22-26 September 2021

INTRODUCTION

Increased use of MRI and improved technique provides the opportunity to detect small pituitary lesions. In a few studies performed only in children there were no cases with further growth of microadenoma during repeated MRI (1-4). There is no consensus on recommendations of the optimal clinical and radiological follow-up of a microadenoma in children.

AIM

We aimed to investigate the growth potential of pituitary microadenoma and cystic lesion <10mm in children, and to evaluate how reproducible the measurements were on magnetic resonance imaging (MRI).

PITUITARY MICROADENOMA IN CHILDHOOD - IS FOLLOW-UP WITH DIAGNOSTIC IMAGING NECESSARY?

Camilla Borghammar¹ MD, Ashkan Tamaddon² MD, Eva-Marie Erfurth³ MD PhD, Pia C Sundgren^{2, 4, 5} MD PhD, Peter Siesjö⁶ MD PhD, Margareta Nilsson^{2*} MD PhD

¹Lund University, Skåne University Hospital, Institution of Clinical Sciences, Department of Pediatrics, Pediatric Endocrinology, Lund, Sweden, ²Lund University, Skåne University, Skåne University, Skåne University, Institution of Clinical Sciences, Department of Endocrinology, Lund, Sweden, ⁴Lund University, Institution for Clinical Sciences, Department of Radiology, Lund, Sweden, ⁵Lund University, Lund Bioimaging Center, Lund, Sweden, ⁶Lund University, Skåne University Hospital, Institution of Clinical Sciences, Department of Neurosurgery, Lund, Sweden *Joint last author



FACULTY OF MEDICINE

RESULTS

In all, 74 children had a non-functioning microadenoma, probable microadenoma (a solid lesion less distinct) or cystic lesion, median age 12 years (range 3 - 17). Of those 55 underwent repeated MRI, median number 3 (range 2-7), with a median follow-up time of 37 months (range 4 – 189). An additional 12 children with prolactinoma, median age 16 years (range 12 – 17), underwent repeated MRI, median number 5.5 (range 2 – 11), median follow-up time 73.5 months (range 26 – 120). In one child the cystic lesion grew and hormonal deficits developed, in another child a prolactinoma increased in size due to problems with compliance to medication. None of the nonfunctioning microadenoma, probable microadenoma or cyst increased significantly in size during follow-up. Two reviewers agreed that no lesion could be identified in 38/269 MRI exams, in 51/231 (22%) there was disagreement about lesion location. They also disagreed in 34/460 (7%) MRI measurements to an extent that would have been considered progression in lesion size (>2mm). The inter-observer agreement was less accurate with the high field strength (3T), especially for probable microadenoma.

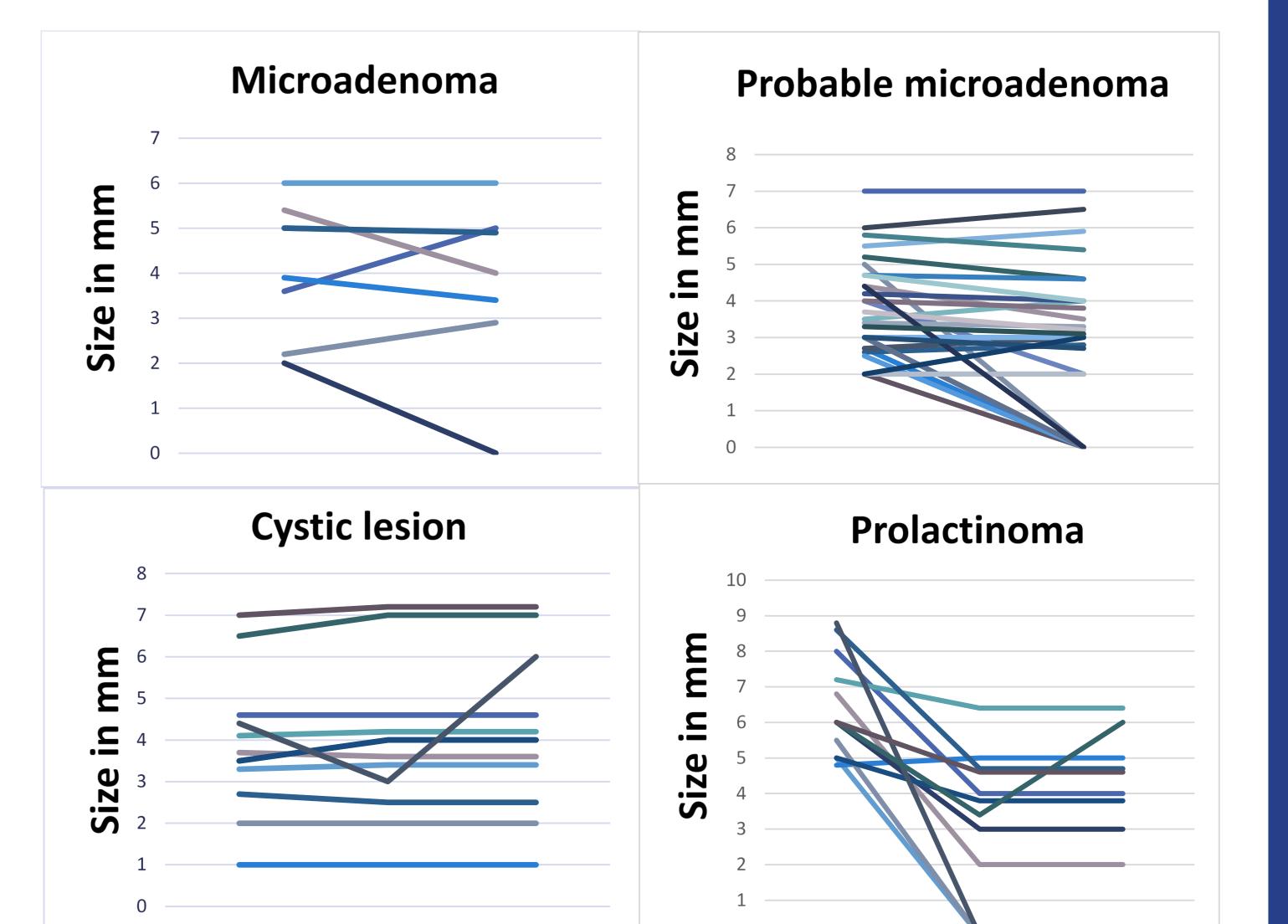
Table 1. Demographics, number of MRI scans and follow up time in children with pituitary microadenoma (including probable), cystic lesion and prolactinoma.

Microadenoma (including probable) and cystic lesion n=74	Prolactinoma n=12
12 (3-17)	16 (12-17)
43 (58%)	8 (67%)
48(65%)	12 (100%)
4(5%)*	7 (58%)*
1 (1%)	1 (8%)
32 (43%)	12 (100%)
2 (1-7)	5.5 (2-11)
37 (4-189)**	73.5 (26-120)
	(including probable) and cystic lesion n=74 12 (3-17) 43 (58%) 48(65%) 4(5%)* 1 (1%) 32 (43%) 2 (1-7)

*Normal result of ophthalmological examination in all cases. **For those 55 with repeated MRI

Table 2. Presenting symptoms pituitary microadenoma (including probable) and cystic lesion. **Presenting symptom** n = 74 **Precocious puberty** 27 **Growth disturbances** 22 **Delayed puberty** Headache Fatigue **Pubertal arrest** Pituitary insufficiency Stereotype movements Focal epilepsy Hyperprolactinemia (risperidone) Secondary amenorrhea

Figure 1. Size at initial and last MRI, as measured by an experienced neuroradiologist, for non-functioning pituitary microadenoma (n=8), probable microadenoma (n=35), cystic lesion (n=12), prolactinoma (n=12).



METHOD

Children included were under 18 years at first pituitary MRI, diagnosed with a microadenoma (microadenoma producing ACTH, GH or TSH excluded) or cyst < 10mm. Children with prolactinoma were included for testing of measurement accuracy. Pituitary lesion size at first and last MRI was reviewed by a neuroradiologist. All individual MRI examinations were re-evaluated by two radiologists.

CONCLUSIONS

Non-functioning pituitary microadenoma or cystic lesion in children, with no hormonal disturbances, show small clinically irrelevant size variations that do not require intervention or extended follow-up. We suggest for smaller pituitary lesions < 4 mm no regular MRI follow up, and for lesions 4-9 mm a follow up MRI after 24 months, since there still is limited knowledge of the natural cause and risk of long term growth in these conditions.

REFERENCES

1. Souteiro P RM, Santos-Silva R, Figueiredo R, Costa C, Belo S, Castro-Correia C, Carvalho D, Fontoura M. Pituitary incidentalomas in paediatric age are different from those described in adulthood. Pituitary. 2019;22(2):124-8 2. Pedicelli S, Alessio P, Scire G, Cappa M, Cianfarani S. Routine screening by brain magnetic resonance imaging is not indicated in every girl with onset of puberty between the ages of 6 and 8 years. J Clin Endocrinol Metab. 2014;99(12):4455-61 3. Derrick Kristina GW, Gensure Robert. Incidence and outcomes of pituitary microadenomas in children with short stature/growth hormone deficiency. Horm Res Paediatr. 2018;90(3):151-60. 4. Thaker VV, Lage AE, Kumari G, Silvera VM, Cohen LE. Clinical Course of Nonfunctional Pituitary Microadenoma in Children: A Single-Center Experience. J Clin Endocrinol Metab.2019;104(12):5906-12.

ACKNOWLEDGEMENTS

Research nurse Lena Rollof is gratefully acknowledged for assistance and support during the study, and Thomas Wiebe, Associate professor, Paediatric oncology, MD, PhD, for contribution of data.

CONTACT INFORMATION

E-mail: camilla.borghammar@med.lu.se

