An Unusual Association: Arnold Chiari Deformity and Meniere's Disease

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Abstract: Rare and fairly unknown, Arnold Chiari deformity is defined by the abnormally low position of the cerebellar tonsils that engage through the foramen magnum. Its association with the triad of tinnitus-hypoacusia-vertigo causes an authentic Meniere’s disease worth discussing. We report an unusual association of Arnold Chiari deformity with Meniere’s disease. A 46-years-old patient was diagnosed with left Meniere’s disease in 1994 on the classical diagnostic triad and the mode of progression: rotatory vertigo evolving by iterative crises; Intermittent buzzing tinnitus; left perception deafness with notion of wadded left ear. For a decade (1994 to 2015), he has been put under hygieno-dietary measures, vestibular re-education and medical treatment. The evolution was marked by the worsening of vertigo becoming progressively incapacitating, as well as an aggravation of the left deafness and persistence of tinnitus. Audiometry highlighted a severe endocochlear left perception deafness. The videoystagmography revealed a well-compensated left vestibular deficit. Cerebral magnetic resonance imaging (MRI) revealed a cerebellar tonsils’ ptosis through the foramen magnum. The diagnosis of Arnold-Chiari deformity associated with Meniere’s disease was then retained. The patient received a surgical left labyrinthectomy in 2015. The clinical course was uneventful, marked by the disappearance of vertigo twenty four months later. Association of Meniere’s disease and Arnold Chiari deformity is rare and must be diagnosed. These two pathologies being manifested by peripheral vertigo, a meticulous clinical and Para clinical examination is necessary to guide the diagnosis. Cerebrospinal fluid flow and pressure anomaly due to Arnold Chiari malformation can truly impact labyrinthine physiology, which explains the correlation between these two entities.

Keywords: Vertigo, Meniere's Disease, Arnold Chiari

1. Introduction

Arnold Chiari disease is due to a mismatch between the size of the skull and that of the brain leading to a malposition of the cerebellum. Rare and fairly unknown, it is defined by the abnormally low position of the cerebellar tonsils that engage for more than 5mm [1] through the foramen magnum itself malformed. It can go undetected throughout life or be revealed by very different symptoms [2, 3] the most frequent being: posterior headaches, the mosalgesic and tactile sensitivity disorders and audio-vestibular disorders.

In the other hand Meniere's disease is a chronic affection of internal ear whose reported prevalence rates have varied widely from 3.5/100 000 to 513/100 000 according to different authors [4]. Its etiology is uncertain. It has an unpredictable evolution, marked by recurrences of episodes of triad tinnitus-hypoacusia-vertigo known in authentic Meniere’s disease. It is a potentially an incapacitating affliction due to impact of vertigos on patient’s personal, social and professional life. Frequency and evolution of vertigo crisis are randomly: 6 to 11 crisis per year [5]. The diagnosis is primarily based on the clinical history. Clinicians also utilize various tests to confirm the diagnosis based on a consensual guidelines. [6-8]. Peripheral vertigo is common to Chiari deformity and Meniere’s disease, their semiological analysis is fundamental to the treatment. We report the case...
of a patient whose main symptom is vertigo and who presents the diagnostic criteria of both pathologies (Arnold-Chiari and Meniere). Our goal was to report the unusual association of these two pathologies and to expose the therapeutic conduct.

2. Case Report

![Audiometric exploration of the patient.](image)

*Figure 1. Audiometric exploration of the patient.*
A 46-years-old man, married with two children, was referred in 2015 to our university hospital center for incapacitating vertigo evolving for more than ten years. He has a history of smoking (20 Packs-year), dorsal scoliosis, a syringomyelia surgery and an appendectomy. The diagnosis left Meniere’s disease was made in 1994 based on the classic diagnostic triad and the mode of progression: rotatory vertigo evolving by iterative crises of more than 20 minutes (20 minutes to 6 hours) being repeated 2 to 3 times per week; Intermittent buzzing tinnitus; left Sensorineural hearing loss (SNHL) with notion of wadded left ear. For ten years (1994-2015), he was treated by hygienic and dietary measures (low sodium diet, alcohol, tobacco, and nicotine restriction; and adequate sleep); vestibular re-education and medical treatment (Acetyl-Leucine central anti-vertigo; Acetazolamide osmotic diuretic; Anxiolytics; Betahistine vasodilators). Evolution under this treatment was marked by the aggravation of vertigo becoming progressively incapacitating, as well as a worsening of the left deafness and persistence of tinnitus. An audio-vestibular check-up was done again as soon as he was admitted in August 2015. Audiometry were concordant and showed a severe endocochlear left perception deafness (average hearing loss of 70 dB predominant for high frequencies, and recruitment) [Figure 1].

Ipsilateral auditory evoked potentials were altered [Figure 2].

Evoked otolithic potentials confirmed the deficit on the left [Figure 3].
Videonystagmography showed well-compensated left vestibular areflexia [Figure 4].
Cerebral magnetic resonance imaging (MRI) revealed a 9.2 mm cerebellar tonsils’ ptosis through the foramen magnum [Figure 5].

The diagnosis of Arnold-Chiari deformity associated with Meniere’s disease was retained. Due to the severity of left hypoacousia and incapacitating vertigo, the patient received a left surgical labyrinthectomy in 2015. The evolution was favorable, marked by disappearance of vertigo and an expected left iatrogenic cophosis. Post-operative follow-up was of twenty four months.

3. Discussion

Type 1 Arnold Chiari deformity is defined as a cerebellar tonsils hernia through the foramen magnum of more than 5 millimeters. It is characterized by a malposition of the cerebellum that ends up into the spinal canal often associated with syringomyelia in 40% to 80% of cases [9, 10]. Its diagnosis is confirmed by magnetic resonance imaging especially on sagittal sections. When it becomes symptomatic, it is often revealed by a non-specific neurological syndrome including occipital headaches [11, 12] and sensory thermo-algesic disorders, balance disorders [13], and more rarely sleep disorders [14]. Its standard treatment is neurosurgical, based on the posterior pit decompression. This surgery is not without risk. [15].

The association with audio-vestibular disorders is not uncommon [16]. And the symptomatology can be menieriform [16]. In our clinical case the intensity of vertigo and the absence of other neurological disorders are not attributable to Arnold Chiari deformity.

The challenge for clinical medicine at this stage is not to ignore an associated Meniere’s disease that would require a specific treatment. Careful clinical and Paraclinical analysis is essential for diagnosis [17]. Recruitment on audiometry reinforces this diagnosis. It advocates for treatment of Meniere’s disease. The evolution was spread over more than a decade in two successive stages: iterative vertigo at the beginning then chronic and incapacitating vertigo. Labyrinthectomy and vestibular neurotomy are the most effective surgical procedures in case of incapacitating vertigo when it comes to Meniere’s disease. Trans-mastoid surgical labyrinthectomy presents less risk of complication than vestibular neurotomy [18]. We chose it because of the incapacitating vertigo and lack of effective hearing. In case audition needs to be preserved, the risk of iatrogenic cophosis due to labyrinthectomy makes endolymphatic sac decompression surgery a preferable option [19] or corticosteroids intratympanic injection [20]. In any diagnosed case of Meniere’s disease, it’s highly recommended to perform an MRI of the posterior pit and of the cervico-occipital hinge in both axial and sagittal planes in order to exclude other secondary causes of Chiari’s deformity [21].

Strengths and Weaknesses

Major strength of our case is the rare association between these two pathologies: Chiari’s deformity and Meniere’s disease. The article emphasizes the diagnostic process based on interrogatory, clinical examination then paraclinical tests. Indeed, taking care of vertigo crisis is sometimes tough, due to symptoms’ complexity and various etiologies. The main weakness of any case report remains its isolation, and we can’t jump to scientific conclusions. But this case stresses the importance of this possible association and arouses lector’s conscience on a meticulous diagnosis process.

4. Conclusion

Chiari deformity remains asymptomatic for a long time and discovery is often random. No sign is specific but the posterior cervicalgia, headache and balance disorders are very evocative. This clinical case draws attention to the vertiginous syndrome that can simulate a revelation mode. Clinical and Para clinical analysis led to the conclusion of an associated Meniere’s disease and adequate care could be provided. This clinical case draws our attention to the very wide field of etiologies for vertigo. Every clinician should remain critical when encountering vertiginous patients.

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