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CLINICO - EPIDEMIOLOGICAL STUDY IN PATIENTS WITH NEUROCYSTICERCOSIS AT TERTIARY CARE HOSPITAL IN VIZIANAGARAM, ANDHRA PRADESH.



Neurology	-id do
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ABSTRACT

INTRODUCTION: Neurocysticercosis is the most common parasitic disease of Nervous system, with a prevalence of 50 million people worldwide, 50,000 deaths worldwide annually. Clinical manifestations of Neurocysticercosis are varied due to individual differences in the number, size, and topography of lesions and in the severity of the host's immune response to the parasites It is the major cause of young and adult onset epilepsy in tropical countries.

OBJECTIVES: To study the Demographic characteristics and Clinical profile of Neurocysticercosis.

MATERIALS AND METHOD: This is a Prospective Type of study in tertiary care centre for a duration of one year from August 2018 to july 2019 To study the Demographic characteristics, Clinical profile of Neurocysticercosis. Based on revised criteria for the diagnosis of Neurocysticercosis, cases were separated into Definitive Cases, and Probable Cases of Neurocysticercosis. Diagnostic criteria of neurocysticercosis was done based on Histological confirmation of parasite from biopsy of brain or spinal cord lesion. Cystic lesion showing the scolex on CT or MRI. Direct visualization of subretinal parasites on fundoscopy. Positive serum immunoelectrotransfer blot(EITB) for the detection of anticysticercal antibodies. Resolution of intracranial cystic lesions after therapy with albendazole. Spontaneous resolution of small single enhancing lesion. Clinical manifestations suggestive of NCC(Neurocysticercosis). Positive CSF enzyme linked immunosorbent assay (ELISA)

RESULTS: In the present study, Maximum number of patients encountered are in the age group of 21 to 30 yrs (31.11%), followed by 41-50 yrs(20.00%), and the mean age of cases was 32.8yrs. Of which 62% are male and 38% are female. 93% of cases have clinical presentation of seizures. The Most common site Of lesion on Ct is in parietal lobe 58%, 31% frontal lobe, 24% multiple lobes. Pattern of seizures is simple partial in 51% cases,29% gtcs 11% complex partial . No of lesion seen On imaging are single in 60% cases, multiple lesions in 40% cases. All the 45 patients are Non-vegetarians, and only 8 patients (17.77%) were pork eaters.

CONCLUSIONS: Neurocysticercosis is the most common parasitic infection of the brain. Most common clinical manifestation is Seizures Commonly presenting as Ring Enhancing lesion. All cases of young and adult onset Epilepsy in Tropical countries should be evaluated for Neurocysticercosis.

KEYWORDS

NCC, Neurocysticercosis

I. INTRODUCTION

Neurocysticercosis is the most common parasitic disease of Nervous system, with a prevalence of 50 million people worldwide.

Neurocysticercosis has been estimated to cause at least 50,000 deaths worldwide annually.

The geographic distribution of cysticercosis is wide, with high prevalence reported from Mexico, Central and South America, India and Sub-Saharan Africa.

Clinical manifestations of Neurocysticercosis are varied due to individual differences in the number, size, and topography of lesions and in the severity of the host's immune response to the parasites.

It is the major cause of young and adult onset epilepsy in tropical countries.

In India, Neurocysticercosis has been identified as a cause of 2-2.6% unselected cases of Seizures.

II.AIMS AND OBJECTIVES

To study the - Demographic characteristics and

Clinical profile of Neurocysticercosis.

III. MATERIALS AND METHODS TYPE OF STUDY: Prospective study.

PLACE OF STUDY:

Maharajah institute of medical sciences. (A Tertiary care hospital.)

DURATION OF STUDY: One year, from August 2018 to July 2019.

INCLUSION CRITERIA

The Diagnosis was based on Clinical and Radiological features. Based on revised criteria for the diagnosis of Neurocystecercosis, cases were separated into

- Definitive Cases, and
- Probable Cases of Neurocysticercosis.

DIAGNOSTIC CRITERIA OF NEUROCYSTICERCOSIS ABSOLUTE CRITERIA:

- Histological confirmation of parasite from biopsy of brain or spinal cord lesion.
- 2. Cystic lesion showing the scolex on CT or MRI.
- 3. Direct visualization of subretinal parasites on fundoscopy.

MAJOR CRITERIA:

- 1. Lesions highly suggestive of NCC on CT or MRI (cyst without scolex,enhancing or calcified lesion)
- Positive serum immunoelectrotransfer blot(EITB) for the detection of anticysticercal antibodies.
- Resolution of intracranial cystic lesions after therapy with albendazole.
- 4. Spontaneous resolution of small single enhancing lesion

MINOR CRITERIA:

- 1. Lesions compatible with NCC on neuroimaging studies.
- 2. Clinical manifestations suggestive of NCC.
- 3. Positive CSF enzyme linked immunosorbent assay (ELISA)

EPIDEMIOLOGICAL CRITERIA:

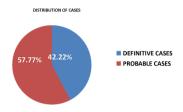
- . Evidence of household contact with Taenia solium infection.
- 2. Individuals coming from or living in an area where cysticercosis is endemic
 - . History of frequent travel to disease endemic areas.

EXCLUSION CRITERIA:

Patients with

- Tuberculosis,
- HIV reactive.
- Malignancy,
- known case of Neurocysticercosis on medication.

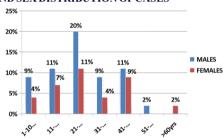
IV. RESULTS



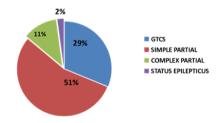
Clinical Presentation Of NCC

Clinical Presentation	Number	Percentage
Seizures	42	93.33%
Headache	08	17.77%
Raised ICT	03	06.66%
Mimicking Stroke	04	08.88%
Cranial Nerve Palsies	03	06.66%
Incoordination	04	08.88%
Dementia	04	08.88%

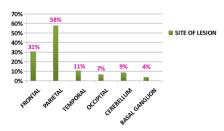
AGE AND SEX DISTRIBUTION OF CASES



SEIZURES PATTERN



DISTRIBUTION OF LESIONS ON CT



V. DISCUSSION AGE

In the present study, maximum number of patients(31.11%) is encountered in the age group of 21 to 30 yrs, followed by 41-50 yrs(20.00%), and the mean age of cases was 32.8yrs.

In Kuruvilla (2001) reported that the majority of the cases were in the age range of 24-62yrs, with a mean age of 35.2yrs.

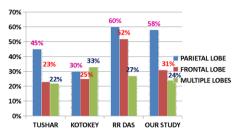
Tushar B Patil (2010) reported that the majority of the cases were in the age group of 21-30 yrs.

DIETARY HABITS:

All the 45 patients were Non-vegetarians, and only 8 patients (17.77%) were pork eaters.

In Kuruvilla's study also all patients were non-vegetarians, and 36% of them were pork eaters.

SITES OF LEISON ON CT



VI. CONCLUSION

Neurocysticercosis is the most common parasitic infection of the

Most common clinical manifestation is Seizures

Commonly presenting as Ring Enhancing lesion as.

We recommend that all cases of young and adult onset Epilepsy in Tropical countries should be evaluated for Neurocysticercosis.

VII. REFERENCES

- Fisher RS, Acevedo C, Arzimanoglou A, Bogacz A, Cross JH, Elger CE, Engel JJr, Forsgren L, French JA, Glynn M, Hesdorffer DC, Lee BI, Mathem GW, Moshé SL, Perucca E, Scheffer IE, Tomson T, Watanabe M, Wiebe S. ILAE official report: a practical clinical definition of epilepsy. Epilepsia. 2014 Apr; 55(4):475-82. doi:10.1111/epi.12550.Epub2014Apr14
- Ngugi AK, Bottomley C, Kleinschmidt I, Sander JW, Newton CR.Estimation of the burden of active and life-time epilepsy: A meta-analytic approach. Epilepsia 2010;51:883-90
- 3. Agarwal A. Social classification: The need to update in the present scenario. Indian J Community Med 2008;33:50 1
- Radhakrishnan K, Pandian JD, Santhoshkumar T, Thomas SV, Deetha TD, Sarma PS, et al. Prevalence, knowledge, attitude, and practice of epilepsy in Kerala, South India. Epilepsia 2000; 41:1027-35.
- Das SK, Biswas A, Roy T, Banerjee TK, Mukherjee CS, RautDK, et al. A random sample survey for prevalence of major neurological disorders in Kolkata. Indian J Med Res 2006: 124:163-72
- 6. Banerjee TK, Ray BK, Das SK, Hazra A, Ghosal MK, Chaudhuri A, et al. A longitudinal study of epilepsy in Kolkata, India. Epilepsia 2010;51:2384-91. Mani KS, Rangan G, Srinivas HV, Kalyanasundaram S, Narendran S, Reddy AK. The
- Yelandur study: A community-based approach to epilepsy in rural South India-epidemiological aspects. Seizure 1998;7:281-8
- Pandey S, Singhi P, Bharti B. Prevalence and treatment gap in childhood epilepsy in a
- north Indian city: A community-based study. J Trop Pediatr 2014;60:118-23. Raina SK, Razdan S, Nanda R. Prevalence of neurological disorders in children less than 10 years of age in RS Pura town of Jammu and Kashmir. J PediatrNeurosci 2011; 6:103-
- Banerjee TK, Hazra A, Biswas A, Ray J, Roy T, Raut DK, et al. Neurological disorders in
- children and adolescents. Indian J Pediatr 2009; 76:139-46. Shah PA, Shapoo SF, Koul RK, Khan MA.Prevalence of epilepsy in school-going children (6-18 years) in Kashmir Valley of NorthwestIndia. J Indian Med Assoc 2009;
- Shaji S, Verghese A, Promodu K, George B, Shibu VP. Prevalenc of priority psychiatric disorders in a rural area in kerala. Indian J Psychiatry 1995;37:91-6.
- Nandi DN, Banerjee G, Chowdhury AN, Banerjee T, Boral GC, Sen B. Urbanization and mental morbidity in certain tribalCommunities in West Bengal. Indian J Psychiatry 1992; 34:334-9
- 14. Banerjee T, Mukherjee SP, Nandi DN, Banerjee G, Mukherjee A, Sen B, et al. Psychiatric morbidity in an urbanized tribal (santal) community-a field survey. Indian J Psychiatry 1986; 28:243-8
- Gourie-Devi M, Gururaj G, Satishchandra P, Subbakrishna DK. Prevalence of 15.
- Gourne-Devi M, Gururaj G, Satistichandra F, Suboakrishina DK. Prevalence of neurological disorders in Bangalore, India: Acommunity-based study with a comparison between urban andrural areas. Neuroepidemiology 2004; 23:261-8

 Gourie-Devi M, Gururaj G, Satishchandra P, SubbakrishnaDK. Prevalence of neurological disorders in Bangalore, India: Acommunity-based study with a comparison between urban annural areas. Neuroepidemiology 2004;23:261-8
- Joseph N, Kumar GS, Nelliyanil M. Pattern of seizure cases in tertiary care hospitals in Karnataka state of India. Ann Indian AcadNeurol 2013;