

Beaumont Health

Beaumont Health Scholarly Works and Archives

Conference Presentation Abstracts

Neurosurgery

12-5-2022

Responsive Neurostimulation (RNS) of the Centromedian Nucleus of the Thalamus for the Treatment of Drug Resistant Idiopathic Generalized Epilepsy

Angelique Manasseh

Hannah Guider

Brooklynn Bondy

Ayman Haykal

Nour Baki

See next page for additional authors

Follow this and additional works at: https://scholarlyworks.beaumont.org/neurosurgery_confabstract



Part of the [Neurology Commons](#), and the [Surgery Commons](#)

Authors

Angelique Manasseh, Hannah Guider, Brooklynn Bondy, Ayman Haykal, Nour Baki, Ashleigh Terrell, Michael Staudt, Christopher Parres, and Andrew Zillgitt

Abstracts

Responsive Neurostimulation (RNS) of the Centromedian Nucleus of the Thalamus for the Treatment of Drug Resistant Idiopathic Generalized Epilepsy

Abstract number : 3.179

Submission category : 3. Neurophysiology / 3E. Brain Stimulation

Year : 2022

Submission ID : 2204627

Source : www.aesnet.org

Presentation date : 12/5/2022 12:00:00 PM

Published date : Nov 22, 2022, 05:25 AM

Authors :

Angelique Manasseh, DO – Beaumont Health; Hannah Guider, DO – Neurology – Beaumont Health; Brooklynn Bondy, DO – Neurology – Beaumont Health; Sanjay Patra, MD – Neurosurgery – Spectrum Health; David Burdette, MD – Neurology – Spectrum Health; Ayman Haykal, MD – Neurology – Beaumont Health; Nour Baki, M – Neurology – Beaumont Health; Ashleigh Terrell, NP-C – Neurology – Beaumont Health; Michael Staudt, MD – Neurosurgery – Beaumont Health; Christopher Parres, MD – Neurology – Beaumont Health; Andrew Zillgitt, DO – Neurology – Beaumont Health

Rationale: Idiopathic generalized epilepsy (IGE) accounts for approximately one-third of all epilepsies. The majority of people with IGE achieve seizure-freedom from treatment with anti-seizure

medications (ASM). However, up to 25% of people with IGE continue to experience seizures despite adequate ASM trials. In patients with drug resistant IGE, neuromodulation with responsive neurostimulation (RNS) in the bilateral centromedian nucleus (CMN) of the thalamus may be an effective treatment option.

Methods: Five patients with drug resistant IGE were evaluated through the Beaumont Adult Comprehensive Epilepsy Center and underwent an epilepsy presurgical evaluation including video-EEG monitoring in the epilepsy monitoring unit, brain MRI, and neuropsychology evaluation. Cases were presented individually at the weekly multidisciplinary patient management conference, and a consensus was made to offer RNS implantation within the bilateral CMN of the thalamus. Four patients underwent RNS implantation through Spectrum Health, while 1 patient was implanted with RNS at Beaumont Health. Following RNS implantation, electrocorticography (ECoG) data were reviewed by the treating epileptologists, and adjustments to detection and therapy settings were implemented to optimize seizure control. Retrospective review was then completed for patient seizure outcomes. Quality of life (QoL) was assessed at clinic visits with a subjective questionnaire in which patients were asked if their QoL was much better, better, same, worse, or much worse after RNS implantation.

Results: Patient demographics are presented in Table 1. The average age of epilepsy onset was 13 years old (range, 2-38 years old) and the average at RNS implantation was 36.6 years old (range, 25-51 years old). The average number of ASM trials was 6 (range, 2-10). At the time of implantation, patients were taking, on average, 3 ASM (range, 2-4). The average follow-up after RNS implantation was 18 months (range, 8-32 months). There were no operative or perioperative complications. At last follow-up, all 5 patients reported at least a 75% reduction in the frequency of disabling seizures (Table 2). Two patients remained free of GTCS following RNS implantation. On a subjective QoL questionnaire, all patients reported QoL was “much better” or “better” after RNS implantation.

Conclusions: Drug resistant IGE is common and, in the past, surgical treatment options have been limited. In this case series, 5 patients with IGE underwent RNS implantation within the bilateral CMN of the thalamus. There were no operative or perioperative complications. All 5 patients experienced at least a 75% reduction in the frequency of disabling seizures, and 2 patients remained free of GTCS following RNS implantation. In addition, QoL was improved after RNS implantation. Responsive neurostimulation within the bilateral CMN of the thalamus may be a safe and efficacious treatment option for people with drug resistant IGE.

Funding: Not applicable
Neurophysiology

Table 1. Patient demographics

| Patient | Sex | Age | Age at onset | Onset | PHs | Initial Seizure | Seizure Types | ERL | RHI | RCS |
|---------|-----|-----|--------------|-------|-----|-----------------|-----------------|---------------------|-----|-------------|
| 1 | F | 36 | 2 | Y | M | FE, MSB | ABS, JMs, MTCs | 2 Hr P5W, 5W, 6W | NL | JME |
| 2 | F | 25 | 5 | N | M | ABS | ABS, JMs, CTCS | 2-4 Hr P5W-SW | NL | CAZEME |
| 3 | F | 23 | 5 | N | Y | AJG | AJG, CTCS | 2-4 Hr P5W-SW, CTEs | NL | FR-LEN, JAG |
| 4 | M | 31 | 28 | N | Y | GTCS | GTCS (JAG) | 4-8 Hr P5W-SW, CTEs | NL | BUJTS |
| 5 | F | 22 | 18 | N | M | ABS | AJG, CTCS (JAG) | 2-4 Hr P5W-SW | NL | JAG + JME |

Abbreviations: ABS, Absence; AJG, Atonic Jerk; CAZEME, Childhood Absence Epilepsy; CTEs, Childhood Temporal Epilepsy; CTCS, Childhood Complex Tonic Clonic Seizure; ERL, Epilepsy Response Latency; ERG, Epilepsy Response Graph; FE, Focal Epilepsy; GTCS, Generalized Tonic Clonic Seizure; JAG, Juvenile Absence Epilepsy; JME, Juvenile Myoclonic Epilepsy; JMs, Juvenile Myoclonic Seizures; LCM, Lennox-Gastaut Syndrome; MTCs, Myoclonic Tonic Clonic Seizures; MSB, Myoclonic Seizure; NL, No Lesion; PHs, Perinatal Hypoxia; P5W, Postnatal 5 Weeks; P5W-SW, Postnatal 5 Weeks - Seizure Worsening; RCS, Response to Carbamazepine; SW, Seizure Worsening; Y, Young Onset; N, Not; M, Male; F, Female; Y, Yes; N, No.

Table 2. Patient outcomes

| Patient | Previous ASM | Current ASM | Follow-up | Outcome |
|---------|---|--------------------|-----------|--|
| 1 | CGE, TPM, LGS, ZNS, LCM, LGS, RGS, CBZ, LEV | CGE, RGS, LGS, VPA | 22 months | 75-80% reduction in disabling seizures |
| 2 | LGS, LEV | LTC, ZNS | 24 months | 96-97% reduction in disabling seizures |
| 3 | VPA, LGS, LEV, TPM, PEG, LGS | REX, LGS, ESP, ZNS | 15 months | Seizure free (100% reduction) |
| 4 | VPA, TPM, RGS, LCM | CGE, LGS, LEV | 11 months | 95-97% reduction in disabling seizures |
| 5 | CGE, VPA, LEV, TPM, LGS, LGS, RGS, CGE, LCM | LCM, TPM, LGS | 5 months | 75-80% reduction in disabling seizures |

Abbreviations: ASM, Antiepileptic Medication; CBZ, Carbamazepine; CGE, Childhood Generalized Epilepsy; CTEs, Childhood Temporal Epilepsy; CTCS, Childhood Complex Tonic Clonic Seizure; ERG, Epilepsy Response Graph; FE, Focal Epilepsy; GTCS, Generalized Tonic Clonic Seizure; JAG, Juvenile Absence Epilepsy; JME, Juvenile Myoclonic Epilepsy; JMs, Juvenile Myoclonic Seizures; LCM, Lennox-Gastaut Syndrome; MTCs, Myoclonic Tonic Clonic Seizures; MSB, Myoclonic Seizure; NL, No Lesion; PHs, Perinatal Hypoxia; P5W, Postnatal 5 Weeks; P5W-SW, Postnatal 5 Weeks - Seizure Worsening; RCS, Response to Carbamazepine; SW, Seizure Worsening; Y, Young Onset; N, Not; M, Male; F, Female; Y, Yes; N, No.