



## STUDY ON DEMOGRAPHIC AND CLINICAL PROFILE OF NEUROCYSTICERCOSIS

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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 Profile and Clinical profile of Neurocysticercosis.

**MATERIALS AND METHOD:** This is a Prospective Type of study in tertiary Centre for a duration of one yr from June 2016 to May 2017. To study the Demographic Profile, 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 funduscopy. 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. 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 32.8 yrs. Of which 62% are male and 38% are female. 93% of cases have clinical presentation of seizures.

**STUDY ON DEMOGRAPHIC AND CLINICAL PROFILE OF NEUROCYSTICERCOSIS**

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 :****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 Profile, 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 June 2017 to May 2018.

**Inclusion Criteria**

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

Definitive Cases, and  
Probable Cases of Neurocysticercosis.

**DIAGNOSTIC CRITERIA OF NEUROCYSTICERCOSIS**

Absolute criteria:

1. 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 funduscopy.  
Major criteria:

1. Lesions highly suggestive of NCC on CT or MRI (cyst without scolex, enhancing or calcified lesion)

2. Positive serum immunoelectrotransfer blot (EITB) for the detection of anticysticercal antibodies.

3. 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 :

- 1.Evidence of household contact with Taenia solium infection.
- 2.Individuals coming from or living in an area where cysticercosis is endemic.

3.History of frequent travel to disease endemic areas.

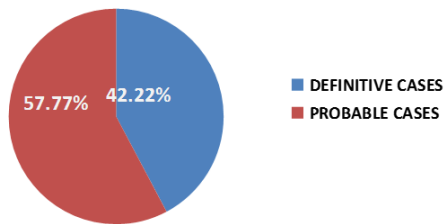
**Exclusion Criteria:**

Patients with

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

**IV. RESULTS**

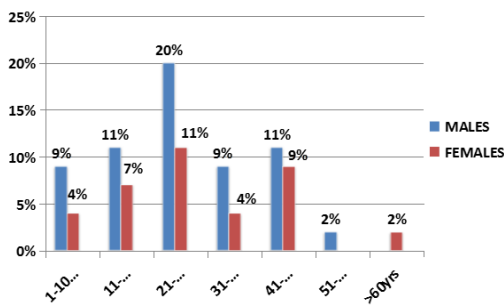
DISTRIBUTION OF CASES



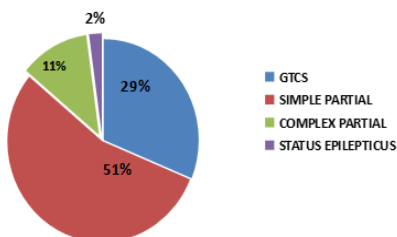
**Clinical Presentation OfNCC**

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%

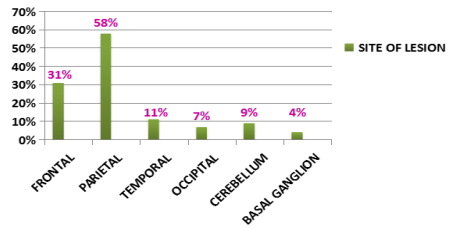
**AGE AND SEX DISTRIBUTION OF CASES**



**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.

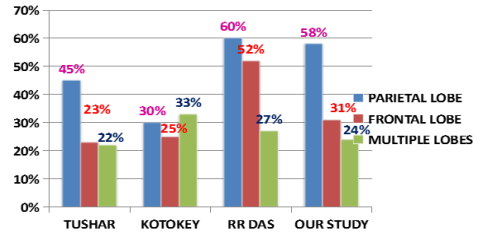
In Kuruvilla (2001) reported that the majority of the cases were in the age range of 24-62yrs, with a mean age of 35.2yrs.

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**



**VII. REFERENCES**

1. Fisher RS, Acevedo C, Arzimanoglou A, Bogacz A, Cross JH, Elger CE, Engel J Jr, Forsgren L, French JA, Glynn M, Hesdorffer DC, Lee BI, Mather 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. Epub 2014 Apr 14
2. 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
4. 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.
5. 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.
7. 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
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.
9. 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; 6:103-5.
10. 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.
11. Shah PA, Shapoo SF, Koul RK, Khan MA. Prevalence of epilepsy in school-going children (6-18 years) in Kashmir Valley of Northwest India. *J Indian Med Assoc* 2009; 107:216-8.
12. Shaji S, Verghese A, Promodu K, George B, Shibu VP. Prevalence of priority psychiatric disorders in a rural area in Kerala. *Indian J Psychiatry* 1995;37:91-6.
13. Nandi DN, Banerjee G, Chowdhury AN, Banerjee T, Boral GC, Sen B. Urbanization and mental morbidity in certain tribal Communities 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.
15. Gourie-Devi M, Gururaj G, Satishchandra P, Subbakrishna DK. Prevalence of neurological disorders in Bangalore, India: A community-based study with a comparison between urban and rural areas. *Neuroepidemiology* 2004; 23:261-8
16. Gourie-Devi M, Gururaj G, Satishchandra P, Subbakrishna DK. Prevalence of neurological disorders in Bangalore, India: A community-based study with a comparison between urban and rural areas. *Neuroepidemiology* 2004;23:261-8
17. Joseph N, Kumar GS, Nelliyanil M. Pattern of seizure cases in tertiary care hospitals in Karnataka state of India. *Ann Indian AcadNeurol* 2013;