extent of destruction, and the basic principle was that the screw should not be close to the local lesion). The C-arm fluoroscopy pedicle screw was in a good position. One side was selected as the channel for lesion and cement removal, while a temporary connecting rod was placed on the other side. The articular joints, lamina, pedicle, and other structures of the destroyed vertebrae were exposed through the compartment of muscles between the longis and multifidus muscles. Piezosurgery was used to remove one side of the lamina, pedicle, facet joint, and other structures. The dural sac and corresponding nerve roots were protected, the destroyed vertebral body and bone cement were removed in piecemeal, and the necrotic tissue around the destroyed vertebral body was completely removed. The tissues at the junction of necrosis and normal tissues were selected for bacterial culture and pathological examination. The surgical area was cleaned repeatedly, and the autogenous ribs or iliac bones were selected for intervertebral bone grafting according to the level of the infected vertebral body; then, bilateral connecting rods and transverse connectors were placed (Fig. 1,Fig.2). A plasma drainage tube was placed in the surgical area, and the surgical incision was closed. After the surgery, the patient was sent to the intensive care unit and transferred to the general ward after the condition was stable.

After the surgery, the surgical area was flushed through the plasma drainage tube using sterile normal saline for 7–10 days. If pathogenic microorganisms were detected before the surgery, antibiotics were selected based on the drug sensitivity test after the surgery and infused intravenously for 2 weeks and orally for 6 weeks. If no evidence of pathogenic microorganisms was found before the surgery, antibiotics were empirically injected with higher bone tissue distribution. After detecting pathogenic microorganisms, the antibiotics were adjusted according to the results of the drug sensitivity test. If pathogenic microorganisms could not be cultivated, empirical intravenous infusion of antibiotics with a higher bone tissue distribution was used for 2 weeks and oral administration for 6 weeks. The conservative treatment mainly involved the use of antibiotics, and the principles of antibiotic use were the same as that of surgical treatment.

Statistical analysis

Statistical analysis was performed using the commercial software package SPSS 19.0 (SPSS, IL, USA). All results for continuous variables were presented as mean ± standard deviation, while those for categorical variables were expressed as n.

Results

From January 2010 to July 2020, 2267 patients underwent VA at 4 hospitals in the region. Further, 19 had PSVA, and only 7 underwent biopsy simultaneously, of which 6 showed negative results in pathological examination, and 1 was detected with chronic low-virulence infection of the vertebral body. The patient's medical history, inflammation indicators, and imaging examinations revealed that four were highly suspected of having suppurative spondylitis before undergoing VA. Among the 19 patients, 5 were men and 14 were women, with an average age of 73.0 (59–90) years. Besides osteoporosis, 18 patients had other comorbidities, among which hypertension and diabetes were the most common. Kyphoplasty was performed in 12 and vertebroplasty in 7 patients. Before VA, the CRP level, ESR, and other inflammatory indexes were completely normal in seven patients (Table 1).
The average time interval between VA and diagnosis of PSVA was 96.4 (14–373) days. According to Abdelrahman's et al. [10] diagnostic criteria, 4 patients had an acute infection (within 1 month) and 15 had a chronic infection (more than 1 month). Further, 10 patients had fever or chills during the course of the disease. After the occurrence of PSVA, the CRP level, ESR, white blood cell (WBC) count, and neutrophil (Neu) count were significantly higher than the normal values in patients. The VAS score reached 7.6 (610) points. Among the 19 patients with PSVA, 7 developed neurological deficits (Table 2).

Among the 19 patients with PSVA, 14 completed the follow-up, with an average follow-up time of 14.3 (33 ± 3.5) months. Sixteen patients finally chose surgery due to unbearable severe pain, neurological deficits, and spinal instability. The surgical method was sPVRIF. All patients had a smooth surgery. The surgical time was 175.0 ± 16.8 (155–210) min, and the intraoperative blood loss was 465.6 ± 166.0 (300–900) mL. After surgical treatment, one patient died of refractory septic shock, one died of prostate cancer, and two still needed to live in a wheelchair. The daily activities of the remaining patients significantly improved after the surgery. The VAS score at the last follow-up was 1.8 (13) points. Among the patients treated conservatively, one died of refractory septic shock and one needed to use a walker for assistance in completing daily activities. The patient who gave up treatment eventually died. A total of 12 of the 19 patients showed evidence of pathogenic microorganisms, among which the most common pathogenic bacteria were *Gluconococcus* and *Streptococcus* (Table 3).

**Discussion**

VA can immediately rebuild the stability of the spine and effectively relieve pain through the insertion and support between the bone cement and the fractured end of the vertebral body. However, bone cement leakage, refracture, bone cement toxicity, and PSVA are still important complications that restrict the wide clinical application of VA. The incidence of PSVA is low, but the consequences are very serious, even threatening the lives of patients. Since Yu et al. [5] first reported the PSVA, different scholars have summarized the characteristics, clinical manifestations, diagnostic methods, and treatment principles of PSVA. The bone cement currently used in clinic has no biological activity, and the polymerization of bone cement monomer leads to an increase in temperature inside the vertebral body. The bone tissue that is burnt and necrotic often forms an isolation zone between the bone cement and the normal bone tissue. The aforementioned factors are all important reasons why it is challenging to cure PSVA using conservative treatment. Patients with PSVA often suffer from uncomfortable severe pain, local kyphosis, and neurological deficits. Hence, it is difficult to achieve satisfactory results with antibiotics alone. Surgical treatment has become an indispensable choice for such patients. However, in-depth and systematic research on the surgical method of such patients lacks due to the small sample size. The scholars in the past often recommended the use of anterior debridement combined with posterior internal fixation for the treatment of PSVA to completely remove the lesions, pus, and bone cement.[3,6,10,17-20]. However, such patients often have multiple diseases due to their advanced age, and multiple-organ dysfunction throughout the body can hardly tolerate such large surgical trauma. Since the 21st century, most spinal diseases could be completed via a single approach with an in-depth understanding of the pathophysiological mechanism of spinal diseases, anatomical structure of the spinal column, highly developed spinal surgical instruments and increasing proficiency of surgical techniques. This study was performed on 19 patients with PSVA, which is the largest sample used so far. All surgical patients in this study were treated with sPVRIF. The surgical time was 175.0 ± 16.8 (155–210) min and the intraoperative blood loss was 465.6 ± 166.0 (300–900) mL, which were significantly lower than those reported in previous studies. This was mainly due to the simultaneous completion of lesion removal and internal fixation via a single surgical approach. In addition, piezosurgery is vital in the resection of the vertebral body and bone cement. Among the 16 patients with PSVA, 14 completed the last follow-up. The daily activities of the other patients significantly improved, except for two patients who still needed to be in a wheelchair after the surgery. Only one patient died of postoperative refractory septic shock, and the mortality rate was significantly lower than that reported by Abdelrahman. Besides the improvement in the surgical method, the following improvements were made during the perioperative period: (1) Patients’ CRP, ESR, Neu, and other inflammatory indicators were dynamically observed besides following the basic principles of antibiotic use. The types of antibiotics used to avoid bacterial imbalance and double infections were dynamically adjusted based on the results of the drug sensitivity test. (2) The stability of the patient’s internal environment was maintained, and the nutritional status was improved. The plasma protein level of such patients was maintained at more than 35 g/L, and continuous maintenance of the hemoglobin level more than 100 g/L was essential to enhance the patient's disease resistance. (3) The surgical area for each patient was routinely flushed for 7–10 days after the surgery to reduce the concentration of local pathogens and inflammatory mediators. (4) The patients’ oral and perineal care was strengthened to prevent urinary tract and lung infections.
Previous studies [10,12,16-17] reported that the incidence of PSVA was 0%–1.6%, and the incidence in this study was 0.83%. Although the incidence was low, it resulted in catastrophic consequences to patients. Previous scholars [21] suggested the use of bone cement mixed with tobramycin for VA, while some recommended the use of perioperative intravenous prophylactic antibiotics, to prevent the occurrence of such complications in high-risk patients. However, the clinical efficacy of the aforementioned methods still requires large-sample prospective comparative studies for validation. In this study, neither cement-loaded antibiotics nor systemic perioperative prophylactic intravenous antibiotics were used; instead, a single intraoperative prophylactic dose of a first-generation cephalosporin was used. The experience was as follows: (1) If the levels of inflammation indicators were elevated before the surgery, pulmonary infection and urinary tract infection needed to be carefully checked. If a clear infection of other parts was detected, it was recommended to perform VA 2 weeks after the infection was cured. (2) If infectious diseases of the vertebral body could not be ruled out, it was recommended to give priority to conservative treatment after 2 weeks and re-examine the magnetic resonance imaging and computed tomography of the fractured vertebral body. If it was an infectious disease, the progress of vertebral body disease was often found at this time. During conservative treatment, a biopsy of the destroyed vertebral body was performed. (3) Routine radionuclide bone scintigraphy was recommended for patients initially diagnosed with OVF.

Obtaining etiological evidence is key to the treatment of infectious diseases. Patients with PSVA often start using antibiotics before obtaining pathogenic evidence, and it is generally difficult to cultivate pathogenic microorganisms. In addition, Vats HS et al. [13] reported that polymerase chain reaction (PCR) increased the detection rate of pathogenic microorganisms. Elderly people have long-term oral usage of multiple drugs to treat other basic diseases and lack personal hygiene and health protection knowledge, leading to significant changes in the bacterial spectrum of infectious diseases. Hence, routine urine culture, sputum culture, and blood culture are recommended. Blood for blood culture should be withdrawn before using antibiotics and during chills. In addition, three blood samples from both sides of the human body for blood culture are required to increase the detection rate of pathogenic microorganisms. While searching for pathogenic evidence, attention should be paid to rare pathogenic microorganisms, such as mycobacteria, fungi, and anaerobic bacteria.

This study was novel in reporting the clinical efficacy of sPVRIF in the treatment of PSVA. However, it had some limitations. This study was a single-center retrospective study and lacked comparative findings. Also, the follow-up time of this study was short. Further, since most patients in this study required oral administration of multiple drugs (such as nonsteroidal anti-inflammatory and analgesic drugs) for other comorbidities, it was impossible to use a unified standard to evaluate the fusion of intervertebral body grafts after the surgery.

**Conclusions**

PSVA after VA is a severe complication and can even threaten the lives of patients. The possibility of vertebral infection should be ruled out before VA. Once diagnosed with suppurative spondylitis, pathogenic microorganisms should be detected through blood culture, bacterial culture of necrotic tissue at the lesion site, and PCR. The use of sPVRIF combined with the standardized antibiotic application is one of the effective treatment methods if the patient has intolerable severe pain, neurological deficits, and spinal stability damage.

**Abbreviations**

CRP, C-reactive protein; ESR, erythrocyte sedimentation rate; Neu, neutrophil; OVF, osteoporotic vertebral fractures; PCR, polymerase chain reaction; PMMA, polymethylmethacrylate; PSVA, pyogenic spondylitis after vertebral augmentation; sPVRIF, single posterior debridement, vertebral body resection, and intervertebral bone graft fusion and internal fixation; VA, vertebral augmentation; VAS, visual analog scale; WBC, white blood count.

**Declarations**

**Ethical approval and consent to participate**
The study protocol was approved by the Ethics Committee of the Affiliated Hospital of Southwest Medical University. All patients provided written informed consent prior to their inclusion in this study.

Consent for publication

Not applicable

Availability of data and materials

Data will be available upon request to the first author ZS.

Competing interests

The authors declare that they have no competing interests.

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Authors’ contributions

ZS and WQ conceived the original study and developed the protocol together with WS, Statistical advice was provided by XS and YJ. ZS wrote the manuscript. All authors read and approved the final manuscript.

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Tables

Table 1 Demographics and comorbidity of patients and information on primary intervention
| Cases | Level(s) | Comorbidities | Type of intervention | CRP (mg/L) | ESR (mm/h) | WBC (10^9/L) | VAS | Neu (%) | Biopsy |
|-------|----------|---------------|---------------------|------------|------------|--------------|-----|---------|--------|
| 1     | T9,L2    | COPD, DM      | KP                  | 7.2        | 28         | 7.6          | 6   | 49      | Negative |
| 2     | L3       | HTN, IHD      | VP                  | NA         | NA         | NA           | NA  | NA      | NA      |
| 3     | T10,T11,T12 | HTN, CIS    | KP                  | 10.7       | 46         | 6.4          | 7   | 62      | NA      |
| 4     | L4       | HF, Af        | VP                  | 8.3        | 19         | 6.7          | 5   | 63      | Chronic inflammation |
| 5     | L2       | HTN, DM, UTI  | KP                  | 21.3       | 44         | 7.8          | 6   | 74      | NA      |
| 6     | L1,L2    | DM, CIS       | KP                  | 17.6       | 36         | 8.2          | 6   | 68      | NA      |
| 7     | T11      | Malnutrition, RI | KP              | 11.4       | 29         | 5.4          | 5   | 59      | NA      |
| 8     | T10,T11  | Obesity, HTN  | KP                  | 13.6       | 33         | 5.9          | 4   | 62      | NA      |
| 9     | L3,L4    | COPD, CIS     | VP                  | NA         | NA         | NA           | NA  | NA      | NA      |
| 10    | T8,T12   | DM, Malnutrition | KP               | 8.3        | 16         | 6.2          | 6   | 63      | Negative |
| 11    | L1       | None          | KP                  | 27.9       | 67         | 4.6          | 5   | 41      | Negative |
| 12    | T12      | SLE           | KP                  | 24.4       | 58         | 5.0          | 6   | 43      | NA      |
| 13    | L1,L2    | PCa (chemotherapy) | VP               | 19.6       | 41         | 3.9          | 5   | 39      | NA      |
| 14    | T11      | Obesity, CIS  | VP                  | 10.0       | 21         | 6.5          | 5   | 60      | Negative |
| 15    | T10,L1   | HTN, CIS      | KP                  | 8.1        | 18         | 7.1          | 6   | 67      | Negative |
| 16    | T11,T12  | UTI, DM       | VP                  | 20.4       | 39         | 8.5          | 6   | 79      | NA      |
| 17    | L1       | DM, HTN       | VP                  | 11.3       | 20         | 8.2          | 6   | 69      | NA      |
| 18    | L3,L4    | COPD, RI      | KP                  | 8.3        | 17         | 7.1          | 4   | 63      | NA      |
| 19    | T12,L1,L2 | Malnutrition, Decubitus | KP               | 7.0        | 42         | 8.0          | 6   | 70      | NA      |

CRR: C-reactive protein; ESR: erythrocyte sedimentation rate; WBC: white blood cell; VAS: visual analog scale; Neu: neutrophil ratio; COPD: chronic obstructive pulmonary disease; DM: diabetes mellitus; KP: Kyphoplasty; VP: vertebroplasty; HTN: hypertension; IHD: ischemic heart disease; NA: not available; CIS: cerebral ischemic stroke; HF: heart failure; Af: atrial fibrillation; UTI: urinary tract infection; RI: renal insufficiency; SLE: systemic lupus erythematosus; PCa: prostatic cancer.

Table 2 Clinical, Laboratory, preoperative data of reoperation.
Cases | Reoperation after infection(d) | Fever/Chill | Neurologic deficits | CRP (mg/L) | ESR (mm/h) | WBC (10^9/L) | VAS | Neu (%)
---|---|---|---|---|---|---|---|---
1 | 67 | + | None | 45.7 | 62 | 11.4 | 7 | 77
2 | 103 | + | None | 81.3 | 59 | 9.8 | 7 | 81
3 | 14 | - | None | 109.7 | 48 | 13.5 | 8 | 89
4 | 88 | - | Paraparesis | 35.6 | 109 | 10.9 | 9 | 82
5 | 97* | - | Radiculopathy | 40.2 | 77 | 9.6 | 9 | 79
6 | 135 | + | Paraparesis | 18.9 | 68 | 11.1 | 7 | 80
7 | 373 | + | None | 22.7 | 60 | 9.0 | 6 | 69
8 | 26* | - | Paraparesis | 173.7 | 114 | 15.2 | 7 | 91
9 | 54 | - | None | 40.9 | 95 | 8.3 | 9 | 83
10 | 61 | + | None | 73.8 | 87 | 10.5 | 8 | 74
11 | 108 | - | None | 36.9 | 124 | 11.7 | 7 | 79
12 | 157 | + | Radiculopathy | 18.7 | 108 | 9.5 | 6 | 71
13 | 19 | - | None | 142.3 | 77 | 14.6 | 7 | 88
14 | 182* | - | Paraparesis | 40.6 | 84 | 10.8 | 9 | 82
15 | 71 | + | None | 87.7 | 79 | 12.3 | 8 | 87
16 | 114 | - | None | 39.0 | 56 | 10.4 | 7 | 70
17 | 93 | + | Paraparesis | 45.5 | 38 | 7.9 | 6 | 69
18 | 21 | + | None | 109.0 | 103 | 10.8 | 10 | 78
19 | 49 | + | None | 92.5 | 81 | 14.1 | 8 | 89

*Conservative treatment; □Abandoning therapy; CRP,C-reactive protein; ESR, erythrocyte sedimentation rate; WBC, white blood count; VAS, visual analog scale; Neu, neutrophil ratio.

Table 3 Operative and postoperative data
| Cases | Treatment method | Operative time(min) | Blood loss(ml) | Causative | Follow-up(month) | Outcome   | VAS follow-up  |
|-------|------------------|---------------------|----------------|-----------|------------------|-----------|---------------|
| 1     | Operation        | 165                 | 500            | No organism | 9.2              | Normal    | 2             |
| 2     | Operation        | 190                 | 400            | Enterococcus faecalis | 5.3          | Normal    | 1             |
| 3     | Operation        | 185                 | 350            | Non-tuberculous mycobacterium | Died           | NA        | NA            |
| 4     | Operation        | 210                 | 450            | Micromonomonas | 17.5          | Wheelchair | 2             |
| 5     | Conservative treatment | NA         | NA             | NA        | Died            | NA        | NA            |
| 6     | Operation        | 180                 | 400            | No organism | 23.5            | Use a walker | 3             |
| 7     | Operation        | 170                 | 300            | MRSE      | 8.5              | Normal    | 2             |
| 8     | None             | NA                  | NA             | NA        | Died            | NA        | NA            |
| 9     | Operation        | 160                 | 350            | Tread fungus | 33.5          | Use a walker | 1             |
| 10    | Operation        | 160                 | 300            | S.albus   | 23.0             | Normal    | 1             |
| 11    | Operation        | 210                 | 650            | Streptococcus | 12.0          | Normal    | 2             |
| 12    | Operation        | 155                 | 300            | No organism | 7.5             | Wheelchair | 2             |
| 13    | Operation        | 165                 | 400            | S.aureus  | 15.5             | Died(cancer) | NA            |
| 14    | Conservative treatment | NA         | NA             | NA        | 24.0             | Use a walker | 3             |
| 15    | Operation        | 180                 | 900            | Enterococcus faecalis | 3.0          | Normal    | 1             |
| 16    | Operation        | 160                 | 700            | S.aureus  | NA               | NA        | NA            |
| 17    | Operation        | 165                 | 400            | No organism | 8.0             | Normal    | 2             |
| 18    | Operation        | 170                 | 500            | Streptococcus | NA            | NA        | NA            |
| 19    | Operation        | 175                 | 550            | Escherichia coli | 10.0         | Normal    | 1             |

VAS, visual analog scale; NA, not available; MRSE, methicillin resistant Staphylococcus epidermidis; S.albus, Staphylococcus albus; S.aureus, Staphylococcus aureus.