Interest of routine MR spectroscopic techniques for differential diagnosis between radionecrosis and progression of brain tumor lesions

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ABSTRACT
Objectives: The main objective of the study is to assess the feasibility and reproducibility of routine MRS to assist in the differential diagnosis between post-radiation necrosis and tumor progression. The secondary objective is to evaluate the accuracy of the method.
Method: An additional sequence of MRS was added to the standard protocol routinely used for patient follow-up. To assess discomfort a control group was formed. The time required to perform MRS and analysis of results, and data about artefacts and technical limitations were collected. MRS results analyzed independently by two neuroradiologists were compared. The diagnostic accuracy of MRS was calculated using a composite reference standard.
Results: The experimental group included 38 patients, the control group 41. The discomfort felt during the examination is not significantly different between the groups. The average quality of SRM is rated as low. The frequency of cerebral radionecrosis is 13 % based on the reference standard used, 54 % and 46 % based on MRS results for the two observers. The additional time is 19,5 min. There is strong inter-observer agreement. The sensitivity and specificity of MRS are respectively for the diagnosis of radionecrosis of 60 % and 45 % (PPV = 16 %, NPV = 87 %), for the diagnosis of tumor tissue of 25 % and 94 % (PPV = 80 %, NPV = 57 %).
Conclusion: MRS is probably not applicable in routine clinical practice; however, in view of our results and the literature, in selected cases, it could be a support in the diagnosis of radionecrosis or brain tumor progression. Radionecrosis is probably underestimated.

1. Introduction
The annual incidence of primitive brain tumors in Europe, according to the RARECARE project, is 5–6 per 100,000 [1]. The annual incidence of secondary tumors in the USA based on autopsies is 83.5/100,000. Between 8.5 % and 9.6 % of patients with major neoplasia who are known to give frequent brain metastases (breast neoplasia, colorectal, kidney, lung, and melanoma) are affected; this is probably underestimated and the incidence is increasing [2]. The treatment of primitive and secondary brain tumors involves the use of systemic therapies, surgery, and radiation therapy alone or simultaneously. Radiotherapy treatments and radiosurgery can cause secondary effects. Two of these effects pose a differential diagnostic problem in brain imaging: pseudo progression, observed in early delayed phases, and radionecrosis, observed in late delayed phases [3]. Differentiation of these entities from tumor progression is a challenge in magnetic resonance imaging (MRI). Diagnosis can be determined by combining diffusion, perfusion, magnetic resonance spectroscopy (MRS) techniques, as by comparing the area, volume, and edges of the lesions in T1 and T2 [4]. Recent literature suggests that the combination of multiple magnetic resonance imaging methods reduces the possibility of lesion misinterpretation, however prospective and longitudinal studies are required to validate multiparametric MRI [5,6]. Post-therapeutic residual or recurrent tumor in patients with primitive or secondary brain tumors are major criteria for survival [7], and accurate and rapid diagnosis is very important for proper management. Routine use of MRS maybe could benefit the patient. Our study intend to evaluate this point. Several studies [8–11] have assessed the usefulness of this method, in most cases with limited cohorts, and have not evaluated the feasibility in routine. The main objective of our work is to assess the feasibility of using MRS in routine use for the differential diagnosis between necrosis and tumor progression of brain lesions after radiotherapy or radiosurgery.
2. Materials and methods

We conducted a non-randomized prospective study that added a sequence of Chemical Shift Imaging (CSI) in MRS to usual patient’s follow-up. Patients were included from June 2019 to February 2020. The inclusion criteria were: adult patients with at least one primary or secondary brain tumor lesion and treated with radiation alone or combined with surgery and/or systemic treatments. The criteria for exclusion were: patient under 18 years of age, contraindications to MRI or contraindication to gadolinium injection. MRS preparation, acquisition and time for analysis were measured, artefacts and technical limitations of the method were registered. The MRS sequence performed is a CSI multivoxel, with an echo time (TE) of 135 ms, centered on the area where the tumor(s) were located on the previous examination if it was no longer visible or properly demarcated on the current examination. A control group has been constituted with adult patients benefiting from post-contrast brain MRI in the context of assessment of an extracranial primary cancer. The perception of the exam was assessed by participants in both groups using a numerical visual scale of discomfort calibrated from 0 to 10. The expected sample size was at least 30 patients per group.

The examinations were performed with 3 T and 1.5 T devices with administration of routine dose of Gadolinium (gadoterate at dose 0.2 mL/kg or gadobutrol at 0.1 mL/kg dose). Two radiologists analyzed the standard images and then the MRS sequence. For the analysis of the latest, the reviewers completed a dedicated report form. Spectrum quality was assessed qualitatively according to baseline disturbance on a scale of 0–3 (0 = uninterpretable 1 = baseline very disturbed 2 = baseline moderately perturbed 3 = very well).

The results of morphological sequences were compared with the results of MRS to identify a possible complementary contribution. When available, results of a composite reference standard (including all or part of the following: PET-CT methionine, PET-CT FDG, pathology, successive MRI scans, or clinical evolution) have been considered. For the interpretation of MRS sequences, each observer selected one or more voxels in the available volume and measured the values and ratios of the

Fig. 1. Right temporo-occipital lesion, radionecrosis. A Axial T1 FAT-SAT with Gadolinium showing a contrast enhancement. B voxel chosen. C Voxel’s spectrum suggesting a radionecrosis (Cho/Cr = 1.44-Cho/NAA = 3.56-Cho/Lip + Lac = 0.29).
following metabolites: choline (Cho), creatine (Cr); N-Acetyl-Aspartate (NAA), Lipid (Lip), lactate (Lac). The thresholds for ratios of these metabolites have been established in comparison with studies whose populations like our cohort [10–14]. The Cho/NAA > 1.5 and Cho/Cr > 2 ratios have been selected as thresholds for indicating tumor tissue and the Cho/(lactate + lipid) < 0.3 ratio for radionecrosis (Figs. 1, 2). Other threshold values have been tested to improve the diagnostic accuracy of MRS. A 5-point score was established, as described in Table 1.

The interobserver reproducibility assessment was evaluated by the kappa test. The hypothesis test for the assessment of a statistically significant difference in discomfort during the brain MRI examination with and without MRS is performed using a numerical visual scale and the Mann-Whitney test for independent samples. Univariate analysis was performed for comparing the two groups. Sensitivity and specificity of MRS was calculated using the composite reference standard defined.

3. Results

The group of patients receiving the spectroscopic sequence consists of 38 individuals, Primitive tumors found are listed in Table 2.

The average time between cerebral irradiation and MRS is 1 year and 11 months, in only 3 cases (8 %) this time is less than 3 months; it is between 3 months and 1 year for 8 patients (21 %).

The control group consists of 46 patients, 5 of whom were excluded due to no tumor context. The characteristics of the two groups were compared by univariate analysis (Table 3). This analysis does not show statistically significant differences in age, sex, tumor pathology, corticoids, symptoms, and history of brain surgery.

Fig. 2. Left insular lesion, glioblastoma. A Axial T1 FAT-SAT with Gadolinium showing a contrast enhancement. B voxel chosen. C spectrum indicating tumor tissues (Cho/Cr = 1,57–Cho/NAA = 2,64–Cho/Lip + Lac = 1,13).
Table 1
MRS 5-points score.

| Score | MRS (thresholds successively established) | MORPHOLOGICAL | Global MRI score |
|-------|------------------------------------------|---------------|-----------------|
| 5 - Highly probable tumor recurrence | Cho/NAA > 1.5(1.5-1.1) AND Cho/Chr > 2 (1.5-1.3) Cho/(Lip + Lac) > 0.3(0.6) | tumor tissue | 3 |
| 4 - Suspicion of tumor recurrence | Cho/NAA > 1.5(1.5-1.1) OR Cho/Chr > 2 (1.5-1.3) Cho/(Lip + Lac) > 0.3(0.6) | tumor tissue | 3 |
| 3 - Undetermined | Cho/NAA > 1.5(1.5-1.1) OR Cho/Chr > 2 (1.5-1.3) AND Cho/(Lip + Lac) < 0.3(0.6) | Radiocnosis is possible | 2 |
| 2 - High probability of radionecrosis | Cho/NAA < 1.5(1.5-1.1) OR Cho/Chr < 2 (1.5-1.3) AND Cho/(Lip + Lac) < 0.3(0.6) | Radiocnosis is possible | 2 |
| 1 - Healthy brain | Cho/NAA < 1.5(1.5-1.1) AND Cho/Chr < 2 (1.5-3) AND Cho/(Lip + Lac) > 0.3(0.6) | Healthy brain | 1 |

Table 2
Primitive tumors.

| Primary tumors | n (%) |
|----------------|-------|
| Lung (3 SCLC, 12 ADC, 1 SCC) | 16(42) |
| Breast | 10(26) |
| Melanoma | 6(16) |
| Cerebral (2 glioblastoma, 1 ependymoma, 1 astrocytoma) | 4(10.5) |
| Clear cell renal carcinoma | 1(2.75) |
| Extragonadal germ cell tumor | 1(2.75) |

For the discomfort experienced during the examination, the Mann-Whitney test shows no statistically significant difference between the two groups, with an average assessment of discomfort on a scale of one to ten values 1 (0–4) for the control group and 2 (0–4) for the experimental group.

For the experimental group, one examination resulted in a technical failure (Fig. 2) demonstrate the usefulness of MRS for the follow-up of brain tumors and differential diagnosis between pseudo progression, radionecrosis and neoplastic tissue. However, the concomitant presence of tumoral and necrotic tissue is possible [22], and molecular imaging such as MRS can confirm this entity. Considering literature data, MRS scores, and the relative quality of reference standard protocols [18], Studies addressing only metastatic lesions versus radionecrosis are small, with varying results and metabolites ratio thresholds, Huang et al. [10] showed a sensitivity of 33 % and a specificity of 100 %, or Menoux et al. [11] 92 % and 56 % respectively. In the meta-analysis of Wang et al., the prevalence of radionecrosis varies from 14 % to 45 % depending on the studies [8].

4. Discussion

The technique of multi-voxel MRS (Spectroscopic Imaging or CSI) seemed to be the most suitable for our study, allowing the study of metabolites distribution in a larger region compared to single-voxel technology, however, this technique is more sensitive to metal artefacts. Its acquisition time is longer, and it performs less well in several brain regions: posterior fossa anterior part of the temporal lobe, frontal lobe [15]. In our study, 56 % of the lesions are in these regions. The localization of lesions is probably one of the factors that has influenced the quality of MRS. Other techniques of improvement, such as the reduction of the shim field of view (FOV) and the focus on the part of lesion to be examined may be considered [16,17]. The training of operator for this technique should be improved if it is to be used in routine.

The additional acquisition time of 14 min, the reading time of 5–6 min, and the failure rate of 10% argue against routine use of MRS. The single voxel technique has not been explored. It allows for a shorter acquisition time, a better homogeneity of the field, but requires a precise knowledge of the part of lesion to be studied [15]. The use of the two techniques in context could improve the time and quality limitations highlighted here. The reproducibility of brain MRI sequences between MRI system manufacturers is good despite the absence of international standardized protocols [18]. Studies addressing only metastatic lesions versus radionecrosis are small, with varying results and metabolites ratio thresholds, Huang et al. [10] showed a sensitivity of 33 % and a specificity of 100 %, or Menoux et al. [11] 92 % and 56 % respectively. In the meta-analysis of Wang et al., the prevalence of radionecrosis varies from 14 % to 45 % depending on the studies [8].

Spectroscopy in oncological brain imaging has an interest in the follow-up and diagnosis of both primary and secondary tumor lesions. In gliomas, it allows for the molecular identification of certain subtypes of gliomas [19], it also allows correlation to the Ki67 antigen expression [20], and an increase of more than 20 % choline is associated with tumor progression [21]. About the choice of thresholds, our aim was to improve the sensitivity for radionecrosis and the specificity for tumor tissue, in addition the specificity of 90 % and the 80 % VPP for tumor tissue detection (Fig. 2) demonstrate the usefulness of MRS for the follow-up of brain tumors and differential diagnosis between pseudo progression, radionecrosis and neoplastic tissue. However, the concomitant presence of tumoral and necrotic tissue is possible [22], and molecular imaging such as MRS can confirm this entity. Considering literature data, MRS scores, and the relative quality of reference standard, the prevalence of radionecrosis in our cohort could be above 13 %.

Recent literature suggests that the combination of multiple magnetic resonance imaging and advanced imaging methods reduces the possibility of lesion misinterpretation and increase sensitivity, specificity and accuracy to differentiate tumor progression and treatment-related changes however prospective and longitudinal studies are required to validate and standardize multiparametric methods [6,23], the aim of our study was to assess the feasibility of using MRS in routine use when...
there was an enhancing lesion in the area that was previously treated, most of the time for the follow-up of brain metastatic disease. A systematic multiparametric approach was not performed and perfusion and diffusion data are not available for all patients.

Radiological investigations involve anxiety in oncological patients and not, the anxiety activates the autonomic nervous system causing hyperactivity, harmful to the quality of radiological examinations [24], in this study the addition of a sequence of MRS has not shown a difference on comfort of patients during the examination.

5. Conclusion

MRS is probably not a clinical routine technique due to the methodological difficulties, failure rate and the extra time required. However, for selected cases, considering the results of the literature, the inter-observer concordance, the small impact on comfort of patients, MRS could add arguments for the diagnosis of cerebral radionecrosis, and detection of persistent or recurrent brain tumor. Radionecrosis is probably underestimated. Standardized metabolites thresholds and reports for MRS remain to be defined.

CRediT authorship contribution statement

Federico De Lucia, Yolene Lefebvre, Marc Lemort: Conceptualization, Methodology, Data curation, Writing – original draft, Visualization, Investigation, Writing – review & editing. Marc Lemort, Yolene Lefebvre: Supervision.

Table 3
Univariate analysis of the two groups.

| Variables                        | Experimental group | Control group | p-value |
|----------------------------------|--------------------|---------------|---------|
| Age (years)                      | 56.5(13)           | 58 (13)       | 0.65**  |
| Sex                              |                    |               |         |
| Female                           | 23 (60)            | 22 (54)       | 0.54**  |
| Male                             | 15 (40)            | 19 (46)       |         |
| Cerebral surgery                 |                    |               |         |
| Yes                              | 10 (26)            | 5 (12)        | 0.11**  |
| No                               | 28 (74)            | 36(88)        |         |
| Orthopnea                        |                    |               |         |
| Yes                              | 4 (10.5)           | 4 (10,5)      | 1**     |
| No                               | 34 (89,5)          | 37 (89.5)     |         |
| MRI                              |                    |               |         |
| Yes                              | 1 (3)              | 1 (3)         | 1***    |
| No                               | 37 (97)            | 40 (97)       |         |
| MRI 3 T                          | 19 (50)            | 14 (34)       | 0.15**  |
| MRI 1.5 T                        | 19 (50)            | 27 (66)       |         |
| Neurological symptoms            |                    |               |         |
| Yes                              | 6 (16)             | 11(29)        | 0.17**  |
| No                               | 32 (84)            | 30(71)        |         |
| Neoplasias                       |                    |               |         |
| Pulmonary                        | 16 (42,1)          | 19 (46,3)     | 0.06*** |
| Breast                           | 10 (26,3)          | 8 (19,5)      |         |
| Melanoma                         | 6 (15,8)           | 5 (12,2)      |         |
| Glioma                           | 4 (10,5)           | 0 (0)         |         |
| Other                            | 2 (5,3)            | 9 (22)        |         |
| Discomfort (EVN)                 | 20(4–)             | 1(0–4)        | 0.84****|

Table 4
Results based on reference standard and MRS SCORE (with the first set thresholds initially established) for each observer.

| Conclusion   | MRS SCORE | Observer1 (%) | Observer2 (%) | Reference standard (%) |
|--------------|-----------|---------------|---------------|------------------------|
| Radionecrosis | 2         | 20/37(54)     | 17/37(46)     | 5/38(13)               |
| Tumor        | 3–4.5     | 6/37(16)      | 5/37(14)      | 25/38(66)              |
| Healthy brain | 1         | 11/37(30)     | 15/37(40)     | 8/38(21)               |

Table 5
The best sensitivity and specificity of MRS with different thresholds, calculated using the composite reference standard for the diagnosis of radionecrosis and tumor tissue.

| Thresholds | Radionecrosis | Tumoral tissue |
|------------|---------------|----------------|
| 1) Cho/NAA = 1.5 Cho/Cr = 2 | Se = 60 % | Se = 25 % |
| Cho/(Lip + Lac) = 0.3 | Sp = 45 % | Sp = 94 % |
| VPP = 16 % | VPP = 80 % |
| VPN = 87 % | VPN = 57 % |
| Se = 75 % | Se = 31 % |
| 2) Cho/NAA = 1.5 Cho/Cr = 1.5 | Sp = 54 % | Sp = 89 % |
| Cho/(Lip + Lac) = 0.6 | VPP = 20 % | VPP = 71 % |
| VPP = 93 % | VPP = 59 % |
| Se = 67 % | Se = 62.5 % |
| 3) Cho/NAA = 1.1 Cho/Cr = 1.3 | Sp = 55 % | Sp = 64 % |
| Cho/(Lip + Lac) = 0.6 | VPP = 14 % | VPP = 67 % |
| VPP = 96 % | VPP = 60 % |
Ethical Statement

The study was approved by the Jules Bordet Institute ethics committee under the number CE3010.

The study was conducted in accordance with the principles embodied in the Declaration of Helsinki and in accordance with local statutory requirements.

All participants gave written informed consent to participate in the study.

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Conflict of Interest

The authors have no conflict of interest.

References

[1] E. Crocetti, A. Trama, C. Stiller, et al., Epidemiology of glial and non-glial brain tumors in Europe, Eur. J. Cancer 48 (10) (2012) 1532–1542, https://doi.org/10.1016/j.ejca.2011.12.013.

[2] E. Taboure, L. Bouchet, A. Carpentier, Epidemiologie des metastases cerebrales et tumeurs neurovasculaires, Bull. Cancer 100 (2013) 57–62, https://doi.org/10.1684/bdc.2012.1681.

[3] J. Fink, D. Born, M.C. Chamberlain, Radiation necrosis: relevance with respect to treatment of primary and secondary brain tumors, Curr. Neurol. Neurosci. Rep. 12 (3) (2012) 276–285, https://doi.org/10.1007/s11910-012-0258-7.

[4] A. Raimbault, X. Cazals, M.A. Lauvin, et al., Radionecrosis of malignant glioma and cerebral metastasis: a diagnostic challenge in MRI, Diagn. Interv. Imaging 95 (10) (2014) 985–1000, https://doi.org/10.1016/j.diii.2014.06.013.

[5] N. Verma, M.C. Cowperthwaite, M.G. Burnett, et al., Differentiating tumor recurrence from treatment necrosis: a review of neuro- oncologic imaging strategies, Neuro Oncol. 15 (5) (2013) 515–534, https://doi.org/10.1093/neo/mct307.

[6] V. Sawlani, N. Davies, M. Patel, et al., Evaluation of response to stereotactic radiosurgery in brain metastases using multiparametric magnetic resonance imaging and a review of the literature, Clin. Oncol. 31 (1) (2019) 41–49, https://doi.org/10.1016/j.clon.2018.09.003.

[7] M.K. Bucci, A. Maity, A.J. Janss, et al., Near complete surgical resection predicts a favorable outcome in pediatric patients with nonbrainstem, malignant gliomas: results from a single center in the magnetic resonance imaging era, Cancer 101 (4) (2004) 817–824, https://doi.org/10.1002/cncr.20042.

[8] W. Wang, Y. Hu, P. Lu, et al., Evaluation of the diagnostic performance of magnetic resonance spectroscopy in brain tumors: a systematic review and meta-analysis, PLoS One 9 (11) (2014), e112577, https://doi.org/10.1371/journal.pone.0112577.

[9] M.T. Chuang, Y.S. Liu, Y.S. Tsai, et al., Differentiating radiation-induced necrosis from recurrent brain tumor using MR perfusion and spectroscopy: a meta-analysis, PLoS One 11 (1) (2016), e0141458, https://doi.org/10.1371/journal.pone.0141438.

[10] J. Huang, A.M. Wang, A. Shetty, et al., Differentiation between intra-axial metastatic tumor progression and radiation injury following fractionated radiation therapy or stereotactic radiosurgery using MR spectroscopy, perfusion MR imaging or volume progression modeling, Magn. Reson. Imaging 29 (7) (2011) 993–1001, https://doi.org/10.1016/j.mri.2011.04.004.

[11] I. Menoux, G. Noel, D. Antoni, et al., PET scan and NMR spectroscopy for the differential diagnosis between brain radiation necrosis and tumour recurrence after stereotactic irradiation of brain metastases: place in the decision tree, Cancer Radiother. 21 (5) (2017) 389–397, https://doi.org/10.1016/j.crrrad.2017.03.003.

[12] E. Matsunae, J.R. Fink, J.K. Rockhill, et al., Distinction between glioma progression and post-radiation change by combined physiologic MR imaging, Neuoradiology 52 (4) (2010) 297–306, https://doi.org/10.1007/s00234-009-0613-9.

[13] M.R. Anbarloui, S.M. Ghodsi, A. Khoshnevisan, et al., Accuracy of magnetic resonance spectroscopy in distinction between radiation necrosis and recurrence of brain tumors, Iran. J. Neurol. 14 (1) (2015) 29–34.

[14] T. Kimura, K. Sako, T. Gotoh, et al., In vivo single-voxel proton MR spectroscopy in brain lesions with ring-like enhancement, NMR Biomed. 14 (6) (2001) 339–491, https://doi.org/10.1002/nbr.711.

[15] P.B. Barker, A. Bizi, N. De Stefano, et al., Pulse sequences and protocol design, in: P.B. Barker, A. Bizi, N. De Stefano, et al. (Eds.), Clinical MR Spectroscopy: Techniques and applications, Cambridge University Press, Cambridge, 2010, pp. 19–33.

[16] J. Stockmann, L. Wald, Neuroimage In vivo B0 field shimming methods for MRI at 7 T, Neuroimage 168 (2018) 71–87, https://doi.org/10.1016/j.neuroimage.2017.06.013.

[17] B. Van Den Bergen, C. Van Den Berg, D. Klop, et al., SAR and power implications of different RF shimming strategies in the pelvis for 7 T MRI, Magn. Reson. Imaging 30 (1) (2009) 194–202, https://doi.org/10.1016/j.mri.2008.09.012.

[18] M. Pavazan, M. Mikkelien, A. Berrington, et al., Comparison of multivendor single-voxel MR spectroscopy data acquired in healthy brain at 26 sites, Radiology 295 (1) (2020) 171–180, https://doi.org/10.1148/radiol.2020191037.

[19] G. Oz, J. Alger, P. Barker, et al., Clinical proton MR spectroscopy in central nervous system disorders, Radiology 270 (3) (2014) 658–679, https://doi.org/10.1148/radiol.13113051.

[20] R. Guellivan, C. Mensel, H. Duffau, et al., Proton magnetic resonance spectroscopy predicts proliferative activity in diffuse low-grade gliomas, J. Neurooncology. 87 (2) (2008) 181–187, https://doi.org/10.1007/s12075-007-9508-y.

[21] G. Tedeschi, N. Lundbom, R. Raman, et al., Increased choline signal coinciding with malignant degeneration of cerebral gliomas: a serial proton magnetic resonance spectroscopy imaging study, J. Neurosurg. 87 (4) (1997) 516–524, https://doi.org/10.3171/jns.1997.87.4.0316.

[22] S. Ritu, V. Surjith, J. Rojymon, et al., Radiation necrosis in the brain: imaging features and differentiation from tumor recurrence, Radiograpges 32 (5) (2012) 1343–1359, https://doi.org/10.1148/rg.325125006.

[23] D. Qin, G. Yang, H. Jing, et al., Tumor progression and treatment-related changes: radiological diagnosis challenges for the evaluation of post treated glioma, Cancers 14 (2022) 3771, https://doi.org/10.3390/cancers14153771.

[24] G. Lo Re, R. De Luca, F. Muscarneri, et al., Relationship between anxiety level and radiological investigation. Comparison among different diagnostic imaging exams in a prospective single-center study, Radiol. Med. 121 (10) (2016) 763–768, https://doi.org/10.1007/s11547-016-0664-z.