

Available online at www.sciencedirect.com

ScienceDirect

Journal of the Chinese Medical Association 77 (2014) 451–452

www.jcma-online.com

Editorial

Risk factors of meningioma



Meningioma is a common pathology in the central nervous system (CNS). It arises from the arachnoid cap cells of the arachnoid villi in the meninges, which envelop the brain and spinal cord. Therefore, a meningioma can occur anywhere from the vertex to the spine. It is often named according to its anatomical location. For example, a meningioma abutting the sphenoid bone between the frontal and temporal lobes is called a sphenoid ridge meningioma. A meningioma can be malignant, but it is usually benign and causes few symptoms until it reaches a large size. Because of the slow rate of tumor growth, a meningioma that is found incidentally may require no treatment other than periodic follow up. For symptomatic meningioma, an intradural extra-axial neoplasm can be managed with surgical resection when it produces a significant mass effect or can be managed with radiosurgery if its maximal diameter is <3 cm. With the advances in modern image technology (e.g., magnetic resonance images), contemporary neurosurgical practice usually obtains good results in the treatment of a meningioma.

The risk factors of meningioma remain elusive. Epidemiology studies have demonstrated a higher prevalence rate of meningioma in women.¹ Type 2 neurofibromatosis (NF) is frequently associated with multiple meningiomas, whereas type 1 NF is not associated with an increased risk of meningioma.² The most common genetic mutations involved in meningiomas are inactivation of the NF 2 gene on chromosome 22q. There are several other genes that may be involved in the development of meningioma such as the *AKT1*, *MNI*, and *PTEN* genes.^{3,4} Environmental causes, ionizing radiation, and exogenous hormones have also been correlated with intracranial meningioma in several previous studies.^{5–8} Despite these reports, the actual etiology and risk factors of meningioma in the CNS remain uncertain.

In this issue of the *Journal of the Chinese Medical Association*, there is an interesting article entitled “Risk of meningioma in patients with head injury: A nationwide population-based study”, by Dr. Kuan and colleagues.⁹ The authors should be commended for utilizing a large data analysis to investigate the association between head injury and meningioma.⁹ They compared 75,292 patients with head injury to a matched cohort without head injury (as the control). During the follow-up of approximately 430,000 person-years, they found 17 meningioma cases in the head injury cohort and 14 meningioma cases in the control cohort. Therefore, the authors concluded that head injury is an unlikely cause of meningioma.

However, there are several caveats to the study. The follow-up period in this cohort was relatively short and variable (e.g., an average of 5 years and ranging from 31 days to 10 years). Considering the fact that a meningioma is a slow-growing tumor, a longer follow-up is required to establish a correlation between it and various etiologies. The incidence rates of meningioma were as low as 3.23 per 100,000 person-years reported in the current study. With such low incidence rates, a case-controlled study design would be more appropriate to elucidate the causal effect of head injury. Furthermore, the heterogeneity of the head injury cohort is a significant concern. It is reasonable to infer that concussion, contusional brain hemorrhage, and diffuse axonal injury cause a great variety of effects in oncogenesis. These confounding factors would limit the power of this study. Further investigation is encouraged to explore the causes of meningioma and other oncology in the CNS.

Conflicts of interest

The author declares that there are no conflicts of interest to declare related to the subject matter or materials discussed in this article.

References

1. Wiemels J, Wrensch M, Claus EB. Epidemiology and etiology of meningioma. *J Neurooncol* 2010;**99**:307–14.
2. Goutagny S, Kalamarides M. Meningiomas and neurofibromatosis. *J Neurooncol* 2010;**99**:341–7.
3. Lekanne Deprez RH, Riegman PH, Groen NA, Warringa UL, van Biezen NA, Molijn AC, et al. Cloning and characterization of MN1, a gene from chromosome 22q11, which is disrupted by a balanced translocation in a meningioma. *Oncogene* 1995;**10**:1521–8.
4. Staal FJ, van der Luijt RB, Baert MR, van Drunen J, van Bakel H, Peters E, et al. A novel germline mutation of PTEN associated with brain tumours of multiple lineages. *Br J Cancer* 2002;**86**:1586–91.
5. Claus EB, Calvocoressi L, Bondy ML, Schildkraut JM, Wiemels JL, Wrensch M. Dental x-rays and risk of meningioma. *Cancer* 2012;**118**:4530–7.
6. Claus EB, Calvocoressi L, Bondy ML, Wrensch M, Wiemels JL, Schildkraut JM. Exogenous hormone use, reproductive factors, and risk of intracranial meningioma in females. *J Neurosurg* 2013;**118**:649–56.
7. Krampla W, Newrkla S, Pfisterer W, Jungwirth S, Fischer P, Leitha T, et al. Frequency and risk factors for meningioma in clinically healthy 75-year-old patients: results of the Transdanube Ageing Study (VITA). *Cancer* 2004;**100**:1208–12.
8. Longstreth Jr WT, Phillips LE, Drangsholt M, Koepsell TD, Custer BS, Gehrels JA, et al. Dental X-rays and the risk of intracranial meningioma: a population-based case-control study. *Cancer* 2004;**100**:1026–34.

9. Kuan AS, Chen YT, Teng CJ, Wang SJ, Chen MT. Risk of meningioma in patients with head injury: A nationwide population-based study. *J Chin Med Assoc* 2014;**77**:457–62.

Jau-Ching Wu*

Department of Neurosurgery, Neurological Institute, Taipei Veterans General Hospital, Taipei, Taiwan, ROC

National Yang-Ming University School of Medicine, Taipei, Taiwan, ROC

*Dr. Jau-Ching Wu, Department of Neurosurgery, Neurological Institute, Taipei Veterans General Hospital, 201, Section 2, Shih-Pai Road, Taipei 112, Taiwan, ROC.
E-mail address: jauching@gmail.com