

Acta medica Lituanica 2021. Vol. 28. No 2, pp. 285–297

# Added value of 3D-DRIVE and SWI Magnetic Resonance Imaging Sequences in Intraventricular Neurocysticercosis (IVNCC): An Institutional Experience from Northeast India

### Deb K. Boruah\*

Department of Radio-diagnosis, Lakhimpur Medical College, Assam, India

### **Bidyut Bikash Gogoi**

Department of Pathology, Assam Medical College, Dibrugarh, Assam, India

## Kuntal Kanti Das

Department of Neurosurgery, Sanjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, Uttar Pradesh, India

### Kalyan Sarma

Department of Neuroradiology, AIIMS, New Delhi, India

### Pranjal Phukan

Department of Radiodiagnosis, North Eastern Indira Gandhi Regional Institute of Health & Medical Sciences, India

### **Binoy Kumar Singh**

Department of Neurosurgery, North Eastern Indira Gandhi Regional Institute of Health & Medical Sciences, India

### Karuna Hazarika

Department of Radio-diagnosis, Tezpur Medical College, Assam, India

### Awadhesh Jaiswal

Department of Neurosurgery, Sanjay Gandhi Postgraduate Institute of Medical Sciences, Lucknow, Uttar Pradesh, India

**Abstract.** *Background:* Prompt diagnosis and early treatment institution are important in intraventricular neurocysticercosis (IVNCC) as compared to the parenchymal or racemose form because it is associated with a poorer patient prognosis. Intraventricular neurocysticercosis is often missed on CT scan or conventional cranial magnetic resonance imaging because of similar density or signal intensity of cysticercus lesion with cerebrospinal fluid. Thestudy aims to evaluate the added value of 3D-DRIVE and SWI MRI sequences in isolated intraventricular cysticercosis with acute neurological presentation.

*Methods and Materials*: This retrospective study was carried out on diagnosed 10 patients with isolated intraventricular neurocysticercosis (IVNCC) presented to a tertiary care hospital with an acute onset of symptoms or acute neurological deficit between June 2019 to May 2021. Relevant neurological examination, CSF analysis, a serological test of neurocysticercosis and MRI scan of the brain were performed.

**Result:** Tenpatients of isolated intraventricular neurocysticercosis (3 males and 7 females) having 3 pediatric and 7 adults were included in this study sample. The common neurological complications of the isolated intraventricular neurocysticercosis in this study are observed as obstructive hydrocephalus in 8(80%) patients and ependymitis in 7(70%) patients. IVNCC with distinctly visualized scolex (visibility score 2) identified in 2(20%) patients in T2WI, 8 (80%) patients in 3D-DRIVE and 3(30%) patients in SWI sequences. The cyst

Received: 15/08/2021. Revised: 30/09/2021. Accepted: 20/10/2021

Copyright © 2021 Deb K. Boruah, Bidyut Bikash Gogoi, Kuntal Kanti Das, Kalyan Sarma, Pranjal Phukan, Binoy Kumar Singh, Karuna Hazarika, Awadhesh Jaiswal. Published by Vilnius University Press. This is an Open Access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

<sup>\*</sup> Corresponding author: Deb K. Boruah, Department of Radio-diagnosis, Lakhimpur Medical College, Assam, India 787051. Email: debpal8028@gmail.com

wall of IVNCC was distinctly visualized (visibility score 2) in 1(10%) patient in T2WI, 8(80%) patients in 3D-DRIVE and 6(60%) patients in SWI sequence.

*Conclusion:* Heavily T2-weighted steady-state and SWI sequences should be added to routine MRI sequences that helps to identify IVNCC and should be used in patients with unexplained hydrocephalus, especially in endemic regions of Neurocysticercosis.

**Keywords:** 3DT2W-driven equilibrium radiofrequency reset pulse (DRIVE), magnetic resonance imaging, susceptibility weighted imaging (SWI), intraventricular neurocysticercosis (IVNCC)

# Papildoma 3D DRIVE ir SWI (svertinio jautrumo vaizdavimo) magnetinio rezonanso tomografijos sekų nauda intraventrikulinės neurocisticerkozės (IVNCC) atveju. Šiaurės Rytų Indijos įstaigų patirtis

**Santrauka.** *Problema.* Intraventrikulinės neurocisticerkozės (IVNCC) atveju, kitaip nei sergant parenchimine ar subarachnoidine forma, svarbu ligą greitai diagnozuoti ir taikyti ankstyvą gydymą, nes nuo to priklauso pasveikimo prognozė. Intraventrikulinė neurocisticerkozė dažnai nenustatoma atliekant KT tyrimą ar įprastą kaukolės magnetinio rezonanso tomografiją, nes cisticerkinio pažeidimo bei galvos ir stuburo smegenų skysčio tankis ir signalo intensyvumas yra panašūs. Tyrimo tikslas – įvertinti papildomą 3D DRIVE ir SWI MRT sekų naudą ūminės neurologinės izoliuotos intraventrikulinės cisticerkozės atveju.

*Metodai ir medžiaga.* Atliktas retrospektyvus 10 pacientų, kuriems diagnozuota izoliuota intraventrikulinė neurocisticerkozė (IVNCC), atvykusių į trečio lygio ligoninę 2019 m. birželio mėn. – 2021 m. gegužės mėn. pasireiškus ūminiams simptomams ar ūminiam neurologiniam deficitui. Atlikta atitinkama neurologinė apžiūra, galvos ir stuburo smegenų skysčio tyrimas, serologinis neurocisticerkozės tyrimas ir galvos smegenų MRT.

**Rezultatas.** Tyrime dalyvavo dešimt pacientų, sergančių izoliuota intraventrikuline neurocisticerkoze (3 vyrai ir 7 moterys, 3 vaikai ir 7 suaugusieji). Šio tyrimo metu nustatytos izoliuotai intraventrikulinei neurocisticerkozei būdingos neurologinės komplikacijos – obstrukcinė hidrocefalija 8(80 %) pacientams ir ependimos uždegimas 7(70 %) pacientams. IVNCC su aiškiai matomu skoleksu (2 matomumo balas) nustatytas 2 (20 %) pacientams taikant T2WI, 8 (80 %) pacientams taikant 3D DRIVE ir 3 (30 %) pacientams taikant SWI sekas. IVNCC cistos sienelės aiškiai matomos (2 matomumo balas) 1 (10 %) pacientui taikant T2WI, 8 (80 %) pacientams taikant SWI sekas.

*Išvada.* T2WI ir SWI sekos turėtų būti atliekamos su įprastomis MRT sekomis, siekiant nustatyti IVNCC, ir turėtų būti atliekama pacientams, kuriems dėl neaiškių priežasčių pasireiškia hidrocefalija, ypač tuose regionuose, kuriuose siaučia neurocisticerkozės endemija.

**Raktažodžiai:** 3DT2W (DRIVE), magnetinio rezonanso tomografija, svertinis jautrumas (SWI), intraventrikulinė neurocisticerkozė (IVNCC)

### Introduction

The incidence of intraventricular neurocysticercosis (IVNCC) was reported as 7.3% to 61.3% in neurocysticercosis [1, 2]. The fourth ventricle is the most common site for intraventricular neurocysticercosis (43–70%), followed by lateral ventricle (11–43%), third ventricle (1–29%) and aqueduct of Sylvius (7–9%) [1, 3, 4, 5, 6]. About 38% of patients of IVNCC presented with rapid neurological detoriation and acute neurological deficit and even sudden death [7].

In intraventricular neurocysticercosis, the larvae of Taeniasolium reach the cerebral ventricles via the choroid plexus and causes symptoms from CSF flow obstruction toependymitis, ventriculitis, or both [1, 2].

Most of the IVNCC is usually occult or missed on CT scan owing to attenuation characteristics of cysticercus cyst fluid equal to those of CSF. Neurocysticercus lesions in the CSF spaces are usually missed on conventional MRI sequences. Sometime post-gadolinium MRI images are not able to characterize the IVNCC because the signal intensity of the cyst is similar to that of CSF [8]. It has been shown that heavily T2-weighted high-resolution steady-state sequences like 3D-DRIVE/3D-CISS/FIESTA better delineates the cysticercus lesions in cisternal spaces, cerebral sulci or within the ventricle [9]. The 3D-DRIVE sequence is a 3D T2-weighted driven equilibrium radio-frequency reset pulse. CISS is a heavily T2W constructive interference in steady-state sequenceand FIESTA is a fast imaging employing steady-state acquisition.

These heavily T2-weighted images increase the conspicuity of the lesion within a CSF space. So it is able to identify the cyst wall or scolex, which are not otherwise seen on conventional MRI [10].

Susceptibility weighted imaging (SWI) sequence is a 3D gradient-echo sequence with both magnitude and phase information, it separately and in combination provides additional information in identifying the calcified scolex of intraventricular cysticercus lesion in comparison to the conventional MRI sequences [11, 12].

Advanced MRI techniques can be able to identify the adherence of IVNCC into the wall of the ventricle or able to detect ventriculitis, ependymitis and obstructive hydrocephalus. Identification of adherent IVNCC is of utmost importance MRI imaging, as even surgical resection sometimes fails to treat it, and in such situation it may need a CSF diversion with or without cyst excision [1, 13].

Intraventricular Neurocysticercosis (IVNCC) had a poorer prognosis than other forms of Neurocysticercosis [14]. The active viable stage of Intraventricular Neurocysticercosis doesn't produce a reaction to the host, however, it may produce noncommunicating hydrocephalus.With the death of intraventricular larva, the host reaction is induced in the form of ependymitis, ventriculitis and meningoencephalitis.

The common alarming symptoms of IVNCC include acute onset of headache, vomiting, decreased visual acuity, altered mental status and even death [1, 4, 5]. Isolated intraventricular neurocysticercosis, especially in the fourth ventricle, can cause mechanical obstruction of CSF flow subsequently resulting in hydrocephalus [4, 5]. The inflammatory process in IVNCC varies according to the location of the parasite [15]. MRI with newer advanced techniques is necessary to locate the IVNCC prior to surgery, as sometime IVNCC are mobile and migrated within the ventricular system.

The study aims to evaluate the added value of 3D-DRIVE and SWI MRI sequences in acute neurological complications of isolated intraventricular cysticercosis.

### **Methods and Materials**

*Patient selection:* The case records of the 10 patients were retrospectively analyzed from June 2019 to May 2021. All patients of isolated intraventricular neurocysticercosis (IVNCC) presented with features of raised intracranial pressure with the onset of symptoms of headache, vomiting or acute neurological deficit. This retrospective study was approved by the institutional ethics review committee.

#### Inclusion criteria:

1. Only isolated intraventricular neurocysticercosis with acute clinical presentation.

### **Exclusion criteria:**

1. Intraventricular neurocysticercosis along with a parenchymal or racemose form of neurocysticercosis.

*MRI Protocol:* All 10 patients underwent an MRI scan of the brain, using a 1.5 T MR scanner, Philips Ingenia (Philips Medical System, The Netherlands). MRI scans of the brain was done using a dedicated 32 channel head coil. Conventional MRI sequences include axial T1WI, T2WI, FLAIR (fluid-attenuated inversion recovery), DWI (diffusion-weighted imaging), sagittal T1WI and coronal T2WI. 3D-DRIVE (3D T2-weighted driven equilibrium radiofrequency reset pulse) and SWI (susceptibility-weighted imaging) sequences were obtained, followed by post-gadolinium T1WI sequences in all three planes. The parameters of the various MRI sequences used are shown in Table 1.

| MRI sequence                                       | TE(ms)  | TR(ms)     | Matrix    | Field of<br>view<br>(FOV) | Slice<br>thickness<br>(mm) | Flip<br>angle | Others                                  |
|--|---------|------------|-----------|---------------------------|----------------------------|---------------|---|
| T2W axial  | 90-110  | 3500-4500  | 512       | 220-250                   | 5                          | 90°           |   |
| T1W axial  | 10-15   | 450-650    | 512       | 220-250                   | 5                          | 90°           |   |
| FLAIR axial  | 100-140 | 9000-11000 | 512       | 220-250                   | 5                          | 90°           | TI=2500-2800ms                          |
| DWI axial  | 9-10    | 3000-4000  | 160 × 100 | 220-250                   | 5                          | 90°           | b-value =<br>1000s/mm <sup>2</sup>      |
| SWI axial  | 10-24   | 50-60      | 512       | 220-250                   | 2                          | 20°           |   |
| T1W-sagittal                                       | 10-15   | 450-650    | 512       | 220-250                   | 4                          | 90°           |   |
| T2W-coronal  | 90-110  | 3500-4500  | 512       | 220-250                   | 4                          | 90°           |   |
| 3D-DRIVE   | 8-10    | 1500       | 512       | 140-160                   | 1.1                        | 90°           |   |
| Post-contrastT1W_SPIR<br>axial, coronal & sagittal | 10-15   | 450-650    | 512       | 220-250                   | 4                          | 90°           | I.V. Gadolinium<br>1ml/kg<br>bodyweight |

Table 1 showing parameters used in various MRI sequences

*MRIEvaluation*: MRI images were evaluated for the location and size of the intraventricular neurocysticercosis. The presence of a scolex was looked for in the T1WI, T2WI, 3D-DRIVE and SWI images. Presence of cyst wall or wall calcification was also looked for in SWI and 3D-DRIVE sequences. The maximum wall thickness of the intraventricular cyst was measured in the 3D-DRIVE sequence. The pattern of post-contrast enhancement of the IVNCC, associated ependymitis, meningitis, hydrocephalus and entrapment of ventricle were also observed.

*Visibility score of Intraventricular neurocysticercosis (IVNCC)*: The visibility of the wall and scolex of the intraventricular neurocysticercosis was categorized on a 3-point scale from 0 to 2. Score 0 means "notdetected", 1 means "probablyseen" and 2 means "distinctlyseen". To know the diagnostic performance of T2W, 3D-DRIVE and SWI sequences in the detection of intraventricular neurocysticercosis, this visibility score was calculated.

*Neurological assessment of intraventricular neurocysticercosis(IVNCC)*: The details of clinical and neurological examinations performed in all patients.

*Follow up and final diagnosis:* Craniotomy with microsurgical cyst excision was done in 2 patients and neuroendoscopic cyst excision in another 4 patients, and histopathological confirmation of cysticercus lesions were confirmed. Three patients were confirmed having IVNCC on subsequent follow up MRI scans on basis of regression in size or complete healing of cysticercus lesion, with initial positive IgM antibodies against cysticercus antigens measured by ELISA technique both in CSF and Serum serology. Another 1 patient was diagnosed with IVNCC only basis of initial MRI imaging findings of intraventricular cyst with positive IgM antibodies against cysticercus antigens both in CSF and Serum.

All 10 patients received anticysticercal therapy with albendazole. Patients were followed up for a period of 6 months to 18 months.

**Statistical analysis**: All statistical analysis was performed using Statistical Package for Social Science (SPSS, version 16). Data were presented in terms of percentage and mean. A Chi-square test is done to find out the sensitivity and specificity of T2W, 3D-DRIVE and SWI sequences.

# Results

The study sample comprised of 10 patients (n=3 males and n=7 females) with a mean age of  $37.2 \pm 22.7$  [SD] years and a male: female ratio of 1:2.3. Five (50%) patients presented with acute onset of headache and vomiting, 3(30%) with altered consciousness and neurological deterioration, 1(10%) patient had a seizure and another 1(10%) patient had visual disturbance. Intraventricular neuro-cysticercosis lesions were identified within the lateral ventricle in 5(50%) patients (Figure 1 and2), fourth ventricle in 2(20%) patients (Figure 3 and 4) and third ventricle in 1(10%) patient (Figure 5). Two (20%) patients had intraventricular neurocyticercus lesions in more than one ventricle. Five (50%) patients had only solitary intraventricular neurocysticercus lesions, 1(10%) patient had two neurocysticercus lesions, 2 (20%) patients had three neurocysticercus lesions (Figure 5) and another 2(20%) patients had more than three intraventricular neurocysticercus lesions (Figure 3).



Figure 1. Brain magnetic resonance imaging of 52 years female with headache and vomiting. (A-C) Axial T2WI images show asymmetrical hydrocephalus with entrapment of temporal horn of the right lateral ventricle with narrowing / adhesion in the trigone of the lateral ventricle (arrow) with surrounding periventricular CSF seepage. (D) Heavily T2-weighted image shows an intraventricular neurocysticercus lesion with a T2 hypointense scolex within (arrow). (E) Axial susceptibility-weighted image shows nodular blooming of the calcified scolex within the thin wall IVNCC (arrow). (F) Axial post-gadolinium T1W image shows thin smooth peripheral cyst wall enhancement with the smooth ependymal enhancement of the asymmetrically dilated lateral ventricle.

Figure 2. Brain magnetic resonance imaging of 50 years female with acute severe headache with acute neurological status and semicomatose state. (A-C) Axial T2WI, FLAIR and T1WI images show a thin-walled T2 hyperintense cystic lesion in the frontal horn of the right lateral ventricle occluding the right-sided foramen of Monroe(arrow) with marked supratentorial hydrocephalus with periventricular CSF seepage. (D-F) Axial and sagittal post-gadolinium T1W images show smooth peripheral cyst wall enhancement (arrow) with a small enhancing mural nodule inferiorly (arrow in image F).



Figure 3. Brain magnetic resonance imaging of 25 years male with headache. (A-B) Axial T2WI and FLAIR images show distension and dilatation of the fourth ventricle with minimal periventricular CSF seepage. Variable signal intensity cystic lesions are seen within the fourth ventricle on the FLAIR image (arrow). (C-D) Axial and coronal heavily T2W images show multiple thin-walled hyperintense cystic lesions (arrows) with less hyperintense CSF signal within the 4th ventricle with variable sizes irregular nodules within. (E-F) Axial and sagittal T1W post-contrast image shows thin smooth enhancement of the intraventricular cyst (arrow) with enhancing irregular nodules. Post-contrast enhancement is also seen along the ependymal lining of the 4th ventricle, in posterior and inferior aspects.



**Figure 4.** Brain magnetic resonance imaging of 13 years male with severe headache, vomiting and acute neurological deterioration. (A-B) Axial T2WI images show marked distension and dilatation of the fourth ventricle with marked periventricular CSF seepage and supratentorial hydrocephalus. (C) Axial FLAIR image shows cystic lesions within the fourth ventricle (arrow). (D-E) Axial heavily T2W images show thin-walled hyperintense cystic lesions within the 4<sup>th</sup> ventricle with an oval to elongated appearing cystic lesion with scolex in the right foramen of Luschka (arrow). (F-H) Axial and sagittal T1W post-contrast image shows a thin peripheral enhancing cystic lesion with an irregular mural nodule in the right foramen of Luschka (arrow) and thin smooth enhancement of the intraventricular cysts. Nodular enhancement was seen in midline inferior margin of 4<sup>th</sup> ventricle near to foramen of Magendie (arrow).



**Figure 5.** Brain magnetic resonance imaging of 80 years male with headache and visual disturbance. (A) Sagittal T2WI image shows a thin-walled T2 hyperintense cysticercus lesion in the fourth ventricle (arrow) causing supratentorial hydrocephalus. (B-C) Axial and sagittal T1W post-contrast images show cysticercus lesions within the 3rd ventricle (white arrow in image B), trigone of the left lateral ventricle (yellow arrow) and fourth ventricle (arrow in image C). (D-E) Neuroendoscopic post-excision cysticercus specimen shows thin-walled cystic lesions. (F) HPE image shows sucker (long arrow) and cuticle (short arrow) of the cysticercus lesion.

The mean diameter of the largest intraventricular cysticercosis was 18.2±6.9 mm [SD] and the mean wall thickness of intraventricular neurocysticercosis was 1.3±0.23 mm [SD].

Very thin smooth cyst wall enhancement was observed in 9(90%) patients (Figure 1 and 2) and irregular peripheral rim-like wall enhancement with an irregular nodule in 1(10%) patient. Ventricular ependymitis with smooth to irregular ependymal lining enhancement was observed in 7(70%) patients (Figure 3). Presence of hydrocephalus detected in 8(80%), where MRI revealed asymmetrical enlargement of lateral ventricle with periventricular CSF seepage in 4(40%) patients (Figure 1), dilatation of  $3^{rd}$  ventricle in 2(20%) patients and ballooned  $4^{th}$  ventricle in another 2(20%) patients (Figure 4).

**T2WI, DRIVE and SWI visibility score of the scolex of intraventricular neurocysticercosis** (*IVNCC*): In 10patients with intraventricular neurocysticercosis, the scolex was distinctly visualized (visibility score 2) in 2(20%) patients on T2WI, 8(80%) patients on 3D-DRIVE (Figure 1) and 3(30%) on SWI sequence (Figure 1). For identification of scolex in intraventricular neurocysticercosis with a visibility score of 2, the 3D-DRIVE sequence had a sensitivity of 80% followed by 30% with SWI and 20% with T2W images. The poorly visualized scolex of intraventricular neurocysticercosis (visibility score 1) noted in 2 (20%) patients on 3D-DRIVE, 5(50%) patients on SWI and 2(20%) on T2W sequence [Table 2].

| MRI sequence | Visibility score | Cyst wall | Scolex | p-value (chi-square) |
|--------------|------------------|-----------|--------|----------------------|
|              | Score 2          | 1         | 2      |                      |
| Axial T2WI   | Score 1          | 4         | 2      | 0.323                |
|              | Score 0          | 5         | 6      |                      |
| 3D-DRIVE     | Score 2          | 8         | 8      |                      |
|              | Score 1          | 2         | 2      | 0.002                |
|              | Score 0          | 0         | 0      |                      |
| SWI          | Score 2          | 6         | 3      |                      |
|              | Score 1          | 4         | 5      | 0.329                |
|              | Score 0          | 0         | 2      |                      |

**Table 2.** Visibility score of cyst wall and scolex of intraventricular neurocysticercosis on T2WI, 3D-DRIVE and SWI sequences in 10 patients.

**T2WI, DRIVE and SWI visibility score of the wall of intraventricular neurocysticercosis:** Of 10patients with IVNCC, the cyst wall was distinctly visualized (visibility score 2) in 1 (10%) patient on T2WI, 8 (80%) patients on 3D-DRIVE (Figure 1 and 2) and 6 (60%) patients on SWI sequence. For identification of wall of IVNCC with a visibility score of 2, the 3D-DRIVE sequence had a sensitivity of 80% followed by 60% with SWI and 10% with T2W images. Cyst wall calcification was identified in 1(10%) patient on SWI. Poorly visualized wall of intraventricular neurocysticercosis (visibility score 1) noted in 2(20%) patients on 3D-DRIVE, 4(40%) patients on SWI and 4(40%) on T2W sequences [Table 2].

Four (40%) patients were treated conservatively with cysticidal drugs only and 6 (60%) patients were treated surgically followed by cysticidal drugs. Craniotomy with micro-surgical resection was done in 2 patients with 4<sup>th</sup> ventricular neurocysticercosis and endoscopic approach in another 4 patients with lateral and 3<sup>rd</sup> ventricular neurocysticercosis. On follow-up, 8(80%) patients show complete healing of the intraventricular neurocysticercosis, 1 (10%) patient showed disease recurrence and another 1 (10%) patient died.

# Discussion

Acute onset of headache, vomiting, altered mental status and visual disturbance can occur in intraventricular neurocysticercosis (IVNCC) [16]. Acute onset of hydrocephalus can lead to sudden death due to brain stem herniation, displacement or distortion[17]. This sudden onset of symptoms is associated with changes in patient's head position, because acute ventricular obstruction can occur in Bruns' syndrome [18, 19]. It occurs due to intermittent CSF flow obstruction by the ball-valve movement of intraventricular cysts. Intraventricular neurocysticercosis occurs in approximately 20% of patients of neurocysticercosis worldwide [20]. It carries high mortality rate due to acute onset of hydrocephalus and neurological detoriation. The viable intraventricular neurocysticercosis is often freely mobile and may lodge in ventricular foramina like foramen of Monroe, Luschka, Magendie or aqueduct of Sylvius, and lead to acute obstructive hydrocephalus [21]. Isolated fourth IVNCC is associated with ependymitis, arachnoiditis and ventriculitis [21].

CT scan fails to identify IVNCC that do not deform the ventricle because of same density of cyst with CSF, cyst wall and scolex not visible or cyst wall not shows any abnormal wall enhancement[22, 23, 20, 21]. MRI can able to identify IVNCC in approximately 80% of cases [24] with a classical intraventricular cyst with scolex [25].

MRI appearance of an intraventricular cystic lesion with T1-weighted hyperintense or T2-weighted hypointense scolex within are considered as pathognomonic for intraventricular neurocysticercosis and these characteristics are usually not detected by CT scan. Conventional MRI sequences routinely fails to identify the scolex or cyst wall. Hence additional newer advanced MRI sequences like 3D-DRIVE and SWI act as a solving tool in the identification and characterization of intraventricular cystic lesions [10].

The common differential diagnosis of intraventricular NCC included colloid cyst, ependymal cyst, choroidal plexus cyst, intraventricular epidermoid and arachnoid cyst. And to differentiate these intraventricular cystic lesions newer advanced MRI techniques are necessary [Table 3].

 Table 3. Common differential diagnosis of isolated intraventricular neurocysticercosis (IVNCC) on MR imaging.

|                                | IV NCC  | Ependymal<br>Cyst  | Choroid<br>plexus<br>cyst  | Intraven-<br>tricular<br>epidermoid<br>cyst  | Intraven-<br>tricular<br>arachnoid<br>cyst   | Colloid cyst   | Intraven-<br>tricular<br>Cystic<br>neoplasm  |
|--------------------------------|---|--|--|--|--|--|--|
| Age                            | - Any age<br>- Common-<br>est in 15-40<br>years   | - Young adult<br>- around 3 <sup>rd</sup><br>decade          | - Common in<br>neonates<br>- Uncommon<br>in adults   | 4-6 <sup>th</sup> decade   | Any age  | 3-4 <sup>th</sup> decade   | Adult<br>- Cystic as-<br>trocytoma<br>Children<br>- cystic me-<br>dulloblas-<br>toma   |
| Clinical<br>presen-<br>tations | headache,<br>vomiting,<br>decreased<br>visual acuity,<br>altered men-<br>tal status and<br>even death | - asympto-<br>matic<br>- symptomat-<br>ic in larger<br>cyst  | - Usually<br>asymptomatic<br>- Symptomat-<br>ic in larger<br>cysts   | - asympto-<br>matic<br>- headache<br>- cognitive<br>deficit<br>- larger pro-<br>duces mass<br>effect, hydro-<br>cephalus | - usually<br>asymptomatic<br>- symptomat-<br>ic with larger<br>cyst<br>- headache,<br>hydrocepha-<br>lus<br>- hemorrhage<br>can occur<br>within a cyst | - usually<br>asymptomatic<br>- Headache<br>- rarely pre-<br>sent with<br>acute ob-<br>structive hy-<br>drocephalus<br>-sudden<br>death[10] | - asympto-<br>matic<br>- Headache<br>- acute ob-<br>structive hy-<br>drocephalus   |
| Location                       | fourth ventri-<br>cle followed<br>by lateral,<br>third ven-<br>tricle and<br>aqueduct of<br>sylvius   | - common in<br>frontal horn<br>of lateral ven-<br>tricle[10] | Located<br>in choroid<br>plexus in pos-<br>terior body<br>of lateral ven-<br>tricle.<br>Usually bilat-<br>eral[10] | 4 <sup>th</sup> ventricle<br>followed by<br>3 <sup>rd</sup> and lateral<br>ventricle                                     | - common in<br>lateral ventri-<br>cle followed<br>by 3 <sup>rd</sup> and 4 <sup>th</sup><br>ventricle  | anterior 3 <sup>rd</sup><br>ventricle fol-<br>lowed by near<br>to foramen of<br>Monroe   | 4 <sup>th</sup> ventricle  |
| Conven-<br>tional<br>MRI       | - similar to<br>CSF<br>- Slightly<br>higher signals<br>on FLAIR<br>compared to<br>CSF.                | Similar to<br>CSF signals<br>in all MRI<br>sequences         | Thin-walled<br>cyst showing<br>slightly to<br>moderately<br>hyperintense<br>to CSF on<br>FLAIR                     | Slightly<br>higher signals<br>on FLAIR<br>compared to<br>CSF.  | Similar to<br>CSF signals<br>in all MRI<br>sequences   | Hypointense<br>on T2W and<br>hyperintense<br>on T1W  | - central T2<br>hyperintense<br>with periph-<br>eral T2 iso to<br>hypointense<br>irregular wall<br>or eccentric<br>solid appear-<br>ing compo-<br>nent |

|                    | IV NCC                      | Ependymal<br>Cyst           | Choroid<br>plexus<br>cyst  | Intraven-<br>tricular<br>epidermoid<br>cyst                          | Intraven-<br>tricular<br>arachnoid<br>cyst | Colloid cyst                            | Intraven-<br>tricular<br>Cystic<br>neoplasm                         |
|--------------------|-----------------------------|-----------------------------|--|--|--|---|---|
| DWI                | No diffusion<br>restriction | No diffusion<br>restriction | 2/3 <sup>rd</sup> showed<br>diffusion<br>restriction             | Shows diffu-<br>sion restric-<br>tion with<br>variable ADC<br>value  | No diffusion<br>restriction                | No diffusion<br>restriction             | Peripheral<br>wall or solid<br>component<br>showed re-<br>striction |
| Scolex             | Present                     | Absent                      | Absent   | Absent   | Absent                                     | Absent                                  | Absent  |
| Wall               | Smooth thin<br>wall         | Smooth wall                 | Smooth or<br>irregular   | Thin or may<br>not seen  | Thin or im-<br>perceptible                 | Not seen                                | Thick and<br>irregular en-<br>hanced walls                          |
| Hydro-<br>cephalus | Common                      | Uncommon                    | - uncommon<br>- larger lesion<br>may cause<br>hydrocepha-<br>lus | - Uncommon<br>- Rarely<br>larger lesion<br>causes hydro-<br>cephalus | Uncommon                                   | - acute hy-<br>drocephalus<br>is common | Common  |

The colloid cyst typically showed T1-weighted hyperintensities and was classically located in the anterior third ventricle or foramen of Monroe region. CT scan shows hyperdensity within the colloid cyst. Ependymal cyst is difficult to differentiate from an adherent intraventricular NCC, but ependymal cyst wouldn't show scolex. The common location of ependymal cyst is in the frontal horn of lateral ventricle, where the cystis located near tothe foramen of Monroe region causing obstructive hydrocephalus. The choroid plexus cyst is usually asymptomatic and confines in the posterior body of the lateral ventricle. Intraventricular epidermoid cyst showed typical diffusion restriction without any scolex. Intraventricular arachnoid cyst suppressed signals on FLAIR images with an absence of scolex.

IVNCC withoutabnormal post-contrast enhancement should be treated surgically with either neuroendoscopic or open surgery for cyst excision. Those IVNCC that showa peripheral rim-like post-contrast enhancement with or without adjacent ependymitis should probably undergo the CSF flow diversion with VP shunt or partial cyst excision with the CSF diversion. IVNCC showing rim-like post-contrast enhancement denotes ependymitis; if it is surgically resected, patient is likely to develop hydrocephalus after surgery.

The surgical treatment options for intraventricular NCC depend on the clinical presentation, location and stage of IVNCC. Usual endoscopic or open microsurgical removal of intraventricular NCC should be considered if there is a CSF obstruction or mass effect, or fourth ventricular cyst.

Various recent literatures show application of various newer MRI sequences for identification and localization of IVNCC [26, 27]. Conventional MRI sequences are lesssensitive for identificationof IVNCC, however the previous study of Singh et al. [26] identified scolex on FLAIR and T1W images in 3 out of 4 patients of IVNCC, and T2W fails to identify the same. Govindappa SS et al. [10] found more accuracy of 3D-CISS than of conventional MRI sequences for identification of IVNCC. However, Robbani I et al.[28] and Mont'Alverne Filho FE et al. [9] found more accuracy of 3D-SPGR (spoiled gradient recalled echo) overthat of conventional MRI sequences for identification of IVNCC. Table 4 shows few review literature of intraventricular neurocysticercosis.

| S/N | Study/year                                    | No of<br>patient<br>of<br>IVNCC | Location<br>of IVNCC   | Mean/<br>Median<br>age(yrs.) | CEMRI<br>enhancement<br>Pattern  | Complication   | Treatment  |
|-----|---|---------------------------------|--|------------------------------|--|--|--|
| 1.  | Cuetter AC et<br>al. / 2002(1)                | 18                              | 2-isolated IVNCC<br>With extra-ventric-<br>ular NCC-16   | Mean = 35                    | -  | Hydrocepha-<br>lus-18  | Medical treat-<br>ment -8<br>Surgical Treat-<br>ment -10 |
| 2.  | Citow JS et<br>al. / 2002(3)                  | 30                              | 30-IVNCC<br>LV-5(16.7%)<br>3V-5(16.7%)<br>4V-21(70%)<br>With extra-ventric-<br>ular NCC-   | Mean= 36                     | 17-no enhance-<br>ment<br>14- peripheral<br>wall enhance-<br>ment of the<br>IVNCC                        | 29   | Surgical<br>Treatment -30                                |
| 3.  | Kumar R et<br>al. / 2008(29)                  | 11                              | 11-IVNCC<br>LV-4(36.4%)<br>FOM-2(18.2%)<br>Aqueduct-1(9%)<br>4V-4(36.4%)<br>With extra-ventric-<br>ular NCC-0                                    | Mean=<br>14.2 ±2.98          | -  | Hydrocepha-<br>lus-7   | Surgical<br>Treatment -11                                |
| 4.  | Mont'Alver-<br>neFilho FE et<br>al. / 2011(9) | 7                               | IVNCC-8<br>4V-4(50%)<br>3V-2(25%)<br>LV-2(25%)<br>With extra-ven-<br>tricular NCC-7(4<br>patient had paren-<br>chymal and 3 had<br>racemose NCC) | Mean=<br>39 ±10              | 3(37.5%)-<br>showed periph-<br>eral rim-like or<br>nodular contrast<br>enhancement                       | -  | -  |
| 5.  | Nash TE et<br>al. / 2018(30)                  | 23                              | LV-11(36.7%)<br>3V-2(6.7%) Aque-<br>duct-1(3.3%)<br>4V-16(53.3%)<br>With extra-<br>ventricular NCC-<br>17(73.9%)                                 | Median =<br>31.8             | -  | -Hydrocepha-<br>lus-17 (73.9%)<br>-Ventriculi-<br>tis-7(30.4%)<br>-Entrapment<br>of lateral ven-<br>tricle-2(8.7%) | -Cyst excised<br>-14(60.9%)<br>-VP shunt<br>-10(43.5%)   |
| 6.  | Present study                                 | 10                              | LV-5(50%)<br>3V-1(10%)<br>4V-2(20%)<br>2 patients - had<br>IVNCC in more<br>than one ventricle<br>With extra-ventric-<br>ular NCC-0              | 37.2 ± 22.7                  | 9- very thin<br>smooth pe-<br>ripheral wall<br>enhancement<br>1-thick irregular<br>wall enhance-<br>ment | Hydrocephalus<br>-8(80%)<br>Ependymitis<br>-7(70%)<br>Entrapment<br>of lateral ven-<br>tricle-1(10%)               | Surgical treat-<br>ment-6<br>Medical treat-<br>ment -4   |

Table 4. Few review literature of intraventricular NCC on magnetic resonance imaging (MRI).

**Limitation:** Due to exclusion of patients with presence of intraventricular NCC along with the parenchymal form of NCC as well as IVNCC with racemose form of cysticercosis, the sample is limited to only isolated IVNCC in our study, and the sample size decreases. Still we tried to findout the visibility scores of T2W, 3D-DRIVE and SWI sequences for detection of wall and scolex of intraven-

tricular NCC. Therefore, a larger study sample size is needed to confirm these added values of various MRI sequences in diagnosis, detection of cyst wall and scolex in isolated intraventricular NCC.

# Conclusion

Newer heavily T2-weighted MRI sequences (like 3D-DRIVE/FIESTA/CISS) and SWI improve the sensitivity of detection and localization of intraventricular neurocysticercosis, and further aid in guiding the management and neuroendoscopic cyst excision and further improvement of patients suffering from this disease. In a patient with an unexplained obstructive hydrocephalus especially from endemic regions of neurocysticercosis these newer MRI sequences may be added to conventional MRI sequences

### References

- 1. Cuetter AC, Andrews RJ. Intraventricular neurocysticercosis: 18 consecutive patients and review of the literature. Neurosurg Focus 2002; 12: e5.DOI: 10.3171/foc.2002.12.6.6
- 2. Sinha S, Sharma BS. Intraventricular neurocysticercosis: a review of current status and management issues. Br J Neurosurg 2011;26: 305-309.DOI: 10.3109/02688697.2011.635820
- 3. Citow JS, Johnson JP, McBride DQ, Ammirati M. Imaging features and surgery-related outcomes in intraventricular neurocysticercosis.Neurosurg Focus 2002; 12: e6.DOI: 10.3171/foc.2002.12.6.7.
- 4. Goel RK, Ahmad FU, Vellimana AK, Suri A, Chandra PS, Kumar R, et al. Endoscopic management of intraventricular neurocysticercosis. J ClinNeurosci 2008; 15: 1096-1101.DOI: 10.1016/j.jocn.2007.10.004
- 5. Husain M, Jha DK, Rastogi M, Husain N, Gupta RK. Neuroendoscopic management of intraventricular Neurocysticercosis (NCC). ActaNeurochir 2007; 149: 341-346.DOI: 10.1007/s00701-006-1059-z.
- Del Brutto OH: Clues to prevent cerebrovascular hazards of cysticidal drug therapy. Stroke 1997; 28: 1088. PMID: 9158657
- 7. Zee CS, Segall HD, Destian S, Ahmadi J, Apuzzo ML. MRI of intraventricular cysticercosis: Surgical implication. J Comput Assist Tomo1993;17:932 –9.DOI: 10.1097/00004728-199311000-00015.
- Lucato LT, Guedes MS, Sato JR, et al. The Role of Conventional MR Imaging Sequences in the Evaluation of Neurocysticercosis: Impact on Characterization of the Scolex and Lesion Burden. Am J Neuroradiol. 2007; 28: 1501-1504.DOI: 10.3174/ajnr.A0623.
- Mont'Alverne Filho FE, Machado LR, Lucato LT, Leite CC. The role of 3D volumetric MR sequences in diagnosing intraventricular neurocysticercosis: Preliminary results. Arg Neuro Psiquiatr2011;69:74-8.DOI: 10.1590/ s0004-282x2011000100015.
- 10. Govindappa SS, Narayanan JP, Krishnamoorthy VM, Shastry CH, Balasubramaniam A, Krishna SS. Improved detection of intraventricular cysticercal cysts with the use of three dimensional constructive interference in steady state MR sequences. AJNR Am J Neuroradiol2000;21:679-84.PMID: 10782777
- 11. Verma A, Awasthi R, Prasad KN, Soni N, Singh K, Paliwal VK et al. Improved detection of parenchymal cysticercal lesions in neurocysticercosis with T2\*-weighted angiography magnetic resonance imaging. AcadRadiol2012;19:958-64.DOI: 10.1016/j.acra.2012.03.019
- 12. Neyaz Z, Patwari SS, Paliwal VK. Role of FIESTA and SWAN sequences in diagnosis of intraventricular neurocysticercosis. Neurol India. 2012; 60: 646-647.DOI: 10.4103/0028-3886.105205
- 13. Sinha S, Sharma BS. Intraventricular neurocysticercosis: a review of current status and management issues. Br J Neurosurg. 2012; 26(3): 305-309.DOI: 10.3109/02688697.2011.635820
- 14. DeGiorgio CM, Houston I, Oviedo S, Sorvillo F. Deaths associated with cysticercosis. Report of three cases and review of the literature. Neurosurg Focus 2002; 12: e2.DOI: 10.3171/foc.2002.12.6.3.
- 15. Fleury A, Escobar A, Fragoso G, Sciutto E, Larralde C. Clinical heterogeneity of human neurocysticercosis results from complex interactions among parasite, host and environmental factors. Trans R Soc Trop Med Hyg 2010; 104: 243-250.DOI: 10.1016/j.trstmh.2010.01.005
- 16. Garcia HH, Coyle CM, White AC. Cysticercosis Tropical Infectios Diseases Principles Pathogens and Practice 3rd Edition. Elsevier Saunders Philadephia. 2011; 815-823

- 17. Goel RK, Ahmad FU, Vellimana AK, et al. Endoscopic management of intraventicularneurocysticercosis. J ClinNeurosci. 2008; 15(10): 1096-1101.DOI: 10.1016/j.jocn.2007.10.004.
- Campbell BR, Reynoso D, White AC Jr. Intraventricular Neurocysticercosis and Bruns' Syndrome: A Review. J Rare Dis Res Treat. 2017;2(2): 1-5.http://www.rarediseasesjournal.com/articles/intraventricular-neurocysticercosis-and-bruns-syndrome-a-review.pdf
- 19. Torres-Corzo J, Rodriguez-dellaVecchia R, Rangel-Castilla L. Bruns syndrome caused by intraventricular neurocysticercosis treated using flexible endoscopy. J Neurosurg2006;104:746-8.DOI: 10.3171/jns.2006.104.5.746
- 20. Garcia HH, Nash TE, Del Brutto OS. Clinical symptoms diagnosis and treatment of neurocysticercosis. Lancet Neurol. 2014; 13: 1202-1215, DOI: 10.1016/S1474-4422(14)70094-8
- 21. Colli BO, Pereira CU, Assirati Junior JA, Machado HR. Isolated fourth ventricle in neurocysticercosis: pathophysiology, diagnosis, and treatment. Surg Neurol 1993;39: 305–310.DOI: 10.1016/0090-3019(93)90010-x
- 22. Kanamalla US, Ibarra RA, Jinkins JR. Imaging of cranial meningitis and ventriculitis. Neuroimaging Clin N Am 2000;10: 309–331.PMID: 10775954
- 23. Zee CS, Go JL, Kim PE, et al: Imaging of neurocysticercosis. Neuroimaging Clin N Am 2000;10:391–407.PMID: 10775958
- 24. Chang KH, Lee JH, Han MH, et al: The role of contrast enhanced MR imaging in the diagnosis of neurocysticercosis.AJNR1991;12:509–512. PMID: 2058504
- 25. Del Brutto OH, Wadia NH, Dumas M, et al: Proposal of diagnostic criteria for human cysticercosis and neurocysticercosis. J Neurol Sci 1996;142:1–6.DOI: 10.1016/0022-510x(96)00130-x
- 26. Singh S, Gibikote SV, Shyamkumar NK. Isolated fourth ventricular cysticercus cyst: MR imaging in 4 cases with short literature review. Neurology India 2003;51:394-396.PMID: 14652451
- 27. Wallin MT, Kurtzke JF. Neurocysticercosis in the United States: review of an important emerging infection. Neurology 2004;63:1559-1564.DOI: 10.1212/01.wnl.0000142979.98182.ff
- Robbani I, Razdan S, Pandita K. Diagnosis of intraventricular cysticercosis by magnetic resonance imaging: improved detection with three-dimensional spoiled gradient recalled echo sequences. AustralasRadiol 2004; 48:237-239.DOI: 10.1111/j.1440-1673.2004.01279.x
- 29. Kumar R, Kalra S.K, Mahapatra A.K. Intraventricular Neurocysticercus Cyst(s) in Indian Children PediatrNeurosurg 2008;44:118–123.DOI: 10.1159/000113113.
- 30. Nash T E, Ware J M, Mahanty S. Intraventricular Neurocysticercosis: Experience and Long-Term Outcome from a Tertiary Referral Center in the United States. Am. J. Trop. Med. Hyg., 98(6), 2018:1755–1762. DOI:10.4269/ajtmh.18-0085