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– Technical Note

Noninvasive MR Cisternography with Fluid-Attenuated Inversion Recovery and 100% Supplemental O₂ in the Evaluation of Neurocysticercosis

Flávio Braga, Antônio J. Rocha, Hélio R. Gomes, Guinel Hernandez Filho, Carlos J. Silva, and Ricardo B. Fonseca

Summary: We describe an MR protocol for the noninvasive imaging of the subarachnoid space, which we use in patients with suspected neurocysticercosis in this space. It consists of a fluid-attenuated inversion recovery sequence performed 5 minutes after the continuous inhalation of 100% O_2 with a resultant increase in the signal intensity of the CSF that leads to a greater conspicuity of cyst walls in relation to the cortex and the extraventricular CSF.

Initial trials have shown the safety of intrathecal administration of gadopentetate dimeglumine in humans (1–3). The administration of paramagnetic contrast agents by the intrathecal route, however, has not yet been approved by regulatory agencies in most countries, including the United States Food and Drug Administration.

On the other hand, artifactual CSF hyperintensity on fluid-attenuated inversion recovery (FLAIR) images has been shown in patients ventilated with 100% O_2 (4–6). We currently use this approach as a noninvasive cisternography technique that allows the detection of cystic lesions in the subarachnoid space, because detection of such lesions is frequently difficult with conventional MR imaging sequences. In neurocysticercosis patients, the resultant hyperintensity of the CSF increases the conspicuity of lesions in the cisterns, sulci, and along the cortical surface.

Among the main complications of neurocysticercosis in the subarachnoid space are hydrocephalus, meningitis, and arteritis (7). The prompt detection of lesions in the extraventricular CSF spaces may help in the diagnosis and management of neurocysticercosis, cases in which medical treatment over surgical approaches is favored (8, 9), and the prevention of complications.

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Technique

Initially, a FLAIR sequence (1.0 T; TR, 11,000 ms; TE, 140 ms; TI, 2600 ms) is obtained with the patient breathing room air. Next, 100% O_2 is delivered for 5 minutes by using a nonbreathing face mask connected to a ventilator, after which another identical FLAIR sequence is obtained. Images are evaluated with a window width of 950 and window level of 650.

It is essential that the mask fit the face well to prevent room air from entering the ventilator system (4, 5), because high concentrations of O_2 are needed to increase the signal intensity of the CSF. Supplemental oxygen at 50% (4) or 60% (6) will not cause the intended hyperintensity in the CSF on FLAIR sequences.

Discussion

FLAIR is very sensitive for the detection of parenchymal lesions. The acquisition of FLAIR sequences before and after the inhalation of 100% O₂ may improve the detection of subtle cisternal and extraventricular subarachnoid space lesions because of the resultant hyperintensity of the CSF. In neurocysticercosis endemic areas, this noninvasive MR cisternography may resolve the not uncommon diagnostic dilemma of a patient (symptomatic or not) with subtle cisternal widening or subtle superficial cortical lesions.

This method enables both detection of neurocys-

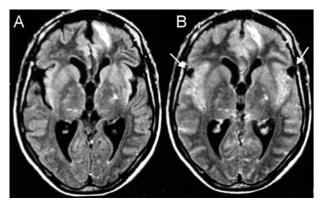


Fig 1. A 40-year-old man with parenchymal neurocysticercosis and suspected subarachnoid involvement.

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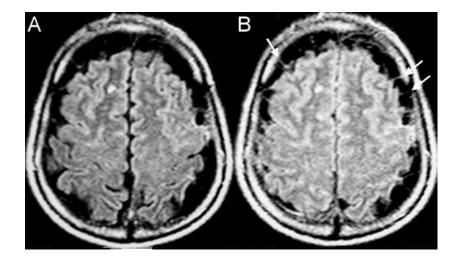
A, MR FLAIR image (1.0 T; TR, 11,000 ms; TE, 140 ms; TI, 2600 ms) shows no extraaxial lesions.

B, FLAIR image after 100% O₂ for 5 minutes shows two small cysts in the Sylvian fissures confirming racemous neurocysticercosis.

Fig 2. A 57-year-old man with multiple vesicular lesions in the brain paren-chyma.

A, FLAIR image shows prominent CSF space along the cerebral convexities.

B, FLAIR image after 100% O_2 for 5 minutes does not show the expected increased signal intensity of the CSF due to the presence of multiple clustered extraaxial cysts. The walls of the cysts (*arrows*) are better visualized after 100% O_2 .



ticercosis lesions in the extraventricular CSF spaces not evident on conventional FLAIR images (Fig 1) and the capacity to rule out the presence of such lesions within the subarachnoid space. Superficial parenchymal lesions may also become more evident (Figs 2 and 3). It should be noted that arachnoid cysts

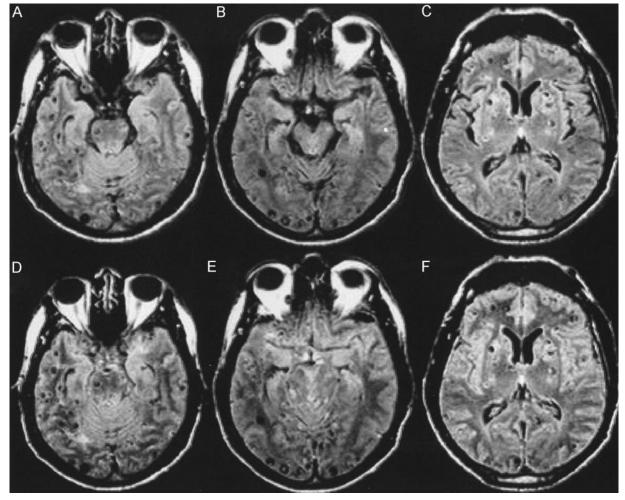


Fig 3. A 34-year-old male patient with known neurocysticercosis.

A, *B*, and *C*, FLAIR images show multiple vesicular lesions in the parenchyma, mainly along the cortical surfaces. The perimesencephalic cisterns are prominent and contain hyperintense foci that could represent the expected flow artifact or cystic lesions with scoleces (*B*).

D, *E*, and *F*, FLAIR after 100% O_2 shows increased signal intensity in the sulci and basal cisterns, allowing the visualization of a greater number of cortical lesions in the frontal, temporal, and occipital lobes. Also, the increased signal intensity of the CSF confirms that there is no cystic lesion in the perimesencephalic cisterns (*E*).

could also become more prominent with this technique. Therefore, one should take into account epidemiologic data and other clinical, laboratory, and imaging findings such as presence of a cyst with a scolex, other cysts, or multiple parenchymal lesions (calcified or not) to diagnose neurocysticercosis (10).

The development of hyperintensity in the CSF after the administration of O_2 is likely due to an increase in the concentration of oxyhemoglobin in the blood, with a paramagnetic effect in the CSF (5, 11). We did not observe an improvement in the detection of intraventricular lesions with this method. The increase of signal intensity of the CSF is less pronounced or even absent within the ventricles when compared with the cortical sulci, fissures, and cisterns probably because of the combination of a less prominent vasculature along the ventricular walls and the dilution of oxyhemoglobin in the ventricles (5).

The evaluation of other types of lesions may also benefit from this technique. For instance, diffusionweighted images allow the diagnosis of epidermoid cysts but are not able to delineate the contours of an arachnoid cyst, which would be possible with MR cisternography. Another theoretical application would be the better demonstration of pathologic superficial vessels, such as an arteriovenous malformation, or suspected superficial cortical lesions.

Conclusion

Noninvasive MR cisternography is a new, easy, low-cost technique that can be used in the evaluation of cortical and extraventricular subarachnoid lesions, such as those of neurocysticercosis.

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Erratum

Article **The Sphenoparietal Sinus of Breschet: Does It Exist? An Anatomic Study** AJNR 25:112–120, January 2004. The sentence on page 112, paragraph 1 which reads "Cruveilhier observed that book II, plate 3 could not be found or confirmed and does not correspond to Breschet's atlas (1)." should read "The reference given by Cruveilhier, that is 'book II, plate3' could not be found or confirmed, and does not correspond to Breschet's atlas (1)."

A misspelling of oligodengrogliomas occurs in the title of the article Glial Tumor Grading and Outcome Prediction Using Dynamic Spin-Echo MR Susceptibility Mapping Compared with Conventional Contrast-Enhanced MR: Confounding Effect of Elevated rCBV of Oligodendroglimoas AJNR 25:214–221, February 2004. The correct title should be Glial Tumor Grading and Outcome Prediction Using Dynamic Spin-Echo MR Susceptibility Mapping Compared with Conventional Contrast-Enhanced MR: Confounding Effect of Elevated rCBV of Oligodendrogliomas.

The authors and their affiliations are incorrectly listed in the article Noninvasive MR Cisternography with Fluid-Attenuated Inversion Recovery and 100% Supplemental O_2 in the Evaluation of Neurocysticercosis, AJNR 25:295–297, February 2004. The authors and their respective affiliations should be listed as:

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Re: Retraction Request: AJNR 23: 868–870, May 2002

Dear Editor:

We request that the manuscript entitled, "Paradoxical Cerebral Arterial Embolization of Cement During Intraoperative Vertebroplasty: Case Report," published in the American Journal of Neuroradiology in May 2002 be retracted. We have discovered several factual errors that we believe warrant its retraction from publication.

We apologize for this inconvenience; however, we believe the integrity of the scientific literature requires the requested action.

Respectfully Submitted,

Joseph Eskridge, M.D. Professor, Dept of Radiology and Neurological Surgery Gavin W. Britz, M.D. Assistant Professor, Dept of Neurological Surgery

The above request for retraction has been reviewed and concurred with by the following University of Washington School of Medicine representatives:

Andrew A. Ziskind, M.D. Associate Vice President for Clinical Specialty Programs Associate Dean for Clinical Affairs

Richard Ellenbogen, M.D. Chair, Dept of Neurological Surgery Norman Beauchamp, M.D. Chair, Dept of Radiology