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Short communication

Outcomes, management, and potential mechanisms of interleaving deep brain stimulation settings



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A R T I C L E I N F O

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ABSTRACT

Introduction: DBS is a therapeutic option for patients with Parkinson disease (PD), tremor and dystonia. In patients who experience suboptimal clinical results with conventional programming (monopolar, double monopolar or bipolar settings), interleaved pulses can sometimes be used to provide differential therapeutic benefits with the possibility of fewer side effects. Interleaving allows a clinician to define two "programs" that automatically alternate. The goal of this paper is to 1) present clinical scenarios where DBS interleaving was used across two clinics to provide improved symptom control in three patients with suboptimal results from conventional programming; 2) address the potential mechanisms of interleaving; and 3) provide practical tips on the use of interleaving.

Methods: Three patients were formally compared for therapeutic benefit on interleaved and conventional parameter settings.

Results: Interleaving is most likely to be useful in two clinical scenarios: 1) different contacts are beneficial for specific symptoms, but each at a different stimulation amplitude; or 2) symptoms are resolved incompletely, and further voltage increase is limited by side effects. The factors underpinning the differences in outcomes with interleaving are unknown but may be highly dependent on specific symptoms and to electrode positioning. Interleaving is a relatively new programming platform and there is no data to demonstrate long-term benefits.

Conclusions: Interleaving is a tool that may augment outcomes, and possibly obviate the need for surgical revisions, although in our experience across two large centers it has been effective for only a small number of patients.

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1. Introduction

Chronic deep brain stimulation (DBS) therapy emerged in the late 1980's for the treatment of tremor and in the early 1990's for advanced Parkinson disease (PD). Although DBS technology has not changed fundamentally since its inception, the latest commercially available neurostimulator upgrades of Activa PC, Activa RC and Activa SC (Medtronic, Inc., Minneapolis, MN) offer increased programming functionality with an interleaving feature.

Interleaving refers to activating two stimulation programs that rapidly alternate from pulse to pulse between each other (Fig. 1). Each of the two interleaving programs can specify the active electrode contact(s), pulse width and pulse amplitude. The frequency for each program must be the same and it is limited by the device up to 125 Hz. One or both electrodes can be interleaved, but no more than two programs can be used for each electrode [1].

Interleaving has been suggested as a strategy when conventional programming techniques fail to achieve desired results. Undesirable results may or may not be due to factors such as suboptimal positioning of the lead. When clinicians struggle to program a DBS device they often employ various additional techniques including bipolar settings, double monopolar or tripolar settings, and alternative pulse widths and frequencies to manipulate the electrical field. Interleaving provides an alternative method to alter the shape of the electrical fields, and possibly maximize benefits. The purpose of this paper is to explain the concept of interleaving, hypothesize on mechanism(s) of action, illustrate its use in three patients, and offer practical tips for its use.

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2. Methods

Three patients with PD were implanted with bilateral subthalamic (STN) DBS electrodes at two expert DBS institutions. All three patients underwent standard clinical programming, but achieved suboptimal results with conventional techniques. They maintained stable clinical benefit for at least 3 months on the interleaved settings. In all patients, interleaving was used in only one hemisphere. UPDRS examination (off medications) was performed for seven stimulation conditions detailed in Table 1, in a single session, in a non-blinded, non-randomized manner, allowing at least 5 min for each stimulation setting to take effect and 30 min for the initial stimulation washout. The aim was to compare stimulation effects using the same parameters, but different electrode configurations.

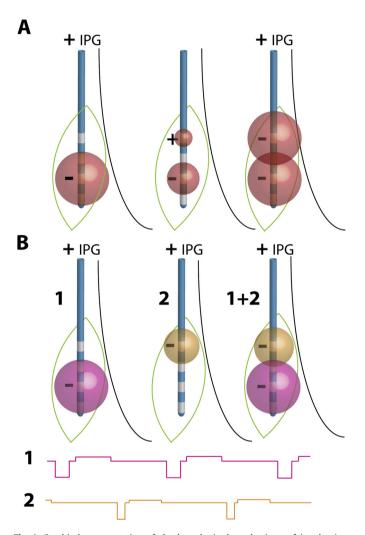


Fig. 1. Graphical representation of the hypothesized mechanisms of interleaving. Spheres represent estimated volume of tissue activated during various DBS programming settings [8]. In this example green outline represents subthalamic nucleus (STN) and black curve is the corticospinal tract. Top section (A) demonstrates conventional settings: monopolar stimulation at contact 1 (left) does not stimulate sufficiently large area of the nucleus and does not provide complete resolution of symptoms. Changing to bipolar setting (middle) provides some additional activation around contact 3, but it is overall insufficient because bipolar stimulation results in a narrower field. Double monopolar (right) provides sufficient coverage of the nucleus but also stimulates corticospinal tract causing side effects. Bottom section (B) illustrates interleaved programming: user specifies two programs which are applied interchangeably. Bottom tracings show shape and timing of the stimulation pulses applied to each contact (large negative pulse activates neurons, while small positive afterpulse recovers charge). For each program, user defines specific contacts, amplitude and pulse width, although frequency is limited up to 125 Hz by the device. This allows greater freedom in selecting different stimulation settings for the two contacts while still benefiting from monopolar field. In this example, interleaving provides sufficient stimulation of the nucleus while avoiding the corticospinal tract. Small overlap area experiences stimulation at twice the frequency since it is seeing pulses from both programs, but clinical utility of this phenomenon is unknown [2].

3. Results

3.1. Case 1

A 61 year-old man with tremor-predominant PD for 11 years suffered from dyskinesia and motor fluctuations with off periods characterized by tremor, marked deterioration of speech and slowing of gait. Bilateral STN DBS was performed with resultant suboptimal tremor control despite multiple programming sessions. On the post-operative MRI, the left electrode was found to be medial to the STN (contact 2 was 7 mm lateral, 1 mm posterior and 1 mm ventral to the midcommisural point (MCP)). The left STN single monopolar setting only partially improved the tremor. Double monopolar setting at the amplitude and pulse width necessary to control tremor could not be tolerated due to diplopia. Interleaving allowed a dorsal contact to be set to higher amplitude and pulse width, while the contact below was set to a tolerable lower amplitude and pulse width. Together, the two contacts provided complete tremor resolution without side effects.

3.2. Case 2

A 65 year-old man with PD for 11 years had tremor, rigidity and bradykinesia that was significantly worse on the left side. He was implanted with bilateral STN DBS due to frequent wearing off and tremor that did not completely respond to high doses of levodopa. On post-operative MRI, his right DBS lead was medial and posterior to the optimal target in the STN (contact 10 was 10.5 mm lateral. 4.5 mm posterior and 4 mm ventral to the MCP). The right STN single monopolar stimulation did not provide sufficient tremor and/or bradykinesia improvement. Double monopolar settings at 90 µs resulted in diplopia and paresthesias which limited further increases in parameters. A double monopolar setting at 60 µs and a slightly higher amplitude improved tremor, but did not further improve bradykinesia. Interleaved settings with similar amplitude and pulse width at the two contacts improved both tremor and bradykinesia, in a seemingly synergistic manner beyond what either double monopolar setting could achieve. The patient did well on this setting for 4 months, but later reported dysarthria which resolved when switched off the interleaved setting, and he therefore was programmed back to a double monopolar setting at 3.0 V and 60 µs which improved his speech but worsened other symptoms. The patient had also been receiving vocal fold injection augmentation for dysphonia, and his last injection was 9 months prior to the reemergence of dysarthria which may have biased his overall outcome.

3.3. Case 3

A 58 year-old man with PD for 16 years had symptoms including stiffness, dystonia and tremor. Over time he developed dyskinesia, wearing off and frequent levodopa dose failures. He underwent staged bilateral STN DBS. On post-operative CT/MRI fusion, his right DBS lead was well placed in the STN (contact 2 was 13 mm lateral, 0 mm posterior and 1 mm ventral to the MCP). The right STN single monopolar programming at contact 2 improved tremor and caused only rare dyskinesias, but worsened nighttime rigidity, which resulted in a reported sleep quality deterioration. Single monopolar DBS at contact 1 improved sleep by decreasing stiffness, but increased daytime dyskinesia. Dyskinesias were present even with elimination of entacapone and controlled release levodopa. Interleaved settings using contacts 1 and 2 decreased daytime dyskinesias and improved nighttime mobility and sleep by patient report. Clinically, contact 2 had an antidykinetic effect even though it was not located in the zona incerta where this can be expected.

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