RESEARCH

Differentiating between non-functioning pituitary macroadenomas and sellar meningiomas using ADC

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Abstract

Introduction and aim: It is difficult to distinguish between non-functioning pituitary macroadenomas (NFPMA) and sellar meningiomas because of their overlapping imaging manifestations on routine MRI, especially in cases of meningiomas growing into the saddle. Here, we aimed to differentiate between these two tumors using apparent diffusion coefficient (ADC) values and MRI characteristics.

Methods: A total of 60 NFPMA and 52 sellar meningioma cases confirmed by the pathological analysis were retrospectively reviewed. All patients were examined via routine MRI and diffusion-weighted imaging (DWI) before undergoing surgery. The clinical characteristics, MRI characteristics, and max ADC (ADC_{max}), average ADC (ADC_{mean}), and minimum ADC (ADC_{min}) values were compared between the two tumors via Chi-square test and two sample t-tests. Receiver operating characteristic (ROC) curve and binary logistic regression analyses were conducted to determine the discrimination ability.

Results: The ADC_{max}, ADC_{mean}, and ADC_{min} values were significantly higher in NFPMA compared to sellar meningiomas ($P < 0.001$ for all). Among ADC values, ADC_{max} demonstrated good performance with an AUC of 0.896 (95% CI, 0.823–0.969) and accuracy of 88.7%. A cut-off value of $0.97 \times 10^{-3} \text{ mm}^2/\text{s}$ was used for ADC_{max} for differentiation between tumors. A combination of ADC_{max} values and clinicoradiological features showed the best discrimination ability for differential diagnosis between the two tumors, with an AUC of 0.981 (95% CI, 0.958–1.000) and accuracy of 96.9%.

Conclusion: A combination of ADC_{max} and clinicoradiological features demonstrates good discrimination ability and high accuracy for differentiation between NFPMA and sellar meningiomas, and is a potential quantitative tool to aid in the selection of surgical techniques.

Key Words

- ▶ ADC
- ▶ DWI
- ▶ meningiomas
- ▶ MRI
- ▶ pituitary macroadenoma

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Introduction

Pituitary adenomas and meningiomas are the most common benign tumors in the CNS (1). Pituitary adenomas arise from the pituitary gland and account for

10–15% of all intracranial tumors (2, 3). Non-functioning pituitary adenomas account for 14–54% of pituitary adenomas and lack biochemical evidence of hormone

secretion (4). Tumors with a diameter of greater than or equal to 10 mm are considered macroadenomas (2). In contrast, sellar meningiomas arise from adjacent dural surfaces, including the dura surrounding the pituitary fossa, diaphragm sellae, and sella turcica, and represent approximately 5–10% of all cranial meningiomas (5). When suprasellar or parasellar meningiomas grow downward into the pituitary fossa, which is particularly observed in combined tuberculum and diaphragm sellae meningiomas (Type B) (6) and pituitary macroadenomas that arise within the sella turcica and extended superiorly, differential diagnosis via routine MRI becomes difficult (5, 7). However, precise preoperative distinction between these two tumors may aid the determination of a suitable surgical planning. Most sellar meningiomas may require a craniotomy and Dzhindzhikhadze *et al.*, have reported that a transpalpebral approach may be followed, whereas most supra- and intrasellar macroadenomas are resected via the trans-sphenoidal route (8, 9, 10, 11).

Currently, some studies have reported on the identification of these two tumors with traditional methods (e.g., CT and MRI) (5, 8, 12). Wu *et al.* (12) reported that quantitative dual-energy CT imaging, iodine content, slope of HU curve, and mean CT value may be valuable parameters for diagnostic differentiation between sellar meningiomas and pituitary adenomas. MRI was mainly performed as the examination of choice for the diagnosis of sellar lesions (13). Some MRI characteristics, such as visibility of the pituitary gland, contrast enhancement, and sellar enlargement can help differentiate pituitary macroadenomas from sellar meningiomas (8). However, the aforementioned methods sometimes may not lead to a confirmed etiological diagnosis because of the uncertainty of the origin of a lesion (13). Diffusion-weighted imaging (DWI) can noninvasively provide information concerning high signal-intensity regions as areas of increased cellularity and tissue microstructures by measuring water diffusion within a tissue (14, 15, 16), which is closely associated with membrane integrity and cell density (15, 17). The diffusion of water molecules is determined by a high cellularity, a small extracellular space, and a high nuclear-to-cytoplasmic ratio (18). The apparent diffusion coefficient (ADC) is a parameter that is derived from DWI, and represents the average diffusion for each voxel. Accordingly, the ADC values in brain tumors inversely associated with cellularity and the ratio of nuclear area to cytoplasm (18). Previous studies have reported that the ADC value or ratio plays an important role in the differentiation of brain tumors, including benign and atypical/malignant meningiomas and solitary fibrous

tumors/hemangiopericytomas and meningiomas (16, 19, 20, 21). However, few studies have assessed the role of ADC in the differentiation of non-functioning pituitary macroadenomas (NFPMAs) from sellar meningiomas.

Therefore, in this study, we compared the clinical and MRI characteristics and ADC values between NFPMAs and sellar meningiomas, and determined whether ADC values can improve the diagnostic accuracy of these two tumors.

Materials and methods

Patients

For this retrospective analysis, ethical approval was obtained from the Institutional Review Board of Lanzhou University Second Hospital (Lanzhou, China), and the need for informed consent was waived. We analyzed 60 patients with NFPMAs (24 females and 36 males; age, 52.6 ± 11.2 years) and 52 patients with sellar meningiomas (43 females and 9 males; age, 52.3 ± 8.53 years) confirmed surgically and pathologically at our institute between January 2014 and December 2019. All patients were consecutively admitted to the hospital and underwent routine MRI and DWI within one week before surgery. The images of each patient were of good quality and without artifacts. Patients showing grossly macrocystic and/or hemorrhagic changes ($n=5$ in the NFPMAs group) on measuring ADC values and those for whom ADC values could not be measured ($n=3$ in the NFPMAs group and $n=7$ in the sellar meningioma group) because of air-tissue interface artifacts were excluded. Accordingly, ADC values of 52 patients with NFPMAs and 45 patients with sellar meningiomas were measured. In all, there were 65 cases of reduced vision, 21 cases of blurred vision, 13 cases of headache and dizziness, 4 cases of epilepsy, and 9 cases of other manifestations.

Imaging protocol

In this study, the Siemens Verio 3.0 T superconducting MR scanner (Germany) was used, and 112 patients underwent T1WI, T2WI, DWI, and contrast-enhance MRI before surgery. The parameters for MRI included the following: T1WI (TR/TE=550/11 ms), slice thickness=5.0 mm, layer spacing 1.5 mm, matrix=256 × 256, field of view (FOV)=260 mm × 260 mm; T2WI (TR/TE=2200/96 ms), echo chain length=8, and number of excitation (NEX)=2. The parameters for DWI (SEPI sequence) were as follows: frequency selection fat suppression technology,

(TR/TE=4000/100 ms), slice thickness=5 mm, layer spacing 1.5 mm, matrix=256 × 192, FOV=260 mm × 260 mm, and diffusion gradients applied in three orthogonal directions (b value=0, 1000 s/mm²). Axial, sagittal, and coronal enhanced T1WI were obtained using Gd-DTPA (Bayer Schering Pharma AG) at 0.1 mmol/kg at a rate of 3.0 mL/s.

Image analysis and processing

Two neuroradiologists (readers 1 and 2, with 12 and 15 years of experience in brain MRI diagnosis, respectively) blindly evaluated the patients data twice within 15 days. In case of a difference in opinions, agreements were reached after discussion between the two. MRI characteristics including maximal diameter, T2W signal, enhancement features, hemorrhage/cyst/necrosis-related changes, DWI signal, sella enlargement, dural tail sign, waist sign (the core of the tumor is thinner than the sides), carotid artery encircling, depression of saddle, pituitary stalk visibility, and pituitary gland visibility were observed and recorded.

ADC maps were automatically generated after DWI. The ADCs of the solid part of the tumors were calculated from the original images. On a continuous-level ADC map, the free-form marker tool was used to manually delineate each slice and measure ADCmean. Subsequently, the regions of interest (ROIs, size 10–20 mm²) were manually placed to cover solid tumor areas and avoid hemorrhagic or cystic lesions, with 6–8 ROIs per level, and ADCmax and ADCmin values were determined. Then, the averages of the ADCmax and ADCmin values of both neuroradiologists were calculated.

Statistical analysis

SPSS 23.0 software was used for statistical analysis. Chi-square test was used to compare the MRI characteristics (tumor variables). Two-sample t tests were performed for the comparisons of age, maximal diameter, and ADC value. Then, variables were included in the multivariate logistic regression analysis. For all statistical analyses, a P value less than 0.05 indicated a statistically significant difference. Differences between the ADC parameters calculated by the two radiologists were evaluated by the intraclass correlation (ICC) test. An ICC value of greater than 0.75 indicated excellent consistency, and average values were calculated. The receiver operating characteristic (ROC) curves and binary logistic regression analysis were used to differentiate between the diagnostic accuracy and optimal thresholds.

Results

Patient clinical characteristics

NFPMAs were diagnosed in 24 females and 36 males, whereas sellar meningiomas were diagnosed in 9 females and 43 males ($P < 0.001$). These findings indicating a statistically significant difference in the incidence between sexes. No significant differences were observed in terms of age between the two disease groups (52.6 ± 11.2 vs 52.3 ± 8.53 years; $P=0.145$), as shown in Table 1.

MRI characteristics and ADC value analysis

After univariate analysis, significant differences were observed in terms of T2W signals, enhancement features, hemorrhage/cysts/necrosis-related changes, sella enlargement, waist sign, depression of saddle, pituitary stalk visibility, and pituitary gland visibility between the NFPA and sellar meningioma groups in routine MRI, all $P < 0.05$. These two groups were similar in terms of the following parameters: maximal diameter, dural tail sign, DWI signal, and carotid artery encircling. Subsequently, the above nine significant MR features were reduced to five independent discrimination factors (including enhancement features, hemorrhage/cysts/necrosis-related changes, depression of saddle, and pituitary gland visibility) by multivariate logistic regression analysis. The aforementioned data are presented in Table 1. Typical MRI-features of these two tumors are shown in Supplementary Figs 1 and 2 (see section on [supplementary materials](#) given at the end of this article).

The ICCs were calculated to evaluate the agreement of the ADC parameters measured by two radiologists, and all values >0.75 , reflecting good agreement. The ADCmax (1.125 ± 0.243 vs $0.853 \pm 0.166 \times 10^{-3}$ mm²/s), ADCmean (1.053 ± 0.249 vs $0.816 \pm 0.168 \times 10^{-3}$ mm²/s), and ADCmin (0.976 ± 0.270 vs $0.778 \pm 0.172 \times 10^{-3}$ mm²/s) values were significantly higher in the NFPA group compared to the sellar meningioma group (all $P < 0.001$), as shown in Table 2.

ROC curve analysis

As shown in ROC curve analyses (Table 3), the ADCmax indicated a relatively high discrimination ability with an AUC of 0.896 (95% CI, 0.823–0.969), accuracy of 88.7%, sensitivity of 82.7%, and specificity of 93.30%; an optimal cutoff value of 0.97×10^{-3} mm²/s was used for discriminating NFPMAs from sellar meningiomas. For clinicoradiological

Table 1 The clinical and magnetic resonance imaging features of non-functioning pituitary macroadenomas and sellar meningioma.

Clinicoradiological features	NFPMA (n = 60)	Sellar meningioma (n = 52)	Univariate analysis (P value)	Multivariate analysis (P value)	Odds ratio (95% CI)
Age (years, mean $\pm$ s.d.)	52.6 $\pm$ 11.2	52.3 $\pm$ 8.53	0.145	–	–
Sex (male/female)	36/24	9/43	0.001 ^a	0.018	4.482 (1.300–15.454)
Maximal diameter (cm)	2.97 $\pm$ 1.09	3.18 $\pm$ 1.21	0.347	–	–
T2W signal-high	50 (83.3%)	29 (55.8%)	0.001 ^a	–	–
Obvious enhancement feature	20 (33.3%)	41 (78.8%)	<0.001 ^a	0.006	0.184 (0.055–0.616)
Hemorrhage/cysts/necrosis changes	42 (70%)	13 (25%)	<0.001 ^a	0.002	6.846 (2.029–23.092)
DWI signal-high	19 (31.7%)	18 (34.6%)	0.151	–	–
Sella enlargement	55 (91.7%)	25 (48.1%)	<0.001 ^a	–	–
Dural tail sign	26 (43.3%)	32 (61.5%)	0.061	–	–
Corset sign	28 (46.7%)	9 (17.3%)	0.001 ^a	–	–
Encircle carotid artery	20 (33.3%)	14 (26.9%)	0.539	–	–
Depression of saddle	49 (81.7%)	19 (36.5%)	<0.001 ^a	0.016	4.976 (1.343–18.438)
Visibility of the pituitary stalk	11 (18.3%)	21 (40.4%)	0.012 ^a	–	–
Visibility of the pituitary gland	3 (5%)	27 (51.9%)	<0.001 ^a	0.003	0.075 (0.014–0.416)

A Student's *t*-test was used to compare the difference in age, while the chi-square test was used to compare the difference in other clinicoradiological features.

^a*P* < 0.05.

NFPMA, non-functioning pituitary macroadenomas.

features, including sex, enhancement features, hemorrhage/cysts/necrosis-related changes, depression of saddle, and pituitary gland visibility, a diagnostic efficacy was achieved with an AUC of 0.934 (95% CI, 0.890–0.978), accuracy of 87.5%, sensitivity of 88.5%, and specificity of 81.7%. Thus, on the combined use of ADCmax values and clinicoradiological features, the discrimination ability was improved to an AUC of 0.981 (95% CI, 0.958–1.000), accuracy of 96.9%, sensitivity of 95.6%, and specificity of 94.2%, which was indicative of the best identification ability and highest accuracy, as shown in Fig. 1.

Discussion

This was a preliminary study combining ADCmax and clinicoradiological features for the distinction between

Table 2 Comparison of ADCmax value, ADCmean value, and ADCmin value between non-functioning pituitary macroadenomas and sellar meningioma.

Parameter ($\times 10^{-3}$ mm ² /s)	NFPMA (n = 52)	Sellar meningioma (n = 45)	P value
ADCmax value	1.125 $\pm$ 0.243	0.853 $\pm$ 0.166	<0.001 ^a
ADCmean value	1.053 $\pm$ 0.249	0.816 $\pm$ 0.168	<0.001 ^a
ADCmin value	0.976 $\pm$ 0.270	0.778 $\pm$ 0.172	<0.001 ^a

A Student's *t*-test was used to compare the difference in ADC value.

^a*P* < 0.001.

ADC, apparent diffusion coefficient; NFPMA, non-functioning pituitary macroadenomas.

NFPMA and sellar meningiomas, which were identified based on the MRI to date. In this study, we showed that the use of both ADCmax and clinicoradiological features had the best discrimination ability and highest sensitivity, and correctly identified the two tumors in 96% of all cases; therefore, this method can be applied to patients who underwent MRI before surgery.

In our study, sex was identified to be a significant factor in subgroup analysis. Females (82.69%) were prone to develop meningiomas, and females accounted for 40% of NFPMA cases, which was similar to the results reported in previous studies (8, 22, 23). The clinical manifestations of both tumors were similar, mainly showing reduced vision, blurred vision, headache, and dizziness.

MRI is the main examination method for intracranial tumors. Our present findings demonstrated significant differences in some MRI characteristics between NFPMA and sellar meningiomas, which may be associated with their pathophysiological changes. In this study, the proportion of high signals in T2WI in NFPMA was higher compared to that in sellar meningiomas, which may indicate that the former is more prone to hemorrhages, cysts, and necrosis-related changes (70% in this study) than the latter. Most sellar meningiomas showed more obvious, uniform enhancement (78.8% in this study), whereas most NFPMA indicated heterogeneous, relatively poor enhancement, which was consistent with the results presented in previous studies (8, 24, 25, 26). We found that the proportion of NFPMA with

Table 3 Diagnostic performance of different parameters for differentiating non-functioning pituitary macroadenomas from sellar meningioma.

Observed parameters	AUC	ACC (%)	SEN (%)	SPE (%)	Cut off
ADCmax	0.896 (0.823–0.969)	88.7	82.7	93.3	0.97 ($\times 10^{-3}$ mm ² /s)
ADCmix	0.787 (0.789–0.947)	80.4	71.2	88.9	0.86 ($\times 10^{-3}$ mm ² /s)
ADCmean	0.868 (0.692–0.883)	86.6	84.6	86.7	0.89 ($\times 10^{-3}$ mm ² /s)
Clinicoradiological features	0.934 (0.890–0.978)	87.5	88.5	81.7	0.3108
Clinicoradiological features and ADCmax	0.981 (0.958–1.000)	96.9	95.6	94.2	0.5103

ACC, accuracy; AUC, area under receiver operating characteristic curve; NFPMAs, nonfunctioning pituitary macroadenomas; SEN, sensitivity; SPE, specificity.

waist signs was significantly higher compared to that of sellar meningiomas, which may be closely associated with the soft texture of NFPMAs. The tumor is blocked when growing through the saddle septum, which leads to waist signs; however, sellar meningiomas are firm in

texture. Furthermore, sella enlargement, depression of saddle, and non-visibility of the pituitary stalk and gland were more commonly encountered in NFPMAs. Together, these features proved the different origins and growth patterns of the two tumors. Sellar meningiomas (except those arising from the dura surrounding the pituitary fossa) first grow above the saddle, then break through the saddle septum and extend downward the saddle. However, NFPMAs first grow in the saddle, spherically enlarging the saddle, cause depression of saddle, and then break through the saddle septum, thereby extending onto the saddle. The aforementioned features showed a strong association with the identification of these two tumors and can be selected to be as predictive factors.

DWI has evolved as a functional MRI modality that utilizes the changes in water molecule diffusion motion to provide information about the tumor microenvironment (27). No statistical differences in DWI signals were observed between these two tumors, which may be because there are many factors affecting the DWI signals. These factors mainly include diffusion sensitivity coefficient (b) and T2 penetration effect, particularly air-tissue interface artifacts from the skull base in our study, among others (28). ADC provides important additional information beyond that provided by MRI (18, 29). The main advantage of using ADC is its ability to extract quantitative information and enable assessment of viable cellularity (28, 30). In this study, we observed that the ADC values were significantly higher in NFPMAs compared to in sellar meningiomas ($P < 0.001$), and ADCmax indicated a relatively higher discrimination ability than ADCmin and ADCmean, with an AUC of 0.896 (optimal cutoff value, 0.97×10^{-3} mm²/s) for distinguishing NFPMAs from sellar meningiomas. These findings, agreed with previously published reports (31), and are possibly due to the wider range of ADC values in pituitary adenomas that tend to be non-homogeneous compared to homogeneous meningiomas. Although low ADC values indicated restricted diffusion of water

Figure 1

(A) ADCmax, ADCmean, and ADCmin values for differential diagnosis of non-functioning pituitary macroadenomas and sellar meningiomas. Compared with ADCmean and ADCmin values, ADCmax value demonstrated a relatively high discriminating ability, with an AUC of 0.896. (B) Compared with the clinicoradiological features alone, combining ADCmax and clinicoradiological features indicated the best identification ability, with an improved AUC of 0.981. ADC, apparent diffusion coefficient.

identified in meningiomas, some authors have reported that the ADC of pituitary adenomas was not significantly different from that of meningiomas (32, 33). This may be related to the small sample size of studies reported in the past.

After analyzing the ADC values, we found that it may be associated with the ratio of extracellular and intracellular gaps and the different cell types (34). NFPMA demonstrated fibrous tissue hyperplasia and loosely arranged cells, sellar meningioma cells were closely connected by intercellular cell junctions, with increased intracellular complex protein molecules, and intercellular areas are packed by numerous cytoplasmic processes (34, 35). Therefore, compared with NFPMA cells, sellar meningioma cells may have more intracellular space and less extracellular space, thereby limiting the diffusion of water. The reason for different ADC values may be explained based on these histological differences.

ADCmax and clinoradiological features together performed better than ADCmax and clinoradiological features alone, resulting in an AUC of 0.981, accuracy of 96.9%, sensitivity of 95.6%, and specificity of 94.2%, and significantly improved the identification ability between NFPMA and sellar meningiomas. The results showed that ADC combined with clinoradiological features, as a noninvasive method, provided more diagnostic information, thereby aiding in the accurate diagnosis of these two tumors before surgery and help in selecting an appropriate surgical plan.

This study has several limitations. First, sellar and parasellar lesions tend to cause limitations in ADC value measurement because of susceptibility artifacts, such as air-tissue interface artifacts. Second, this is a single-center study, and multi-center studies will be needed to further validate the results. Finally, multimodal MRI techniques such as perfusion-weighted imaging (PWI) and susceptibility-weighted imaging (SWI), were not included in this study and will have to be explored in the future.

Conclusion

In conclusion, ADCmax values and clinoradiological features are useful for distinguishing NFPMA from sellar meningioma. Moreover, combining ADCmax and clinoradiological features may further improve this discrimination ability.

Supplementary materials

This is linked to the online version of the paper at <https://doi.org/10.1530/EC-20-0434>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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