

INDIAN JOURNAL OF PRACTICAL PEDIATRICS

- IJPP is a quarterly subscription journal of the Indian Academy of Pediatrics committed to presenting practical pediatric issues and management updates in a simple and clear manner
- Indexed in Excerpta Medica, CABI Publishing, Scopus

Vol.22 No.1	JAN MAR. 2020
Dr.S.Thangavelu	Dr.T.L.Ratnakumari
Editor-in-Chief	Executive Editor

CONTENTS

TOPIC OF INTEREST - "NEUROLOGY "

Febrile fits	5
- Kumaresan G	
Stroke in children	9
- Vykuntaraju K Gowda, Balamurugan Nagarajan	
Acute disseminated encephalomyelitis in children	21
- Naveen Sankhyan, Priyanka Madaan, Chandana Bhagwat	
Metabolic encephalopathies	26
- Sangeetha Yoganathan, Bidkar Sayli Umakant	
Acute flaccid paralysis beyond polio - A case based approach	
- Mohammed Kunju PA, Ahamed Subir H, Merin Eapen	
Newer interventions in epilepsy management	46
- Ramalakshmi Ramiah	
Neuroimaging	55
- Leema Pauline	
Diagnostic neurophysiology in children	
- Lakshminarayanan Kannan	
Genetic testing in neurological disorders - Radiogenomics	78
- Sheffali Gulati, Sonali Singh, Rahul Sinha	

Journal Office and address for communications: Dr. S.Thangavelu, Editor-in-Chief, Indian Journal of Practical Pediatrics, 1A, Block II, Krsna Apartments, 50, Halls Road, Egmore, Chennai - 600 008. Tamil Nadu, India. Tel.No. : 044-28190032 E.mail : ijpp_iap@rediffmail.com

NEWS AND NOTES	85
CLIPPINGS	8,25,45,54,70,77,91,106,108
ADVERTISMENTS	111,112
- Cheran B	
Pediatrician in delivery room	107
MEDICOLEGAL MATTERS	
- Vijayalakshmi G, Balaji S, Raveendran J	
The dilated collecting system - 2	105
RADIOLOGY	
- Nair MKC, Leena Sumaraj, Swapna S, Sajitha Ja	asmine JR
Academic success - Student support and guida	nce 100
ADOLESCENCE	
- Mani Ram Krishna, Mohammed Farooq Kunde	
A clinical approach to syncope	92
GENERAL ARTICLE	
- Jeeson C Unni, Ranjit Baby Joseph	
Oxygen as a prescription	86
DRUG PROFILE	
Indian Journal of Practical Pediatrics	2020;22(1) : 2

FOR YOUR KIND ATTENTION

- * The views expressed by the authors do not necessarily reflect those of the sponsor or publisher. Although every care has been taken to ensure technical accuracy, no responsibility is accepted for errors or omissions.
- * The claims of the manufacturers and efficacy of the products advertised in the journal are the responsibility of the advertiser. The journal does not own any responsibility for the guarantee of the products advertised.
- * Part or whole of the material published in this issue may be reproduced with the note "Acknowledgement" to "Indian Journal of Practical Pediatrics" without prior permission.

- Editorial Board

Published by Dr. S.Thangavelu, Editor-in-Chief, IJPP, on behalf of Indian Academy of Pediatrics, from 1A, Block II, Krsna Apartments, 50, Halls Road, Egmore, Chennai - 600 008. Tamil Nadu, India and Printed by Mr. D.Ramanathan, at Alamu Printing Works, 9, Iyyah Street, Royapettah, Chennai-14.

FEBRILE FITS

*Kumaresan G

Abstract: Febrile fits is a common condition seen in day to day practice. The diagnosis is mainly clinical and there is limited indications for investigations. EEG often adds to confusion and is best avoided. The role of genetics is being recognised. Intracranial infections, febrile myoclonus, epileptic syndromes presenting initially as febrile seizures are to be considered in the differential diagnosis. Long term anti-convulsants should be avoided except in rare situations. Intermittent prophylaxis with clobazam is useful in reducing recurrences and parental anxiety.

Keywords: Febrile fits, Prognosis, Genetics, Differential diagnosis, Intermittent clobazam.

Points to Remember

- Febrile fits is a benign age related, self limiting condition.
- Clinical observation to exclude other conditions is most important than investigations.
- Early therapy to stop on-going seizure is important.
- *Hippocampal abnormalities can be both cause and effect of febrile fits in different situations.*
- Intermittent therapy is useful to minimise recurrences and parental anxiety.
- Restrict use of continuous anti-epileptic drugs.

References

- Byeon JH, Kim GH, Eun B. Prevalence, Incidence, and Recurrence of Febrile Seizures in Korean Children Based on National Registry Data. J Clin Neurol 2018; 14(1): 43-47.
- 2. Nelson KB, Ellenberg JH. Prognosis in children with febrile seizures. Pediatrics 1978; 61:720-727.
- 3. Verity CM, Butler NR, Golding J. Febrile convulsions in a national cohort followed up from birth. I-Prevalence and recurrence in the first five years of life. Br Med J (Clin Res Ed). 1985; 290(6478):1307-1310.
- 4. Vandenberg. Convulsive diseases in children. Pediatr Res 1969; 3:298-304.
- Meyer A, Falconer MA, Beck E. Pathological findings in temporal lobe epilepsy. J Neurol Neurosurg Psychiatry 1954; 17:276-285.
- Fernández G, Effenberger O, Vinz B, Steinlein O, Elger CE, Döhring W, et al Hippocampal malformation as a cause of familial febrile convulsions and subsequent hippocampal sclerosis. Neurology 1998; 50:909-917.
- Van Landingham KE, Heinz ER, Cavazos JE, Lewis DV. Magnetic resonance imaging evidence of hippocampal injury after prolonged focal febrile convulsions. Ann Neurol 1998; 43:413-426.
- Lewis DV, Shinnar S, Hesdorffer DC, Bagiella E, Bello JA, Chan S, et al. Hippocampal sclerosis after febrile status epilepticus: The FEBSTAT study. Ann Neurol 2014; 75:178-185.

email: jayalakshmi_kumaresan@hotmail.com

Professor of Pediatric Neurology (Retd.),
ICH & HC, Madras Medical College
Chennai.

- NIH Consensus Development Conference summary. Febrile seizures-long-term management of children with fever associated seizures. J Tenn Med Assoc 1981; 74(1):62-65.
- Mikati MA, Rahi AC. Febrile seizures From molecular biology to clinical practice. Neurosciences 2005; 10(1): 14-22.
- Chauhan U, Shanbag P, Mallad V. Febrile myoclonus: a missed clinical diagnosis. Indian J Pediatr 2013; 80(11):972-973.
- Khair AM, Elmagrabi D. Febrile seizures and febrile seizure syndromes: an updated overview of old and current knowledge. Neurol Res Int 2015. http://dx.doi.org/10.1155/ 2015/849341.
- Kimia AA. Utility of lumbar puncture for first febrile fits among Children of 6-18 months. Pediatrics 2009; 6:123-126.
- 14. Arzimanoglou A, Aicadi J, Guerrini R. Aicardiïs epilepsy in children. Lippincott Williams & Wilkins 2004; 223.
- 15. Winsley Rose, Chellam Kripakaran, Julies Xavier Scott. Intermittent Clobazam therapy in febrile seizures. Indian J Pediatr 2005; 72:31-33.
- Mohamed Sani Barghat AzzaKamal Al Shahwy Pediatric neurology 2019; 101:33-38.

STROKE IN CHILDREN

*Vykuntaraju K Gowda ** Balamurugan Nagarajan

Abstract: Pediatric stroke is an acute cerebrovascular event that occurs in children after 28 days of life up to 18 years of age. Pediatric stroke results in significant morbidity and mortality. Ischemic stroke can be due to arterial ischemia or venous sinus thrombosis. Hemorrhagic stroke is either due to non-traumatic, intra-parenchymal hemorrhage or subarachnoid hemorrhage. In young children, the symptoms could be non-specific. Stroke like conditions are very common, hence neuroimaging is mandatory for all cases of suspected stroke. Clinical awareness and recognition is crucial for diagnosis to ensure prompt management for better outcome.

Keywords: Stroke, Children, Pediatric stroke.

email: drknvraju08@gmail.com

Points to Remember

- Consider stroke in any child presenting with acute onset hemiparesis or focal deficit, change in mental status, headache, seizure or speech disturbance.
- MRI brain with MRA is the investigation of choice, but it is recommended at least that a CT brain is performed within one hour of arrival at hospital in every child. MR venography is done if cerebral venous sinus thrombosis is suspected.
- Aspirin has to be started in all cases of arterial ischemic stroke as soon as possible in the absence of contraindications except arterial dissection, cardio-embolic stroke and hyper-coagulable states.
- In CVST as well as in AIS caused by arterial dissection, cardio-embolic stroke and hypercoagulable states, anticoagulation using LMWH (enoxaparin) / un-fractionated heparin or oral warfarin is used.
- Role of thrombolysis in pediatric age group is controversial, however, there is growing evidence and looks promising.

- Sacco RL, KasnerSE, Broderick JP, Caplan LR, Connors JJ, Culebras A, et al. An updated definition of stroke for the 21st century: a statement for healthcare professionals from the American Heart Association/ American Stroke Association. Stroke 2013; 44:2064–2089.
- Ferriero DM, Fullerton HJ, Bernard TJ, Billinghurst L, Daniels SR, DeBaun MR,et al. Management of stroke in neonates and children: a scientific statement from the American Heart Association/American Stroke Association. Stroke 2019; 50:51-96.
- Kirton A, Deveber G, Pontigon AM, Macgregor D, Shroff M. Presumed perinatal ischemic stroke: vascular classification predicts outcomes. Ann Neurol 2008; 63:436–443.
- 4. Kirton A, Deveber G. Life after perinatal stroke. Stroke. 2013; 44:3265–3271.
- 5. Golomb MR, MacGregor DL, Domi T, Armstrong DC, McCrindle BW, Mayank S, et al. Presumed pre- or

^{*} Associate Professor of Pediatric Neurology

 ^{**} Senior Resident, Department of Pediatric Neurology, Indira Gandhi Institute of Child health, Bengaluru.

perinatal arterial ischemic stroke: risk factors and outcomes. Ann Neurol 2001; 50:163-168.

- Agarwal N, Johnston SC, Wu YW, Sidney S, Fullerton HJ. Imaging data reveal a higher pediatric stroke incidence than prior US estimates. Stroke 2009; 40:3415-3421.
- Lehman LL, Khoury JC, Taylor JM, Yeramaneni S, Sucharew H, Alwell K, et al. Pediatric stroke rates over 17 years: report from a population-based study. J Child Neurol 2018; 33:463-467.
- Mallick AA, Ganesan V, Kirkham FJ, Fallon P, Hedderly T, McShane T, et al. Childhood arterial ischaemic stroke incidence, presenting features, and risk factors: a prospective population-based study. Lancet Neurol 2014; 13(1):35–43.
- Goldenberg NA, Bernard TJ, Fullerton HJ, Gordon A, deVeber G. International Pediatric Stroke Study Group. Antithrombotic treatments, outcomes, and prognostic factors in acute childhood-onset arterial ischaemic stroke: a multi-centre, observational, cohort study.Lancet Neurol 2009; 8(12):1120-1127.
- Amlie-Lefond C, Guillaume S, Fullerton H. Recent developments inchildhood arterial ischemic stroke. Lancet Neurol 2008; 7:425-435.
- Kossorotoff M, Chabrier S, Tran Dong K, Nguyen The Tich S, Dinomais M. Arterial ischemic stroke in nonneonate children: Diagnostic and therapeutic specificities.Rev Neurol (Paris). 2019 Jun 7. pii: S0035-3787(18)30968 10.1016/j.neurol.2019.03.005.[Epub ahead of print]
- Kim T, Oh CW, Bang JS, Kim JE, Cho WS. Moyamoya disease: treatment and outcomes. J Stroke.2016; 18(1): 21-30.

- 13. Suzuki J, Takaku A. Cerebrovascular "moyamoya" disease. Disease showing abnormal net-like vessels in base of brain. Arch Neurol 1969;20:288-299.
- Gowda VK, Manjeri V, Srinivasan VM, Sajjan SV, Benakappa A. Mineralising angiopathy with basal ganglia stroke after minor trauma. Case series including two familial cases. J Pediatr Neurosci 2018; 13:448-454.
- 15. Yang FH, Wang H, Zhang JM, Liang HY.Clinical features and risk factors of cerebral infarction after mild head trauma under 18 months of age. Pediatr Neurol 2013; 48:220-226.
- Lingappa L, Varma RD, Siddaiahgari S, Konanki R. Mineralising angiopathy with infantile basal ganglia stroke after minor head trauma. Dev Med Child Neurol 2014;56:78-84.
- Pedneuroaiims: E-learningmodules. Cerebral Palsy and Other Neurodevelopmental Disorders. https:// pedneuroaiims.chalopadho.com/s/classroom/1/chapter/7 (Accessed on Nov 19, 2019)
- Stroke in childhood: Clinical guideline for diagnosis, management and rehabilitation (2017). Royal College of Pediatrics and Child Health. http://www.rcpch.ac.uk/ stroke-guideline (Accessed on November 18, 2019).
- Bhatia K, Kortman H, Blair C, Parker G, Brunacci D, Ang T et al. Mechanical thrombectomy in pediatric stroke: systematic review, individual patient data meta-analysis, and case series. J Neurosurg Pediatr. 2019 Aug 9:1-14. doi: 10.3171/2019.5.PEDS19126. [Epub ahead of print]
- 20. Schapkaitz E, Sherman GG, Jacobson BF, Haas S, Buller HR, Davies V, et al. South African Society of Thrombosis and Haemostasis. Paediatric anticoagulation guidelines. S Afr Med J 2012; 102: 171-175.

ACUTE DISSEMINATED ENCEPHALOMYELITIS IN CHILDREN

*Priyanka Madaan **Chandana Bhagwat ***Naveen Sankhyan

Abstract: Acute disseminated encephalomyelitis is a demyelinating inflammatory disorder characterized clinically by acute-onset polyfocal neurologic deficits and encephalopathy and fluffy white matter lesions radiologically. Antecedent factors include infections, vaccinations and others. Usually, a monophasic illness, recurrences should raise a suspicion of a relapsing disorder such as myelin oligodendrocyte glycoprotein associated demyelination or multiple sclerosis. Investigations are warranted to rule out other causes of encephalopathy. Management includes immunomodulation with high dose pulse methylprednisolone, intravenous immunoglobulin and / or plasmapheresis. Prognosis depends on the recovery after the acute stage and the risk of recurrent demyelination.

Keywords: Demyelination, Acute disseminated encephalomyelitis, Myelin oligodendrocyte glycoprotein associated demyelination, Neuroinflammation.

- * Senior Research Associate
- ** Senior Resident

*** Additional Professor, Pediatric Neurology Unit, Department of Pediatrics, Advanced Pediatrics Center, Post Graduate Institute of Medical Education and Research, Chandigarh. email: drnsankhyan@yahoo.co.in

Points to Remember

- ADEM is an inflammatory disorder of the brain, characterized by acute-onset polyfocal neurologic deficits and encephalopathy.
- Fluffy white matter demyelinating lesions are the typical MRI findings.
- ADEM is monophasic, but it may be the first presentation of related inflammatory disorders such as MOG associated demyelination.
- Treatment includes immunomodulation with pulse steroids resulting in a brisk improvement in most children.
- *IVIG* is initiated if there is no clinical improvement within seven days of completing pulse steroids.

- 1. Krupp LB, Banwell B, Tenembaum S. International Pediatric MS Study Group. Consensus definitions proposed for pediatric multiple sclerosis and related disorders. Neurology 2007; 68(16 suppl 2):S7-S12.
- Krupp LB, Tardieu M, Amato MP, Banwell B, Chitnis T, Dale RC, et al. International Pediatric Multiple Sclerosis Study Group criteria for pediatric multiple sclerosis and immune-mediated central nervous system demyelinating disorders: revisions to the 2007 definitions. Mult Scler 2013; 19:1261-1267.
- Cole J, Evans E, Mwangi M, Mar S. Acute Disseminated Encephalomyelitis in Children: An updated review based on current diagnostic criteria. Pediatr Neurol 2019, article in press. doi:10.1016/j.pediatrneurol.2019.06.017.
- 4. Pohl D, Alper G, Van Haren K, Kornberg AJ, Lucchinetti CF, Tenembaun S. Acute disseminated encephalomyelitis: Updates on an inflammatory CNS syndrome. Neurology 2016; 87:S38-S45.
- 5. Pavone P, Pettoello-Mantovano M, Le Pira A, Giardino I, Pulvirenti A, GiugnoR, et. al. Acute disseminated encephalomyelitis: a long-term prospective study and metaanalysis. Neuropediatrics 2010; 41(6):246-255.
- Tenembaum S, Chitnis T, Ness J, Hahn JS. Acute disseminated encephalomyelitis. International Pediatric MS Study Group. Neurology 2007; 68(16 suppl 2):S23-S36.

- Karussis D, Petrou P. The spectrum of post-vaccination inflammatory CNS demyelinating syndromes. Autoimmun Rev 2014; 13(3):215-224.
- Hennes EM, Baumann M, Schanda K, Anlar B, Bajer-Kornek B, Blaschek A, et al. Prognostic Relevance of MOG antibodies in children with an acquired demyelinating syndrome. Neurology 2017; 89:900-908.
- 9. Wingerchuk DM, Banwell B, Bennett JL, Cabre P, Carroll W, Chitnis T, et al. International consensus diagnostic criteria for neuromyelitis optica spectrum disorders. Neurology 2015; 85:177-189.
- Huppke P, Rostasy K, Karenfort M, Huppke B, Seidl R, Leiz S, et al. Acute disseminated encephalomyelitis followed by recurrent or monophasic optic neuritis in pediatric patients. MultScler 2013; 19:941-946.
- 11. Mikaeloff Y, Caridade G, Husson B, Suissa S, Tardieu M. Acute disseminated encephalomyelitis cohort study: prognostic factors for relapse. Eur J Paediatr Neurol 2007;11:90-95.
- Hart MN, K M Earle. Haemorrhagic and perivenous encephalitis: a clinical-pathological review of 38 cases. J Neurol Neurosurg Psychiatry 1975; 38:585-591.
- 13. Alper G. Acute disseminated encephalomyelitis. J Child Neurol 2012; 27(11):1408-1425.
- Hacohen Y, Absoud M, Deiva K, Hemingway C, Nytrova P, Woodhall M, et al. Myelin oligodendrocyte glycoprotein antibodies are associated with a non-MS course in children. Neurology - Neuroimmunology

Neuroinflammation 2015: 2. doi:10.1212/nxi.000000000 000081.

- 15. Young NP, Weinshenker BG, Parisi JE, Scheithauer B, Giannini C, Roemer SF, et al. Perivenous demyelination: association with clinically defined acute disseminated encephalomyelitis and comparison with pathologically confirmed Multiple Sclerosis. Brain 2010; 133:333-348.
- Manguinao M, Krysko KM, Maddike S, Rutatangwa A, Francisco C, Hart J, et al. A retrospective cohort study of plasma exchange in central nervous system demyelinating events in children. MultScler Relat Disord 2019; 35: 50-54.
- Llufriu S, Castillo J, Blanco Y, Ramió-Torrentà L, Río J, Vallès M, et al. Plasma exchange for acute attacks of CNS demyelination: Predictors of improvement at 6 months. Neurology 2009; 73:949-953.
- Pradhan S, Gupta RP, Shashank S, Pandey N. Intravenous immunoglobulin therapy in acute disseminated encephalomyelitis. J Neurol Sci 1999; 165:56-61.
- Ayed H, Chaudhary MW, AlBaradie R, Mir A. Use of cyclophosphamide in a child with fulminant acute disseminated encephalomyelitis. Child Neurol Open 2018; 5:2329048X18754631.
- 20. Hacohen Y, Wong YY, Lechner C, Jurynczyk M, Wright S, Konuskan B, et al. Disease course and treatment responses in children with relapsing myelin oligodendrocyte glycoprotein antibody-associated disease. JAMA Neurol 2018; 75:478-487.

METABOLIC ENCEPHALOPATHIES

*Sangeetha Yoganathan *Bidkar Sayli Umakant

Abstract: The etiologies of metabolic encephalopathy are often diverse in children. Encephalopathy could result from lack of glucose, vitamin cofactors or oxygen and end organ failure. Inborn errors of metabolism, hypoglycemia, dyselectrolytemia, endocrine disorders and Reye syndrome are the reported causes of metabolic encephalopathies in children and adolescents. The clinical manifestations, biochemical parameters and radiological findings vary according to the etiology. Early diagnosis and management lead to reversal of symptoms and can prevent long-term neurological sequelae.

Keywords: Metabolic encephalopathy, Inborn error of metabolism, Osmotic demyelination syndrome, Hepatic encephalopathy, Uremic encephalopathy.

 Post Doctoral Fellow in Pediatric Neurology, Department of Neurological Sciences, Christian Medical College, Vellore, Tamil Nadu.

email: doc_ys@yahoo.co.in

Points to Remember

- Metabolic encephalopathy should be suspected in any child with altered consciousness after excluding CNS infection, structural disorders, toxin ingestion and trauma.
- Organ or system failure like hepatic encephalopathy, hypoxia, dyselectrolytemia and endocrine dysfunction are responsible for metabolic encephalopathy.
- Underlying etiologies are diverse which can be narrowed down by recognizing the clinical clues.
- Management includes acute stabilization and specific measures based on etiology including organ support.

- 1. Perugula ML, Lippmann S. Encephalopathy or Psychosis? Innov Clin Neurosci 2016: 13; 41-42.
- 2. Angel MJ, Young GB. Metabolic encephalopathies. Neurol Clin 2011; 29:837-882.
- Parke, JT. Acute encephalopathies. In: McMillan JA, Feigin RD, De Angelis C, Jones MD (eds), Oski's Pediatrics. Principles and Practice, 4th edn. Philadelphia: Lippincott, Williams & Wilkins 2006; p2258.
- 4. Chiriboga CA. Acute toxic-metabolic encephalopathy in children. Available from:https.//www.uptodate.com/ contents/acute-toxic-metabolic-encephalopathy-in-children/.Accessed on 7th January, 2020.
- Butterworth RF. Metabolic Encephalopathies. In: Siegel GJ, Albers RW, Brady ST, Price DL (eds), Basic Neurochemistry: Molecular, Cellular and Medical Aspects. 7th edn Boston: Elsevier 2006; pp593-594.
- 6. King JD, Rosner MH. Osmotic demyelination syndrome. Am J Med Sci 2010; 339:561-567.
- Zunga PM, Farooq O, Dar MI, Dar IH, Rashid S, Rather AQ, et al. Extra pontine osmotic demyelination syndrome. Ann Neurosci 2015; 22:51-53.
- 8. Gandhi K. Approach to hypoglycemia in infants and children. Translational Pediatrics 2017; 6:408.
- 9. Saudubray JM, Nassogne MC, de Lonlay P, Touati G. Clinical approach to inherited metabolic disorders in neonates: an over-view. Semin Neonatol 2002; 7:3-15.

[#] Professor of Pediatric Neurology,

- Saudubray JM, Cazrola AG. Clinical approach to inborn errors of metabolism in Pediatrics. In: Saudubray JM, Baumgartner MR, Walter J editors. Inborn metabolic diseases diagnosis and treatment. 6th edn. Heidelberg: Springer Berlin 2016; pp33-57.
- 11. Lallas M, Desai J. Wernicke encephalopathy in children and adolescents. World J Pediatr 2014; 10:293-298.
- Arya R, Gulati S, Deopujari S. Management of hepatic encephalopathy in children. Postgrad Med J 2010; 86: 34-41.
- 13. Ryan JM, Shawcross DL. "Hepatic encephalopathy". Medicine 2011; 39:617-620.
- 14. Dara N, Sayyari AA, Imanzadeh F. Hepatic encephalopathy: early diagnosis in pediatric patients with cirrhosis. Iran J Child Neurol 2014; 8:1-11.
- 15. Ferenci P, Lockwood A, Mullen K, Tarter R, Weissenborn K, Blei AT, et al. Hepatic encephalopathydefinition, nomenclature, diagnosis, and quantification: final report of the working party at the 11th World Congresses of Gastroenterology, Vienna, 1998. Hepatology 2002; 35:716-721.
- 16. Bajaj JS, Cordoba J, Mullen KD, Amodio P, Shawcross DL, Butterworth RF, et al. International Society for Hepatic Encephalopathy and Nitrogen Metabolism (ISHEN). Review article: the design of clinical trials in hepatic encephalopathy-an International Society for Hepatic Encephalopathy and Nitrogen Metabolism (ISHEN) consensus statement. Aliment Pharmacol Ther 2011; 33:739-747.
- Chapman J, Arnold JK. Reye Syndrome. [Updated 2019 Jan 17]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2019 Jan. Available from: https://www.ncbi.nlm.nih.gov/books/NBK526101/. Accessed on 1st January, 2020.
- Centers for Disease Control and Prevention. Reye syndrome: 1990 clinical case definition. 1990. Available at http://www.cdc.gov. Accessed on 1st January, 2020.
- 19. Hurwitz ES, Nelson DB, Davis C, Morens D, Schonberger LB. National surveillance for Reye syndrome: a five-year review. Pediatrics 1982; 70:895-900.
- Zemaitis MR, Foris LA, Chandra S, Bashir K. Uremia. [Updated 2019 Jul 5]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2019 Jan. Available from: https://www.ncbi.nlm.nih.gov/books/NBK441859. Accessed on 2nd January, 2020.
- 21. Arnold R, Issar T, Krishnan AV, Pussell BA. Neurological complications in chronic kidney disease. JRSM cardiovascular disease. 2016;5:2048004016677687.
- Hinchey J, Chaves C, Appignani B, Breen J, Pao L, Wang A, et al. A reversible posterior leukoencephalopathy syndrome. N Engl J Med 1996; 334:494-500.
- 23. Lamy C, Oppenheim C, Meder JF, Mas JL. Neuroimaging in posterior reversible encephalopathy syndrome. J Neuroimaging 2004; 14:89-96.

- 24. Yu J. Endocrine disorders and the neurologic manifestations. Ann Pediatr Endocrinol Metab 2014; 19:184-190.
- EL-Mohandes N, Huecker MR. Pediatric Diabetic Ketoacidosis. [Updated 2019 Apr 1]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2019 Jan Available from: https://www.ncbi.nlm.nih.gov/ books/NBK470282/.Accessed on 3rd January, 2020.
- 26. Adeyinka A, Kondamudi NP. Hyperosmolar Hyperglycemic Nonketotic Coma (HHNC, Hyperosmolar Hyperglycemic Nonketotic Syndrome) [Updated 2019 Jun 3]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2019 Jan-. Available from: https:// www.ncbi.nlm.nih.gov/books/NBK482142/. Accessed on 3rd January, 2020.
- 27. Schiess N, Pardo CA. Hashimoto's encephalopathy. Ann N Y Acad Sci 2008; 1142:254-265.
- 28. Lee J, Yu HJ, Lee J. Hashimoto encephalopathy in pediatric patients: Homogeneity in clinical presentation and heterogeneity in antibody titers. Brain Dev. 2018; 40: 42-48.
- 29. Tucci V, Sokari T. The clinical manifestations, diagnosis, and treatment of adrenal emergencies. Emerg Med Clin North Am 2014; 32:465-484.
- 30. Bowden SA, Henry R. Pediatric adrenal insufficiency: diagnosis, management, and new therapies. International Journal of Pediatrics. 2018;2018.
- Kunze K. Metabolic encephalopathies. J Neurol 2002; 249:1150-1159.
- 32. Berisavac II, Jovanoviæ DR, Padjen VV, Ercegovac MD, Stanarèeviæ PD, Budimkiæ-Stefanoviæ MS, et al. How to recognize and treat metabolic encephalopathy in Neurology intensive care unit. Neurol India 2017; 65:123-128.
- Babanrao SA, Prahladan A, Kalidos K, Ramachandran K. Osmotic myelinolysis: Does extrapontinemyelinolysis precede central pontine myelinolysis? Report of two cases and review of literature. Indian J Radiol Imaging 2015; 25:177-183.
- Zuccoli G, Santa Cruz D, Bertolini M, Rovira A, Gallucci M, Carollo C, Pipitone N. MR imaging findings in 56 patients with Wernicke encephalopathy: nonalcoholics may differ from alcoholics. AJNR Am J Neuroradiol 2009; 30:171-176.
- Emeksiz S, Kutlu NO, Çaksen H, Alkan G, Yýkmaz HÞ, Tokgöz H. Posterior reversible encephalopathy syndrome in children: a case series. Turk Pediatri Ars 2016; 51:217-220.
- Kim DM, Lee IH, Song CJ. Uremic Encephalopathy: MR Imaging Findings and Clinical Correlation. AJNR Am J Neuroradiol 2016; 37:1604-1609.
- Rovira A, Alonso J, Córdoba J. MR imaging findings in hepatic encephalopathy. AJNR Am J Neuroradiol 2008; 29:1612-1621.

Indian Journal of Practical Pediatrics

- Singh P, Goraya JS, Gupta K, Saggar K, Ahluwalia A. Magnetic resonance imaging findings in Reye syndrome: case report and review of the literature. J Child Neurol. 2011; 26:1009-1014.
- 39. Lin CC. [EEG manifestations in metabolic encephalopathy]. Acta Neurol Taiwan 2005; 14:151-161.
- 40. Kaplan PW. The EEG in metabolic encephalopathy and coma. J Clin Neurophysiol 2004; 21:307-318.
- 41. Faigle R, Sutter R, Kaplan PW. The electroencephalography of encephalopathy in patients with endocrine and metabolic disorders. J Clin Neurophysiol 2013; 30:505-516.
- 42. Crocker JF, Bagnell PC. Reye's syndrome: a clinical review. Can Med Assoc J 1981; 124:375-382, 425.
- 43. Price A, Losek J, Jackson B. Hyperglycaemic hyperosmolar syndrome in children: Patient characteristics, diagnostic delays and associated complications. J Paediatr Child Health 2016; 52:80-84.

ACUTE FLACCID PARALYSIS BEYOND POLIO- A CASE BASED APPROACH

*Mohammed Kunju PA **Ahamed Subir H ***Merin Eapen

Abstract: Acute flaccid paralysis is a complex clinical syndrome that requires immediate and careful evaluation for making a diagnoses. Each case of acute flaccid paralysis is an emergency from both clinical as well as public health perspective. The precise knowledge of the etiology, underlying pathophysiology and concurrent changes have profound implications in the treatment and prognosis. With the eradication of polio, Gullian Barrie Syndrome has become the major acute flaccid paralysis. Seasonal occurrence of Gullain Barrie Syndrome with spurt of viral fever is also seen. However, the clinical features of polio must be taught to the younger residents since imported or vaccine associated polio can still occur. Better usage of Magnetiic resonance imaging scanning will help in establishing the diagnosis. Acute management of such patient with acute flaccid paralysis due to different causes in intensive care unit has become a necessity. Based on severity IVIg for Gullain Barrie Syndrome and Methyl prednisolone for transverse myelitis are now accepted protocols. We are still in the process of consolidating the eradication of polio by the endgame strategy from 2019-2023.

Keywords: Acute flaccid paralysis, Guillain-Barre Syndrome, Lower motor neuron localization, Transverse myelitis

- * Professor and Head, Dept of Pediatric Neurology, Medical College, Trivandrum
- ** Assistant Professor, Dept of Neurology, MES Medical College, Perinthalmanna, Malappuram
- *** Senior Resident Dept of Pediatric Neurology, Medical College, Trivandrum email : drpamkunju@gmail.com

Points to Remember

- Clinical features of polio must be taught to the younger residents as imported or vaccine associated polio can still occur
- GBS requires prompt diagnosis and management and it is the major AFP now because of the spurt of various viral infections.
- Transverse myelitis (TM) and traumatic neuritis are the other common causes of AFP.
- TM with long segment involvement may be mistaken for GBS because of lack of sensory level and prolonged spinal shock which may be due to enterovirus related TM, or NMO (neuromyelitis optica).
- In TM preservation of dorsal column (joint position sensation) → Anterior cord syndrome → Anterior spinal artery occlusion
- Rabies can present with features of GBS.

- Kasper DL, Fauci AS, Hauser SL, Longo DL, Jameson JL, Loscalzo J (eds). Harrisons Principles of Internal Medicine. 19th Ed. New Delhi: Mc Graw Hill; 2015.
- Campell WW. DeJong's The Neurologic Examination. 7th Ed. Philadelphia. Lippincot Williams Wilkins; 2012.
- Daroff RB, Jankovic J, Mazziotta JC, Pomeroy SL (editors). Bradley's Neurology in Clinical Practice. 7th Ed. Amsterdam Elsevier; 2015.
- World Health Organization. Acute flaccid paralysis. [Internet] 2013. Available at https://www.hpsc.ie > acute flaccid paralysis afp > guidance > File, 14207, en. Last accessed on January, 2020.
- Suresh S, Forgie S, Robinson J. Non-polio Enterovirus detection with acute flaccid paralysis: A systematic review. J Med Virol 2018; 90(1):3-7.
- Messacar K, Schreiner TL, Van Haren K, Yang M, Glaser CA, Tyler KL, et al. Acute flaccid myelitis: A clinical review of US cases 2012-2015. Ann Neurol 2016; 80(3):326-338.

Indian Journal of Practical Pediatrics

- Krishnan C, Kaplin AI, Pardo CA, Kerr DA, Keswani SC. Demyelinating disorders: update on transverse myelitis. Curr Neurol Neurosci Rep 2006; 6:236-243.
- Maloney JA, Mirsky DM, Messacar K, Dominguez SR, Schreiner T, Stence NV. MRI findings in children with acute flaccid paralysis and cranial nerve dysfunction occurring during the 2014 enterovirus D68 outbreak. AJNR Am J Neuroradiol 2015; 36(2):245-250.
- 9. Suresh S, Rawlinson WD, Andrews PI, Stelzer-Braid S. Global epidemiology of nonpolio enteroviruses causing severe neurological complications: A systematic review and meta-analysis. Rev Med Virol 2019; e2082-2087.

NEWER INTERVENTIONS IN EPILEPSY MANAGEMENT

*Ramalakshmi Ramiah

Abstract: Epilepsy is a global issue affecting about 70 million people among the world population. Nearly 80% of them live in low and middle-income countries with limited resources. Although highly advanced treatment is available in some countries, up to 90% of people with epilepsy are not adequately treated or are not treated with conventional antiepileptic therapy in resource limited countries.

This review will highlight a few of the newer advances in management of epilepsy in children. They include pharmacological interventions, ketogenic diet, early genetic diagnosis and newer model multi-disciplinary team management of children with epilepsy.

Keywords: Epilepsy, Treatment, Advances.

 * University Hospital of Coventry and Warwickshire NHS Trust, Coventry CV2 2DX, United Kindgom.
email: ramlucuk@googlemail.com

Points to Remember

- Epilepsy in children can differ from epilepsy in adults both in seizure type and epilepsy syndrome.
- Medical management of epilepsy is complex and has to be tailored to the individual patient. Monotherapy is generally preferred.
- If the monotherapy fails, it is considered preferable to try alternative monotherapy.
- Children who continue to have seizures on monotherapy are prescribed a long term second drug in addition.
- Pharmacotherapy with newer drugs and nonpharmacological therapy like ketogenic diet useful in certain resistant epilepsy.
- Genetic testing aids in diagnosis.

- World Health Organization. Epilepsy. Fact sheet. No. 999. 2012. Updated February 2016. http://www.who.int/ mediacentre/factsheets/fs999/en/. Accessed on 15th Jan, 2020.
- French JA. Refractory epilepsy: clinical overview. Epilepsia 2007; 48 Suppl 1:3-7
- NICE Epilepsies: diagnosis and management Clinical guideline [CG137] Published date: January 2012 Last updated: October 2019. 16th Jan, 2020.
- 4. French JA, Krauss GL, Wechsler RT, Wang XF, DiVentura B, Brandt C, et al. Perampanel for tonic-clonic seizures in idiopathic generalized epilepsy: a randomized trial. Neurology 2015; 85:950-957.
- Steinhoff BJ, Ben-Menachem E, Ryvlin P, Shorvon S, Kramer L, Satlin A, et al. Efficacy and safety of adjunctive perampanel for the treatment of refractory partial seizures: a pooled analysis of three phase III studies. Epilepsia 2013; 54:1481-1489.
- NDA 202834: FDA Approved Labeling Text for FYCOMPA [online]. Available at: http:// www.accessdata.fda.gov/drugsatfda_docs/label/2012/ 202834lbl.pdf.Accessed on 15th Jan, 2020.
- 7. Nichol K, Stott C, Jones N, Gray RA, Bazelot M, Whalley BJ. The proposed multimodal mechanism of action of cannabidiol (CBD) in epilepsy: modulation of

intracellular calcium and adenosine-mediated signaling (P5.5-007). Neurology 2019; 92(15 Supplement) P5.5-007.

- Devinsky OE, Marsh D, Friedman D, Thiele E, Laux L, Sullivan J, et al. Cannabidiol in patients with treatment resistant epilepsy: an open label interventional trial. Lancet Neurol2016; 15:270-278.
- 9. Rosenberg EC, Louik J, Conway E, Devinsky O, Friedman D.Quality of Life in Childhood Epilepsy in pediatric patients enrolled in a prospective, open-label clinical study with cannabidiol. Epilepsia 2017; 58:e96-e100.
- Thiele EA, Marsh D, French JA, Mazurkiewicz-Beldzinska M, Benbadis SR, Joshi C, et al. Cannabidiol in patients with seizures associated with Lennox-Gastut Syndrome (GWPCARE4); a randomized, double blind, placebo controlled phase 3 trial. Lancet 2018;391(10125) :1085-1096.
- 11. GW Pharmaceuticals Announces Positive Phase 3 Pivotal Study Results for Epidiolex[®] (cannabidiol) in the Treatment of Dravet Syndrome.
- 12. NICE guidelines Cannabidiol with clobazam for treating seizures associated with Dravet syndrome December 2019.
- 13. Nice guidelines Cannabis-based medicinal products [D] Evidence review for epilepsy 2019.
- 14. Neal EG, Chaffe H, Schwartz RH, Lawson MS, Edwards N, Fitzsimmons G, et al. The ketogenic diet for the treatment of childhood epilepsy: a randomised controlled trial. Lancet Neurol 2008; 7(6):500-506.
- 15. Lambrechts DA, de Kinderen RJ, Vles JS, de Louw AJ, Aldenkamp AP, Majoie HJ. A randomized controlled trial of the ketogenic diet in refractory childhood epilepsy. Acta Neurol Scand 2017; 135(2):231-239.
- Scholl-Burgi S, Holler A, Pichler K, Michel M, Haberlandt E, Karall D. Ketogenic diets in patients with inherited metabolic disorders. J Inherit Metab Dis 2015; 38(4):765-773.

- 17. Neal EG, Chaffe HM, Edwards N, Lawson MS, Schwartz RH, Cross JH. Growth of children onclassical and medium-chain triglyceride ketogenic diets. Pediatrics 2008; 122(2):e334-e340.
- Dressler A, Trimmel-Schwahofer P, Reithofer E, Gröppel G, Mühlebner A, Samueli S, et al. The ketogenic diet in infants – Advantages of early use. Epilepsy Research 2015; 116:53-58.
- Winawer MR, Shinnar S. Genetic epidemiology of epilepsy or what do we tell families? Epilepsia2005; 46(suppl 10):24-30.
- 20. Ottman R, Hauser WA, Barker-Cummings C, Lee JH, Risch N. Segregation analysis of cryptogenic epilepsy and an empirical test of the validity of the results. Am J Hum Genet 1997; 60:667-675.
- 21. Ostrander BE, Butterfield RJ, Pedersen BS, Farrell AJ, Layer RM, Ward A, et al. Whole-genome analysis for effective clinical diagnosis and gene discovery in early infantile epileptic encephalopathy. NPJ genomic medicine. 2018; 3(1):1-0.
- 22. England MJ, Liverman CT, Schultz AM, Strawbridge LM. Epilepsy across the spectrum: Promoting health and understanding: [Internet]. Inst Med Comm Public Heal Dimens Epilepsies. Washington, DC: NatlAcad Press; 2012.
- 23. Meads C, Burls A, Bradley P. Systematic reviews of specialist epilepsy services. Seizure 2002; 11:90-98.
- 24. Bradley PM, Lindsay B, Fleeman N. Care delivery and self management strategies for adults withepilepsy. Cochrane Database Syst Rev 2016; 2:CD006244.
- 25. Hawasli AH, Bandt SK, Hogan RE, Werner N, Leuthardt EC. Laser ablation as treatment strategy for medically refractory dominant insular epilepsy: therapeutic and functional considerations. Stereotact Funct Neurosurg 2014; 92(6):397-404.

NEUROIMAGING

*Leema Pauline

Abstract: *Availability of neuroimaging facilities has made* the evaluation of neurological problems easier in the last few decades. Computed tomography scan of brain is the initial choice in very sick children because of its wider availability, faster turnaround time and lower cost. Cranial ultrasonography is an important modality in the follow up of infants in the postnatal period, particulary in the evaluation of hypoxic ischemic encephalopathy, subependymal- periventricular- intraventricular hemorrhage and hydrocephalus. It is used as a point of care investigation by neonatologists. Absence of radiation exposure and precision makes magnetic resonance imaging the modality of choice in emergency situations. But the disadvantages are the need for sedation or brief anaesthesia, longer procedural time and cost. But benefits outweigh the disadvantages and additional tools like Magnetic resonance angiography, Magnetic resonance venography and Magnetic resonance spectroscopy add precious information for further evaluation.

Keywords: *Neuroimaging, Cranial ultrasonography, CT, MRI, MRV, MRA, MRS.*

email: leemapauline@rediffmail.com

Points to Remember

- Neuroimaging is an invaluable tool in the evaluation of neurological problems.
- Cranial ultrasonography is the most frequently used neuroimaging modality in the perinatal period, particulary in the evaluation of hypoxic ischemic encephalopathy, subependymal- periventricularintraventricular hemorrhage and hydrocephalus.
- *CT* brain is the first imaging modality in unstable patients as it is widely available for emergencies, has shorter imaging time and lower cost. However, CT is generally suboptimal for imaging of structures in the posterior fossa and brain stem.
- MRI is an indispensable tool in diverse CNS problems such as developmental anomalies, infections, neurocutaneous syndromes, demyelination and metabolic disorders.
- Constraints with MRI are the need for sedation in young infants and its contraindication in the presence of metallic devices and implants.
- MRA, MRV and MRS are additional facilities useful in identifying vascular and metabolic pathology.

- 1. Camprodon JA, Stern TA. Selecting neuroimaging techniques: a review for the clinician. Prim Care Companion CNS Disord 2013; 15(4):PCC.12f01490. doi:10.4088/PCC.12f01490.
- Shehadi WH. Contrast media adverse reactions: occurrence, recurrence, and distribution patterns. Radiology 1982; 143(1):11-17.
- Yousem DM, Grossman RI. Techniques in neuroimaging. In: Yousem DM, Grossman RI, editors. Neuroradiology: The Requisites. 3rd ed. Philadelphia, PA: Mosby; 2010.
- 4. Koeller KK, Shih RY. Viral and Prion Infections of the Central Nervous System: Radiologic-Pathologic Correlation. Radiographics 2017; 37:199-233.
- 5. Shankar SK, Mahadevan A, Kovoor JM. Neuropathology of viral infections of the central nervous system. Neuroimaging Clin N Am 2008; 18(1):19-39.

Professor of Pediatric Neurology, ICH & HC, Madras Medical College, Chennai.

- Mohan S, Jain KK, Arabi M, Shah GV. Imaging of meningitis and ventriculitis. Neuroimaging Clin N Am 2012; 22(4):557-583.
- Fukui MB, Williams RL, Mudigonda S. CT and MR imaging features of pyogenic ventriculitis. AJNR Am J Neuroradiol 2001; 22(8):1510-1516.
- Theron S, Andronikou S, Grobbelaar M, Steyn F, Mapukata A, du Plessis J. Localized basal meningeal enhancement in tuberculous meningitis. Pediatr Radiol 2006; 36(11): 1182-1185.
- da Rocha AJ, Maia ACJr, Ferreira NP, do Amaral LL. Granulomatous Diseases of the Central Nervous System. Top Magn Reson Imaging 2005; 16:155-187.
- Patkar D, Narang J, Yanamandala R, Lawande M, Shah GV. Central nervous system tuberculosis: pathophysiology and imaging findings. Neuroimaging Clin N Am 2012; 22(4):677-705.
- Wasay M, Kheleani BA, Moolani MK, Zaheer J, Pui M, Hasan S,et al. Brain CT and MRI findings in 100 consecutive patients with intracranial tuberculoma. J Neuroimaging 2003; 13:240-247.
- Gupta RK, Jena A, Sharma A, Guha DK, Khushu S, Gupta AK. MR imaging of intracranial tuberculomas. J Comput Assist Tomogr 1988; 12:280-285.
- Rajshekhar V, Haran RP, Prakash GS, Chandy MJ. Differentiating solitary small cysticercus granulomas and tuberculomas in patients with epilepsy. Clinical and computerized tomographic criteria. J Neurosurg 1993; 78:402-407.
- Noujaim SE, Rossi MD, Rao SK, Cacciarelli AA, Mendonca RA, Wang AM, Coelho FH. CT and MR imaging of neurocysticercosis. AJR Am J Roentgenol 1999; 173(6):1485-1490.
- 15. Herron J, Darrah R, Quaghebeur G. Intra-cranial manifestations of the neurocutaneous syndromes. Clin Radiol 2000; 55(2):82-98.
- Huisman TA, Wisser J, Martin E, Huch RK, Marincek B. Fetal magnetic resonance imaging of the central nervous system. Eur Radiol 2002; 12:1952-1961.
- Batura A, Sakarya ME. Congenital brain abnormalities: Pictorial essay. Eastern Journal of Medicine 2015; 20: 11-19.

- Simon EM, Barkovich AJ. Holoprosencephaly: new concepts. Magn Reson Imaging Clin North Am 2001; 9:149-164.
- Poretti A, Blaser SI, Lequin MH, Fatemi A, Meoded A, Northington FJ, et al. Neonatal neuroimaging findings in inborn errors of metabolism. J Magn Reson Imaging 2013; 37(2):294-312.
- 20. Barkovich AJ. An approach to MRI of metabolic disorders in children. J Neuroradiol 2007; 34(2):75-88.
- 21. Tenembaum S, Chamoles N, Fejerman N.Acute disseminated encephalomyelitis: a long-term follow-up study of 84 pediatric patients.Neurology 2002; 59(8): 1224-1231.
- 22. Dutra BG, da Rocha AJ, Nunes RH, Maia Júnior ACM. Neuromyelitis Optica Spectrum Disorders: Spectrum of MR Imaging Findings and Their Differential Diagnosis. RadioGraphics 2018; 38(1):169-193.
- 23. Dos Passos GR, Oliveira LM, da Costa BK, Apostolos-Pereira SL, Callegaro D, Fujihara K, et al. MOG-IgG-Associated Optic Neuritis, Encephalitis, and Myelitis: Lessons Learned From Neuromyelitis Optica Spectrum Disorder. Front NeuroL 2018; 9:217.
- 24. Khalaf A, Iv M, Fullerton H, Wintermark M. Pediatric Stroke Imaging. Pediatric Neurology 2018; 86:5-18.
- 25. Mirsky DM, Lefond CA, Krishnan P, Laughlin S, Lee S, Lehman L, et al. Pathways for Neuroimaging of Neonatal Stroke. Pediatr Neurol 2017; 69:11-23.
- 26. Wasay M, Azeemuddin M. Neuroimaging of cerebral venous thrombosis. J Neuroimaging 2005; 15:118-128.
- 27. Ford K, Sarwar M. Computed tomography of dural sinus thrombosis. AJNR Am J Neuroradiol 1981; 2:539-543.
- Bianchi D, Maeder P, Bogousslavsky J, Schnyder P, Meuli RA. Diagnosis of cerebral venous thrombosis with routine magnetic resonance: an update. Eur Neurol 1998; 40:179 -190.
- 29. Saposnik G, Barinagarrementeria F, Brown RD Jr, Bushnell CD, Cucchiara B, Cushman M, et al. Diagnosis and management of cerebral venous thrombosis: a statement for healthcare professionals from the American Heart Association/American Stroke Association. Stroke 2011; 42:1158-1192.

DIAGNOSTIC NEUROPHYSIOLOGY IN CHILDREN

*Lakshminarayanan Kannan

Abstract: Neurophysiological tests are an extension of the clinical examination and should be interpreted in the overall clinical context. In children they are often misinterpreted due to lack of experience. Normal EEG does not rule out epilepsy and an abnormal EEG per se is not diagnostic of epilepsy. EEG is not a confirmatory test but rather supports clinical diagnosis of epilepsy. Routine interictal EEG does not distinguish between epilepsy and epilepsy mimics. Nerve conduction study and needle electromyography are performed infrequently in the current era of genetic diagnosis.

Keywords: Pediatric electroencephalogram, Video-electroencephalogram, Electroencephalogram in pediatric intensive care unit, Electromyography.

Points to Remember

- Epilepsy is a clinical diagnosis and EEG is not a confirmatory test.
- Inter ictal EEG does not distinguish between seizures and seizure mimics.
- Some epileptiform patterns in EEG have low association with seizures in children.
- Video-EEG is the gold standard to confirm psychogenic non-epileptic attacks.
- Genetic testing is the investigation of choice in suspected cases of muscular dystrophy or spinal muscular atrophy; nerve conduction and EMG do not add value in these conditions.

- Smith SJM. EEG in the diagnosis, classification, and management of patients with epilepsy. J Neurol Neurosurg Psychiatry 2005; 76(Suppl II): ii2–ii7. doi: 10.1136/ jnnp.2005.069245.
- 2. Herman ST, Abend NS, Bleck TP, Chapman KE, Drislane FW, Emerson RG, et al. Consensus Statement on Continuous EEG in Critically III Adults and Children, Part I: Indications. J Clin Neurophysiol 2015; 32:87-95.
- Jain P, Sharma S, Sharma A, Goel S, Jose A, Aneja S. Efficacy and safety of oral triclofos as sedative for children undergoing sleep electroencephalogram: An observational study. J Pediatr Neurosci 2016; 11:105-108. doi:10.4103/ 1817-1745.187622.
- Kuratani J, Pearl PL, Sullivan LR, Riel-Romero RMS, Cheek J, Stecker MM, et al. American Clinical Neurophysiology Society Guideline 5: Minimum Technical Standards for Pediatric Electroencephalography. Neuro diagn J 2016; 56:266-275. doi: 10.1080/21646821.2016. 1245568.
- Elewa MK, Mostafa MA. The subclinical epileptiform discharges among nonepileptic cerebral palsy patients. Egypt J Neurol Psychiatry Neurosurg 2016; 53:268-273. Available from: http://www.ejnpn.eg.net/text.asp?2016/53/ 4/268/202379
- 6. So EL. Interictal epileptiform discharges in persons without a history of seizures: What do they mean? J Clin Neurophysiol 2010; 27:229-238.

Pediatric Neurologist and Epileptologist, Rainbow Children's Hospital, Chennai.
email: dr kln@yahoo.co.in

- Riquet A, Lamblin M, Bastos M, Bulteau C, Derambure P, Vallée L, et al. Usefulness of video-EEG monitoring in children. Seizure 2011; 20:18-22. Available from https:// doi.org/10.1016/j.seizure.2010.09.011
- Pitt MC. Nerve conduction studies and needle EMG in very small children. European Journal of Paediatric Neurology 2012; 16:285-291. https://doi.org/10.1016/ j.ejpn.2011.07.014
- 9. Hyllienmark L, Ludvigsson J, Brismar T. Normal values of nerve conduction in children and adolescents. Electroencephalography and Clinical Neurophysiology/ Electromyography and Motor Control 1995; 97:208-214.
- Heise CO, Siqueira MG, Martins RS, Gherpelli JL. Clinical-Electromyography Correlation in Infants with Obstetric Brachial Plexopathy. The Journal of Hand Surgery 2007; 32:999-1004. https://doi.org/10.1016/j. jhsa.2007.05.002
- 11. Mathur N, Dhawan R. An alternative strategy for universal infant hearing screening in tertiary hospitals with a high delivery rate, within a developing country, using transient evoked oto-acoustic emissions and brainstem evoked response audiometry. J Laryngol Otol 2007; 121:639-643. doi:10.1017/S0022215106004403.
- 12. Odom JV, Bach M, Brigell M, Holder GE, McCulloch DL, Tormene AP, et al. ISCEV standard for clinical visual evoked potentials (2009 update). Doc Ophthalmol 2010; 120:111-119. doi: 10.1007/s10633-009-9195-4.
- Suppiej A, Gaspa G, Cappellari A, Toldo I, Battistella PA. The Role of Visual Evoked Potentials in the Differential Diagnosis of Functional Visual Loss and Optic Neuritis in Children. J Child Neurol 2011; 26:58-64. https://doi.org/ 10.1177/0883073810373947.

GENETIC TESTING IN NEUROLOGICAL DISORDERS - RADIOGENOMICS

*Sheffali Gulati **Sonali Singh **Rahul Sinha

Abstract: *"Radiogenomics" is a new emerging field which* correlate imaging features with genotype of disease. It has potential to play an important role in medicine, particularly neuro-oncology, metabolic and neurodegenerative disorders. Genetic testing confirms the diagnosis, helps in prognostication, predicts the risk of recurrence. But selecting the right genetic test is of prime concern and is difficult. Radiological finding or radio phenotype is useful in selecting the genetic tests needed. Role of radiogenomics is bidirectional. It predicts genotype on the basis of radiological phenotype and vice versa. So it plays an important role in bridging the gap between phenotype and genotype. It will further help the clinician to go for targeted sequencing of the gene which will be cost effective and time saving. For example the pediatric tumours which are studied on the basis of radiogenomics are medulloblastoma and glioblastoma. New generation genomic sequencing is very useful in sequencing the DNA at faster rate and lower cost. Whole genome sequencing, whole exome sequencing, clinical exome sequencing and target gene panel sequencing are various tests which are available. Radiogenomics is an important tool to indicate the likely genotype and directs towards the right genetic investigation.

Keywords: *Whole genome sequencing, Whole exome sequencing, Gene panel sequencing, Radio genomics.*

* Chief,

Child Neurology Division, Center of Excellence and Advanced Research on Childhood Neurodevelopmental Disorders

** Senior Resident, D.M Pediatric Neurology, Department of Pediatrics, AIIMS, New Delhi.

email:sheffaligulati@gmail.com

Points to Remember

- "Radiogenomics" is a new emerging field which correlates the imaging characteristics of a disease (radiophenotype), with its (genotype) genetic or molecular features.
- Next generation genetic sequencing is a technological advance which enable sequencing of million base pairs of DNA at a faster rate and lower cost.
- Whole genome sequencing, whole exome sequencing, clinical exome sequencing and target gene panel sequencing are various tests which are available.
- Knowledge about the various features of these tests is essential to decide the specific test needed.
- Good clinical phenotyping remains the corner stone of deciding the appropriate genetic test and increasing likelihood of a positive yield. But radiophenotype helps bridging the gap between clinical phenotype and genotype, by helping to select the type of genetic test is needed.

- Balmaceda C, Critchell D, Mao X, Cheung K, Pannullo S, DeLaPaz RL, et al. Multisection 1H magnetic resonance spectroscopic imaging assessment of glioma response to chemotherapy. J Neurooncol 2006; 76(2):185-191.
- Yates LR, Gerstung M, Knappskog S, Desmedt C, Gundem G, Van Loo P, Aas T, et al. Subclonal diversification of primary breast cancer revealed by multiregion sequencing. Nat Med 2015; 21(7):751e9.
- 3. Lambin P, Rios-Velazquez E, Leijenaar R, Carvalho S, van Stiphout RG, Granton P, et al. Radiomics: extracting more information from medical images using advanced feature analysis. Eur J Cancer 2012; 48(04):441-446
- 4. Shashi V, McConkie-Rosell A, Rosell B, Schoch K, Vellore K, McDonald M, et al. The utility of the traditional medical genetics diagnostic evaluation in the context of next-generation sequencing for undiagnosed genetic disorders. Genet Med 2014; 16:176-182.
- Venkatesh HS, Chaumeil MM, Ward CS, Haas-Kogan DA, James CD, Ronen SM. Reduced phosphocholine and hyperpolarized lactate provide magnetic resonance

biomarkers of PI3K/Akt/mTOR inhibition in glioblastoma. Neuro Oncol 2012; 14(3):315-325.

- Sudhakar SV, Muthusamy K, Arunachal G, Shroff M. Genomics and radiogenomics in inherited neurometabolic disorders – a practical primer for pediatricians. Indian J Pediatr 2019.
- Kruer MC, Boddaert N. Neurodegeneration with brain iron accumulation: a diagnostic algorithm. SeminPediatr Neurol 2012; 19:67-74.
- Kim JH, Kim HJ. Childhood X-linked adrenoleukodystrophy: clinical-pathologic overview and MR imaging manifestations at initial evaluation and follow-up. Radiographics 2005; 25:619-631.
- Provenzale JM, Peddi S, Kurtzberg J, Poe MD, Mukundan S, Escolar M. Correlation of neurodevelopmental features and MRI findings in infantile Krabbe'sdisease. Am J Roentgenol 2009; 192:59-65.
- 10. Van der Knaap MS, Valk J. Magnetic Resonance of Myelination and Myelin Disorders. Berlin:Springer 2005.
- Nunes J, Loureiro S, Carvalho S, Pais RP, Alfaiate C, Faria A, et al. Brain MRI findings as an important diagnostic clue in glutaric aciduria type1.Neuroradiol J 2013; 26:155-161.
- 12. Vasquez ML, RenaultIZ. Understanding genetics in neuro imaging. Neuroimag Clin N Am 2015; 25:1-16.
- Steenweg ME, Salomons GS, Yapici Z, Uziel G, Scalais E, Zafeiriou DI, et al. L-2-hydroxyglutaric aciduria: pattern of MR imaging abnormalities in 56 patients. Radiology 2009; 251:856-865.
- Menkes, J. H. Kinky hair disease. Pediatrics 1972; 50: 181-182.
- D'Arco F, Alves CA, Raybaud C, Chong WKK, Ishak GE, Ramji S, et al. Expanding the distinctive neuroimaging phenotype of ACTA2 mutations. Am J Neuroradiol 2018; 39:2126-2123.
- Van der Knaap MS, Salomons GS. Leukoencephalopathy with Brain Stem and Spinal Cord Involvement and Lactate Elevation. 2010May25[Updated2015Feb12]. In:AdamMP, Ardinger HH, Pagon RA, et al., editors. GeneReviews® [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2018.
- 17. Biesecker LG, Green RC. Diagnostic clinical genome and exome sequencing. N Engl J Med 2014; 370:2418-2425.

DRUG PROFILE

OXYGEN AS A PRESCRIPTION

*Jeeson C Unni **Ranjit Baby Joseph

Abstract: Oxygen is one of the most common life saving drugs widely used all over the world. It is often needed in many of the acute respiratory emergencies such as pneumonia and asthma where there is a risk of hypoxia. Long term oxygen therapy may be indicated in chronic respiratory conditions such as bronchopulmonary dysplasia (BPD), cystic fibrosis (CF), sleep disordered breathing, interstitial lung disease and pulmonary hypertension. Since medical grade oxygen is classified as a drug with specific biochemical and physiologic actions, with a distinct range of effective doses and well-defined adverse effects at high doses, oxygen needs to be prescribed like any other medication with specifications regarding dose, duration and method of delivery.

Keywords: Oxygen, Prescription, Delivery devices.

** Senior Specialist, Sr. Assc. Consultant, Aster Medcity, Kochi.

email: jeeson1955@gmail.com

Points to Remember

- Oxygen must be considered a medication that warrants a documented prescription before administration, except in emergency situations where written prescription is not mandatory to initiate the therapy
- The prescription should include indication, target saturation range, mode of delivery and flow rate.
- Choice of oxygen delivery device is based on clinical decision.

- Partington JR. A short history of chemistry, 3rd edn. New York: Dover Publications; 1989:p90.
- 2. Hess J. Oxygen unit for premature and very young infants. Am J Dis Child 1934; 47: 916-917.
- Weinberger B, Laskin DL, Heck DE, Laskin JD. Oxygen toxicity in premature infants. Toxicol Appl Pharmacol 2002; 181(1): 60-67.
- O'Driscoll BR, Howard LS, Davison AG. British Thoracic society. Guideline for emergency oxygen use in adult patients. Thorax. 2008; 63: Suppl VI; 45-47.
- National health service 2009. Oxygen safety in hospitals. Available from http://www.nrls.npsa.nhs.uk/alerts/? entryid45=62811. Last accessed on 31st January 2011.
- 6. Pilcher J, Beasley R. Acute use of oxygen therapy. Aust Prescr. 2015; 38 (3): 98-100.
- Weaver LK. Hyperbaric oxygen therapy for carbon monoxide poisoning. Undersea Hyperb Med 2014; 41(4):339-354.
- 8. Airway disease, obstructive. In BMJ group and the royal pharmaceutical society of Great Britain. British National Formulary 71, 2006;pp216-217.
- Perrone S, Bracciali C, Di Virgilio N, Buonocore G. Oxygen Use in Neonatal Care: A Two-edged Sword. Front Pediatr. 2017; 4: 143. Published 2017 Jan 9. doi:10.3389/ fped.2016.00143
- 10. Lindford AJ, Tehrani H, Sassoon EM, O'Neill TJ. Home oxygen therapy and cigarette smoking: a dangerous practice. Ann Burns Fire Disasters. 2006; 19(2):99-100.
- Adde FV, Alvarez AE, Barbisan BN, Guimarães BR. Recommendations for long-term home oxygen therapy in children and adolescents. JPediatr (Rio J) 2013; 89(1):6-17. doi: 10.1016/j.jped.2013.02.003.

^{*} Editor-in-Chief, IAP Drug Formulary

- Inwald D, Roland M, Kuitert L, McKenzie SA, Petros A. Oxygen treatment for acute severe asthma. BMJ 2001; 323(7304):98-100.
- 13. Tang AW. A practical guide to anaphylaxis. Am Fam Physician 2003; 68:1325-1332.
- 14. Tasker RC. Emergency treatment of acute seizures and status epilepticus. Arch Dis Child 1998; 79:78-83.
- 15. Gérardin M, Couec ML, Grall-Bronnec M, Feuillet F, Wainstein L, Rousselet M, et al. PHEDRE trial protocolobservational study of the prevalence of problematic use of Equimolar Mixture of Oxygen and Nitrous Oxide (EMONO) and analgesics in the French sickle-cell disease population. BMC Psychiatry 2015; 15:281.
- Hönemann C, Hagemann O, Doll D. Inhalational anaesthesia with low fresh gas flow. Indian J Anaesth 2013; 57(4):345-350.
- 17. Cope C. The Importance of Oxygen in the Treatment of Cyanide Poisoning. JAMA. 1961; 175(12):1061-1064.
- Hardavella G, Karampinis I, Frille A, Sreter K, Rousalova I. Oxygen devices and delivery systems. Breathe 2019; 15: 108-116.
- Nippers I, Sutton A. Oxygen therapy: professional compliance with national guidelines. Br J Nurs 2014; 23(7):382-386.
- Derek S. Wheeler, Hector R. Wong. Management of the pediatric airway. Pediatric Critical Care Medicine: Basic Science And Clinical Evidence. Springer; 2007;pp47-48.
- 21. Peter C, Poets CF. Prescription of home oxygen therapy to infants in Germany. Biol Neonate 2001; 80:148-151.
- 22. Walsh BK, Smallwood CD. Pediatric Oxygen Therapy: A Review and Update. Respiratory Care 2017; 62(6): 645-661.
- Myers TR, American Association for Respiratory Care. AARC clinical practice guideline: selection of an oxygen delivery device for neonatal and pediatric patients: 2002 revision and update. Respir Care 2002; 47(6):707-716.
- World Health Organisation. Oxygen therapy for acute respiratory infections in young children in developing countries. Geneva: World Health Organisation, 1993: WHO/ARI 93.28. Available at https://apps.who.int/iris/ handle/10665/60985. Last accessed on 15th February, 2020.
- Guilfoile T, Dabe K. Nasal catheter oxygen therapy for infants. Respir Care1981; 26:35-40.

- 26. Udani S. Oxygen therapy. Principles of pediatric and neonatal emergencies, 3rd edn. Jaypee publications; 2011; 52-57.
- 27. Uygur P, Oktem S, Boran P, Tutar E, Tokuc G. Low- versus high-flow oxygen delivery systems in children with lower respiratory infection. Pediatr Int 2016; 58(1):49-52.
- 28. Gupta S, Donn SM. Continuous positive airway pressure: Physiology and comparison of devices. Semin Fetal Neonatal Med 2016; 21(3):204-211.
- Pinto VL, Sharma S. Continuous Positive Airway Pressure (CPAP) [Updated 2019 May 11]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2020 Jan-. Available from: https://www.ncbi.nlm.nih.gov/books/ NBK482178/ Last accessed on 20th February,2020.
- Mikalsen IB, Davis P, Øymar K. High flow nasal cannula in children: a literature review. Scand J Trauma Resusc Emerg Med 2016; 24:93.
- 31. Dani C, Pratesi S, Migliori C, Bertini G. High flow nasal cannula therapy as respiratory support in the preterm infant. Pediatr Pulmonol 2009; 44(7):629-634.
- 32. Holleman-Duray D, Kaupie D, Weiss MG. Heated humidified highflow nasal cannula: use and a neonatal early extubation protocol. J Perinatol 2007; 27(12):776-781.
- 33. Marino BS, Tabbutt S, MacLaren G, Hazinski MF, Adatia I. Atkins DL, et al. Cardiopulmonary Resuscitation in Infants and Children With Cardiac Disease. A Scientific Statement From the American Heart Association. Circulation 2018; 137:e691-e782.
- 34. Smith LL. Mechanism of paraquat toxicity in lung and its relevance to treatment. Hum Toxicol 1987; 6:31-36.
- Cersosimo RJ, Matthews SJ, Hong WK. Bleomycin pneumonitis potentiated by oxygen administration. Drug Intell Clin Pharm 1985; 19:921-923.
- 36. Abdo WF, Heunks LM. Oxygen-induced hypercapnia in COPD: myths and facts. Crit Care 2012; 16(5):323.
- 37. Perrin K, Wijesinghe M, Healy B, Wadsworth K, Bowditch R, Bibby S, et al. Randomised controlled trial of high concentration versus titrated oxygen therapy in severe exacerbations of asthma. Thorax 2011; 66:937-941.
- 38. Jobe AH, Kallapur SG. Long term consequences of oxygen therapy in the neonatal period. Semin Fetal Neonatal Med 2010; 15(4):230-235.

GENERAL ARTICLES

A CLINICAL APPROACH TO SYNCOPE

*Mani Ram Krishna **Mohammed Farooq Kunde

Abstract: Syncope is a common clinical complaint that a pediatrician encounters in the outpatient clinic or in the emergency room. The causes of syncope include autonomic disturbances, cardiovascular causes and neurological problems. Autonomic syncope is the commonest and is usually benign. Cardiovascular causes can potentially be life threatening and it is important to recognize them and refer these children to an appropriate specialist in a timely fashion. It is possible to identify the cause of syncope in most patients with a detailed history and physical examination. In this chapter, we list the causes of syncope and attempt to provide a clinical approach that will permit accurate triage of patients with syncope by a pediatrician.

Keywords: *Syncope, Neurocardiogenic syncope, Cardiac arrhythmia, Fainting, Convulsive syncope.*

** Resident in Pediatric Cardiology, Department of Pediatric Cardiology, Amrita Institute of Medical Sciences, Kochi. email: mann_comp@hotmail.com

Points to Remember

- Syncope is an important clinical problem in the pediatric age group attending ED.
- There are three groups of causes, autonomic instability which is usually benign and more common, cardiac cause which are serious but less common and others including neurological causes (e.g. convulsive syncope) which are rare.
- The differentiation is usually possible by a detailed history, physical examination and basic investigation like ECG.
- ECG is an inexpensive investigation that probably has the highest diagnostic yield in the evaluation of syncope.
- CT has the lowest diagnostic yield and can be replaced by a good focused neurological examination.
- Unnecessary investigations can be avoided and diagnostic yield can be increased if the pediatrician meticulously takes the history and performs the clinical examination.

- Shen WK, Sheldon RS, Benditt DG, Cohen MI, Forman DE, Goldberger ZD, et al. 2017 ACC/AHA/HRS Guideline for the Evaluation and Management of Patients With Syncope: Executive Summary: A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines and the Heart Rhythm Society. J Am Coll Cardiol 2017; 70(5):620-663.
- McHarg ML, Shinnar S, Rascoff H, Walsh CA. Syncope in childhood. Pediatr Cardiol 1997; 18(5):367-371.
- Anderson JB, Czosek RJ, Cnota J, Meganathan K, Knilans TK, Heaton PC. Pediatric syncope: National Hospital Ambulatory Medical Care Survey results. J Emerg Med 2012; 43(4):575-583.
- Kanjwal K, Calkins H. Syncope in children and adolescents. Cardiac Electrophysiology Clinics 2013; 5:443-455.
- Grubb BP. Clinical practice. Neurocardiogenic syncope. N Engl J Med 2005; 352(10):1004-1010.

^{*} Clinical Assistant Professor

- 6. Behere SP, Weindling SN. Inherited arrhythmias: The cardiac channelopathies. Ann Pediatr Cardiol 2015; 8(3):210-220.
- 7. Sheldon R. How to differentiate syncope from seizure. Card Electrophysiol Clin 2013; 5(4):423-431.
- 8. Strickberger SA, Benson DW, Biaggioni I, Callans DJ, Cohen MI, Ellenbogen KA, et al. AHA/ACCF Scientific Statement on the evaluation of syncope: from the American Heart Association Councils on Clinical Cardiology, Cardiovascular Nursing, Cardiovascular Disease in the Young, and Stroke, and the Quality of Care and Outcomes Research Interdisciplinary Working Group; and the American College of Cardiology Foundation: in collaboration with the Heart Rhythm Society: endorsed by the American Autonomic Society. Circulation 2006; 113(2):316-27.
- 9. Hurst D, Hirsh DA, Oster ME, Ehrlich A, Campbell R, Mahle WT, et al. Syncope in the Pediatric Emergency Department - Can We Predict Cardiac Disease Based on History Alone? J Emerg Med 2015; 49(1):1-7.
- Colman N, Bakker A, Linzer M, Reitsma JB, Wieling W, Wilde AA. Value of history-taking in syncope patients: in whom to suspect long QT syndrome? Europace 2009; 11(7):937-943.
- Johnson ER, Etheridge SP, Minich LL, Bardsley T, Heywood M, Menon SC. Practice variation and resource use in the evaluation of pediatric vasovagal syncope: are pediatric cardiologists over-testing? Pediatr Cardiol 2014; 35(5):753-758.
- 12. Tretter JT, Kavey RE. Distinguishing cardiac syncope from vasovagal syncope in a referral population. J Pediatr 2013; 163(6):1618-1623 e1.
- Goble MM, Benitez C, Baumgardner M, Fenske K. ED management of pediatric syncope: searching for a rationale. Am J Emerg Med 2008; 26(1):66-70.

ADOLESCENCE

ACADEMIC SUCCESS - STUDENT SUPPORT AND GUIDANCE

*Nair MKC **Leena Sumaraj ***Swapna S ***Sajitha Jasmine JR

Abstract: Academic success is paramount in any school/ college program. The faculty spent a lot of time to teach many subjects, but often do not guide the students how to study. Poor scholastic performance of school going children is a problem that affect many parents. *The predictor variables for poor scholastic performance* were; not studying daily lessons, poor concentration in studies, lower education status of father and unhappy family. Early guidance and support may stop students experiencing a cycle of failure. Key to supporting struggling students is to identify reasons for poor performance. Many students lack basic academic skills and do not know how to learn effectively. The transition from school to professional education may affect students emotionally, socially and academically. Recall of information is essential for successful performance in examinations. Better recall can be achieved by time management of study periods and regular systematic learning. Prepare a revision timetable and set out what topics, subjects you want to cover each day.

Keywords: Academic success, Scholastic performance, Student support, Enhancing memory.

Points to Remember

- In addition to teaching individual subjects, every faculty should guide the students on method of study.
- This is implemented by organizing student support and guidance program with specific strategies.
- * Formerly Vice Chancellor, Kerala University of Health Sciences and Director
- ** Senior Research Coordinator, CDC
- *** Developmental therapist, NIMS-Spectrum - Child Development Research Centre (CDRC), Kerala. email: cdcmkc@gmail.com

- Faculty should identify reasons for student's failure, and offer specific help.
- *Time management by balancing study and leisure time activities.*
- Students should be taught about preparing for different type of questions namely multiple choice, short answers and essay.
- Organizing the time, practicing memory enhancing methods, taking balanced diets, controlling the stress, discussing with teachers, combined study and organized revision will help succeed in exam.

- Nair MKC, Paul MK, Padmamohan J. Scholastic performance of adolescents. Indian J Pediatr 2003; 70(8):629-631.
- Krishnakumar P, Jisha AM, Sukumaran SK, Nair MKC. Developing a model for resource room training for slow learners in normal schools. Indian J Psychiatry 2011; 53:336-339.
- 3. Nair MKC, Paul MK, John R. Prevalence of depression among adolescents. Indian J Pediatr 2004; 71:523-524.
- Study skills Guide 2011, Australian College of Applied Psychology. Available from: http://currentstudents.acap. edu.au/assets/Student-Services/Study-Skills-Guide-2011. pdf Last accessed on Januray 2020.
- Study Skills Guide, Disability Support Service, University of Newcastle upon Tyne. Available from: file:///C:/Users/ micro/Downloads/study-skills-guide-ncl-ac-uk%20(1).pdf Last accessed on December 2019.
- Jayashree K, Mithra PP, Nair MKC, Unnikrishnan B, Pai K. Depression and Anxiety Disorders among Schoolgoing Adolescents in an Urban Area of South India. Indian J Community Med 2018; 43(Suppl 1):S28-S32.
- Russell PS, Nair MKC, Chandra A, Subramaniam VS, Bincymol K, George B, et al. Suicidal behavior in Anxiety Disorders among adolescents in a rural community population in India. Indian J Pediatr 2013; 80Suppl 2: S175-80. doi: 10.1007/s12098-013-1122-8. Epub 2013 Sep 24.