

Get Clarity On Generics

Cost-Effective CT & MRI Contrast Agents

 **FRESENIUS
KABI**

[WATCH VIDEO](#)

AJNR

This information is current as
of August 31, 2025.

Introduction of Ultra-High-Field MR Imaging in Infants: Preparations and Feasibility

K.V. Annink, N.E. van der Aa, J. Dudink, T. Alderliesten, F. Groenendaal, M. Lequin, F.E. Jansen, K.S. Rhebergen, P. Luijten, J. Hendrikse, H.J.M. Hoogduin, E.R. Huijting, E. Versteeg, F. Visser, A.J.E. Raaijmakers, E.C. Wiegers, D.W.J. Klomp, J.P. Wijnen and M.J.N.L. Benders

AJNR Am J Neuroradiol published online 30 July 2020
<http://www.ajnr.org/content/early/2020/07/30/ajnr.A6702>

Introduction of Ultra-High-Field MR Imaging in Infants: Preparations and Feasibility

K.V. Annink, N.E. van der Aa, J. Dudink, T. Alderliesten, F. Groenendaal, M. Lequin, F.E. Jansen, K.S. Rhebergen, P. Luijten, J. Hendrikse, H.J.M. Hoogduin, E.R. Huijting, E. Versteeg, F. Visser, A.J.E. Raaijmakers, E.C. Wiegers, D.W.J. Klomp, J.P. Wijnen, and M.J.N.L. Benders



ABSTRACT

BACKGROUND AND PURPOSE: Cerebral MR imaging in infants is usually performed with a field strength of up to 3T. In adults, a growing number of studies have shown added diagnostic value of 7T MR imaging. 7T MR imaging might be of additional value in infants with unexplained seizures, for example. The aim of this study was to investigate the feasibility of 7T MR imaging in infants. We provide information about the safety preparations and show the first MR images of infants at 7T.

MATERIALS AND METHODS: Specific absorption rate levels during 7T were simulated in Sim4life using infant and adult models. A newly developed acoustic hood was used to guarantee hearing protection. Acoustic noise damping of this hood was measured and compared with the 3T Nordell hood and no hood. In this prospective pilot study, clinically stable infants, between term-equivalent age and the corrected age of 3 months, underwent 7T MR imaging immediately after their standard 3T MR imaging. The 7T scan protocols were developed and optimized while scanning this cohort.

RESULTS: Global and peak specific absorption rate levels in the infant model in the centered position and 50-mm feet direction did not exceed the levels in the adult model. Hearing protection was guaranteed with the new hood. Twelve infants were scanned. No MR imaging-related adverse events occurred. It was feasible to obtain good-quality imaging at 7T for MRA, MRV, SWI, single-shot T2WI, and MR spectroscopy. T1WI had lower quality at 7T.

CONCLUSIONS: 7T MR imaging is feasible in infants, and good-quality scans could be obtained.

ABBREVIATIONS: dB(A) = A-weighted decibels; -50-mm FH = 50 mm from the isocenter in feet direction; +50-mm FH = 50 mm from the isocenter in head direction; SAR = specific absorption rate

Infants who are admitted to the neonatal intensive care unit are at risk of delayed or impaired neurodevelopmental outcome due to brain injury, cerebral malformations, and genetic or metabolic disorders.^{1,2}

MR imaging is the criterion standard to assess brain development, malformations, and injury in infants.³ The first neonatal 3T field strength MR imaging scans were reported in 2004,⁴ and 3T scanners are now routinely used by many centers. 3T MR imaging has several advantages compared with 1.5T MR imaging. The quality of the MR images improved because of the increased SNR, leading to higher spatial resolution, improved susceptibility contrast, and increased chemical shift dispersion leading to

improved quality of MR spectroscopy.^{5,6} The increased SNR in neonatal 3T MR imaging also led to shorter acquisition times.⁶

In adults, the introduction of ultra-high-field MR imaging provided new opportunities, further improving the spatial resolution at 7T compared with 3T when the same acquisition times were used.⁷ This feature provided additional anatomic information.⁸ At 7T, the sensitivity to susceptibility is strongly increased, enabling

The study was funded by the Dr. C.J. Vaillant Foundation. The PhD position of the first author Kim Annink is supported by the Horizon 2020 Research and Innovation Programme of the European Union (grant agreement No 667224, ALBINO).

Paper previously presented as a poster at: Congress of the Joint European Neonatal Societies/Eastern Society for Pediatric Research Conference, September 17–21, 2019; Maastricht, the Netherlands; and in part, at: International Society for Magnetic Resonance in Medicine Workshop on Ultrahigh Field Magnetic Resonance, March 31 to April 3, 2019, Dubrovnik, Croatia.

Please address correspondence to Manon J.N.L. Benders, MD, Department of Neonatology, UMC Utrecht Brain Center, University Utrecht, Lundlaan 6, 3508AB Utrecht, the Netherlands; e-mail: m.benders@umcutrecht.nl

Indicates open access to non-subscribers at www.ajnr.org

Indicates article with supplemental on-line table.

<http://dx.doi.org/10.3174/ajnr.A6702>

Received December 16, 2019; accepted after revision May 19, 2020.

From the Departments of Neonatology (K.V.A., N.E.v.d.A., J.D., T.A., F.G., M.J.N.L.B.), and Paediatric Neurology (F.E.J.), University Medical Center Utrecht Brain Center, and the Departments of Radiology (M.L., P.L., J.H., H.J.M.H., E.R.H., E.V., F.V., A.J.E.R., E.C.W., D.W.J.K., J.P.W.), and Otorhinolaryngology and Head and Neck Surgery (K.S.R.), University Medical Center Utrecht, University Utrecht, Utrecht, the Netherlands.

K.V. Annink and N.E. van der Aa shared first authorship.

J.P. Wijnen and M.J.N.L. Benders shared last authorship.

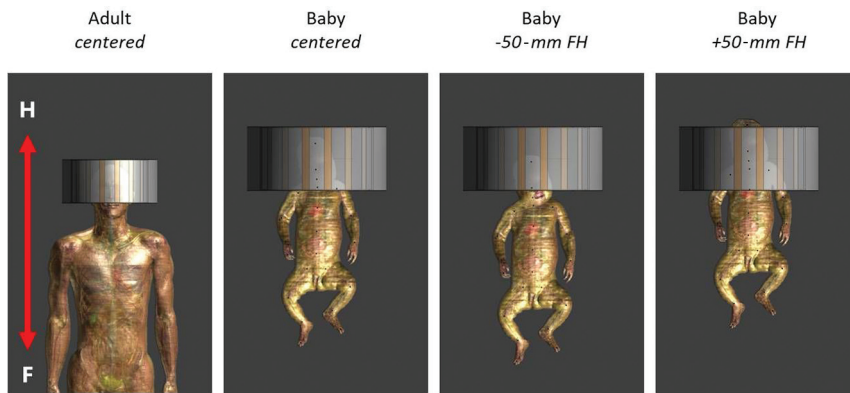


FIG 1. Different positions of the infant model in the radiofrequency coil. On the left, the adult head and torso are shown. They are used as a reference.

the diagnosis of microbleeds and visualization of microvasculature. Due to the increased chemical shift dispersion at 7T, additional metabolite peaks could be detected with MR spectroscopy.^{5,8-11} Currently, accessibility of 7T MR imaging scanners in adult research is increasing rapidly, and 7T MR imaging is more often used for clinical purposes. The initial safety concerns of using 7T MR imaging in adults have been addressed in the past decades. The largest safety concern was an increase in body temperature because of the higher local and global specific absorption rate (SAR) for the same B_1^+ at 7T. However, an increase in body temperature has not been reported in ultra-high-field MR imaging in adults or children.^{5,12} Besides the increased SAR, a higher static magnetic field increases the risk of attracting ferromagnetic objects, which can be prevented with screening for ferrometal before MR imaging.¹³ It also can potentially influence biologic systems, such as cardiac and neurophysiological responses, but harmful effects have never been described in follow-up studies in infants.¹³ Sensory symptoms such as vertigo, headaches, and an iron taste due to the varying gradient field⁵ were reported by patients undergoing 7T MR imaging; however, only 5% rated these symptoms as very unpleasant.¹⁴ Acoustic noise protection should be guaranteed during the MR imaging, similar to 3T MR imaging.

While studies at 7T have now been shown to be safe in adults,^{14,15} the literature about safety in children is scarce. In a study by Harris et al,¹⁶ 42 children between 5 and 10 years of age underwent 7T MR imaging, which was well-tolerated and safe in all children. The Food and Drug Administration approved 7T MR imaging in infants of 1 month and older.¹⁷ The limit of the main static field in neonates is 4T MR imaging. SAR limits are the same for adults and neonates.¹⁷

We initiated a feasibility pilot study in infants. This study shows the first MR images of infants at 7T and provides information about the safety preparations.

MATERIALS AND METHODS

Preparation: SAR Simulation

Using a radiofrequency simulation, we investigated whether MR imaging protocols could be translated from the adult to the infant

brain without exceeding the SAR limits. Finite-difference time-domain simulations were performed using Sim4Life (Zurich Med Tech) to evaluate transmit efficiency and radiofrequency safety limits of the setup, assuming full decoupling of the receiver coils. The geometry and electrical circuits of the NOVA head coil (Nova Medical) for 7T were implemented in Sim4Life. The simulations were performed on a virtual infant model (Charlie, 2 months of age, 4.3 kg) of the Integrated Taxonomic Information System virtual family¹⁸ in different positions in the coil (Fig 1), using the head coil in quadrature mode.

The same simulations were performed on adult heads (male Duke, female Ella) of the virtual family¹⁸ as a reference. Local SAR levels (10-g average) for 1-W input power and global SAR levels (average SAR over the whole head) were calculated. Also, the average SAR per B_1^2 was calculated as the average SAR over the whole head divided by the average B_1^2 in a central section in the brain. The peak local SAR was defined as the highest SAR in the whole infant.

SAR simulations were validated by comparing simulations and measurements of B_1 maps of a spherical phantom and power measurements for data scaling (data not shown).

Preparation: Acoustic Noise Protection

At 3T MR imaging, acoustic noise protection is guaranteed by Alpine Muffy Baby (Alpine Hearing Protection), Natus MiniMuffs (Natus Medical), and a hood for acoustic noise protection,¹⁹ respectively, leading to 6.4- to 31.6-dB, 7-dB, and 4- to 13.6-dB reduction. A prototype of the hood (190 cm long) for noise protection that fits in the 7T MR scanner was developed using a layer of 5-cm foam (EASYfoam TC2; EASY Noise Control).

A test setup with a dummy MR imaging bore (old 7T MR imaging bore) was made in a sound-isolated booth to test the attenuation of acoustic noise with different hoods and no hood. We conducted all sound-level measurements using a sound-level meter (B&K type 2250; Brüel & Kjær). A microphone (B&K, type 4189) was placed in the isocenter of the dummy bore to record the sound volumes in A-weighted decibels 9dB(A). Four speakers (Yamaha MSP5A; Hamamatsu) were positioned around this test setup: one on each end of the dummy bore and one on each side (Fig 2). The speakers were separately calibrated on an acoustic noise level of 55 dB(A). The acoustic noise was measured with the 3T hood, the 7T hood, and without a hood, with the acoustic noise coming from the speakers at both ends, both sides, and from all 4 speakers.

We measured the attenuation of the Alpine Muffy Baby and Natus MiniMuffs using the B&K type 2250 G4 SLM and B&K Artificial Ear type 4153. A stimulus (AudioNigma; Decos) of 80 dB(A) was sent to an artificial ear.

Study Population

Clinically stable infants, between term-equivalent age and the corrected age of 3 months were included in this pilot study.

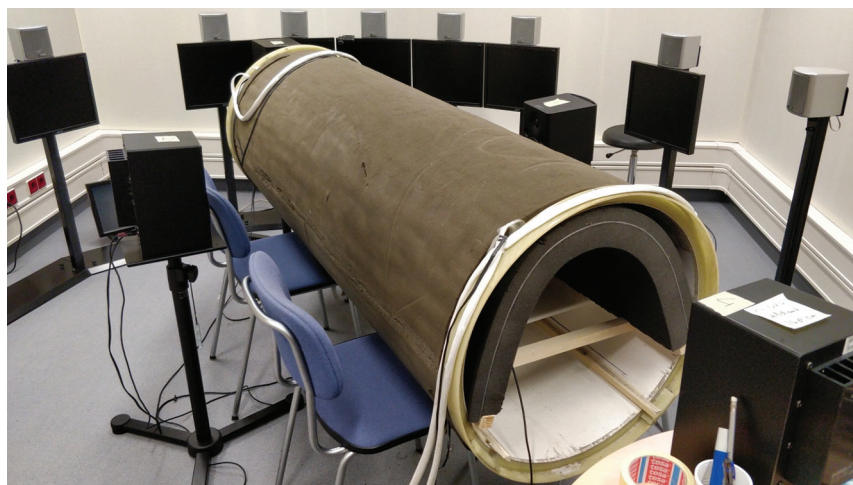


FIG 2. Setup for measuring acoustic noise at the MR imaging table in the presence of the hood. This test setup consisted of a dummy bore with dimensions similar to those of the 7T MR system with a 10-mm plastic plate of polyoxymethylene to create an MR imaging table. The speakers producing the sound are positioned at 28-cm distance around the dummy bore to mimic the sound produced by the MR imaging scanner.

Table 1: Global and peak SAR levels

	Duke Centered	Ella Centered	Charlie Centered	Charlie −50- mm FH	Charlie +50- mm FH
Global SAR levels					
Average SAR for 1 Watt input power (W kg^{-1})	0.066	0.069	0.062	0.050	0.075
Average SAR per B_1^2 ($\text{W kg}^{-1} \mu\text{T}^{-2}$)	0.462	0.465	0.289	0.454	0.474
Average B_1^+ in central section for 1 Watt input power (μT) ^a	0.379	0.385	0.466	0.333	0.398
Peak local SAR levels					
Peak local SAR (10-g averaged) for 1 Watt input power (W kg^{-1})	0.435	0.398	0.321	0.213	0.487
Peak local SAR (10-g averaged) per B_1^2 ($\text{W kg}^{-1} \mu\text{T}^{-2}$)	3.04	2.63	1.48	1.92	3.08

^a The power optimization procedure of the MR imaging scanner software calibrates the needed input power to achieve a certain B_1 in the subject. This calibration is based on the average B_1^+ in a central section of the subject (brain in this case).

Infants with respiratory support or an intravenous catheter were excluded. They underwent 7T MR imaging immediately after their routine 3T MR imaging scan (both Philips Healthcare Best, The Netherlands). All infants were sedated with chloral hydrate before the 3T MR imaging in combination with the feed-bundle technique, as parts of routine clinical care. An additional dose of chloral hydrate before the 7T MR imaging was not allowed by the medical ethics committee. For neonatal scans of the brain at 7T, the 2-channel transmit 32-channel receive head coil (Nova Medical, Wilmington, MA) was used. The selection of sequences was based on the clinical indication. Details of the scan protocols can be found in the On-line Table. Hearing was protected as described above. Safety parameters were monitored before, during, between, and after the MR images. We monitored heart rate, peripheral oxygen saturation, temperature (core temperature before

and after and abdominal skin temperature during the scans), and comfort scales.²⁰

This study was approved by the medical ethics committee of the University Medical Center Utrecht (NL66198.041.18), and written informed consent was obtained from the parents of all participants.

RESULTS

Preparation: SAR Simulation

The global SAR and peak local SAR of the virtual infant model in the centered position and 50 mm from the isocenter in feet direction (−50-mm FH) did not exceed the SAR of the adult models. However, when infant Charlie was positioned 50 mm from isocenter in the head direction (+50-mm FH), global SAR levels and peak local SAR levels exceeded those of the adult models (by +13% and +12% compared with Duke, respectively) (Table 1).

The 10-g averaged local SAR in the head of the infant model was lower than that in the adult head for all positions. When the infant model was positioned +50-mm FH, the local SAR was highest in the neck/shoulder transitions (Fig 3).

The SAR per B_1^2 was lower in the infant model than in the adult models, meaning that less power is needed to reach the same B_1 , except for the +50-mm FH position.

Preparation: Acoustic Noise Protection

The background noise in the sound booth was 28 dB(A). The 7T hood attenuated the acoustic noise by 8.5 dB, and the 3T hood, by 7 dB (Table 2).

Without hearing protection, the artificial ear measured 80 dB(A). The Alpine Muffy Baby reduced the acoustic noise level to 56 dB(A), and the use of only the Natus Minimuffs resulted in 73.4 dB(A). The combination of both, not totally closing the artificial ear, led to an acoustic noise level of 58 dB(A). If they were both well-placed on the ear using the elastic head band, the level decreased to 47.8 dB(A).

Feasibility of MR Imaging

Twelve infants have been included with a median gestational age of 28.2 weeks (range, 25.0–41.7 weeks), median birth weight of 1127 g (range, 585–4570 g), median postnatal age at MR imaging of 95 days (range, 31–114 days), and a median weight at MR

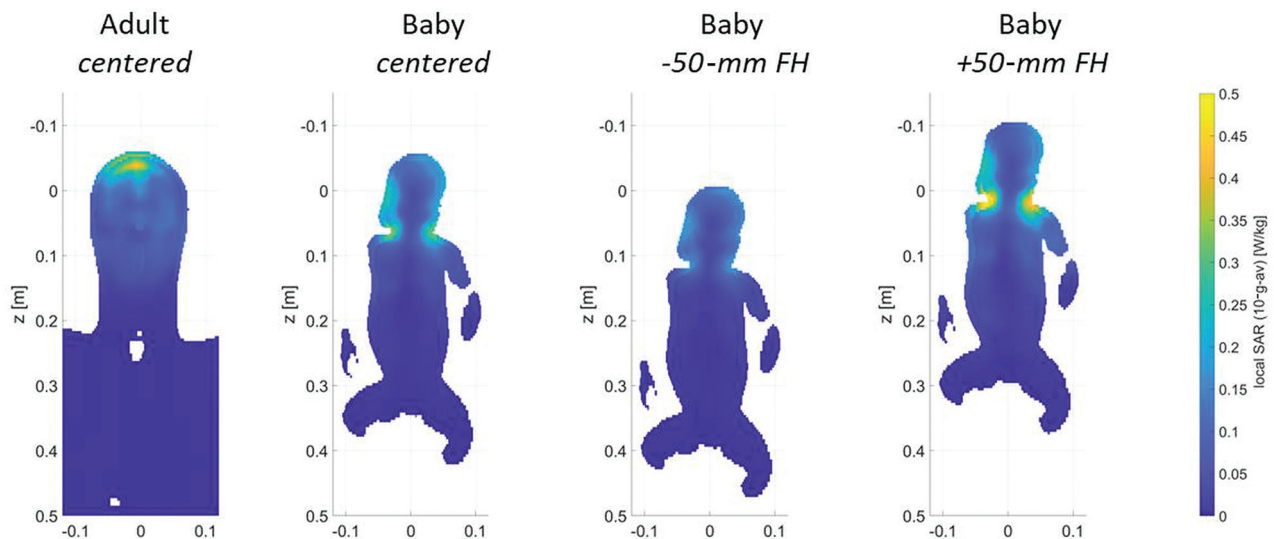


FIG 3. Local SAR levels in adult head (left) and Charlie in the different coil positions. Shifts of an infant in the x and y directions are unlikely because of limited space; therefore, the results are not included in the figure. The SAR values when infant Charlie is positioned 50 mm in the x or y direction are comparable with those in the +50-mm FH position.

Table 2: Acoustic noise levels in decibel(A), with use of no hood, the 3T hood, and the newly developed 7T hood

	Both Ends of the Bore	Sides of the Bore	Sides and Ends of the Bore
No hood for acoustic noise protection	60.0 dB(A)	58.5 dB(A)	62.5 dB(A)
3T Nordell hood for acoustic noise protection	54.0 dB(A)	51.0 dB(A)	55.5 dB(A)
Prototype 7T hood for acoustic noise protection	54.0 dB(A)	43.0 dB(A)	54.0 dB(A)

imaging of 3322 g (range, 2715–6335 g). The clinical indications for the MR imaging scans were the following: MR imaging at term-equivalent age because of preterm birth before 28 weeks' gestation ($n = 7$); MR imaging at term-equivalent age because of white matter injury ($n = 2$); follow-up MR imaging at 3 months' corrected age because of a thalamic hemorrhage ($n = 1$) or hemorrhage in the temporal lobe ($n = 1$); and follow-up MR imaging at 6 weeks because of an arterial ischemic stroke ($n = 1$). Temperature, heart rate, peripheral oxygen saturation, and comfort scales were stable before, during, and after MR imaging. No serious adverse events related to the MR imaging occurred.

MRV at 7T provided good visibility of the different veins and sinuses: The superficial cerebral veins could be followed in detail (Fig 4A, -G).

Also, SWI at 7T was feasible and showed details of the deep venous circulation, i.e., the deep medullary veins (Fig 4B, -H).

Single-shot T2WI at 7T demonstrated good gray and white matter contrast. In 1 patient with a perinatal arterial stroke, perivascular spaces were seen at 7T that were not visible at 3T (Figs 4C, -I). Otherwise, no clinically relevant additional findings were reported by the pediatric neuro radiologist (M.L.) at 7T. The gray and white matter differentiation at T1WI was suboptimal at 7T (Figs 4E, -K).

MRA showed more peripheral arteries at 7T, and little noise was visible at 7T, making it easier to see the thickness and curves of the arteries (Figs 4D, -J).

MR spectroscopy was of improved quality. For example, the patient shown in Fig 4F, -L had an SNR of 19 at 7T compared with 6 at 3T. It was possible to correctly fit more metabolites with a Cramer–Rao Lower Bound of <20% at 7T, such as *N*-acetyl aspartylglutamate, taurine, and glycine.

DISCUSSION

We demonstrated that scanning infants in a 7T scanner is feasible and results in good-quality images. While optimization of the sequences is ongoing, we already demonstrated that some sequences showed more details compared with 3T MR imaging.

Before scanning, simulated SAR levels at 7T were lower in the virtual infant than in adult models.¹⁸ When the infant's head was further in the coil than isocenter, or 5 cm in the x or y direction, SAR levels did exceed the adult situation. Thus, the center position of the infant in the coil is essential. Therefore, the position of the infant's head was constrained in the coil, making it mechanically impossible to put the infant's head farther in the coil than center position. Differences in the SAR due to intersubject variability cannot be completely ruled out. However, previous simulations at 3T showed that different-sized infant models and different positions did not result in major differences in simulated SAR levels.²¹

The SAR simulations had 2 important limitations. The first limitation is that the Sim4Life model of Charlie uses the dielectric values of adults, which might slightly differ from infant dielectric properties. The dielectric values of human infants are unknown

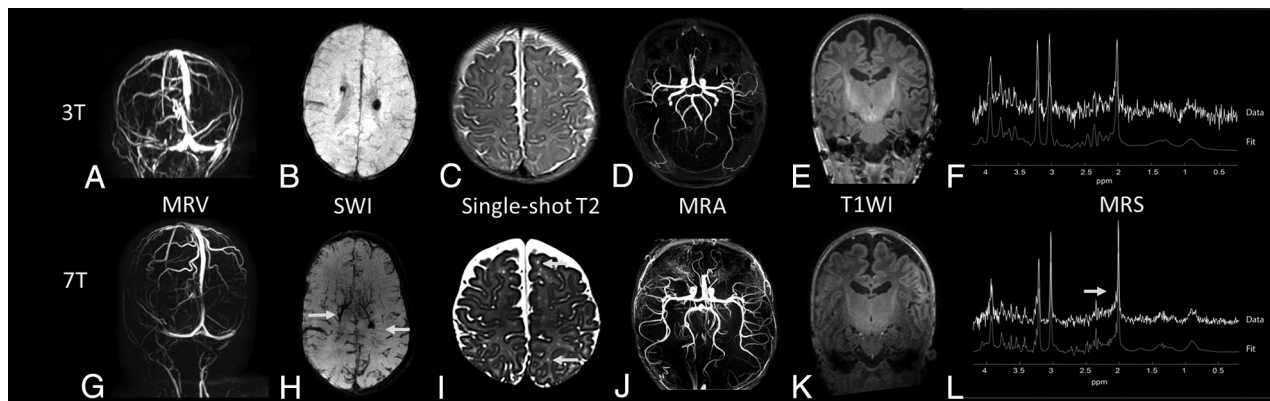


FIG 4. Examples of images of different patients at 3T (upper row) versus 7T MR imaging (lower row). A and G, MRV at 3T and 7T, respectively, of a preterm infant at term-equivalent age. B and H, An SWI at, respectively, 3T and 7T of a preterm infant at term-equivalent age. C and I, The 6-week follow-up single-shot T2-weighted image of a term-born infant with a perinatal stroke. The arrows indicate perivascular spaces that were better visualized at 3T compared with 7T. D and J, A 3-month follow-up MRA at 3T and 7T of a term-born infant with an occipital stroke. E and K, T1WI (MPRAGE) of a preterm-born infant at term-equivalent age. F and L, An MR spectroscopy spectrum at 3T and 7T of a preterm infant at term-equivalent age, both with a comparable ROI in the left basal ganglia/thalami region.

and require further research. In the study of Malik et al,²¹ conversion of adult dielectric values was based on rat data; however, these are not validated. Second, only 1 virtual model of a 2-month-old infant was available, which cannot be completely translated to an infant of term-equivalent age. Head circumference and body composition differed between term-equivalent age and 2 months of age. Malik et al showed that term neonates with smaller head sizes or lower body weight had lower SAR depositions, suggesting that the 2-month-old infant model does not underestimate SAR values. Regarding body composition, Malik et al simulated the effect of fat percentage on SAR depositions in neonates in 2 extreme scenarios: 1 model with only skin and 1 model with a thick layer of pure fat. The model with only skin had 10% higher peak local SAR depositions.²¹ The fat percentage of neonates is lower compared with 2-month-old infants.^{22,23} In the worst case scenario, a neonate might have a higher peak local SAR up to 10% compared with a 2-month-old infant based on fat composition. This will still not exceed the safety limits of the FDA because the 7T MR scanner, Best, the Netherlands has implemented an additional safety factor larger than 2. Furthermore, the global SAR levels of Charlie in the centered position were 6% lower compared with the adult model, and the peak local SAR was 26% lower, leading to an additional safety margin. Adult limits are, therefore, still safe to use.

Another concern might be that the thermoregulation in neonates is immature compared with adults, so the effect of SAR values on body temperature might differ between neonates and adults.²¹ Neonates have less isolating subcutaneous fat and a larger surface-to-body weight ratio, making them more prone to develop hypothermia.^{21,24-26} The risk of high local peak SAR values in neonates is lower compared with adults because less power is needed in neonates to reach the same B_1 . Furthermore, the risk of high local peak SAR is reduced by the above-described safety margins (Table 1).

No MR imaging-related adverse events occurred in the infants scanned at 7T in this pilot study, and comfort scales were stable, which are both indicators that infants did tolerate the higher main static field.

The possible improvements in the quality of SWI and single-shot T2WI are caused by a shorter T2-relaxation time, improved spatial resolution, and increased susceptibility.⁵ These might enable physicians to assess the extent of injury on a microstructural level: diagnosing microbleeds and polymicrogyria and thereby improving the prediction of neurodevelopmental outcome.²⁷⁻²⁹

As expected, the quality of T1WI at 7T was worse compared with 3T in infants. The T1-relaxation time increases at higher field strengths.¹³ Furthermore, the brains of neonates have a relatively high water content, which results in less contrast between white and gray matter. To compensate for this longer T1 relaxation time, the TR can be increased, but this increase leads to longer scanning time, which is also not preferable in neonates.^{6,13} On the other hand, this increased T1-relaxation time enables higher quality angiography, which, in the future, can help, for example, to evaluate small perforator strokes.^{6,10}

For MR spectroscopy, the increased chemical shift dispersion at 7T results in less overlap between the different metabolite peaks; also, the SNR is increased (>2-fold). Of note is that the maximal required B_1 for MR spectroscopy cannot be achieved when the infant is in the -50-mm position in the coil. This can happen if the shoulders do not fit in the head coil when the infant is wrapped in the vacuum mattress. In such cases, MR spectroscopy at 7T has SNR comparable with that of 3T, but with the advantage of less overlap between metabolite peaks. Nevertheless, 7T MR spectroscopy enabled more accurate detection of *N*-acetyl aspartylglutamate, taurine, and glycine and possibly other metabolites such as glutamate, gamma-aminobutyric acid, and myo-inositol, as has been described in adults.^{5,6} This feature could be helpful for the diagnosis of metabolic diseases and neuronal injury but can also provide information about the biochemical development of the neonatal brain.

In the future, 7T might be particularly helpful to answer specific questions about the diagnosis or outcome in, for example, infants with small (perforator) strokes, metabolic diseases, or unexplained neurologic symptoms, i.e., seizures. The clinical

implications and the additional value of 7T in infants should be investigated in larger 7T MR imaging studies. We did not scan preterm neonates; the safety and feasibility of 7T MR imaging in these neonates should also be investigated in the future.

CONCLUSIONS

This pilot study shows, for the first time, that 7T ultra-high-field MR imaging is feasible in infants. Good-quality images could be obtained, with some sequences providing additional details compared with 3T. Positioning of the infant in the isocenter of the coil is important for SAR safety.

ACKNOWLEDGMENTS

We thank Patrick Stroosnijder for helping with the audiology tests and the physician assistants of the neonatal intensive care unit for their help and supervision during the MR imaging scans.

Disclosures: Kim V. Annink—RELATED: Grant: Horizon 2020 grant, grant Dr. C.J. Vaillant Foundation, Comments: The study was funded by the Dr. C.J. Vaillant Foundation. The PhD position of the first author Kim Annink is supported by Horizon 2020 Research and Innovation Programme of the European Union (grant agreement No 667224, ALBINO).^{*} Niek E. van der Aa—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Floris Groenendaal—UNRELATED: Expert Testimony: Court, Comments: expert witness in medicolegal cases of perinatal asphyxia; Grants/Grants Pending: Neurophyxia, Comments: unrestricted grant from Neurophyxia for research on neuroprotection^{*}; Royalties: UpToDate, Comments: royalties for publication.^{*} Erik R. Huijting—UNRELATED: Employment: University Medical Center Utrecht. Fredy Visser—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Alexander J. E. Raaijmakers—UNRELATED: Employment: University Medical Center Utrecht. RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Evita C. Wieggers—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Dennis W.J. Klomp—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Jannie P. Wijnen—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} Manon J.N.L. Benders—RELATED: Grant: Dr. C.J. Vaillant Foundation.^{*} ^{*}Money paid to the institution.

REFERENCES

1. Semple B, Blomgren K, Gimlin K, et al. **Brain development in rodents and humans: identifying benchmarks of maturation and vulnerability to injury across species.** *Prog Neurobiol* 2013;106–107:1–16 [CrossRef Medline](#)
2. Arthur R. **Magnetic resonance imaging in preterm infants.** *Pediatr Radiol* 2006;36:593–607 [CrossRef Medline](#)
3. de Vries L, Benders M, Groenendaal F. **Imaging the premature brain: ultrasound or MRI?** *Neuroradiology* 2013;55:13–22 [CrossRef Medline](#)
4. Rutherford M, Malamateniou C, Zeka J, et al. **MR imaging of the neonatal brain at 3 Tesla.** *Eur J Paediatr Neurol* 2004;8:281–89 [CrossRef Medline](#)
5. Kolk AG, Van Der Hendrikse J, Zwanenburg JJM, et al. **Clinical applications of 7 T MRI in the brain.** *Eur J Radiol* 2013;82:708–18 [CrossRef Medline](#)
6. Ditchfield M. **3 T MRI in paediatrics: challenges and clinical applications.** *Eur J Radiol* 2008;68:309–19 [CrossRef Medline](#)
7. Robitaille PL, Abduljalil AM, Kangarlou A, et al. **Human magnetic resonance imaging at 8T.** *NMR Biomed* 1998;11:263–65 [CrossRef Medline](#)
8. Springer E, Dymerska B, Cardoso PL, et al. **Comparison of routine brain imaging at 3 T and 7 T.** *Invest Radiol* 2017;51:469–82 [CrossRef Medline](#)

9. De Cockler LJ. **Cerebellovascular disease: magnetic resonance imaging.** *J Belgian Soc Radiol* 2016;100:1–6 [CrossRef Medline](#)
10. Park C, Kang C, Kim Y, et al. **Advances in MR angiography with 7T MRI: from microvascular imaging to functional angiography.** *Neuroimage* 2018;168:269–78 [CrossRef Medline](#)
11. Stefanits H, Springer E, Patariaia E, et al. **Seven-Tesla MRI of hippocampal sclerosis an in vivo feasibility study with histological correlations.** *Invest Radiol* 2017;52:666–71 [CrossRef Medline](#)
12. Shellock FG, Cruess JV. **Radiology MR procedures: biologic effects, safety, and patient care.** *Radiology* 2004;232:635–52 [CrossRef Medline](#)
13. Tocchio S, Kline-Fath B, Kanal E, et al. **MRI evaluation and safety in the developing brain.** *Semin Perinatol* 2015;39:73–104 [CrossRef Medline](#)
14. Theysohn JM, Maderwald S, Kraff O, et al. **Subjective acceptance of 7 Tesla MRI for human imaging.** *MAGMA* 2008;21:63–72 [CrossRef Medline](#)
15. Kraff O, Quick HH. **7T: physics, safety, and potential clinical applications.** *J Magn Reson Imaging* 2017;46:1573–89 [CrossRef Medline](#)
16. Harris AD, Singer HS, Horska A, et al. **GABA and glutamate in children with primary complex motor stereotypies: a 1H MRS study at 7T.** *AJNR Am J Neuroradiol* 2016;37:552–57 [CrossRef Medline](#)
17. U.S. Department of Health and Human Services; Food and Drug Administration; Center for Devices and Radiological Health. **Criteria for Significant Risk Investigations of Magnetic Resonance Diagnostic Devices 2014.** <http://www.fda.gov/downloads/RegulatoryInformation/Guidances/ucm126418.pdf>. Accessed September 6, 2019
18. Christ A, Kainz W, Hahn EG, et al. **The Virtual Family: development of surface-based anatomical models of two adults and two children for dosimetric simulations.** *Phys Med Biol* 2010;55:N23–38 [CrossRef Medline](#)
19. Nordell A, Lundh M, Horsch S, et al. **The acoustic hood: a patient-independent device improving acoustic noise protection during neonatal magnetic resonance imaging.** *Acta Paediatr* 2009;98:1278–83 [CrossRef Medline](#)
20. Ambuel B, Hamlett KW, Marx CM, et al. **Assessing distress in pediatric intensive care environments: the COMFORT scale.** *J Pediatr Psychol* 1992;17:95–109 [CrossRef Medline](#)
21. Malik SJ, Beqiri A, Price AN, et al. **Specific absorption rate in neonates undergoing magnetic resonance procedures at 1.5T and 3T.** *NMR Biomed* 2015;28:344–52 [CrossRef Medline](#)
22. Demerath EW, Fields DA. **Body composition assessment in the infant.** *Am J Hum Biol* 2014;26:291–304 [CrossRef Medline](#)
23. Fields DA, Gilchrist JM, Catalano PM, et al. **Longitudinal body composition data in exclusively breast-fed infants: a multicenter study.** *Obesity (Silver Spring)*. 2011;19:1887–91 [CrossRef Medline](#)
24. Sharma A, Ford S, Calvert J. **Adaptation for life: a review of neonatal physiology.** *Anaesthesia & Intensive Care Med* 2011;12:85–90 [CrossRef Medline](#)
25. Plaisier A, Raets MMA, van der Starre C, et al. **Safety of routine early MRI in preterm infants.** *Pediatr Radiol* 2012;42:1205–11 [CrossRef Medline](#)
26. Don Paul JM, Perkins EJ, Pereira-Fantini PM, et al. **Surgery and magnetic resonance imaging increase the risk of hypothermia in infants.** *J Paediatr Child Health* 2018;54:426–31 [CrossRef Medline](#)
27. Brundel M, Heringa SM, de Bresser J, et al. **High prevalence of cerebral microbleeds at 7 Tesla MRI in patients with early Alzheimer's disease.** *J Alzheimers Dis* 2012;31:259–63 [CrossRef Medline](#)
28. De Ciantis A, Barkovich AJ, Cosottini M, et al. **Ultra-high-field MR imaging in polymicrogyria and epilepsy.** *AJNR Am J Neuroradiol* 2015;36:309–16 [CrossRef Medline](#)
29. Cosottini M, Donatelli G, Costagli M, et al. **High-resolution 7T MR imaging of the motor cortex in amyotrophic lateral sclerosis.** *AJNR Am J Neuroradiol* 2016;37:455–61 [CrossRef Medline](#)