UKCGG-National Cancer Genetics MDT meeting: 18-01-2024

Schwannomatosis lead by Oxford Centre for Genomic Medicine

Meeting contact: Dorothy Halliday, Dorothy.halliday@ouh.nhs.uk

Thank you to everyone who attended this session. Below is a summary of the presented cases and relevant publications and resources to refer to.

Presented Cases

Case 1	**Schwannomatosis overview**	Presented by: Dr Dorothy Halliday Oxford Genetics Department Dorothy.halliday@ouh.nhs.uk
	Actions/Outcome:	
Case 2	**Schwannomatosis overview and case of schwannomatosis with CDKN2A**	Presented by: Prof Gareth Evans Manchester Genetics Department Gareth.Evans@mft.nhs.uk
	Actions/Outcome:	
Case 3	**Schwannomatosis with Coffin Sirus features caused by SMARCB1**	Presented by: Dr Deirdre Donnelly Belfast Genetics Department Deirdre.Donnelly@belfasttrust.hscni.net
	Actions/Outcome:	
Case 4	**Schwannomatosis LZTR1**	Presented by: Dr Adam Shaw Guys Genetics Department Adam.shaw@gstt.nhs.uk
	Actions/Outcome:	
Case 5	**Schwannomatosis**	Presented by: Amy Taylor Cambridge Genetics Department amytaylor3@nhs.net
	Actions/Outcome:	
Case 6	**Ring 22 and Phelan McDermid Syndrome**	Presented by: Dr Claire Forde Manchester Genetics Department Claire.Forde@mft.nhs.uk
	Actions/Outcome:	

Next meeting details

Date	Thursday 16th May 2024
Time	12:30 pm-1:45 pm
Theme	Mixed Cases
Leading centre	Glasgow
Contact for cases	Rosemarie Davidson, rosemarie.davidson@ggc.scot.nhs.uk

Please send any questions or ideas for future meetings to Helen Hanson (helen.hanson6@nhs.net)