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Idiopathic Intracranial Hypertension

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Deadline for manuscript submissions:

closed (31 July 2021)

Message from the Guest Editors

Idiopathic intracranial hypertension (IIH) is characterized by raised intracranial pressure (ICP) with no identifiable cause. There is a rising incidence and prevalence in this disease, and it appears that this is related to country-specific prevalence of obesity. It typically affects women of working age, and headache is the predominant morbidity in over 90%. Due to the nature of the disease, 10% have severe visual loss at presentation caused by papilledema, requiring neurosurgical intervention. The complexity of this disorder requires the expertise of a multiprofessional team to provide the best healthcare for these patients.

This Special Issue aims to highlight research from this multiprofessional perspective within the neurosciences. In this Special Issue, "Idiopathic Intracranial Hypertension", the readership will find research that reflects the interplay between basic research and clinical neurology, ophthalmology, and neurosurgery professionals at the leading edge of this growing research field.













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Message from the Editor-in-Chief

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