



CORRESPONDENCE OPEN

Fear and discomfort of children and adolescents during MRI: ethical consideration on research MRIs in children

Pediatric Research _____; <https://doi.org/10.1038/s41390-020-01277-6>

INTRODUCTION

Magnetic resonance imaging (MRI) is the imaging modality of choice for clinical diagnostics and research settings. Despite the fact that unседated MRI without contrast agents is defined as a minimal-risk procedure,^{1–4} some research review boards consider exposure of healthy children and adolescents to magnetic fields of 1.5 or 3 Tesla only for research purposes ethically unjustifiable. According to the US regulations, minimal risk means that “the probability and magnitude of harm or discomfort anticipated in the research are not greater than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or tests”.³ Despite that MRI is a low-risk procedure, it might come along with some burden. Determining the level of burden associated with a study protocol is a difficult endeavor, especially in pediatric research.

Only few studies describe fear and discomfort of children during research MRI.^{5–9} The likelihood of discomfort during clinical MRI is reported to be very low with more than 98% children and adolescents stating no or only minimal fear during clinical MRI.⁵ The same study demonstrated that anxiety levels did not differ between MRI and electroencephalography assessments; however, first-ever MRI was related to significantly higher physiological arousal than repeated MRI.⁵ A functional MRI study compared anxious children to non-anxious children and adults and did not find a difference in their emotional reaction to the MRI.⁶ Children are suggested to experience less discomfort and more enjoyment during MRI than adults.⁷ In older studies, the insertion of a needle for the application of contrast agents, the confined space, loud noise, and the instruction to lay still was suggested to particularly cause anxiety and distress in some children and adolescents.^{8,9} Research projects in developmental neuroscience rarely use contrast agents, particularly not when healthy controls are included. From a psychological point of view, unседated MRI without contrast agents in pediatrics meets the minimal-risk standard.^{4,10} Various strategies exist which help reduce the level of fear and discomfort during MRI in children (see refs. ^{11–13}, and for an overview of such strategies see Table S4). Children showed stronger engagement, more comfort, and more enjoyment than adults⁷ and furthermore rated the MRI as less distressing than their parents did.⁹ These findings highlight the positive side of enrolling children in volunteer-based research MRI.

Likewise, the risk for physical harm in unседated MRI is very low.^{1,4,14} Radiologist’s associations provide useful information on hazards that need to be considered during MRI and for risk assessment.^{15,16} However, they exclude children and do not separate between clinical and research MRI.⁴ Consequently, international controversies remain whether it is ethically justifiable to perform MRI scans in children and adolescents for research purposes only.

The guidelines of swissethics (Swiss Association of Research Ethics Committees) precise that in research procedures with children “not only the objective risk, but also the subjective experience of the minors plays a role”.¹ To avoid experience of discomfort and to deal with a supposedly low risk–benefit ratio, some review boards recommend using, instead of healthy controls, MRI exams from “healthy” children having undergone MRI for other reasons (i.e. suspicion of epilepsy, brain tumor, stroke or others). However, these children do not present a healthy sample, receive different MRI sequences than used for research assessments, and are therefore unsuitable as a control group.

To summarize, the review boards’ recommendation to exclude healthy children from MRI studies does not appear feasible and lacks scientific basis. It is rather a worrying obstruction, which hinders new insights into the developing brain.

The aim of the present study is to provide empirical evidence on fear and discomfort during unседated research MRI without contrast agents in 212 participants aged 7–18 years. We further considered aspects that—based on previous research—are expected to relate to fear and discomfort during MRI such as age,¹⁷ intelligence,¹⁸ parental education level,¹⁹ and cognitive self-control.²⁰ Additionally, this study provides data on the intra-individual longitudinal development of fear and discomfort during MRI between childhood and adolescence. Our findings will be helpful for review boards evaluating study protocols including research MRIs in healthy children and adolescents and will enable review boards to base their decision about the risk–benefit ratio on group-level data and hence on solid grounds.

METHODS

Between 2010 and 2018, participants were scanned by doctoral students in psychology who received an extensive introduction and practical training on how to perform MRI in children. Participants became familiar with the noise inside the scanner via audio presentation and were carefully instructed on the MRI procedure in detail before the scan. Participants, who wished for their parents to be close, were scanned with a parent sitting next to them. In between every MR sequence, the examiner talked to the participants over the intercom to make sure that everything is all right.

To capture the extent of children’s fear and discomfort, all participants completed a self-report rating scale immediately after the MRI, such as suggested in a previous study.²¹ Answers were given on a smiley-based scale (0 = no discomfort/no fear, 1 = almost no discomfort/almost no fear, 2 = a little discomfort/a little fear, 3 = considerable discomfort/considerable fear, 4 = high discomfort/high fear), ranging from a friendly laughing smiley (=0) to a very worried smiley (=4). Fear and discomfort were each assessed on a separate scale.

Parental education was chosen as one of the many possible proxies for socio-economic status and was defined as the highest maternal and paternal education level according to the Swiss education system (no graduation = 1, college = 2, college of

Received: 15 July 2020 Revised: 1 October 2020 Accepted: 2 October 2020
Published online: 20 April 2021

higher education = 3, university degree = 4). Intelligence was assessed using the WISC-IV (see ref. ²¹; NEMO-Study) and the TONI-4 (see ref. ²²; Brainfit-Study). Cognitive self-control was captured using an inhibition task (color word interference task²³). All cognitive assessments and questionnaires were administered on the day of scanning.

In total, 102 healthy controls and 110 patients (7–18 years) were included. All participants were either enrolled in the NEMO-Study²⁴ assessing children born very preterm ($n = 51$) and healthy controls ($n = 49$) or the Brainfit-Study,²⁵ examining survivors of childhood cancer ($n = 59$) and healthy controls ($n = 53$). For inclusion criteria see the corresponding references.

Both studies took place at the Children's University Hospital in Bern, Switzerland; study protocols were approved by the local ethics committee. All parents and participants signed written informed consent prior to enrollment if >14 years (for participants <14 years, the legal guardian signed the consent).

Categorical data (MRI questionnaire, parental education) was analyzed using Mann–Whitney U -test. T -test and Chi-square test was used to analyze group difference concerning age (T -test) and sex (Chi-square). To evaluate the relationship between fear, discomfort and sex, parental education, IQ, and cognitive self-control, Spearman correlation was used. For longitudinal analyses, Wilcoxon signed ranks test was applied to compare fear and discomfort in childhood versus adolescence. The level of significance was set to p 0.05 (two tailed). Bonferroni correction was applied for correlation analyses setting the p value to 0.007.

RESULTS

Mean age at exam ($t = 2.409$, $p = 0.017$), IQ ($t = 2.855$, $p = 0.005$), and parental education ($U = 3517.5$, $p = 0.000$) differed significantly between patients and controls (Table S1). IQ differed significantly between studies, with preterm-born participants (NEMO-Study, $M = 100.88$) presenting significantly lower IQ than childhood cancer survivors (Brainfit-Study, $M = 106.14$, $t = 2.674$, $d.f. = 108$, $p = 0.009$). In controls, IQ did not differ between studies ($t = -0.299$, $d.f. = 94.097$, $p = 0.766$).

Patients and controls did not differ concerning perceived fear ($U = 5193.0$, $p = 0.298$) and discomfort ($U = 5432.0$, $p = 0.671$, Table S2). The majority of participants experienced no or almost no fear (82.1%) and discomfort (74.0%, Fig. 1). In all, 17.5% experienced a little fear and 22.6% a little discomfort, whereas the

rate of considerable fear (0.5%) or discomfort (3.3%) was very low. The maximum score of high fear/discomfort was never given, neither in controls nor in patients. There was a significant sex difference, with discomfort being higher in male than female patients ($U = 1174.0$, $p = 0.036$), whereas the level of fear did not differ between males and females ($U = 1444.0$, $p = 0.703$). In controls, no sex difference occurred in terms of discomfort ($U = 1244.5$, $p = 0.714$) or fear ($U = 1230.5$, $p = 0.631$) during the MRI.

In controls, fear and discomfort was positively associated. In patients, age was negatively associated with discomfort whereas maternal education correlated positively with cognitive self-control and paternal education was positively associated with IQ (Fig. 2).

Longitudinal analysis on the development of fear and discomfort between childhood and adolescence showed that fear during MRI decreased significantly between childhood and adolescence in patients ($z = -2.36$, $p = 0.018$) but not in controls ($z = -1.22$, $p = 0.222$, Table S3). The level of perceived discomfort remained stable between childhood and adolescence in patients ($z = -0.218$, $p = 0.827$) and controls ($z = -0.922$, $p = 0.356$).

DISCUSSION

We provide empirical data on fear and discomfort associated with unседated head MRI in 212 children and adolescents aged 7–18 years (without the use of contrast agents). Self-reported fear and discomfort was very low in participants undergoing research MRI with younger patients reporting more fear than older participants. Older participants are likely to have more experience in handling unfamiliar situations and had more opportunities to internalize possible coping strategies than younger participants and might therefore be less prone to a fearful reaction. Based on the present finding, a general exclusion of minors from MRI research studies due to possible discomfort does not appear justified.

Our results show more discomfort in male than in female patients. The literature concludes that the male and female brain differs in structure and function considerably²⁶ and that males and females use different ways to solve the same problem and converge on the same behavior,²⁷ a fact likely reflected in the sex difference found in our study.

Review boards are often concerned that children do not benefit from participating in neuroimaging research studies and hence presume a low risk–benefit ratio. Our experience and the experience of others⁷ show that MRI is an excellent teaching tool giving children

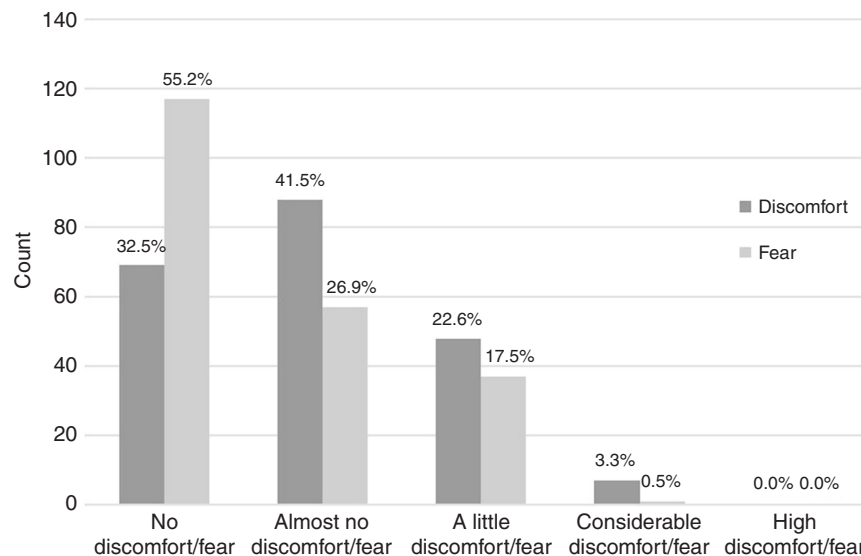


Fig. 1 Fear and discomfort in children. Number and percentages of children experiencing fear and discomfort on a scale between 0 (no fear/discomfort) to 4 (high fear/discomfort).

| | Fear | Discomfort | Age | IQ | Cognitive self-control | Maternal education | Paternal education |
|------------------------|----------------|-----------------|----------------|-----------------|------------------------|--------------------|--------------------|
| Fear | <i>r</i> | 0.277 | -0.165 | 0.125 | -0.010 | -0.017 | 0.042 |
| | <i>p</i> | (0.005*) | (0.097) | (0.808) | (0.924) | (0.867) | (0.690) |
| Discomfort | 0.084 | | -0.126 | 0.229 | 0.002 | 0.032 | 0.242 |
| | (0.383) | | (0.206) | (0.020) | (0.983) | (0.760) | (0.020) |
| Age | -0.076 | -0.196 | | -0.191 | 0.059 | -0.130 | -0.206 |
| | (0.439) | (0.040) | | (0.054) | (0.573) | (0.209) | (0.049) |
| IQ | -0.077 | 0.038 | 0.097 | | -0.112 | 0.081 | 0.078 |
| | (0.424) | (0.695) | (0.314) | | (0.281) | (0.435) | (0.460) |
| Cognitive self-control | 0.112 | -0.160 | 0.267 | 0.219 | | -0.057 | 0.094 |
| | (0.271) | (0.116) | (0.008) | (0.030) | | (0.595) | (0.392) |
| Maternal education | -0.197 | -0.119 | -0.055 | 0.182 | 0.293 | | 0.077 |
| | (0.045) | (0.228) | (0.577) | (0.064) | (0.005*) | | (0.464) |
| Paternal education | -0.051 | -0.048 | 0.042 | 0.305 | 0.061 | 0.615 | |
| | (0.607) | (0.628) | (0.677) | (0.002*) | (0.562) | (0.000*) | |

Fig. 2 Correlation between output measures (fear and discomfort) and age at exam, IQ, cognitive self-control, and parental education ($n = 102$ controls above the diagonal line; $n = 110$ patients below the diagonal line).

the opportunity to interact with modern medical technology, get an insight into potential career choices, and contribute to research. These are striking benefits that are neglected in the discourse on the risk–benefit ratio of pediatric study participants. By all means, we recommend to formulate elaborated research questions with convincing rationales to clarify the necessity of a study and to justify any minor inconveniences experienced by the children for the sake of scientific advancement.

Recruiting healthy children nowadays is challenging and expensive and often forces researchers to focus merely on the study of ill children. To allow for statements about brain development and reorganization after disease, the comparison with healthy children's brains is inevitable. For example, the investigation on resting-state functional networks after stroke in childhood needs normative data of healthy controls²⁸ or studying characteristics of the working memory network in preterm-born children is impossible without the knowledge on the development of the working memory network in healthy children.²⁹ Recently, open access imaging data are available for healthy children and adolescents (i.e. <https://data.rocklandsample.rfmh.org>). These datasets allow for the inclusion of control data serving as normative information when studying patients. However, technical constraints often hinder the comparison of open access imaging data with imaging sequences from the local site.³⁰

Limitations of our study are the following: Children undergoing MRI for clinical reasons might experience more fear and discomfort than participants in our research setting who consented to the study procedure on a voluntary basis and likely exhibit extraordinary motivation to undergo an MRI scan. Additionally, it has been shown that post-MRI anxiety is significantly lower than pre-MRI anxiety,^{10,31,32} pointing towards

a positive bias due to relieving feelings after the MRI scan. Whether specific factors such as the noise of the scanner, the restricted space, the presence of a parent or the parenting style contribute to perceived fear and discomfort remains to be investigated in future studies.

To conclude, our data present evidence that the majority of study participants do not suffer from discomfort during the MRI scan. Restraints from research review boards concerning unneeded research MRI without contrast agents in healthy children and adolescents seem unjustified and need to be carefully reconsidered to not impede the advancement of developmental neuroscience.

AUTHOR CONTRIBUTIONS

Substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data: R.E., R.M., V.S., K.L., and M.S. Drafting the article or revising it critically for important intellectual content: R.E., R.M., K.L., V.S., R.W., and M.S. Final approval of the version to be published: R.E., R.M., K.L., V.S., R.W., and M.S.

ADDITIONAL INFORMATION

The online version of this article (<https://doi.org/10.1038/s41390-020-01277-6>) contains supplementary material, which is available to authorized users.

Competing interests: The authors declare no competing interests.

Patient consent: All parents and participants signed written informed consent prior to enrollment if >14 years (for participants <14 years, the legal guardian signed the consent).

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Regula Everts^{1,2,3}, Raphaela Muri^{2,4}, Kurt Leibundgut³,
Valerie Siegwart^{1,3}, Roland Wiest⁴ and Maja Steinlin¹
¹Division of Neuropediatrics, Development and Rehabilitation,
Children's University Hospital, Inselspital, Bern University Hospital,
University of Bern, Bern, Switzerland; ²Department of Diabetes,
Endocrinology, Nutritional Medicine and Metabolism, Inselspital, Bern
University Hospital, University of Bern, Bern, Switzerland;
³Department of Pediatric Hematology and Oncology, Children's
University Hospital, Inselspital, Bern University Hospital, University of
Bern, Bern, Switzerland and ⁴Institute of Diagnostic and
Interventional Neuroradiology, Inselspital, Bern University Hospital,
University of Bern, Bern, Switzerland
Correspondence: Regula Everts (regula.everts@insel.ch)

REFERENCES

- Swissethics. Leitlinien zur Forschung mit gesunden Kindern und Jugendlichen. 1st edn (2017).
- Behrman, R. E. & Field, M. J. (Eds.). *Ethical Conduct of Clinical Research Involving Children* (National Academies Press, 2004).
- U.S. Department of Health and Human Services. *Protection of Human Subjects*. 45 CFR. (1995).
- Schmidt, M. H., Marshall, J., Downie, J. & Hadskis, M. R. Pediatric magnetic resonance research and the minimal-risk standard. *IRB Ethics Hum. Res.* **33**, 1 (2011).
- Jaite, C. et al. A comparison study of anxiety in children undergoing brain MRI vs adults undergoing brain MRI vs children undergoing an electroencephalogram. *PLoS ONE* **14**, e0211552 (2019).
- Shechner, T. et al. Empirical examination of the potential adverse psychological effects associated with pediatric fMRI scanning. *J. Child Adolesc. Psychopharmacol.* **23**, 357–362 (2013).
- Thomason, M. E. Children in non-clinical functional magnetic resonance imaging (fMRI) studies give the scan experience a "thumbs up". *Am. J. Bioeth.* **9**, 25–27 (2009).
- Marshall, S. P., Smith, M. S. & Weinberger, E. Perceived anxiety of pediatric patients to magnetic resonance. *Clin. Pediatr.* **34**, 59–60 (1995).
- Tyc, V. L., Fairclough, D., Fletcher, B., Leigh, L. & Mulhern, R. K. Children's distress during magnetic resonance imaging procedures. *Child Health Care* **24**, 5–19 (1995).
- Marshall, J., Martin, T., Downie, J. & Maliszka, K. A comprehensive analysis of MRI research risks: in support of full disclosure. *Can. J. Neurol. Sci.* **34**, 11–17 (2007).
- Greene, D. J., Black, K. J. & Schlaggar, B. L. Considerations for MRI study design and implementation in pediatric and clinical populations. *Dev. Cogn. Neurosci.* **18**, 101–112 (2016).
- Hallowell, L. M., Stewart, S. E., de Amorim E Silva, C. T. & Ditchfield, M. R. Reviewing the process of preparing children for MRI. *Pediatr. Radiol.* **38**, 271–279 (2008).
- de Bie, H. M. et al. Preparing children with a mock scanner training protocol results in high quality structural and functional MRI scans. *Eur. J. Pediatr.* **169**, 1079–1085 (2010).
- Dempsey, M. F., Condon, B. & Hadley, D. M. MRI safety review. *Semin. Ultrasound CT MRI* **23**, 392–401 (2002).
- Canadian Association of Radiologists. *CAR Standards for Magnetic Resonance Imaging*. April 2011. <https://car.ca/wp-content/uploads/Magnetic-Resonance-Imaging-2011.pdf> [retrieved 7 July 2020].
- Kanal, E. et al. ACR guidance document for safe MR practices. *AJR Am. J. Roentgenol.* **188**, 1447–1474 (2007).
- William Li, H. C. & Lopez, V. Do trait anxiety and age predict state anxiety of school-age children? *J. Clin. Nurs.* **14**, 1083–1089 (2005).
- Coplan, J. D. et al. The relationship between intelligence and anxiety: an association with subcortical white matter metabolism. *Front. Evol. Neurosci.* **3**, 8 (2012).
- Zhu, Y. et al. Socioeconomic status disparities affect children's anxiety and stress-sensitive cortisol awakening response through parental anxiety. *Psychoneuroendocrinology* **103**, 96–103 (2019).
- Powers, J. P., Moshontz, H. & Hoyle, R. H. Self-control and affect regulation styles predict anxiety longitudinally in university students. *Collabra Psychol.* **6**, (2020).
- Maliszka, K. L., Martin, T., Shiloff, D. & Yu, D. C. Reactions of young children to the MRI scanner environment. *Magn. Reson. Med.* **64**, 377–381 (2010).
- Brown, L., Sherbenou, R. J. & Johnsen, S. K. *Test of Nonverbal Intelligence: TONI-4* (Pro-ed., Austin, TX, 2010).
- Delis, D. C., Kaplan, E. & Kramer, J. H. *Delis-Kaplan Executive Function System* (The Psychological Corporation, San Antonio, TX, 2001).
- Everts, R., Schöne, C. G., Mürner-Lavanchy, I. & Steinlin, M. Development of executive functions from childhood to adolescence in very preterm-born individuals—a longitudinal study. *Early Hum. Dev.* **129**, 45–51 (2019).
- Benzing, V. et al. The Brainfit study: efficacy of cognitive training and exergaming in pediatric cancer survivors—a randomized controlled trial. *BMC Cancer* **18**, 1–10 (2018).
- Cahill, L. Why sex matters for neuroscience. *Nat. Rev. Neurosci.* **7**, 477–484 (2006).
- De Vries, G. J. Minireview: sex differences in adult and developing brains: compensation, compensation, compensation. *Endocrinology* **145**, 1063–1068 (2004).
- Kornfeld, S. et al. Resting-state connectivity and executive functions after pediatric arterial ischemic stroke. *NeuroImage: Clinical.* **17**, 359–367 (2018).
- Mürner-Lavanchy, I. M. et al. White matter microstructure is associated with language in children born very preterm. *NeuroImage Clin.* **20**, 808–822 (2018).
- Biberacher, V. et al. Intra- and interscanner variability of magnetic resonance imaging based volumetry in multiple sclerosis. *Neuroimage* **142**, 188–197 (2016).
- Wechsler, D. *Wechsler Intelligence Scale for Children-WISC-IV* (Pearson, London, 2003).
- Viggiano, M. P. et al. Impact of psychological interventions on reducing anxiety, fear and the need for sedation in children undergoing magnetic resonance imaging. *Pediatr. Rep.* **7**, (2015).



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2020