Trigeminal autonomic cephalalgias: fancy term or constructive change to the IHS classification?

P J Goadsby

A classification based on pathophysiology is a useful aid to differential diagnosis and effective treatment planning

For the neurologist faced with the day to day grind of clinical work a change to terminology may seem like the academics “at it again”. I will try to set out this change and illustrate a physiology that may be attractive to understand, and hopefully one that enhances, clinical practice. Appreciating the physiology of the trigeminal-autonomic reflex can make patients presenting with varying degrees of cranial autonomic activation, such as lacrimation, conjunctival injection, nasal congestion or rhinorrhoea and the like, comprehensible at the bedside.1

The trigeminal autonomic cephalalgias (TACs) is a grouping of headache syndromes recognised in the second edition of the International Headache Society (IHS) classification.2 The term was coined to reflect a part of the pathophysiology that may be attractive to pathologists, notably cranial autonomic activation.3

TACs are classified in section III of the second edition of the classification2 and include cluster headache,4 episodic and chronic paroxysmal hemicrania,5 hemicrania continua6 and autonomic episodic vertigo7 and are independent of cerebral metabolism.8 Cranial parasympathetic autonomic features seen clinically are consistent with cranial parasympathetic activation (lacrimation, rhinorrhoea, nasal congestion, and eyelid oedema) and sympathetic hypofunction (ptosis and miosis). The latter is likely to be a neurapraxia effect of carotid wall swelling13 with cranial parasympathetic activation. Some degree of cranial autonomic sympathometabolism is, therefore, a normal physiological response to cranial nociceptive input.13 Indeed other primary headaches, notably migraine,2 patients with facial pain, such as trigeminal neuralgia,9 would be expected to have cranial autonomic activation, and they do. The distinction between the TACs and other headache syndromes is the degree of cranial autonomic activation, not its presence alone.10 This is why some patients with migraine have minor cranial autonomic activation that leads to the term cluster-migra, when most patients with migraine with cranial autonomic activation.

Permitting trigeminal-parasympathetic activation

What is the basis for the cranial autonomic symptoms being so prominent in the TACs? Is it due to a central disinhibition of the trigeminal-autonomic reflex? Functional imaging studies—positron emission tomography studies in cluster headache11–12 and a functional magnetic resonance imaging (MRI) study in SUNCT syndrome13—has demonstrated ipsilateral posterior hypothalamic activation. Posterior hypothalamic activation seems specific to these syndromes, which is not seen in episodic14–16 or chronic migraine, or in experimental ophthalmic trigeminal distribution head pain.17 There are direct hypothalamic-trigeminal connections17 and the hypothalamus is known to have a modulatory role on the nociceptive and autonomic pathways, specifically trigeminovascular nociceptive pathways.18 Hence, cluster headache and SUNCT syndrome are probably due to an abnormality in the region of the hypothalamus (fig 1) with subsequent trigeminovascular and cranial autonomic activation. Imaging data with paroxysmal hemicrania are keenly awaited. Cranial autonomic features are not invariably linked with trigeminal pain and may persist after lesions of the trigeminal nerve.

Differential Diagnosis of TACs

The TACs need to be differentiated from secondary TAC producing lesions, from...
other primary headaches, and from each other. The differentiation from secondary causes is not a problem if one images patients but can be extremely difficult if one does not. An MRI of the brain with attention to the pituitary fossa and cavernous sinus will detect most secondary causes. It is easy to make an argument given the rarity of paroxysmal hemicrania and SUNCT syndrome that MRI would detect no more than one in 100 cases of lesions in episodic cluster headache and in whom multiple attacks were captured.

For other primary headaches, migraine is the single biggest problem in the differential diagnosis of cluster headache. Migraine can cluster and despite the best intentions of the IHS classification committee short attacks do occur. Cranial autonomic symptoms are well reported, and the neuropeptide changes are the same as in cluster headache. The occurrence of attacks together does not seem to have the seasonal preponderance that is so typical of cluster headache, and this can be a useful differential diagnostic feature. I regard the term cluster-migraine as unhelpful and I am yet to see a convincing case of a distinct biological entity usefully described by this name. The criterion for the effect of movement was added to cluster headache to sharpen the difference with migraine. The committee hoped this would draw attention to the fact that most cluster headache patients feel restless or agitated, whereas most migraine patients are quiescent, as IHS-I recognised.

In clinical practice, this symptom, and the periodicity, are extremely helpful in differential diagnosis. The other feature of cluster headache, and this is a feature of TACs when compared with migraine, is that patients with TACs often complain of unilateral, homolateral photophobia, whereas patients with migraine often complain of bilateral photophobia. Bilateral photophobia in patients with TAC could be speculated to occur in about 25% purely by the chance of them having some migrainous biology. The TACs themselves (table 1) can often be differentiated by their attack

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**Table 1** Clinical features of the trigeminal autonomic cephalalgias (TACs)

<table>
<thead>
<tr>
<th></th>
<th>Cluster headache</th>
<th>Paroxysmal hemicrania</th>
<th>SUNCT syndrome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex (F:M)</td>
<td>1:4</td>
<td>2:1</td>
<td>1:2</td>
</tr>
<tr>
<td>Pain</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Type</td>
<td>Stabbing, boring</td>
<td>Throbbing, boring</td>
<td>Burning, stabbing, sharp</td>
</tr>
<tr>
<td>Severity</td>
<td>Severe to excruciating</td>
<td>Excruciating</td>
<td>Moderate to severe</td>
</tr>
<tr>
<td>Site</td>
<td>Orbit, temple, face</td>
<td>Orbit, temple</td>
<td>Periorbital</td>
</tr>
<tr>
<td>Attack frequency</td>
<td>15–180 minutes</td>
<td>1–40/day</td>
<td>1/day–30/hour</td>
</tr>
<tr>
<td>Duration of attack</td>
<td>5–240 seconds</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>Autonomic features</td>
<td>Yes</td>
<td>Occasional</td>
<td>No</td>
</tr>
<tr>
<td>Migrainous features*</td>
<td>Yes</td>
<td>Yes</td>
<td>No†</td>
</tr>
<tr>
<td>Alcohol trigger</td>
<td>Yes</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Indomethacin effect</td>
<td>–</td>
<td>++</td>
<td>–</td>
</tr>
</tbody>
</table>

*Nausea, photophobia (often ipsilateral to the pain) or phonophobia.
†May have photophobia ipsilateral to the pain.

SUNCT, Short lasting unilateral neuralgiform headache attacks with conjunctival injection and tearing.

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**Table 2** Cluster headache

3.1 Diagnostic criteria:

A At least five attacks fulfilling B–D
B Severe or very severe unilateral orbital, supraorbital and/or temporal pain lasting 15–180 minutes if untreated
C Headache is accompanied by at least one of the following:
   (1) Ipsilateral conjunctival injection and/or lacrimation
   (2) Ipsilateral nasal congestion and/or rhinorrhea
   (3) Forehead and facial sweating
   (4) Ipsilateral eyelid oedema
   (5) Ipsilateral forehead and facial sweating
   (6) Ipsilateral miosis and/or ptosis
   (7) A sense of restlessness or agitation
D Attacks have a frequency from one every other day to eight per day
E Not attributed to another disorder

3.1.1 Episodic cluster headache

Description: Occurs in periods lasting seven days to one year separated by pain free periods lasting one month or more

Diagnostic criteria:

A All alphabetical headings of 3.1
B At least two cluster periods lasting from 7 to 365 days and separated by pain free remissions of one month or more

3.1.2 Chronic cluster headache

Description: Attacks occur for more than one year without remission or with remissions lasting less than one month

Diagnostic criteria:

A All alphabetical headings of 3.1
B Attacks recur for more than one year without remission periods or with remission periods lasting less than one month
length. This is certainly true when comparing cluster headache with SUNCT/short lasting unilateral neuralgiform headache attacks with cranial autonomic symptoms (SUNA). The IHS criteria for TACs does betray an uncomfortable biological naivety with regard to the timing. The A, C, D, E/F criteria are rather similar for each TAC (tables 2–4). It seems neat in some way to have SUNCT be up to four minutes long, paroxysmal hemicrania from two to 30 minutes and cluster headache from 15 minutes onwards. The overlap seems minimal. It almost goes without saying that this must be wrong in absolute terms, biology rarely provides such neat rules, but it does provide a useful way to identify cases of sufficiently similarity to make biologically meaningful studies.

### CHALLENGES FOR THE TACs

The classification and biology of the TACs have come a long way in a short time. The syndromes are well established, and although rare compared with migraine they are sufficiently common, with cluster headache affecting about 0.2% of the population, to demand a neurological and headache specialist’s attention. There are some particular issues of classification that are not currently clear.

**Cluster headache**

A patient with a first attack of cluster headache is now simply classified as cluster headache (3.1). This takes the top-down view—that is, diagnose what you can and fill in the detail as available. Such cases are unsuitable for almost any study except natural history studies where they are ideally the starting point. A similar problem is how to refer to patients who have one type of TAC, typically an episodic form, and then evolve to the chronic form. The old classification differentiated primary from secondary chronic cluster headache depending on whether there was a period of episodic headache first. This argument would apply equally to cluster headache (3.1). The syndromes are well established, and although rare compared with migraine they are sufficiently common, with cluster headache affecting about 0.2% of the population, to demand a neurological and headache specialist’s attention. There are some particular issues of classification that are not currently clear.

**Paroxysmal hemicrania**

The diagnosis of PH by the IHS criteria requires a response to indometacin. This is very difficult. It is not clear what the basis for the indometacin effect is, although it is perfectly clear that the effect is clinically very meaningful (table 5). Patients with PH who are treated with indometacin have an almost unbelievably spectacular resolution. This response seems so distinct that reserving the diagnosis of PH for these patients seems reasonable. Given varying sensitivity to indometacin, we have seen a requirement for a single dose given first thing in the morning of 300 mg indometacin to produce a complete response—perhaps there are unrecognised dosing requirements. There is certainly a timing requirement and again we have seen patients turn off, but only after 10 days at the dose of 275 mg daily.

**SUNCT**

For SUNCT the most immediate challenge must be to define the phenotype properly. We have seen patients who fulfill criteria for SUNA (table 6) but not SUNCT (see table 4). Typically the eye is not red, but we have also seen, for example external auditory canal swelling and periaural flushing as the sole cranial autonomic symptom, as has been reported for PH. It seems possible, given the relative proportion of patients with cluster headache who have lacremitation and conjunctival injection as compared with other cranial autonomic symptoms, that these symptoms are for some reason biologically more likely. This is supported by the same relative changes being seen in experimentally induced headache.

### Table 3 Paroxysmal hemicrania

<table>
<thead>
<tr>
<th>Diagnostic criteria</th>
<th>3.2</th>
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</thead>
<tbody>
<tr>
<td>A At least 20 attacks fulfilling B–D</td>
<td></td>
</tr>
<tr>
<td>B Severe unilateral orbital, supraorbital, or temporal pain lasting 2–30 minutes</td>
<td></td>
</tr>
<tr>
<td>C Headache is accompanied by at least one of the following:</td>
<td></td>
</tr>
<tr>
<td>(1) Ipsilateral conjunctival injection and/or lacrimation</td>
<td></td>
</tr>
<tr>
<td>(2) Ipsilateral nasal congestion and/or rhinorrhea</td>
<td></td>
</tr>
<tr>
<td>(3) Forehead and facial sweating</td>
<td></td>
</tr>
<tr>
<td>(4) Ipsilateral eyelid oedema</td>
<td></td>
</tr>
<tr>
<td>(5) Ipsilateral forehead and facial sweating</td>
<td></td>
</tr>
<tr>
<td>(6) Ipsilateral miosis and/or ptosis</td>
<td></td>
</tr>
<tr>
<td>D Attacks have a frequency above five per day for more than half the time, although periods with lower frequency may occur</td>
<td></td>
</tr>
<tr>
<td>E Attacks are prevented completely by therapeutic doses of indometacin</td>
<td></td>
</tr>
<tr>
<td>F Not attributed to another disorder</td>
<td></td>
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</tbody>
</table>

**Table 4 Short lasting unilateral neuralgiform headache attacks with conjunctival injection and tearing (SUNCT)**

<table>
<thead>
<tr>
<th>Diagnostic criteria</th>
<th>3.3</th>
</tr>
</thead>
<tbody>
<tr>
<td>A At least 20 attacks fulfilling criteria B–E</td>
<td></td>
</tr>
<tr>
<td>B Attacks of unilateral, orbital, supraorbital or temporal stabbing or pulsating pain last 5–240 seconds</td>
<td></td>
</tr>
<tr>
<td>C Pain is accompanied by ipsilateral conjunctival injection and lacrimation</td>
<td></td>
</tr>
<tr>
<td>D Attacks occur with a frequency from three to 200 per day</td>
<td></td>
</tr>
<tr>
<td>E Not attributed to another disorder</td>
<td></td>
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</tbody>
</table>

**Table 5 Effects of treatment on trigeminal autonomic cephalalgias**

<table>
<thead>
<tr>
<th>Cluster headache</th>
<th>Paroxysmal hemicrania</th>
<th>SUNCT syndrome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Indometacin effect</td>
<td>Sumatriptan 6 mg s/c ar</td>
<td>+ +</td>
</tr>
<tr>
<td>Abortive treatment</td>
<td>Oxygen 20 mg nasal insufflation</td>
<td>Nil</td>
</tr>
<tr>
<td>Preventive treatment</td>
<td>Verapamil</td>
<td>Indometacin</td>
</tr>
<tr>
<td></td>
<td>Methysergide</td>
<td>Topiramate</td>
</tr>
<tr>
<td></td>
<td>Lithium</td>
<td>Prednisone</td>
</tr>
</tbody>
</table>
Table 6  Short lasting unilateral neuralgiform headache attacks with cranial autonomic symptoms (SUNA)

<table>
<thead>
<tr>
<th>A 3.3 Diagnostic criteria:</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>A At least 20 attacks fulfilling criteria B–E</td>
<td></td>
</tr>
<tr>
<td>B Attacks of unilateral orbital, supraorbital, or temporal stabbing pain lasting from two seconds to 10 minutes</td>
<td></td>
</tr>
<tr>
<td>C Pain is accompanied by one of:</td>
<td></td>
</tr>
<tr>
<td>(1) Conjunctival injection and/or tearing</td>
<td></td>
</tr>
<tr>
<td>(2) Nasal congestion and/or rhinorrhea</td>
<td></td>
</tr>
<tr>
<td>(3) Eyelid oedema</td>
<td></td>
</tr>
<tr>
<td>D Attacks occur with a frequency of one or more per day for more than half the time</td>
<td></td>
</tr>
<tr>
<td>E Not attributed to another disorder</td>
<td></td>
</tr>
<tr>
<td>A 3.3.1 Episodic SUNA</td>
<td></td>
</tr>
<tr>
<td>Description: SUNA attacks occurring for seven days to one year with pain free intervals longer than one month</td>
<td></td>
</tr>
<tr>
<td>A 3.3.2 Chronic SUNA</td>
<td></td>
</tr>
<tr>
<td>Description: At least two attack periods last seven days to one year separated by remission periods of less than one month (untreated)</td>
<td></td>
</tr>
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</table>

Thus research criteria for a more encompassing syndrome are proposed (see table 6).

CONCLUSION

The TACs represent a great success story in headache. From a classification point of view, the syndromes share much biology so their agglomeration in section III draws attention to them and to the trigeminoparasympathetic reflex. It is highly desirable that headache classification moves to a more biological and pathophysiological basis and the TACs are a step in that direction. The TACs also represent excellent clinical opportunities to take a careful history and offer effective therapy to otherwise highly disabled, suffering patients. Lastly, further investigations of the TACs are bound to illuminate physiological processes whose understanding will be useful to the range of primary headache syndromes.

J Neurol Neurosurg Psychiatry 2004;75:630–635
doi: 10.1136/jnp.2004.036012
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Competing interests: P JG is a Wellcome Trust Senior Research Fellow.

REFERENCES

Cholinesterase inhibitors

Action of cholinesterase inhibitors in patients’ brains

K Herholz

Cholinesterase inhibitors in patients’ brains

Cholinesterase (ChE) inhibitors are the only class of drug that have consistently shown improvement in cognitive function in patients with mild to moderate Alzheimer’s disease. Unfortunately, improvement is generally rather small. Recent clinical trials have been shown to improve cognitive function in patients with mild to moderate Alzheimer’s disease. However, on the other hand, recent studies have suggested more extensive use because the improvement in cognitive function has also been observed in vascular dementia, dementia with Lewy bodies, and Parkinson’s disease with dementia. However, the other hand a recent study in community resident patients with mild to moderate Alzheimer’s disease concluded that benefits were “below minimally relevant thresholds.”

On the background of this confusing situation, studies are particularly welcome that provide clues as to how ChE inhibitors exert their beneficial effect in patients and how we could increase their efficacy. In this issue, such information is provided in a study by Bohnen et al., which measured the actual inhibition of cortical acetylcholine esterase (AChE) activity by donepezil in vivo and studied the correlation of the degree of inhibition with the cognitive effects. Several observations were made that indicate directions for improving therapy.

The inhibition of cortical AChE activity by donepezil at the recommended dose of 10 mg daily was rather low (on average 16–24% depending on cortical regions) and it varied considerably among patients. Although somewhat higher values had been measured with a slightly different tracer by other authors cited in the paper, the inhibition of human cerebral AChE is much less than observed in peripheral blood, which is in contrast to findings in rats. Thus, dosage, pharmacokinetics, or specific binding of the drug to human cerebral AChE appear to be suboptimal, and this had not become evident during preclinical and clinical phases of drug development and testing.

The study also indicates that the degree of cerebral AChE inhibition makes a clinical difference because it was significantly correlated with measures of executive function and attention. This indicates that it could indeed be worthwhile to increase inhibition in selected patients, e.g., by higher dosages in side effects permit. It is expected that similar positron emission tomography (PET) studies will be performed to measure inhibition of cerebral butyrylcholine esterase (BChE) for selection and development of drugs that achieve higher effective levels of acetylcholine by additional inhibition of this degradation pathway.

Another interesting aspect is that the relatively small inhibition effects were observed in temporal and parietal association cortex—structures that are thought to be of pivotal importance for episodic and semantic memory—that did not benefit significantly from treatment in this and other studies. One would wish to see similar studies with other ChE inhibitors to determine whether this is a property of the entire class of drugs.

It is gratifying that such direct in vivo assessments of pharmacological action are happening now, which means that we do not depend solely on large trials with clinical outcome measures that are, of course, of utmost clinical importance but often tell very little about the mechanisms that explain interindividual variation. One can hope that this will ultimately provide rational means to improve treatment of individuals, which is in the primary interest of patients and doctors.

REFERENCES


J Neurol Neurosurg Psychiatry; first published as 10.1136/jnnp.2004.059931 on 16 February 2005. Downloaded from http://jnnp.bmj.com/or November 4, 2023 by guest. Protected by copyright.
Recreational cannabis use: not so harmless!

D Deplanque

Cannabis and stroke

Cannabis is currently the most widely used illicit drug in Western populations. The question of whether or not it should remain prohibited is under debate in many European countries. Possible adverse health effects play an important role in this debate. The classical anxiolytic, sedative, analgesic, and psychedelic properties of cannabis are well known, but it has recently been suggested that it may also induce cerebrovascular disease. Now for the first time, Mateo et al have shown (this issue, see page 435) that a causal relation with cerebrovascular events is highly plausible. The major argument in favour of this is that the events only occurred during periods when the patient was consuming cannabis, and this is a major criterion of adverse drug reaction monitoring. In the present case, the patient was not a regular user and both the arterial abnormalities and the time course of the disease suggest an immuno-allergic vasculitis.

The systemic effects of natural cannabis compounds (Δ9-tetrahydrocannabinol, Δ8-tetrahydrocannabinol, cannabidiol) are primarily mediated by the activation of cannabinoid receptors (CB1 and CB2), which are present in various tissues including the brain, cells of the immune system, blood vessels, and heart. Apart from the vasoactive properties of cannabinoids, it is also possible that a central nervous system vasculitis could result from a dysfunction of the immune system. It remains to be determined whether the immunomodulatory effects of cannabinoids could be involved as a mechanism of cannabis induced vasculitis through their action on CB2 receptors. The very low frequency of such complications may reflect a genetic predisposition in a few individuals. Better identification of patients with cannabis related strokes may lead to epidemiological case-control studies that include genomic investigations. In that recreational cannabis use appears not to be as harmless as was thought, there is a need to improve public information. The therapeutic potential of cannabis and its derivatives should be rigorously evaluated and the benefit to risk ratio taken into account before authorising their medical use.

REFERENCES