Research letter

Sedation for screening MRI in patients with congenital melanocytic naevi under the age of one is a successful, safe and economical firstline approach

DOI: 10.1111/bjd.17263

DEAR EDITOR, A single screening magnetic resonance imaging (MRI) scan of the central nervous system with contrast in patients under age 1 year (ideally < 6 months) is currently recommended as the best predictor of adverse outcome measures in children with multiple congenital melanocytic naevi (CMN).^{1,2} This recommendation is based on using sedation rather than general anaesthesia (GA), however, this practice is

not routine in many departments. Recently, concerns regarding neurodevelopmental effects of GA in animals³ and children^{4,5} have suggested that avoidance of GA is desirable where possible in infants.

We therefore undertook a retrospective analysis of the success of MRI using different modalities of sedation or anaesthesia. Records were reviewed of 247 patients with CMN who had had an MRI attempted, 161 under age 1 year. In total 114 of 247 (46%) were male, 208 of 244 (85%) had brain and whole-spine imaging and 193 of 202 (96%) had contrast injection. The mean and median ages were 2.02 and 0.66 years, respectively (range 0-18.8 years).

Information regarding sedation and anaesthetic was available in 208 of 247. Forty-five of 208 (22%) were preselected

fore sedation Patient assessment	Suitability for sedation	Baseline	Fasting
 Past medical history Medications, allergies and vaccination history Metal implants 	Consider awake scan	 Observations Weight 	 4 h food/milk 2 h clear fluids/breast
ıring sedation	sedationist on the day of the s	e most appropriate protocol sho can. This decision is currently bas ration of scan and sedationist pre	ed on individual patient
< 5 kg	5–12 kg	12–20 kg	> 20 kg
+	+	+	+
'Feed and wrap'	Chloral hydrate 100 mg kg ⁻¹ (max 1 g)	Chloral hydrate 100 mg kg ⁻¹ (max 2 g) + alimemazine 1–2 mg kg ⁻¹ (max 60 mg)	Dexmedetomidine 3 µg kg ⁻¹ loading dose over 10 min
	+	+	+
		effective after 10 min	Then 2 µg kg ⁻¹ h ⁻¹ continuous IV infusion for 30–120 min as required
		+	+
	< 20 kg slowly	100 μg kg ⁻¹ for / in increments kg ⁻¹ or 10 mg)	If sedation ineffective at any point before scan completed
	(agitation,	ical effects uncontrolled cination etc.)	Repeat loading dose dexmedetomidine
		versal 10–20 μg s, in two doses	

Fig 1. Current sedation protocol for paediatric magnetic resonance imaging used at Great Ormond Street Hospital for Children NHS Foundation Trust, London, and for which a 90% success rate was obtained in this cohort. Sedation is also cheaper than general anaesthesia (GA). IV, intravenous. for GA by an experienced sedationist, for example due to comorbidities, although a comparison of the demographic and phenotypic profiles of those who received sedation or GA showed no significant differences. Ten of 208 (5%) were awake, 151 of 208 (73%) were sedated and two of 208 (1 \cdot 0%) underwent the 'feed and wrap' technique, all according to local protocols (Fig. 1). In total 219 of 234 scans (94%) were successfully completed, with the 15 abandoned scans all in the sedation group. This equates to a sedation success rate of 136 of 151 (90%). Regression analysis demonstrated no significant difference in sedation success by age.

The literature supports sedation for paediatric MRI as effective and safe,⁶ with comparable success rates to our results,⁷ although there is no consensus on the methods of sedation.⁸ Common minor adverse events with sedation include vomiting and excessive secretions and apnoea; serious adverse events are very rare,⁶ with none reported here. Sedation is also cheaper than GA, at £462 vs. £770 per patient.⁷

These data strongly support the use of sedation rather than GA as a first-line approach for MRI in infants with multiple CMN, after triage by a sedationist or anaesthetist and with access to anaesthetic support if required.

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References

- 1 Waelchli R, Aylett SE, Atherton D et al. Classification of neurological abnormalities in children with congenital melanocytic naevus syndrome identifies magnetic resonance imaging as the best predictor of clinical outcome. Br J Dermatol 2015; 173:739–50.
- 2 Kinsler VA, O'Hare P, Bulstrode N et al. Melanoma in congenital melanocytic naevi. Br J Dermatol 2017; 176:1131-43.
- 3 Rappaport B, Mellon RD, Simone A, Woodcock J. Defining safe use of anesthesia in children. N Engl J Med 2011; **364**:1387–90.
- 4 Warner D, Zaccariello M, Katusic S et al. Neuropsychological and behavioral outcomes after exposure of young children to procedures requiring general anesthesia: the Mayo Anesthesia Safety in Kids (MASK) study. *Anesthesiology* 2018; **129**:89–105.
- 5 Andropoulos DB, Greene MF. Anesthesia and developing brains implications of the FDA warning. N Engl J Med 2017; **376**: 905–7.
- 6 Cravero JP, Blike GT, Beach M et al. Incidence and nature of adverse events during pediatric sedation/anesthesia for procedures outside the operating room: report from the Pediatric Sedation Research Consortium. Pediatrics 2006; 118:1087–96.
- 7 Bailey MA, Saraswatula A, Dale G, Softley L. Paediatric sedation for imaging is safe and effective in a district general hospital. Br J Radiol 2016; 89:20150483.
- 8 The Royal College of Radiologists. Sedation, analgesia and anaesthesia in the radiology department. Available at: https://www.rcr.ac.uk/sys tem/files/publication/field_publication_files/bfcr182_safe_sedation. pdf (last accessed 16 October 2018).

Funding sources: V.K. is funded by the Wellcome Trust (grant WT104076MA), and the work was supported by the U.K. National Institute for Health Research through Biomedical Research Centres at Great Ormond Street Hospital for Children NHS Foundation Trust and University College London.

Conflicts of interest: none to declare.