Imaging Biomarkers for Adult Medulloblastomas: Genetic Entities May Be Identified by Their MR Imaging Radiophenotype

V.C. Keil, M. Warmuth-Metz, C. Reh, S.J. Enkirch, C. Reinert, D. Beier, D.T.W. Jones, T. Pietsch, H.H. Schild, E. Hattingen, and P. Hau

ABSTRACT

BACKGROUND AND PURPOSE: The occurrence of medulloblastomas in adults is rare; nevertheless, these tumors can be subdivided into genetic and histologic entities each having distinct prognoses. This study aimed to identify MR imaging biomarkers to classify these entities and to uncover differences in MR imaging biomarkers identified in pediatric medulloblastomas.

MATERIALS AND METHODS: Eligible preoperative MRIs from 28 patients (11 women; 22–53 years of age) of the Multicenter Pilot-study for the Therapy of Medulloblastoma of Adults (NOA-7) cohort were assessed by 3 experienced neuroradiologists. Lesions and perifocal edema were volumetrized and multiparametrically evaluated for classic morphologic characteristics, location, hydrocephalus, and Chang criteria. To identify MR imaging biomarkers, we correlated genetic entities sonic hedgehog (SHH) TP53 wild type, wingless (WNT), and non-WNT/non-SHH medulloblastomas (in adults, Group 4), and histologic entities were correlated with the imaging criteria. These MR imaging biomarkers were compared with corresponding data from a pediatric study.

RESULTS: There were 19 SHH TP53 wild type (69%), 4 WNT-activated (14%), and 5 Group 4 (17%) medulloblastomas. Six potential MR imaging biomarkers were identified, 3 of which, hydrocephalus (P = .03), intraventricular macrometastases (P = .02), and hemorrhage (P = .04), when combined, could identify WNT medulloblastoma with 100% sensitivity and 88.3% specificity (95% CI, 39.8%–100.0% and 62.6%–95.3%); WNT-activated nuclear β-catenin accumulating medulloblastomas were smaller than the other entities (95% CI, 5.2–22.3 cm³ versus 35.1–47.6 cm³; P = .03). Hemorrhage was exclusively present in non-WNT/non-SHH medulloblastomas (P = .04; n = 2/5). MR imaging biomarkers were all discordant from those identified in the pediatric cohort. Desmoplastic/nodular medulloblastomas were more rarely in contact with the fourth ventricle (4/15 versus 7/13; P = .04).

CONCLUSIONS: MR imaging biomarkers can help distinguish histologic and genetic medulloblastoma entities in adults and appear to be different from those identified in children.

ABBREVIATIONS: AUC — area under the curve; CE — contrast-enhanced; CMB — classic medulloblastoma; DNMB — desmoplastic/nodular medulloblastoma; SHH — sonic hedgehog; WHO — World Health Organization; WNT — wingless

Medulloblastomas (World Health Organization [WHO] grade IV) rarely occur in adults. According to the United States registry analysis from the Surveillance, Epidemiology, and End-Results data base, incidence rates around 0.6 cases per million have been recorded, which is >50 times lower than the incidence of glioblastoma.1–3 A higher age at diagnosis is a negative prognostic factor for survival, with a median overall survival currently between 7.7 and 9.7 years, provided patients receive the best medical care.4 The 2016 revision of the WHO classification of CNS tumors introduced the concept of an integrative medulloblastoma diagnosis.5 The diagnosis includes 4 histologically and 4 genetically defined entities known to have an influence on the...
course of the disease in both children and adults.\textsuperscript{6–12} The genetic entities that are currently defined are sonic hedgehog (SHH)-activated (and exist with or without TP53 mutation), wingless (WNT)-activated, and non-SHH/non-WNT (Groups 3 and 4) medulloblastomas. The defined histologic groups are classic (CMB), large cell anaplastic, desmoplastic/nodular (DNMB) medulloblastomas and medulloblastoma of extensive nodularity. The exact classification at the earliest possible time point is of great importance to evaluate prognosis and possible targeted therapies.

Radiogenomics is a dynamically evolving field in radiology based on standard diagnostic MR imaging. It seeks to identify so-called MR imaging biomarkers that may predict the genetic profile of a tumor, assuming that the genetic profile is reflected in a distinctive radiophenotype, and can also be of benefit when true genetic analysis is not available.\textsuperscript{13} Only 1 study has been published in a predominantly pediatric cohort, which investigated a radiogenomic approach to differentiate genetically defined medulloblastoma entities.\textsuperscript{14} The authors found that genetic entities were distinguishable by several MR imaging biomarkers such as tumor location or enhancement pattern. Different relative frequencies and varying prognostic influences of the genetic medulloblastoma entities between adult and pediatric cohorts suggest that MR imaging biomarkers identified in pediatric cohorts may be different from those in adult medulloblastoma.\textsuperscript{8–10,12,15,16}

The Multicenter Pilot-study for the Therapy of Medulloblastoma of Adults (NOA-07) is the first prospective trial of an adult medulloblastoma cohort that systematically evaluated radiochemotherapy as the first-line treatment and included, among others, imaging biomarkers in its analysis. The study presented here is dedicated to identifying MR imaging biomarkers that will allow differentiation of medulloblastoma genetic entities based on the entirely adult NOA-07 cohort. Identification of such MR imaging biomarkers may facilitate presurgical tumor assessment and assist in the categorization of the differences between adult and pediatric MR imaging biomarkers.

\textbf{MATERIALS AND METHODS}

\textbf{Patient and Imaging Data}

This trial was registered at ClinicalTrials.gov (NCT01614132) and under the EudraCT number 2007-002560-10 (https://www.clinicaltrialsregister.eu/ctr-search/search?query=2007-002560-10) after approval by the ethics committee of the University of Regensburg, Regensburg, Germany (08-112-05a; substantial amendment of July 1, 2016) and of all participating sites. Between 2008 and 2014, 15 neuro-oncologic centers recruited 30 adult patients older than 21 years of age with suspected medulloblastomas. Medulloblastoma diagnosis was confirmed in all cases. Presurgical CT and MR imaging datasets were retrospectively analyzed involving multiparametric imaging criteria based on T1-weighted, T2-weighted, FLAIR, and contrast-enhanced T1-weighted sequences (available in \(n = 28/30\) as well as apparent diffusion coefficient maps (\(n = 23/30\), On-line Figure)). In 2 of 30 cases, presurgical MR imaging data were not available.

\textbf{Imaging Criteria Definition and Analysis}

Three neuroradiologists (E.H., M.W.-M., V.C.K., with 26, 28, and 6 years of experience, respectively) who were blinded to the histologic and genetic backgrounds of each patient, evaluated noncontrast-enhanced CT scans and MR imaging datasets (1.5T or 3T). Criteria were grouped as follows: 1) classic morphologic imaging characteristics applied to describe brain tumors, 2) Chang medulloblastoma staging criteria and tumor location,\textsuperscript{17} 3) diagnostic criteria of hydrocephalus or brain stem compression, and 4) tumor and edema volumes and respective ratios.

Classic morphologic characteristics of brain tumors were defined as the following: enhancement pattern (strong, weak, nonenhancing), signal homogeneity (homogeneous, inhomogeneous), signal intensity (hypoointense, isointense, hyperintense to gray matter) on nonenhanced T1-weighted and T2-weighted images, the sharpness of tumor margins, and the presence of cyst formation and hemorrhage. The presence of hemorrhage was diagnosed from either additional SWI or T2-weighted gradient-echo sequences or on presurgical CT.

Location criteria were the following: infiltration of the vermis, the hemispheres, or the brain stem/peduncular region, including contact with the lower rhombic lip, the eighth cranial nerve, the fourth ventricle, or the cerebellopontine angle. In addition, the presence of hydrocephalus, brain stem compression, midline shift, supratentorial growth, and multifocality was evaluated.

Medulloblastoma volumes were determined with commercially available software (tumor tracking tool, Intellispace Portal 5.0; Philips Healthcare, Best, the Netherlands) on automatically aligned contrast-enhanced T1WI (CET1WI) and T2-weighted images on the basis of manually defined signal-intensity thresholds. Perifocal edema was quantified on FLAIR datasets. Volume ratios were established as the following: 1) tumor volume ratio, which was defined as the volume fraction of the tumor part with exclusive T2 signal elevation and without the contrast-enhancing volume fraction (T2-weighted volume minus CET1WI volume divided by T2-weighted volume); and 2) edema-tumor ratio, which was defined as edema volume on FLAIR images divided by T2-weighted tumor volume.

\begin{table}[h]
\centering
\begin{tabular}{|l|c|c|}
\hline
\textbf{Histologic entity (No.)} & \textbf{Adult (This Cohort,} \(n = 28\)) & \textbf{Pediatric (Perreault et al.} \(n = 47\)) \\
\hline
\hline
\textbf{In clinic} & \textbf{Median age (range) (yr)} & \textbf{Median age (range) (yr)} \\
\hline
\textbf{Male/female} &  & \\
\hline
\textbf{SHH} & 19 (67.9\%) & 13 (27.7\%) \\
\textbf{WNT} & 4 (14.2\%) & 8 (17.0\%) \\
\textbf{Group 4} & 5 (17.9\%) & 17 (36.2\%) \\
\textbf{Group 3} & 0 (0.0\%) & 13 (27.7\%) \\
\hline
\textbf{Histologic entity (No.)} & \textbf{Classic} & \textbf{Desmoplastic/nodular} \\
\hline
\textbf{Other subgroups} & 13 (46.4\%) & 31 (66.0\%) \\
\textbf{Large cell anaplastic} & 15 (53.6\%) & 4 (8.5\%) \\
\textbf{Desmoplastic/nodular} & 0 (0.0\%) & 10 (21.3\%) \\
\textbf{Classic} & 0 (0.0\%) & 2 (4.2\%) \\
\hline
\end{tabular}
\caption{Adult and pediatric cohorts by histology and genetic status}
\end{table}

Data were compared with that in the cohort of Perreault et al.,\textsuperscript{14} in which 42 of 47 cases were pediatric (Table 1).

\textbf{Table 1: Adult and pediatric cohorts by histology and genetic status}

| Histologic entity (No.) | Adult (This Cohort, \(n = 28\)) | Pediatric (Perreault et al. \(n = 47\)) |
|-------------------------|----------------------------------|---------------------------------------|
| SHH                     | Median age (range) (yr)          | 39.0 (33.4–41.3)                      |
|                         | Male/female                      | 1:1.0                                 |
|                         | Genetic entity (No.) (%)         | 1:1.0                                 |
| SHH                     | 19 (67.9\%)                      | 13 (27.7\%)                           |
| WNT                     | 4 (14.2\%)                       | 8 (17.0\%)                            |
| Group 4                 | 5 (17.9\%)                       | 17 (36.2\%)                           |
| Group 3                 | 0 (0.0\%)                        | 13 (27.7\%)                           |
| Classic                 | 13 (46.4\%)                      | 31 (66.0\%)                           |
| Desmoplastic/nodular    | 15 (53.6\%)                      | 4 (8.5\%)                             |
| Large cell anaplastic   | 0 (0.0\%)                        | 10 (21.3\%)                           |
| Other subgroups         | 0 (0.0\%)                        | 2 (4.2\%)                             |

\textbf{Note:—SHH} indicates sonic hedgehog–activated medulloblastoma; \textbf{WNT}, wingless-activated medulloblastoma.
The criteria of Chang et al.\textsuperscript{17} were determined on the basis of tumor margins on both T2-weighted and CET1-weighted images. Macroscopically visible subependymal/ventricular metastases (Chang M-stage ≥2) were determined on CET1WI. Micrometastases (Chang M-stage 1) were previously investigated in the CSF at the time of diagnosis on the basis of the evaluation of 2 board-certified neuropathologists in all cases.\textsuperscript{17}

**Neuropathologic Evaluation**

Medulloblastoma diagnoses had been confirmed at the German Brain Tumor Reference Center of the German Society for Neuropathology and Neuroanatomy by at least 2 board-certified neuropathologists. Classification was performed with histologic, immunohistochemical, and genetic analyses according to the revised WHO Classification of Tumors of the CNS 2016 as described in current Best Practice Guidelines.\textsuperscript{18} Histologically and genetically defined medulloblastoma entities were available for all 28 patients. In addition, the whole genomic copy number and allelic distribution were analyzed by molecular inversion probe methodology without evidence of amplifications in MYCC, MYCN, or GLI2 genes. One SHH-activated tumor was found to exhibit copy number losses of 9q and chromosome 14, typical for SHH-activated entities. A subgroup of samples was classified according their epigenetic subgroup by methylation arrays (450k array), which confirmed the genetically defined subgroups in all cases.\textsuperscript{19,20} The complete analysis catalogue is listed in the On-line Figure.

Imaging criteria of the comparative pediatric cohort are listed in Table 2 extracted from the earlier publication.\textsuperscript{14}

**Statistics**

Statistical analyses were designed by an independent biomathematician and performed with STATA 14.0 (StataCorp, College Station, Texas). Because no tentative assumptions regarding the relationship between imaging criteria and genetic or histologic status could be made, an exploratory statistical approach to identify possible MR imaging biomarkers involved testing imaging criteria against all other criteria with 2-sided Fisher exact or ANOVA testing (depending on whether categoric or continuous data were handled). Deliberately, no multiple testing correction approach was chosen to not overlook any possible significant associations.\textsuperscript{21} Statistical significance was reached at $P < .05$.

Imaging criteria that were significantly associated with genetic criteria, according to the exploratory tests, were defined as potential imaging biomarkers. These were then assessed regarding their sensitivity and specificity in the differentiation of genetic entities. Evaluation involved both single and combined MR imaging biomarker applications and was based on an area under the curve (AUC) analysis.

To investigate potential differences between adult and pediatric biomarkers, we compared imaging biomarkers identified in the present study with those of a previous study involving a predominantly pediatric cohort.\textsuperscript{14}

**RESULTS**

**Patient Cohort**

The cohort of adult patients with medulloblastoma in this NOA-07 substudy included 11 women and 17 men (detailed overview in Table 1). In this cohort, no SHH-activated TP53–mutated medulloblastomas were found, all were TP53 wild type. Histologically, no cases with large cell anaplastic histology or with extensive nodularity were present. All WNT-activated and non-WNT/non-SHH medulloblastomas were histologically CMBL, while 4 of 15 SHH-activated medulloblastomas were DNMB ($P = .001$).

**Imaging Biomarkers of Genetic Entities and Chromosomal Aberrations**

Six categoric and continuous volumetric imaging criteria could be identified as MR imaging biomarkers to differentiate genetically defined entities of SHH TP53 wild type, WNT-activated, and non-SHH/non-WNT Group 4 medulloblastomas in this cohort (Table 3). Notably, apart from hemorrhage, none of the classic MR imaging criteria used for tumor description, as defined in the “Materials and Methods” section, were identified as suitable MR imaging biomarkers.

The absence of the biomarker “hydrocephalus” was observed in 4 of 4 WNT-activated medulloblastomas. Hence, normally

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**Table 2: Comparative overview of MR imaging biomarkers for adult and pediatric cohorts regarding genetic entity discrimination**

| Adult Cohort Imaging Criteria (n = 28) | P Value | Correspondence | P Value | Pediatric Cohort Imaging Criteria (n = 47)* |
|--------------------------------------|---------|----------------|---------|------------------------------------------|
| Hemorrhage\textsuperscript{b}        | .04*    | No\textsuperscript{b} | NS      | Blood products                           |
| Edema volume\textsuperscript{b}      | .02\textsuperscript{b} | No\textsuperscript{b} | NS      | Edema present                            |
| Hydrocephalus\textsuperscript{b}     | .03\textsuperscript{b} | NA             | NA      | NA                                       |
| Chang M-stage ≥2\textsuperscript{b}  | .02\textsuperscript{b} | NA             | NA      | NA                                       |
| Contact with lower rhombic lip\textsuperscript{b} | .03\textsuperscript{b} | NA             | NA      | NA                                       |
| Chang T-stage > T2 on T2WI\textsuperscript{b} | .01\textsuperscript{b} | NA             | NA      | NA                                       |
| Contact with fourth ventricle\textsuperscript{b} | .02\textsuperscript{b} | Pseudocorrespondence | <.005\textsuperscript{b} | Midline/fourth ventricle\textsuperscript{b} |
| Infiltration of vermis                | .80     | No\textsuperscript{b} | NS      | Enhancement pattern                      |
| Cysts                                 | .82     | Both NS         | NS      | Cyst                                     |
| Contrast-enhancement homogeneity      | .29     | Both NS         | NS      | Enhancement pattern solid                |
| Infiltration of cerebellar hemispheres| .08     | No\textsuperscript{b} | <.005\textsuperscript{b} | Cerebellar hemisphere\textsuperscript{b} |
| Contact with cerebellopontine angle   | 1.0     | No\textsuperscript{b} | <.005\textsuperscript{b} | Contact with cerebellopontine angle\textsuperscript{b} |
| Enhancement pattern                  | .81     | No\textsuperscript{b} | <.005\textsuperscript{b} | Enhancement minimal to none\textsuperscript{b} |
| Sharp margins                        | .63     | No\textsuperscript{b} | .03\textsuperscript{b} | Ill-defined margins\textsuperscript{b} |
| NA                                   | NA      | NA             | NS      | Enhancement pattern, other               |

Note:—NA indicates not available (parameter not analyzed); NS, criterion not significantly unbalancedly distributed, hence not identified a biomarker by Perreault et al.\textsuperscript{14}

* Pediatric cohort: discovery cohort of Perreault et al.\textsuperscript{14} No identical MR imaging biomarkers could be identified among imaging criteria of both cohorts.

**SHH**-activated and non-**SHH**/non-**WNT** medulloblastomas were histologically CMBL, while 4 of 15 **SHH**-activated medulloblastomas were DNMB ($P = .001$).
Table 3: Imaging biomarkers for adult medulloblastoma genetic entity differentiation

| Imaging Biomarker | P Value | SHH (n = 19) | WNT (n = 4) | Group 4 (n = 5) |
|------------------|---------|--------------|-------------|----------------|
| Hydrocephalus    | .03b    | 12/19        | 0           | 4/5            |
| Chang M-stage ≥ 2 (CET1WI) | .02c    | 10/19        | 0           | 1/5            |
| Hemorrhage       | .04a    | 0            | 0           | 2/5            |
| Contact with fourth ventricle | .02b   | 3/19         | 3/4         | 3/5            |
| Contact with lower rhombic lip | .03a   | 0            | 1/4         | 2/5            |
| Chang T-stage ≥ 2 (T2WI) | .01b   | 17/19        | 1/4         | 4/5            |
| Tumor volume on T2WI (median and 95% CI) | .06a   | 30.6 cm³ (5.2–22.3 cm³) | 5.6 cm³ (6.6–7.7 cm³) | 0.04–4.74 cm³ |

Note: SHH indicates sonic hedgehog-activated medulloblastoma (only TP53 wild type in this cohort), WNT, wingless-activated medulloblastoma.

Comparison with Previously Reported MR Imaging Biomarkers for Medulloblastoma Entities in Pediatric Patients

Relative frequencies of histologically and genetically defined medulloblastoma entities were different between both cohorts: None of the MR imaging biomarkers identified in the adult cohort corresponded to those identified in the comparative predominantly pediatric cohort (Table 2). Midline/fourth ventricle contact was a significant biomarker to identify non-SHH/non-WNT medulloblastomas in the pediatric cohort. Midline location of the lesion was not significant in our cohort (criterion “vermis”). Contact with the fourth ventricle was analyzed separately for the present adult cohort and was determined as a biomarker with excess representation in WNT-activated medulloblastomas (Table 3).

DISCUSSION

According to the recent revision of the WHO classification of brain tumors, the definition of genetic and histologic features of medulloblastoma is a prerequisite for a proper diagnosis of this tumor. In both pediatric and adult patients, genetically and histologically defined entities were determined to be relevant for prognosis and patient outcome. Therefore, precise characterization of medulloblastomas can facilitate the design of personalized therapies. Biomarkers based on standard MR imaging techniques have been successfully applied in the diagnosis of numerous brain lesions, including medulloblastomas.

As part of the criteria of Chang et al.,17 MR imaging morphometric analyses are well-established for medulloblastomas and are relevant in the prognosis of both adult and pediatric cohorts. However, entities of adult medulloblastomas seem to behave differently from the same entities in pediatric cohorts. To our knowledge, this is the first study to date that analyzes MR imaging biomarkers for genetic entity discrimination of exclusively adult patients with medulloblastoma in comparison with a pediatric cohort. The imaging criteria used in this study differ slightly from those presented in the study of Perreault et al.14 Because the present...
study was a multicenter study, sequence parameters and image quality varied among the centers; these differences make the analysis more difficult. In detail, the pediatric comparative cohort did not differentiate between 2 of the biomarkers in our cohort (“contact with the fourth ventricle” and “vermis/midline location”) and did not include the Chang criteria or hydrocephalus. Remarkably, no coherent MR imaging biomarkers for the same genetic entities could be identified between adult and pediatric cohorts.

As a main result of our study, WNT-activated genetic entities of adult patients with medulloblastoma could be differentiated from SHH-activated TP53 wild type and non-SHH/non-WNT Group 4 medulloblastomas by a combination of MR imaging biomarkers. Adult WNT-activated tumors were observed to be correlated to several imaging biomarkers, which might be described as their radiophenotype. At the time of tumor detection (and thus of neurologic symptoms), WNT tumors are in contact with the fourth ventricle, but without Chang M-stage $\geq 2$ and are of small volume (lower Chang T-stages), in consequence without hydrocephalus. This finding might suggest that early midline location at the fourth ventricle induces neurologic symptoms, which allow diagnosis despite low tumor size. In contrast, most SHH-activated or non-SHH/non-WNT tumors were larger, multilocular, or focally metastatic and associated with hydrocephalus. Smaller, monofocal tumors can be more easily resected in total. This characteristic may also partially explain a frequently identified higher survival in these patients. Most interesting, an association of WNT-activated medulloblastomas in the lower rhombic lip could only be confirmed for a minority of tumors in this trial, despite the embryologic association of WNT-activation within the lower rhombus.

The combination of 3 of 6 identified imaging biomarkers, hemorrhage, subependymal macrometastases, and hydrocephalus, allowed an optimal discrimination of WNT-activated medulloblastomas from the other 2 genetic entities in our cohort. All MR imaging biomarkers relevant for genetic entity discrimination in this cohort were, however, different from those identified for the comparative pediatric cohort. The pediatric cohort did not differentiate between 2 of the biomarkers in our cohort (contact with the fourth ventricle and vermis/midline location). Contact with the fourth ventricle is, therefore, a pseudo-
consistent biomarker between both cohorts because it was linked to the criterion midline location in the pediatric cohort, which was not identified as a biomarker in our study.

Commonly, medulloblastoma location is stratified into med- dial, cerebellopontine, and hemispheric lesions. However, at- tempts to use location criteria as biomarkers for differentiation of genetic medulloblastoma entities have yielded only inconsistent results.14,29–31 Our cohort also exhibits this inconsistency, be- cause no significant association between genetic entities and tu- mor location was observed. This may suggest that these location categories are not useful as biomarkers in adult medulloblasto- mas. A lack of association between cerebellar location and genetic entity is, to some extent, in conflict with findings that suggested that different medulloblastoma genetic entities had a spatially and cytologically distinct tumorigenesis.2,7,29,32

Classic morphologic tumor criteria such as signal homogene- ity were not significantly associated with genetic entities in our study. This finding was strikingly different from the data of Perreault et al.,14 in which sharpness of tumor margins, enhance- ment pattern, and location categories were highly predictive of genetic entities (though tumor margins later proved not an ade- quate MR imaging biomarker when tested with the validation cohort of the pediatric study). Hemorrhage was the only classic morphologic tumor criterion in our cohort that exclusively oc- curred in Group 4 tumors. However, hemorrhage was also re- ported in WNT-activated medulloblastomas.27,33 Therefore, this MR imaging biomarker should be re-evaluated in a larger inde- pendent cohort.

Considering the substantial amount of genetic and chromo- somal factors investigated in this study, it remains surprising how many of these factors apparently do not result in a particular radiophenotype, especially when considering chromosomal aber- rations. It can thus be postulated that the biologic impact of these factors cannot be quantified by standard MR imaging sequences. Quantitative MR imaging sequences and multifactorial computer-assisted image data analyses may, however, allow a better identifica- tion of biomarkers.34

As a limitation of our study, the comparative pediatric cohort comprised 4 adults (besides 43 children). However, the main limi- tation is the small size of our cohort. All findings regarding im- aging biomarkers must, therefore, be taken as hypothesis-gener- ating and should be tested for their validity in larger cohorts in the future. Different from our comparative pediatric cohort, in the discovery cohort of Perreault et al.,14 no validation cohort was available to test the identified MR imaging biomarkers for adult medulloblastoma genetic entity differentiation in a blinded set- ting. Further research is needed to determine why biomarkers for medulloblastoma entities in pediatric patients differed from adult biomarkers identified in this study and also why classic tumor imaging criteria were usually not relevant as MR imaging bio- markers in this study. Possible reasons can be statistical effects due to a small cohort size, different methods of image assessment, but also a true lack of influence of the genetic entity on medulloblas- toma radiophenotype. An international prospective trial with larger patient numbers is planned and will be an opportunity to elaborate on these open questions.

CONCLUSIONS

Several MR imaging biomarkers could be identified in this cohort of adult patients with medulloblastoma and allowed successful presurgical discrimination of genetically as well as histologically defined medulloblastoma entities. MR imaging biomarkers used in differentiation of medulloblastomas in adults seem distinct from biomarkers in children. Both findings will need to undergo further validation by radiogenomic analyses of larger medul- loblastoma cohorts.

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