Surgical Management of Deep Brain Stimulator Infection without Electrode Removal: Report of Two Cases

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Abstract

Objective Stimulation of the subthalamic nucleus by implanted electrodes (deep brain stimulation [DBS]) is performed to suppress symptoms of Parkinson's disease. However, postoperative wound dehiscence and infection can require removal of the implanted electrode leads. This report describes treatment of intractable unilateral wound infection in two patients without removing the DBS device.

Methods First, components of the DBS system were removed except for the electrode lead and thorough debridement of the infected wound was conducted. Second, the edges of the bone defect left by removal of DBS components were smoothed to eliminate dead space. Subsequently, the electrode lead was covered by using a pericranial-frontalis-muscle flap or a bi-pedicled-scalp flap with good blood supply. Closed intrawound continuous negative pressure and irrigation treatment was conducted for 1 week after the surgery, and then the drain was removed.

Results We treated two patients with wound infection after implantation of DBS electrodes. Case 1 developed a cutaneous fistula and Case 2 had wound dehiscence. After treatment by the method described above, complete wound healing was achieved in both patients.

Conclusion DBS is always associated with a risk of infection or exposure of components and treatment can be very difficult. We successfully managed intractable wound infection while leaving the electrode lead in situ, so that it was subsequently possible to continue DBS for Parkinson's disease.

Keywords ► deep brain stimulation
► electrode lead
► infection
► intractable wound
► cutaneous fistula
► intrawound continuous negative pressure and irrigation therapy
► reconstruction
► pericranial flap
► scalp flap

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Introduction

Deep brain stimulation (DBS) therapy involves electrical stimulation of the subthalamic nucleus and the globus pallidus, and it is widely used for Parkinson’s disease. However, wound infection and dehiscence may occur in rare cases, and such infection can be very difficult to treat, with removal of the electrode leads being necessary. In 2006, we reported the usefulness of a pericranial flap for reconstruction of the dura mater. In 2007, intrawound continuous negative pressure and irrigation treatment (IW-CONPIT) was reported, after which this method has been applied to treat infected and/or intractable wounds at various sites. In 2016, the efficacy of IW-CONPIT using a closed system was reported for controlling infection of all layers of the cranium combined with cerebrospinal fluid leakage. We applied these methods (pericranial flap reconstruction and IW-CONPIT) in two patients with wound infection after implantation of DBS systems (Medtronic Inc., Minneapolis, Minnesota, United States), allowing infection to be managed without removal of the electrode leads so that DBS therapy could subsequently be continued.

Case 1

History and Presentation

A 61-year-old woman developed a cutaneous fistula at 6 months after surgery for placement of DBS electrodes to treat Parkinson’s disease at the Neurosurgery Department of another hospital. The patient was referred to our department because the fistula persisted. At initial examination, a cutaneous fistula was seen in a scar on the forehead. Drainage of pus was observed, and a silicon cap for fixing the electrode was visible inside the fistula.

Operation

After re-opening the wound via the previous incision line, infected granulation tissue was detected around the silicon cap and along the electrode line. The silicon cap was removed, but the electrode lead was left in situ to continue DBS therapy for Parkinson’s disease. Debridement of the infected granulation tissue was performed and the edges of the bone defect left after component removal were smoothed. A pedicled pericranial frontalis muscle flap was harvested from the parietal region on the healthy (right) side, and was used to cover the electrode lead and fill the dead space in the bone defect. The edges of the debrided fistula showed very poor circulation. Because the pericranial flap was placed under the fistula, there was no bone exposure; hence, healing by secondary intention was deemed possible and the fistula was left open without suturing. To prevent postoperative infection, two drains were placed between the pericranial flap and the bone, and closed IW-CONPIT was initiated with saline at 2,000 mL/d after conducting IW-CONPIT for 1 week, the drains were removed since there were no signs of infection.

Postoperative Course

The postoperative course was good and the fistula showed epithelialization by 2 weeks after surgery. There has been no relapse of the fistula or infection during follow-up for 1.5 years. In addition, a depression has not developed at the previous site of the fistula or at the frontalis muscle donor site, with the cosmetic outcome being satisfactory.

Case 2

History and Presentation

A 67-year-old woman with Parkinson’s disease developed wound dehiscence at 4 months after surgery for implantation of DBS electrodes at another hospital. At initial examination, drainage of pus from a cutaneous fistula and exposure of a silicon cap were observed.

Operation

A fusiform skin incision was made to resect the cutaneous fistula, revealing that the old surgical wound was filled with infected granulation tissue. As was done for Case 1, the cap was removed while retaining the electrode lead. Debridement of infected granulation tissue was performed and the edges of the bone defect were smoothed. Then the wound was closed by using a bi-pedical flap raised from the right side, and was continued for 1 week, following which the drains were removed because there were no signs of infection.

Table 1 Summary of patient characteristics and surgical interventions

<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Age (y/Sex)</th>
<th>Time to fistula, (mo)</th>
<th>Infectious agent</th>
<th>Primary intervention</th>
<th>Secondary intervention</th>
<th>Duration of continuous irrigation and negative pressure</th>
<th>Follow-up</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>61/F</td>
<td>6</td>
<td>MRSA</td>
<td>Scalp undermining and pericranial flap</td>
<td>None</td>
<td>7 d</td>
<td>18 mo, no recurrence</td>
</tr>
<tr>
<td>2</td>
<td>67/F</td>
<td>4</td>
<td>MRSA</td>
<td>Bi-pedical scalp flap, right-thigh split-thickness skin graft</td>
<td>None</td>
<td>7 d</td>
<td>19 mo, no recurrence</td>
</tr>
</tbody>
</table>

Abbreviation: MRSA, methicillin-resistant Staphylococcus aureus.
Postoperative Course
At 1 year and 7 months after wound healing, there has been no recurrence of the fistula or infection (Fig. 2J–L). In addition, DBS therapy has been continued for Parkinson’s disease.

Discussion
Parkinson’s disease is caused by degeneration of dopaminergic cells in the substantia nigra of the midbrain, with the consequent lack of dopaminergic stimulation causing motor symptoms such as tremor, muscle rigidity, and akinesia. Anticholinergics and levodopa preparations are used as standard medical treatment, but control of symptoms such as wearing-off and dyskinesia is problematic. DBS therapy was reported by Benabid et al in 1987, and it is a treatment for various neurological diseases that involves electrical stimulation of the globus pallidus, thalamus, or subthalamic nucleus via implanted electrodes. In patients with Parkinson’s disease, DBS therapy can be effective for akinesia and rigidity, as well as for wearing-off symptoms, and it is currently the mainstream surgical treatment for this disease. However, the risk of
infection and exposure always exists when devices are implanted surgically. In principle, infections that do not respond to conservative treatment require removal of the implanted device, making it impossible to continue DBS therapy.

We encountered three problems when managing the patients presented here. First, wound healing was prevented by the presence of the implanted devices and infected granulation tissue. Second, there was a risk of creating dead space under the scalp if the wound was simply closed after debridement of granulation tissue. Third, it was important to retain the electrode leads for future performance of DBS, despite the intractable wound infection.

These three problems were managed as follows: First, the nonessential parts of the DBS system were removed and thorough debridement was performed. Second, the edges of the bone defect were smoothed to promote adherence of the skin flap. Then we covered the electrode lead by using a local flap such as a pericranial flap or bi-pedicled scalp flap with good blood supply. Flaps such as galeal, temporalis fascial, or pericranial flaps are reliable, thin, and supple, as well as having a good arc of rotation and minimal donor site morbidity. These flaps can also be raised in the vicinity of the operative field, which is convenient for the neurosurgeon. In Case 1, we used a thin and flexible pericranial flap to fill the dead space while covering the electrode lead. In Case 2, a bi-pedicled flap with good mobility was transferred freely to broadly cover the bone defect without suturing under tension. Finally, postoperative closed IW-CONPIT was performed with a high volume of saline for infection control. In 1997, vacuum-assisted closure was reported as an effective method for increasing granulation tissue formation in subacute and chronic wounds. However, this method has no suppressive effect on wound infection. On the other hand, IW-CONPIT is designed for infected wounds and can be used to treat intractable wound infection at various sites. The key points of IW-CONPIT are reducing bacteria in the wound by continuous irrigation for 24 hours per day and elimination of dead space by negative pressure. IW-CONPIT can be supplemented by flap transplantation. In our patients, adhesion between a local flap with a good blood supply and the tissues around the fistula was...
promoted by maintenance of negative pressure using IW-CONFIT, resulting in control of wound infection.

If infection occurs following implantation of a DBS device, taking these measures can make it possible to eliminate wound infection while retaining the electrode lead in situ, allowing DBS therapy to be continued.

Conflict of Interest
None.

References