Long-Term Outcomes of Occipital Nerve Stimulation for New Daily Persistent Headache With Migrainous Features

Susie Lagrata, MSc¹; Sanjay Cheema, MBBS, MRCP¹; Laurence Watkins, PhD²; Manjit Matharu, PhD¹

ABSTRACT

Objectives: New daily persistent headache (NDPH) is a subset of chronic headache where the pain is continuous from onset. Phenotypically it has chronic migraine or chronic tension type features. NDPH is considered to be highly refractory. Occipital nerve stimulation (ONS) has been used for treatment of refractory chronic migraine but there are no specific reports of its use for NDPH with migrainous features.

Materials and Methods: Nine patients with NDPH with migrainous features were identified as having had ONS implants between 2007 and 2014 in a specialist unit with experience of using ONS in chronic migraine. Moderate to severe headache days were compared at baseline and follow-up. A positive response was defined as at least 30% reduction in monthly moderate to severe headache days.

Results: Patients had suffered NDPH for a median of 8 years (range 3–16 years) and had failed a median of 11 previous treatments (range 8–15). After a median follow-up of 53 months (range 27–108 months), only a single patient showed a positive response to ONS. At no point did the cohort as a whole show any change in monthly moderate to severe headache days or disability scores.

Conclusion: Our experience suggests that ONS is not effective in the treatment of NDPH with migrainous features even in centers with experience in treating chronic migraine with ONS. The difference in response rates of chronic migraine and NDPH with migrainous features supports the concept of a different pathophysiology to the two conditions.

Keywords: Chronic headache, chronic migraine, new daily persistent headache, occipital nerve stimulation

Conflict of Interest: Laurence Watkins has served on advisory boards for St Jude Medical and Medtronic. Susie Lagrata and Sanjay Cheema have no competing interests. Manjit Matharu serves on the advisory board for Allergan, Novartis, Lilly, and Medtronic and has received payment for the development of educational presentations from Allergan, Abbott, Medtronic, and electroCore.

INTRODUCTION

New daily persistent headache (NDPH) is a headache characterized by the acute onset of head pain that occurs daily from onset and which presents in patients without an increasing frequency of pre-existing headaches. It was first described by Vanast in 1986 as a chronic benign headache with clinical features similar to chronic tension headache where most patients would recover spontaneously within six months (1). Since this original description, multiple reports have described large cohorts of NDPH patients with migrainous features. This has led the International Classification of Headache Disorders version 3 (ICHD 3) listing NDPH as a persistent headache, daily from onset which may be migraine-like, tension-type-like, or have features of both (2). Similarly, subsequent to the original description as a transient condition, multiple case series have recognized NDPH as a prolonged and treatment-resistant headache, especially in those with the migrainous phenotype (3–6).

Occipital nerve stimulation (ONS) is a form of peripheral nerve stimulation targeted at the occipital nerves (predominantly the greater occipital nerve) and involves the implantation of

suboccipital electrodes attached to an implantable pulse generator (IPG). The procedure is performed for the preventative treatment of refractory primary headaches such as chronic migraine

Address correspondence to: Manjit Matharu, PhD, Headache and Facial Pain Group, University College London (UCL) Queen Square Institute of Neurology and The National Hospital for Neurology and Neurosurgery, Queen Square, London WC1N 3BG, UK. Email: m.matharu@uclmail.net

¹ Headache and Facial Pain Group, UCL Queen Square Institute of Neurology and The National Hospital for Neurology and Neurosurgery, London, UK; and

² Department of Neurosurgery, UCL Queen Square Institute of Neurology and The National Hospital for Neurology and Neurosurgery, London, UK

For more information on author guidelines, an explanation of our peer review process, and conflict of interest informed consent policies, please go to <http://www.wiley.com/WileyCDA/Section/id-301854.html>

Source(s) of financial support: There was no funding or financial support for this study.

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made.

and chronic cluster headache. There is currently only a limited amount of controlled evidence to support the use of ONS in refractory chronic migraine although case series and systemic reviews report more promising results (7–11). Our specialist headache unit has previously reported a series of 53 patients with refractory chronic migraine with an average response rate of 45.3% at final follow-up (12). Despite the use of ONS for refractory chronic migraine, there are no specific reports in any of the above publications for the use of ONS in NDPH. This cohort therefore presents the long-term follow-up of nine patients with NDPH with migrainous features with ONS implants.

MATERIALS AND METHODS

Patients

All patients were treated by a single specialist team at the National Hospital for Neurology and Neurosurgery, Queen Square, London, UK. Implants took place between May 2007 and February 2014. Patients fulfilled the ICHD 3 diagnostic criteria for NDPH with migraine features (Box 1) (2). As intractability is not defined for NDPH, the recommendations of Goadsby et al. for the definition of intractable chronic migraine were applied (13).

Patients had been offered ONS as part of standard care for their refractory headache under the National Health Service (NHS). The treatment was carried out under the supervision of our hospital's Clinical Effectiveness Supervisory Group (CESG) and ethics approval for data collection and publication was granted by Northwick Park Hospital Research Ethics Committee, Hampstead, London, UK.

Surgical Procedure

All patients were implanted with bilateral suboccipital leads connected to an IPG. Seven patients were implanted with a Medtronic and two with Abbott (previously St Jude Medical) systems. Octad electrodes were implanted in all.

During surgery, patients were placed into the lateral position and a midline posterior cervical incision made. In earlier cases, the insertion point of the leads was the spinous process of C1,

passing laterally and superiorly, with the procedure conducted using a curved Tuohy needle and an image intensifier. In later cases, the implant level was superior to the nuchal line. In these cases, the lead was passed using a blunt plastic tube to reduce the likelihood of the tip of the lead being tunneled too close to the skin causing erosions. Changes in operative procedure were to reduce adverse events such as unwanted recruitment of neck muscles or erosion of the tip of the lead through the skin. Given that both techniques target the same nerve the implant technique alone would not account for any change in clinical response. Leads were anchored to cervical fascia and then tunneled to a lateral cervical or infraclavicular skin crease intermediate incision. An infraclavicular incision was made forming a pocket for the IPG, to which the distal end of the leads were attached. Silicone sheaths were used to protect lead connections. Topical gentamicin was introduced around the pocket and the incisions closed.

Trial stimulation was not conducted as our unit feels that current evidence does not yet support that the benefit of significantly improved response rates is outweighed by the risks of multiple surgical interventions.

Frequency was initially set at 70 Hz with a pulse width of 450 μ s. During follow-up, the polarity of the electrodes was adjusted to achieve bilateral paresthesia in the greater occipital region. Stimulation settings were recorded at each follow-up. Medications were changed as needed at the discretion of the treating physicians.

Data Collection

All data were collected prospectively and entered onto a clinical database (Microsoft Excel, Microsoft Corporation, Redmond, WA, USA). For the first-year, postimplant patients were seen in clinic every three months and after that every 6–12 months. Data were collected on patient demographics and medication history at time of implant. Throughout follow-up data were collected on moderate to severe headache days, daily pain score and duration, headache disability scores and quality of life scores, acute and preventative medication use, ONS settings, and adverse events.

Headache disability scores included the Migraine Disability Assessment Scores (MIDAS) and Headache Impact Test-6 Scores (HIT-6). These were recorded pre- and postimplant. Quality of life scores used were the Euro-QoL (Euro-QoL 5D index [EQ-5D] and Euro-QoL visual analogue score [EQ-VAS]), Short Form 36 Questionnaires (SF36) both physical (SF36-P) and mental (SF36-M) components, Beck Depression Inventory II (BDI-II), and Hospital Anxiety and Depression Scores (HAD-A and HAD-D). All scores were recorded before and after implant.

Adverse events were recorded as they occurred and categorized as either hardware, biological, or stimulation related. Hardware-related events were recorded if problems occurred involving the device components, biological events if reactions occurred to the device or related to the surgical procedure and stimulation related events recorded if problems were due to stimulation issues (i.e., painful stimulation) (11).

Primary outcome measure was the improvement in mean monthly moderate to severe headache days at final follow-up compared to baseline. A reduction of 30% or more was defined as a positive clinical response. This response rate was chosen in line with our previous paper on ONS in chronic migraine to allow meaningful comparison and as per the International Headache Society clinical trials subcommittee (12,14). Secondary outcome

Box 1. International Classification of Headache Disorders Version 3
Definition of New Daily Persistent Headache.

ICHD-3: New Daily Persistent Headache

Distinct and clearly remembered onset, with pain becoming continuous and unremitting within 24 hours

Present for >3 months

NDPH may have features of either

1. Migraine

- o Headache lasting 4–72 hours,
- o Headache having at least two of the following four characteristics: unilateral location, pulsing quality, moderate to severe intensity, aggravation by routine physical activity
- o During headache presence of at least one of following: nausea and/or vomiting, photophobia, and phonophobia

2. Tension-type headache

- o Headache lasting from 30 minutes to seven days
- o Headache having at least two of the four characteristics: bilateral location, pressing/tightening/nonpulsatile quality, mild–moderate intensity, not aggravated by routine physical activity
- o Both of the following: no nausea or vomiting, only one of photophobia or phonophobia

measures were changes in median daily pain intensity, median daily headache hours, and changes in disability and quality of life scores at final follow-up.

Statistical Analysis

SPSS 24.0 statistical software was used to analyze data. Continuous variables were expressed as median plus range and/or interquartile range (IQR). Categorical variables were summarized as percentages. Median values pre- and postimplant were compared using Wilcoxon signed rank tests. All tests were two-sided with a significance level of 95%.

RESULTS

Demographic Data

Between May 2007 and February 2014, nine patients with NDPH with migrainous features underwent ONS implants. Their demographic data are shown in Table 1. Median age at implant was 42.00 years (range 29–64, IQR 22.00) and four patients were female. The cohort had failed to respond to a median of 11 preventative medications prior to their implant (10–13, IQR 3.00) (Table 2). Two patients reported a previous positive response to a greater occipital nerve block (a more than 50% reduction in headache severity or frequency lasting at least two weeks).

Patient 9 had coexistent chronic cluster headache and NDPH with migraine features but as they were able to differentiate between their two headache types they kept separate diaries to allow us to monitor the response of each phenotype throughout the follow-up.

Patient 9 and patient 3 met the ICHD-3 criteria for the overuse of acute medication prior to implant. Patient 9 was using daily triptans for their coexistent cluster headache and so these could not be stopped. Patient 3 was using daily opiates but had undergone a supervised withdrawal prior to the ONS but failed to show any significant improvement when off of the opiates. This observation excludes medication overuse headache as a cause of symptoms and therefore the patient chose to continue on the painkillers.

Clinical Outcomes

Median follow-up time was 53.00 months (27–108 months, IQR 36.50). At final follow-up, there was zero change in the monthly moderate to severe headache days of the cohort ($p = 0.414$; Table 3). No significant change was seen in the median monthly moderate to severe headache days of the cohort at any time during the follow-up: three-month median change = 0, $p = 0.285$; six-month median change = 0, $p = 0.655$; nine-month median change = 0, $p = 0.144$; 12-month median change = 0, $p = 0.285$; 24-month median change = 0, $p = 0.144$; 36-month ($n = 7$) median change = 0, $p = 0.180$.

A statistically significant reduction was seen across the cohort in median daily pain intensity (–1.00 points on verbal rating scale [VRS], $p = 0.011$); however, this reduction is below the two-point reduction in VRS usually considered clinically significant. There was no significant change seen in median total headache days (median change = 0, $p = 1.00$) or median daily pain duration (median change = –5.00, $p = 1.00$) (Table 3). There was no significant change seen in the headache disability scores (MIDAS median change = –5.00, $p = 0.674$ and HIT-6 median change = 1, $p = 0.446$) and the only quality of life score to show any significant change was BDI-II with a median reduction of 3.00 points ($p = 0.008$) (Table 4).

A clinically significant reduction in moderate to severe headache days was seen in only one patient (patient 5). A positive response was observed within three months of implant and was maintained throughout follow-up while the device was operational. At 18–20 months postimplant, the patient's IPG failed. Battery failure was associated with an increase in headache frequency so that within four months he was back to baseline in terms of his monthly moderate to severe headache days. Following battery replacement, it took a further 6–8 months to regain the previous levels of benefit.

In the patient with coexistent chronic cluster headache and NDPH with migrainous features (patient 9) daily cluster attack frequency fell by more than 50% at final follow-up despite there being no response in their NDPH with migrainous features.

Table 1. Patient Demographics.

	Age (years)	Sex	Duration (years)	Laterality	Monthly moderate-to-severe HA days	Mean daily VRS	Medication overuse	Previous preventatives	Response GONB	Follow-up (months)
1	33	F	13	Bilateral	30	8	No	11	Negative	65
2	62	M	8	Bilateral	30	8	No	13	Negative	49
3	29	F	8	Unilateral	30	5	Yes	11	Negative	64
4	37	M	5	Bilateral	30	8	No	10	Negative	51
5	46	M	6	Unilateral	30	6	No	10	Positive	108
6	42	M	12	Bilateral	30	7	No	11	Negative	89
7	64	F	10	Bilateral	30	6	No	11	Negative	53
8	34	F	16	Bilateral	30	9	No	12	Negative	27
9	49	M	3	Unilateral	30	7	Yes*	12	Positive	32
Median	42	5M: 4F	8	6 Bilateral:	30	7.00	7 No:	11	7 Negative:	53.00
Range	29–64		3–16	3 Unilateral	(0)	5–9	2 Yes	10–13	2 Positive	27–108
(IQR)	(22)		(7)			(2.00)		(3.00)		(36.50)

*Patient using daily triptans for coexistent chronic cluster headache.

F, female; GONB, greater occipital nerve block; HA, headache; M, male; VRS, verbal rating scale.

Table 2. Previous Medications Used for Treatment of NDPH Prior to ONS.

	1	2	3	4	5	6	7	8	9	Median dose
Propranolol	DNK	80 mg	160 mg	-	DNK	80 mg	160 mg	80 mg	40 mg	100 mg
Topiramate	50 mg	-	200 mg	400 mg	100 mg	100 mg	200 mg	DNK	25 mg	153.57 mg
Sodium valproate	900 mg	2000 mg	-	3000 mg	DNK	100 mg	DNK	DNK	200 mg	1040 mg
Pizotifen	2.5 mg	3 mg	DNK	-	DNK	DNK	-	DNK	1.5 mg	2.33 mg
Flunarizine	10 mg	-	10 mg	10 mg	10 mg	-	10 mg	10 mg	-	10 mg
Methysergide	6 mg	-	-	12 mg	8 mg	1 mg	DNK	-	9 mg	7.2 mg
Gabapentin	2700 mg	1800 mg	2700 mg	-	3600 mg	DNK	DNK	1600 mg	1800 mg	2366.66 mg
Pregabalin	-	600 mg	-	600 mg	-	300 mg	-	DNK	400 mg	475 mg
Tricyclic Anti-depressant	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes	
Amitriptyline	DNK	10 mg	50 mg	200 mg	DNK	25 mg	DNK	DNK	100 mg	77 mg
Dosulepin	75 mg	150 mg	-	-	-	75 mg	75 mg	75 mg	-	93.75 mg
Other	DHE	Botox	Botox	DHE	DHE	DHE	DHE	Botox	DHE	
	GONB	DHE	DHE	Indomethacin	GONB	GONB	GONB	GONB	GONB	
		GONB	Indomethacin	GONB			Candesartan	Acupuncture	MCNB	
		MCNB	GONB	MCNB					Indomethacin	
		Co-enzyme Q10	MCNB							

BOTOX, onabotulinum toxin A; DHE, dihydroergotamine; DNK, do not know; GONB, greater occipital nerve block; MCNB, multiple cranial nerve block.

Table 3. Headache Characteristics Before and After Treatment.

	Follow-up (months)	Moderate-to-severe headache days			Mean daily VRS			Mean daily pain duration (hours)			Estimated improvement [†]	Patient recommend [‡]
		Pre-ONS	Post-ONS	% Change	Pre-ONS	Post-ONS	VRS change	Pre-ONS	Post-ONS	% Change		
1 [§]	65	30	30	0	8	8	0	16.00	16.00	0	0	N
2	49	30	30	0	8	4	4	18.00	18.00	0	30	Y
3 [§]	64	30	30	0	5	5	0	15.00	16.00	0	0 (70% prior to battery failure)	Y
4	51	30	30	0	8	7	1	16.00	16.00	0	0	N
5 [¶]	108	30	8	73.3%	6	1	5	17.00	17.00	0	90	Y
6	89	30	30	0	7	7	0	17.00	18.00	0	0	N
7	53	30	29	3.3%	6	7	0	20.00	20.00	0	5	N
8	27	30	30	0	9	6	3	18.00	18.00	0	38	Y
9	32	30	30	0	7	6	1	18.00	18.00	0	50	Y
Median	53.00	30	30	0.0	7	6	1.00	17.00	18.00	0		
(IQR)	(36.5)	(0)	(0.5)	(0.5)	(2)	(2.5)	(3.50)	(2.00)	(2.00)			

* $p < 0.05$.

[†]Patient estimate of improvement at final follow-up.

[‡]Patient asked if they would recommend the procedure to another person with their disorder.

[§]ONS nonfunctioning at time of follow-up, awaiting replacement battery.

[¶]ONS removed at final follow-up.

IQR, interquartile range; ONS, occipital nerve stimulation; VRS, verbal rating scale.

Medication Use

Patient 2 and patient 7 were taking at least one preventative medication at the time of implantation. At final follow-up, patient 2 had stopped all oral medications but was receiving regular occipital nerve blocks. Although they stated they found these beneficial, they showed no change in the median moderate to severe headache days. Patient 7 had stopped their original oral medications by final follow-up (flunarizine and topiramate) but was taking candesartan with no positive effect. Two other patients (patient 4 and patient 9) had started preventative

medications by final follow-up yet reported no change in median monthly to severe headache days.

At both time of implant and final follow-up, the median number of days patients used any type of acute medication was zero with no change ($p = 0.144$). As previously discussed, patient 3 had medication overuse prior to implant in the form of daily opiates. At final follow-up, this patient had stopped all opiates and was using paracetamol less than 15 days a month and so no longer met the criteria for medication overuse by ICHD-3 guidelines (2). Patient 9 with coexistent chronic cluster headache and NDPH

Table 4. Headache Specific Disability, Affect, and Quality of Life Score Changes With ONS.

	Pre-ONS median (IQR)	Post-ONS median (IQR)	Median change (IQR)	<i>p</i> value
	Range	Range	Range	
MIDAS (<i>n</i> = 9)	64.00 (184.00) 8–270	21.00 (256.00) 0–270	–5.00 (66.5) –90 to 105	0.674
HIT-6 (<i>n</i> = 9)	66.00 (6.50) 36–70	65.00 (6.50) 44–82	0.0 (7) –46 to 17	0.446
HAD-A (<i>n</i> = 9)	9.00 (5.50) 4–16	7.00 (3.50) 2–9	1.0 (5.50) –2 to 9	0.107
HAD-D (<i>n</i> = 9)	8.00 (4.50) 2–14	7.00 (4.50) 1–11	2.00 (5.50) –1 to 8	0.176
BDI-II (<i>n</i> = 9)	21.00 (11.50) 6–28	9.00 (10.00) 4–22	3.00 (6.50) 1 to 21	0.008*
SF-36 physical (<i>n</i> = 9)	37.00 (17.75) 20.30–47.20	35.50 (0.25) 20.30–56.20	0.00 (17.35) –19.20 to 11.70	0.499
SF-36 mental (<i>n</i> = 9)	41.70 (21.25) 24.90–53.40	49.80 (8.2) 31.20–61.20	–9.10 (17.3) –20.20 to 6.90	0.063
Euro-QoL (<i>n</i> = 7)	0.76 (0.12) 0.71–0.83	0.79 (0.12) 0.48–0.83	0.00 (0.02) –0.07 to 0.28	0.715
Euro-Scale (<i>n</i> = 7)	50.00 (20.00) 30–80	60.00 (30.00) 20–95	0.0 (30.00) –35.00 to 10.00	0.273

BDI-II, Becks Depression Inventory; HAD-A, Hospital Anxiety and Depression scores-anxiety specific; HAD-D, Hospital Anxiety and Depression scores—depression-specific; HIT-6, Headache Impact Test; IQR, interquartile range; MIDAS, Migraine Disability Assessment Scale; SF-36, Short Form 36.

with migraine features had reduced their triptans from twice daily to twice a week post-ONS implant.

Stimulation Settings

Mean stimulation amplitude was 1.46 V (0.29–3.95), pulse width 449.90 μ s (370–570), and frequency 72.13 Hz (50–140).

Throughout follow-up, six patients had their ONS turned off for a period. Two were due to lack of efficacy and eventual removal, three were due to battery failure, and one due to failure of the battery caused by the patient not charging the battery. In three of these patients, their headaches worsened when the ONS was off despite the fact that only one patient was reporting a positive effect prior to ONS switch off. The mean time to pain from ONS switch off to headache worsening was 9.00 months (6–48 months).

Adverse Events

A total of 15 adverse events were recorded in nine patients and 12 of these events required additional surgical input. One patient (patient 1) suffered an electrode erosion a month after implant which required surgical revision. One patient suffered an infection over a lead connector (patient 5) which resulted in the removal and subsequent reinsertion of the whole ONS system. One patient (patient 7) suffered from three repeated battery failures after only 5–6 months. This led to a revision and replacement of the ONS system with a rechargeable device. No episodes of lead migration or fracture were recorded.

DISCUSSION

Since the initial description of NDPH by Vanast a benign self-limiting headache with tension-type features the clinical view of

the condition has changed somewhat (1). The original diagnostic criteria in the 2004 ICHD-2 is of a headache that within three days of onset is daily and unremitting for more than three months, which has at least two characteristics from: bilateral location, non-pulsatile nature, mild-moderate intensity, not aggravated by movement and which is not accompanied by nausea or vomiting and more than one of photophobia or phonophobia (15). The main differential is of chronic tension-type headache. However, since this time numerous case series have found that migraine features not described in the ICHD-2 classification are common. Robbins et al. found that 56.3% of their cohort of 71 patients with a headache persistent from onset failed to fulfill the ICHD-2 criteria because of their migrainous features (4). Peng et al. phenotyped 92 NDPH in their clinic of which 35.9% reported features of migraine and a majority of the cohort of Li and Rozen described migraine associated symptoms such as nausea (68%), photophobia (66%), and phonophobia (61%) (16,17). The increasing evidence of a wider clinical spectrum to NDPH has led to ICHD-3 including migraine features within the diagnostic criteria allowing a much larger number of patients to be included (2). The prognosis initially thought to be very positive also has been reassessed. In the original Vanast cohort, 73% with NDPH had resolved without treatment within two years of onset (1). Now, it appears that patients with NDPH, especially those with migraine features, appear to have a poorer prognosis than chronic migraine and often prove medically refractory. The cohort of Robbins et al. found that 75.0% of those with NDPH with migraine features had a persistent and continuous headache from onset with a mean duration of 31 months compared to the 77.4% of those with ICHD-2 defined NDPH which had a mean duration of only 18 months (4). Across the cohort as a whole, 63.6% of those with remission did so within 24 months. In the group of Peng et al., after a mean follow-up of 50.1 months post-headache onset, only 26.4% were headache free despite 75/92 patients being treated with preventative drugs (16). A cohort of

30 Japanese patients with NDPH duration of 39.5 months before presentation were treated with various preventative medications (5). Only 27% of these patients reported a good response to treatment, 20% a minimal effect, and 50% reported no effect at all. This information suggests that those patients with NDPH who remit are likely to do so early and that those who do not are unlikely to respond to current treatments.

In terms of current treatments for NDPH, there have been no randomized trials to date. Most specialists will treat NDPH based on the clinical subtype of their headache—migraine or tension-type. There has been some suggestion of specific NDPH treatments such as gabapentin, topiramate, doxycycline (on the basis of a possible infective etiology to NDPH), steroids, and mexiletine (18). None of these agents have shown promise in the clinical setting. Robbins et al. have reported the effects of multiple cranial nerve blocks (MCNBs) on NDPH stating that although 54% had an acute response to the procedure the effect only lasted for 24 hours on average (4). Our units MCNB data showed that only 10% of patients reported a response to the procedure compared to 49% of chronic migraine patients (19).

ONS has been used to treat refractory chronic daily headaches in particular chronic migraine. There are three randomized controlled trials of ONS in chronic migraine and although results are somewhat contradictory, a meta-analysis of them suggests an overall positive response (7–11). The ONSTIM trial (ONS for the treatment of intractable chronic migraine headache) conducted in 2010 included 77 patients with chronic migraine (10). The three-month responder rates (subjects reporting more than 50% reduction in monthly headache days or more than three points reduction in pain intensity) was 39% for the treatment group, 6% for the sham-control group, and 0% for the conventional medical management group. The PRISM (Precision Implantable Stimulator for Migraine) study is still only available in abstract form despite publication in 2009 but it was a negative study reporting no difference in outcomes at three months between sham and active groups (9). The largest randomized controlled trial of ONS in migraine was conducted by Silberstein et al. and involved 157 patients with refractory chronic migraine (11). The primary outcome measure of a 50% or more reduction in average pain intensity after three months of treatment was negative; however, a number of secondary outcomes, including a 30% reduction in pain severity and headache frequency, did show a significant change, suggesting that ONS did have some positive benefit in chronic migraine. A long-term follow-up of this cohort showed that after 52 weeks there was a significant drop in monthly headache days of 6.7 days (± 8.4 days) and that 47.8% of patients reported a more than 50% reduction in pain severity or headache days (8). A meta-analysis of the pooled results of the three studies show that three months of ONS treatment for chronic migraine is associated with a mean reduction of moderate to severe headache days of 2.59 days when compared to sham (7).

There are numerous case series of ONS for chronic migraine including our own units' cohort of 53 patients with medically refractory chronic migraine treated with the same ONS procedure (surgery and programming) as our current NDPH series (12,20). For chronic migraine, our unit reported a clinical response (30% reduction in moderate to severe headache days) in 45.3% of the group with a median follow-up of 42.00 months. Significant reductions also were seen in pain intensity and duration as well as in quality of life scores. These patients were treated in exactly the same way as the NDPH cohort, yet the outcome is strikingly different. Only one patient (11.1%) showed a clinical response

(same definition as for chronic migraine patients) and there was no improvement in the same secondary outcome measures such as quality of life or disability scores. This difference is further compounded by our chronic migraine group being highly refractory and complex having failed a much higher number of medications, having continuous background pains and a number of coexistent headache types.

The poor treatment response of NDPH with migraine features to ONS supports the notion that in those that do not spontaneously resolve early on in the course of the syndrome, their condition will prove highly resistant to treatments otherwise useful in chronic migraine. It also adds to the limited data that nonoral treatments such as nerve blocks appear to be of limited use in the condition. The reason for this emerging difference in treatment response of chronic migraine and NDPH with migraine features poses interesting questions over the underlying pathophysiology of NDPH. Such a dramatic difference in response to oral medications, nerve blocks and now ONS suggests that there may be a very different underlying etiology to chronic migraine and NDPH with migraine features despite the clinical similarities. NDPH as a secondary headache is a concept previously raised in the NDPH literature. Studies have looked at potential trigger events in NDPH such as infections, trauma, stress, and surgical procedures (21). Vanast et al. initially suggested a viral trigger, specifically Epstein Barr virus, which caused a prolonged autoimmune disorder manifesting as chronic headache (22). Rozen and Swidan also explored a possible inflammatory cause of NDPH reporting increased levels of pro-inflammatory cytokines (tumor necrosis factor) in the cerebrospinal fluid of subjects with the syndrome (23). However, findings have not been replicated and methodological issues have cast doubt on the conclusions of Rozen and Swidan's study (24). Other causes speculated to be relevant in NDPH include cervical hypermobility and even intracranial pressure disturbances (25,26). If NDPH is a secondary headache, this may explain why ONS is so much less effective than in chronic migraine as the treatment is not targeting the cause of the secondary headache and therefore, the original driver to create pain is maintained despite attempts to wind down the central pain system.

The main limitation of this study is the small sample size; however, NDPH is a rare disorder with estimated prevalence between 0.03% and 0.1% of the population (27,28), some of whom will respond to medical treatment or spontaneously improve. It is therefore unlikely that large series of patients with treatment refractory NDPH treated with ONS will be reported. This was an uncontrolled open label study with no comparison to placebo or sham-stimulation. Given the rarity of the condition, it is unlikely that a randomized controlled trial of ONS for NDPH will ever be conducted. The response rate of NDPH patients in our group of patients was below that seen in the sham stimulation group in randomized controlled trials of ONS for chronic migraine (11).

Our cohort suggests that ONS is not an effective treatment for NDPH with migraine features even when conducted in specialized and experienced units. If this finding is duplicated in other units, it may be that the treatment should not be offered to these patients due to the significant cost and risk:benefit ratio. The stark differences between response rates of chronic migraine and NDPH with migraine features highlights the potential of different underlying mechanisms of the two clinically similar conditions. There is a need for further studies into NDPH and its etiology and also into new distinctive treatment regimes.

Authorship Statement

Susie Lagrata was responsible for the recruitment of subjects, analysis and interpretation of data, drafting and revision of manuscript. Sanjay Cheema was responsible for the manuscript revision. Laurence Watkins performed surgery and manuscript revision. Manjit Matharu was responsible for the study concept, recruitment of subjects, interpretation of data, and manuscript revision. All authors approved the final version of the manuscript.

How to Cite this Article:

Lagrata S., Cheema S., Watkins L., Matharu M. 2021. Long-Term Outcomes of Occipital Nerve Stimulation for New Daily Persistent Headache With Migrainous Features. *Neuromodulation* 2021; 24: 1093–1099

REFERENCES

1. Vanast WJ, Edmondton M. New daily persistent headaches definition of a benign syndrome. Program abstracts of the twenty-eighth Annual Meeting of the American Association for the Study of Headache June 27, 28 and 29, 1986, Chicago, Illinois U.S.A. *Headache* 1986;26:309–321.
2. Headache Classification Committee of the International Headache Society (IHS). The international classification of headache disorders, 3rd edition (beta version). *Cephalalgia* 2018;38:1–211.
3. Rozen TD. New daily persistent headache: an update. *Curr Pain Headache Rep* 2014;18:431.
4. Robbins MS, Grosberg BM, Napchan U, Crystal SC, Lipton RB. Clinical and prognostic subforms of new daily-persistent headache. *Neurology* 2010;74:1358–1364.
5. Takase Y, Nakano M, Tatsumi C, Matsuyama T. Clinical features, effectiveness of drug-based treatment, and prognosis of new daily persistent headache (NDPH): 30 cases in Japan. *Cephalalgia* 2004;24:955–959.
6. Prakash S, Saini S, Rana KR, Mahato P. Refining clinical features and therapeutic options of new daily persistent headache: a retrospective study of 63 patients in India. *J Headache Pain* 2012;13:477–485.
7. Chen YF, Bramley G, Unwin G et al. Occipital nerve stimulation for chronic migraine—a systematic review and meta-analysis. *PLoS One* 2015;10:e0116786.
8. Dodick DW, Silberstein SD, Reed KL et al. Safety and efficacy of peripheral nerve stimulation of the occipital nerves for the management of chronic migraine: long-term results from a randomized, multicenter, double-blinded, controlled study. *Cephalalgia* 2015;35:344–358.
9. Lipton R, Goadsby PJ, Cady R et al. PRISM study: occipital nerve stimulation for treatment-refractory migraine. *Cephalalgia* 2009;29:30.
10. Saper JR, Dodick DW, Silberstein SD, McCarville S, Sun M, Goadsby PJ. Occipital nerve stimulation for the treatment of intractable chronic migraine headache: ONSTIM feasibility study. *Cephalalgia* 2011;31:271–285.
11. Silberstein SD, Dodick DW, Saper J et al. Safety and efficacy of peripheral nerve stimulation of the occipital nerves for the management of chronic migraine: results from a randomized, multicenter, double-blinded, controlled study. *Cephalalgia* 2012;32:1165–1179.
12. Miller S, Watkins L, Matharu M. Long-term outcomes of occipital nerve stimulation for chronic migraine: a cohort of 53 patients. *J Headache Pain* 2016;17:68.
13. Goadsby PJ, Schoenen J, Ferrari MD, Silberstein SD, Dodick D. Towards a definition of intractable headache for use in clinical practice and trials. *Cephalalgia* 2006;26:1168–1170.
14. Silberstein S, Tfelt-Hansen P, Dodick DW et al. Guidelines for controlled trials of prophylactic treatment of chronic migraine in adults. *Cephalalgia* 2008;28:484–495.
15. Headache Classification Subcommittee of the International Headache Society. The international classification of headache disorders, 2nd edition. *Cephalalgia* 2004;24:1–160.
16. Peng KP, Fuh JL, Yuan HK, Shia BC, Wang SJ. New daily persistent headache: should migrainous features be incorporated? *Cephalalgia* 2011;31:1561–1569.
17. Li D, Rozen TD. The clinical characteristics of new daily persistent headache. *Cephalalgia* 2002;22:66–69.
18. Rozen TD. New daily persistent headache: clinical perspective. *Headache* 2011;51:641–649.
19. Miller S, Lagrata S, Matharu M. Multiple cranial nerve blocks for the transitional treatment of chronic headaches. *Cephalalgia* 2019;39:1488–1499.
20. Magis D, Schoenen J. Advances and challenges in neurostimulation for headaches. *Lancet Neurol* 2012;11:708–719.
21. Rozen TD. Triggering events and new daily persistent headache: age and gender differences and insights on pathogenesis—a clinic-based study. *Headache* 2016;56:164–173.
22. Vanast WJ, Diaz-Mitoma F, Tyrrell DL. Hypothesis: chronic benign daily headache is an immune disorder with a viral trigger. *Headache* 1987;27:138–142.
23. Rozen T, Swidan SZ. Elevation of CSF tumor necrosis factor alpha levels in new daily persistent headache and treatment refractory chronic migraine. *Headache* 2007;47:1050–1055.
24. Saxon A. CSF TNFalpha has not been shown to be elevated in headache patients. *Headache* 2015;55:1266.
25. Rozen TD, Roth JM, Denenberg N. Cervical spine joint hypermobility: a possible predisposing factor for new daily persistent headache. *Cephalalgia* 2006;26:1182–1185.
26. Goadsby PJ, Boes C. New daily persistent headache. *J Neurol Neurosurg Psychiatry* 2002;72:ii6–ii9.
27. Castillo J, Muñoz P, Guitera V, Pascual J, Kaplan Award 1998. Epidemiology of chronic daily headache in the general population. *Headache* 1999;39:190–196.
28. Grande RB, Aaseth K, Lundqvist C, Russell MB. Prevalence of new daily persistent headache in the general population. The Akershus study of chronic headache. *Cephalalgia* 2009;29:1149–1155.

COMMENTS

The use of occipital nerve stimulation in the treatment of new daily persistent headache (NDPH) is of major interest. This study showed that NDPH with migrainous features does not respond to occipital nerve stimulation compared to patients with chronic migraine who do.

Stephen Silberstein, MD
Philadelphia, PA, USA

The response of even clear cut chronic migraine to occipital nerve stimulation (ONS) is very variable. By highlighting one headache subgroup with migrainous features for whom ONS is clearly not beneficial, the authors have provided very useful information to help with patient selection.

Interestingly, we have found that in patients with a history of chronic migraine that progresses from episodic migraine, the presence of headache absolutely every day predicts a poor response to ONS. Those who report at least some headache free days appear to do better. It may be that those with a headache every day have a different pathology, adverse psychological factors, or are simply well above the ceiling of the 'headache days' scale.

James Fitzgerald, MA, BM, BCh, PhD
Oxford, UK