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Creating Informed Interest in Mal De Débarquement Syndrome

To the Editor: Mal de débarquement syndrome (MdDS) is a chronic illness manifested by a persistent false sensation of nonspinning self-motion, such as rocking, swaying, bobbing, or gravitational pull in a particular direction. The case report by Williams et al¹ correctly identifies that MdDS is often complicated by psychiatric comorbidities and that underrecognition of the illness among clinicians poses a significant burden on afflicted patients. Pertinently to the latter point, patients are said to make an estimated average of 7–19 visits to health care professionals before their MdDS diagnosis, which can take several frustrating years and cost thousands of dollars for the visits, diagnostic tests, and travel expenses.^{2,3} Unfortunately however, the authors seem to have overlooked the publication of the expert consensus document on MdDS diagnostic criteria, which predates the case report by 2 years and takes some contrasting positions.⁴ Critically, the document, as previously noted, states that exposure to passive motion temporarily reduces symptoms of MdDS, which is a rather unique qualifier and diagnostic distinction from persistent postural perceptual dizziness.^{4–8} The document also states that clinical structural brain imaging is of low yield in the diagnosis of MdDS, as incidental findings are rather common.^{5,9}

The journal readership should also be made aware that the chance of a positive outcome of MdDS has recently significantly improved with the discovery that a central vestibular mechanism, known as velocity storage, may be maladapted in patients with MdDS.^{10–12} Velocity storage is a major element of the vestibulo-ocular reflex (VOR), underpinned by an interaction between the bilateral vestibular nuclei in the brain stem and the posterior vermal cerebellum.^{13,14} Activated by head rotation, large-field visual motion, or tactile cues for continuous rotation, this mechanism holds an estimate of head rotational velocity in space, as expressed by nystagmus in response to head rotation or optokinetic stimulation (OKS) or during circular locomotion.^{15,16} In addition to serving brain stem reflexes with a working memory–like function for self-motion,¹⁷ velocity storage is thought to contribute to the illusory sensation of self-motion during OKS,¹⁸ as well as the false sensation of self-motion in MdDS. Importantly, a maladapted velocity storage can be corrected by a technique aimed to induce readaptation of the VOR.¹⁰ The original approach that combines OKS and an alternating head tilt maneuver has yielded successful outcomes in many patients with MdDS in multiple settings,^{19–23} and new but related approaches are being developed, including one that uses OKS alone to counter gravitational pull.²⁴

The case report¹ mentions that the patient responded well to an OKS therapy, although potentially informative details of this therapy are missing. Also unclear is whether the possibility of reducing or tapering off amitriptyline initially prescribed to the patient was considered at this point. Although data are lacking as to whether a nonmedication treatment such as VOR readaptation or cognitive-behavioral therapy can lead to reduced use of psychoactive medication previously prescribed for comorbid conditions that were exacerbated by MdDS, benefits of such

medication should be continually evaluated against risks associated with its long-term use.

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