Novel advances in monitoring and therapeutic approaches in IIH
Mitchell, James; Mollan, Susan; Vijay, Vivek; Sinclair, Alexandra

DOI: 10.1097/WCO.0000000000000690

License: Creative Commons: Attribution (CC BY)

Document Version
Publisher's PDF, also known as Version of record

Citation for published version (Harvard):

Link to publication on Research at Birmingham portal

General rights
Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

• Users may freely distribute the URL that is used to identify this publication.
• Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
• Users may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

Take down policy
While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact UBIRA@lists.bham.ac.uk providing details and we will remove access to the work immediately and investigate.

Download date: 15. Aug. 2019
Novel advances in monitoring and therapeutic approaches in idiopathic intracranial hypertension

James L. Mitchell\textsuperscript{a,b}, Susan P. Mollan\textsuperscript{c}, Vivek Vijay\textsuperscript{a,b}, and Alexandra J. Sinclair\textsuperscript{a,c,d}

\section*{Purpose of review}

The current article appraises the recent developments in idiopathic intracranial hypertension (IIH), with particular attention to novel therapeutic avenues and advanced clinical assessment and monitoring with optical coherence tomography and telemetric intracranial pressure devices.

\section*{Recent findings}

The incidence of IIH is increasing. The first consensus guidelines for IIH have been published detailing investigation and management algorithms for adult IIH. Improved understanding, clinical assessment and monitoring are emerging with the use of optical coherence tomography. Intracranial pressure telemetry is providing unique insights into the physiology of raised intracranial pressure in IIH. There are now an increasing number of ongoing clinical trials evaluating weight loss methods and novel targeted therapies, such as 11ß-HSD1 inhibition and Glucagon-like peptide 1 (GLP-1) receptor agonists.

\section*{Summary}

Several studies are evaluating new therapies for IIH. Monitoring techniques are advancing, aiding diagnosis and allowing the clinician to accurately evaluate changes in papilloedema and intracranial pressure.

\section*{Keywords}

glucagon-like peptide 1 receptor agonists, idiopathic intracranial hypertension, intracranial telemetry, novel therapies, optical coherence tomography, raised intracranial pressure

\section*{INTRODUCTION}

Idiopathic intracranial hypertension (IIH) is characterized by increased intracranial pressure (ICP) with no identifiable cause. IIH, also known as pseudotumor cerebri, is a syndrome with the major risk factor of recent weight gain, occurring mainly in overweight women of working age\cite{1,2}. There is a rising incidence in this disease\cite{3} and the incidence appears related to country-specific prevalence of obesity\cite{4}.

In the majority of those presenting with IIH, they will have headache that is progressively more severe and frequent, with a divergence of traditional considerations of a raised ICP headache\cite{5} to a phenotype that is highly variable and commonly mimics migraine\cite{6,7}. Other reported symptoms include transient visual obscurations (unilateral or bilateral darkening of the vision typically lasting seconds), pulsatile tinnitus, back pain, dizziness, neck pain, visual blurring, cognitive disturbances, radicular pain, and horizontal diplopia\cite{2,8,9}. Prognosis is variable as IIH can either be self-limited or have a lifelong chronic course with significant affects on quality of life\cite{10,11}.

In 2018, the first consensus IIH guidance was published\cite{1}. The document was reviewed by a committee of international key opinion leaders and a patient group, which established a James Lind Alliance Priority Setting Partnership for adult IIH\cite{12}. It sets out key diagnostic and management principles. The diagnostic principles of the investigation of papilloedema are to find any underlying treatable cause in a timely manner, protect the vision and ensure timely re-examination when...
Headache

KEY POINTS

- The incidence of IIH is increasing, and clinicians now have consensus recommendations to help guide their management decisions.
- Novel therapeutic targets are emerging for IIH including GLP-1R agonists.
- Optical coherence tomography provides the clinician with accurate assessment of changes in papilloedema.
- For those with problematic intracranial pressure, telemetric ICP monitors are likely to be increasingly utilized.

ADVANCES IN OCULAR IMAGING

Visual monitoring of patients is a key principle of management [1••], in addition to visual field perimeter, optical coherence tomography (OCT) has allowed new observations in papilloedema. OCT is a rapid, reliable, reproducible and noninvasive imaging technique, using reflected light waves to produce high-resolution cross-sectional and 3D representations of retinal structures. Optic nerve head (ONH) OCT measures have been correlated with the modified Frisen grading of papilloedema [16,17]. The noninvasive nature of these techniques make them ideal in follow-up in contrast to lumbar puncture, which is feared by patients [18].

Whenever investigating papilloedema, OCT is useful in the differentiation of pseudopapilloedema from true papilloedema, a key area of misdiagnosis [19,20]. Combining blue autofluorescence and disc volume OCT scanning can highlight buried crystalline drusen clearly (Fig. 1). There is debate regarding OHN drusen that appear de novo in papilloedema. Peripapillary hyperreflective mass-like structures, termed PHOMS (Fig. 2), [21,22] may be nerve fibre in origin [21]. Further work may define their significance.

Standard measurements for papilloedema include peripapillary retinal nerve fibre layer (pRNFL) and ONH volume. These are reliably increased in active IIH compared with controls, and are significantly associated with CSF opening pressure and improve following treatment [23]. OCT has revealed dynamic deformation of the peripapillary retinal pigment epithelium and Bruch’s membrane (pRPE/BM) regressing towards the normal shape with reduction of ICP [24,25]. Deformation in pRPE/BM may be of particular value in evaluating atrophic papilloedema with minimal RNFL swelling, as deflection of pRPE/BM may correlate with disease activity. Macular RNFL thickness has been shown to be significantly reduced compared with controls, reduces over time and is associated with ONH volume measurements at baseline and visual function [26,27].

OCT angiography (OCTA) is a relatively new, noninvasive investigative modality, allowing visualization of ONH vasculature. OCTA is based on detecting differences in amplitude, intensity or phase variance between sequential B-scans taken at the same location of the retina. Early use has been directed at papilloedema with small cohorts, reported on multiple OCT platforms, with differing methodology. The advantage of OCTA is the ability to segment layers and view below retinal haemorrhage, but despite the small field of view, there is high resolution [28•]. Early findings include dilated tortuous OHN capillaries with no vascular dropout in comparison to ischaemic OHN oedema where there is vascular dropout [29]. OCTA may provide useful biomarkers and aide diagnosis as fundus fluorescence angiography, its predecessor, remains an invasive test [30].
FIGURE 1. (a) Fundus photograph of the right optic disc in a patient referred for papilloedema. The optic nerve shape is irregular. Note there is no loss of view of any of the retinal vessels as they run over the border of the optic nerve. (b) Fundus photograph of the left optic disc in a patient referred for papilloedema. (c) Red-free fundus photograph of the right optic disc highlights hyper reflectivity at the optic nerve head. (d) Red-free fundus photograph of the left optic disc highlights hyper reflectivity at the optic nerve head. (e) Blue autofluorescence (BAF) imaging using the Heidelberg Spectralis optic coherence tomography (OCT) imaging. This clearly highlights buried optic disc drusen as a cause of the pseudopapilloedema. (f) BAF
OCT technology is limited by penetration particularly in severe OHN swelling. Measurement of the pRPE/BM deflection is semi-automated and accurately placing landmarks is challenging [25]. Proprietary automated algorithms commonly fail to correctly segment the layers, and manual re-segmentation is recommended [15,31]. Overall, as the different parameters of OCT are investigated, the outstanding advantage of OCT is the provision of a reliable quantitative method for the longitudinal monitoring of IIH patients (Fig. 2).

**Figure 2.** (a) Optical coherence tomography infrared image, in same patient as other figures, shows the cross-sectional cut for (b) with the arrowed line. (b) OCT cross-sectional volume image shows a typical peripapillary hyperreflective ovoid mass-like structure (PHOMS) (arrow). (c) OCT infra-red image, in same patient as other figures, shows the cross-sectional cut for (b) with an arrowed line. Note the reduction in optic nerve head swelling. (d) OCT cross-sectional volume image shows the reduction in the size of the PHOMS (arrow). (e) and (f) Fundus photographs of the right (e) and left (f) eye at baseline showing Frisen grade 4 papilloedema. (g) and (h) Resolution of papilloedema following bariatric surgery seen on fundus photographs. (i) and (j) Resolution of papilloedema seen on OCT as reduction of total retinal thickness at the optic nerve head in right (i) and left (j) eyes. OCT, optical coherence tomography.

**Advances in Intracranial Pressure Telemetry**

The most common method of ICP measurement in IIH remains lumbar puncture, with several well documented negative aspects [32]. Direct
measurement of ICP is either noninvasive (yet to be used routinely) or invasive. Invasive ICP measurement can be performed at various anatomical sites namely intraventricular, intraparenchymal, subarachnoid, subdural, epidural; where there is CSF communication ICP can be measured by lumbar puncture [33]. Basic ICP measurement by external ventricular, lumbar drains or lumbar puncture is made by fluid column; more recently a variety of microsensors have become available that locate in the target tissue. These have been limited by drift phenomena and the requirement for external wiring, thus the risk of infection with use in excess of 72 h.

Telemetric ICP monitors are now available commercially. There are two main systems, Neurovent p-Tel, Raumedic, Helmbrechts, Germany and Sensor Reservoir, Miethke, Potsdam, Germany. The wiring issue is resolved with wireless power and reading utilizing induction technology. The drift issue is solved in both systems by way of an external monitor reading atmospheric pressure and solid-state sensor technology. The two systems differ with Neurovent p-Tel siting a pressure sensor in the brain parenchyma; meanwhile the Miethke system is based on a sensor within a reservoir attached to a ventricular drain. Both devices are readable by external hand-held equipment, both devices are passive in the sense that they have no power or memory integral to the device and so pressure is only recorded with the external equipment in situ.

The major initial application for the Miethke system is in refining valve settings in challenging patients, a recent case series highlights this application, of note the highest frequency of valve adjustments was seen in the IIH cohort [34].

More data is available for the Raumedic p-Tel as it has been available for longer. It sites a solid-state pressure sensor approximately 20 mm into the brain parenchyma, usually within the right frontal lobe. From the largest published series, there is a low complication rate, approximately 6% overall, with seizures affecting 3% and infection in 1.5% [35**]; however, this was in a series of patients with significant structural brain abnormality (for example, hydrocephalus) and the rate of such complications is likely lower in IIH patients. Of note, the UK driving regulations allow resumption of driving 1 week after insertion in patients without complications. The device provides a high degree of accuracy with low drift [36]. Many have been kept in situ beyond the licensed 3-month period [35**,37]; where they have been shown to retain their accuracy with low drift of 2.5 mmHg over a median 241-day implantation period. The device samples at 5 Hz, considerably lower than the wired and Miethke systems, although this is sufficient for waveform analysis [36,38]. The device is capable of long-term recordings for up to 1 week with the present hardware and can be worn by an ambulant patient out with the hospital environment (Fig. 3) [38].

Telemetric ICP monitors have an evolving role in diagnosis and monitoring of several conditions. In IIH, particular roles could include evaluating whether neurosurgical shunt placement is advised in a deteriorating patient developing fulminant disease. Furthermore, it is useful in evaluating whether pressure is pathologically elevated in those with...
minimal ocular features and in shunted patients. Monitors can inform the setting of CSF shunt valves aiming to abrogate low pressure headaches, at present seen in 23% [14]. ICP telemetry may also facilitate the differentiation between raised pressure headaches and migrainous headaches [39].

**NOVEL THERAPEUTICS AND INTRACRANIAL HYPERTENSION**

Acetazolamide is the longest established treatment for IIH. In 2015, following the publication of the first two randomized control trials for medical treatment in IIH [8,40], an updated Cochrane review highlighted that there was insufficient evidence to recommend or reject the efficacy of acetazolamide for treating IIH and insufficient evidence for other drugs used in IIH [41]. Of note, there was no effect of acetazolamide on headache seen in the IIHTT [6]. The common existing drugs used in IIH have been evaluated acutely in vivo at clinically relevant doses, and were not found to significantly reduce ICP, with the exception of topiramate [42]. There is, therefore, an unmet need for novel therapeutic strategies in IIH (Fig. 4).

Disordered CSF dynamics have been suspected to underlie the raised ICP seen in IIH. There are currently no novel drugs targeting the underlying pathogenesis driving IIH, which remains elusive. Novel therapies are currently focussed on reducing ICP through reducing CSF secretion. Ideally novel therapies would also reduce weight as this approach is disease modifying in IIH [43].

The choroid plexus is the principle site of CSF production; this is driven by the net movement of sodium ions (Na⁺) from the blood to the cerebral ventricles, creating an osmotic gradient down, which water moves. Although several channels are involved in this process, the principal channel is the Na⁺/K⁺/ATPase that actively transports Na⁺ into the cerebral ventricle and is the rate-limiting step [44,45].

**11ß-HYDROXYSTEROID DEHYDROGENASE TYPE 1**

11ß-hydroxysteroid dehydrogenase type 1 (11ß-HSD1) is an intracellular enzyme that converts inactive cortisone to the active cortisol. This amplifies local glucocorticoid activity independent of systemic cortisol. 11ß-HSD1 expression and activity has been demonstrated in choroid plexus epithelial cells, along with other key elements of the glucocorticoid signalling pathway [46,47]. Inhibitors have been developed, including AZD4017, originally as potential therapies for diabetes mellitus type 2 and the metabolic syndrome. Glucocorticoid metabolism has been characterized in IIH in relation to therapeutic weight reduction; global 11ß-HSD1 activity decreased with weight loss as measured by urinary glucocorticoid metabolites by gas chromatography/mass spectroscopy [43]. Importantly it was noted that the reduction in ICP significantly correlated with reduction in 11ß-HSD1 activity [47]. Of interest, is that 11ß-HSD1 inhibition reduced intraocular pressure and it has been shown that secretory mechanisms of the ocular ciliary body are akin to that of choroid plexus epithelium [46–48].

11ß-HSD1 inhibitors do not affect systemic glucocorticoid metabolism [49], but would reduce CSF secretion though reducing local cortisol availability in the choroid plexus with subsequent reduction of downstream glucocorticoid receptor-mediated sodium transportation, reduced osmotic gradient and decreased water movement into the cerebral ventricle [46,47]. Conversely systemic administration of glucocorticoids has been found to precipitate intracranial hypertension [50]. The IIH Drug trial (IIHDT), clinicaltrials.gov identifier NCT02017444, has investigated the ability of an 11ß-HSD1 inhibitor to reduce CSF secretion and hence ICP in patients with IIH [51]. IIHDT is the first phase 2 double-blind placebo-controlled trial in IIH. It recently completed recruitment and is expected to report in 2019.

**GLUCAGON-LIKE PEPTIDE 1**

The incretin glucagon-like peptide 1 (GLP-1) is a gut peptide secreted by the distal small intestine in response to food intake [52]. GLP-1 stimulates glucose-dependant insulin secretion and inhibits glucagon release, lowering blood glucose only in the presence of insulin and not resulting in hypoglycaemia [53]. GLP-1 is also synthesized in neurons of the nucleus tractus solaris that project to the hypothalamus [54] and promotes satiety and weight loss [55]. GLP-1 analogues have a clinical role in the management of type 2 diabetes mellitus, as well as for weight loss in obesity. Several GLP-1 agonists have been developed and are now licensed drugs [56]. These include exenatide twice daily, exenatide once weekly, liraglutide, lixisenatide, albiglutide, dulaglutide and most recently semaglutide. Currently, only liraglutide is licensed for weight loss in obesity. They vary in structure and pharmacology, ability to penetrate the blood–brain barrier (BBB) as evidenced by CNS effects. Importantly, the choroid plexus epithelium lies outside the BBB [57].
FIGURE 4. The major ion channels responsible for CSF secretion in the choroid plexus are shown with sites of action of acetazolamide, AZD4017 and exenatide. Cortisone is converted to the active cortisol by 11ß-HSD1, cortisol binds to the GR and MR receptors, which upregulate Na\(^+\) K\(^+\) ATPase activity; AZD4017 inhibits 11ß-HSD1 reducing local availability of cortisol. Exenatide binds and activates GLP-1R stimulating the conversion of ATP to cAMP by AC. cAMP activates PKA, which inhibits the Na\(^+\) H\(^+\) exchanger reducing Na\(^+\) re-absorption and also inhibits the Na\(^+\) K\(^+\) ATPase reducing Na\(^+\) excretion. Carbonic anhydrase catalyzes the conversion of H\(_2\)O and CO\(_2\) to H\(^+\) and HCO\(_3^-\), which is important in the establishment of the osmotic gradient. Both acetazolamide and topiramate inhibit carbonic anhydrase function. AC, adenylate cyclase; AE2, anion exchange protein 2; cAMP, cyclic adenosine monophosphate; CSF, cerebrospinal fluid; CTFR, cystic fibrosis.
GLP-1 also has a diuretic effect by reducing Na\(^+\) re-absorption in the renal proximal tubule, thereby increasing Na\(^+\) and water excretion [58]. Activation of GLP-1R stimulates the conversion of adenosine triphosphate to cyclic adenosine monophosphate (cAMP) by adenylate cyclase. cAMP activates protein kinase A, which inhibits the Na\(^+\)/H\(^+\) exchanger, thus reducing Na\(^+\) re-absorption. Choroid plexus epithelial cell function is inverted compared with renal proximal tubule but with an analogous fluid transport mechanism [59], and as such GLP-1R was investigated as a potential target for conditions with raised ICP. It has been shown that GLP-1 receptor (GLP-1R) is expressed in the human choroid plexus. Treatment with the agonist exendin-4 modulates the GLP-1R in the rat choroid plexus through agonist induced receptor internalization, which was shown to increase cAMP generation and reduce Na\(^+\)/K\(^+\)/ATPase activity. Importantly exendin-4 reduced ICP in conscious rats at clinically relevant doses. There was a 65% reduction in ICP 30 min post dose compared with baseline and a cumulative effect seen with reduction in the ICP to 79.3 and 72.5% of baseline values predose at days 2 and 4, respectively, of the experiment. The action was blocked by intraventricular administration of the GLP-1R antagonist exendin 9-39, suggesting the effect is mediated by the GLP-1R in the brain. Importantly, the effect was also seen in a rat model with markedly raised ICP [60\(^*\)]. The IIH Pressure Trial, ISRCTN12678718, is a double-blinded, placebo-controlled physiology study assessing the effects of exenatide, a GLP-1R agonist, on ICP in a cohort with active IIH and is expected to report this year [39].

There are also developments with surgical management of IIH, two trials are currently ongoing, the IIH Weight Trial [61] (ClinicalTrials.gov Identifier: NCT02124486) is a randomized controlled trial of Bariatric Surgery Versus a Community Weight Loss Programme and opened to recruitment in 2014. The SIGHT trial (ClinicalTrials.gov Identifier: NCT03501966) opened in 2018 and is a triple-arm randomized controlled trial of medical therapy (acetazolamide) vs. medical therapy with Optic Nerve Sheath Fenestration vs. medical therapy with ventriculoperitoneal shunting.

**CONCLUSION**

The incidence of IIH is increasing along with global rates of obesity, a key pathological factor, making IIH research and management increasingly important. The first consensus guidelines for the management of IIH have now been published guiding the clinical management of this condition. Looking to the future, there are several new avenues of clinical therapeutics based on reducing CSF secretion, with GLP-1 receptor agonists appearing promising as they also significantly reduce weight. Advances in OCT technology will continue to improve our diagnosis and monitoring of papilloedema, and may provide unique biomarkers. Early indications show that telemetric ICP monitoring may provide insight into CSF dynamics in IIH and facilitate management decisions.

**Acknowledgements**

None.

**Contributor Statement:** All authors have read and approved the final manuscript.

**Financial support and sponsorship**

None.

**Conflicts of interest**

A.J.S. is an inventor and the University of Birmingham applicant on patent application no. PCT/GB2015/052453 related to this work entitled ‘Elevated intracranial pressure treatment.’

A.J.S. is funded by an NIHR Clinician Scientist Fellowship (NIHR-CS-011-028) and by the Medical Research Council, UK (MR/K015184/1).

**REFERENCES AND RECOMMENDED READING**

Papers of particular interest, published within the annual period of review, have been highlighted as:

- of special interest
- of outstanding interest

1. Mollan SP, Davies B, Silver NC, et al. Idiopathic intracranial hypertension: consensus guidelines on management. J Neurol Neurosurg Psychiatry 2018; 89:1088–1100. Over 20 common questions answered for clinicians on all aspects of IIH and IIH WOP. Consensus-based statements founded by a specialist interest group that included patients, allied health professionals, neurologists, neuroophthalmologists, neurosurgeons, neuroradiologists, and informed by multiple surveys of practice. It was reviewed by four professional bodies in the United Kingdom (UK), an international panel and by IIH UK, a UK-based patient charity for IIH.


1350.7540 Copyright © 2019 The Author(s). Published by Wolters Kluwer, Inc.

www.co-neurology.com

Novel advances in therapeautic approaches in IIH Mitchell et al.

8. Up-to-date review of our understanding of headache in IIH. Presents the latest literature phenotyping the IIH headache as migraine, and highlighting the issue of medication overuse headache in the IIH cohort.
10. The Idiopathic Intracranial Hypertension Treatment Trial (IIH TT) represents the largest randomized, prospectively analyzed cohort of untreated patients with IIH to date. It reported the use of acetazolamide with a low-sodium weight-reduction diet compared with diet alone resulted in modest improvement in visual field function in patients with mild visual loss. The IIHTT also reported improved quality-of-life outcomes at 6 months with acetazolamide and diet group compared with diet alone.
42. Is cerebrospinal fluid shunting in idiopathic intracranial hypertension worthwhile? A 10-year review.