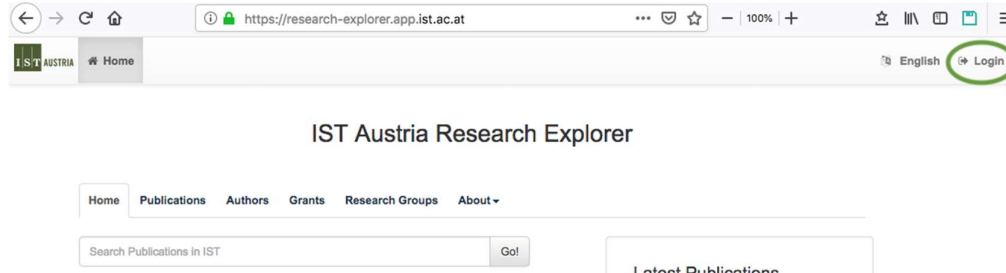


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## CUGC for pontocerebellar hypoplasia type 9 and spastic paraplegia-63

A. Marsh, G. Novarino, P. Lockhart, R. Leventer, European Journal of Human Genetics (2018).

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DOI **10.1038/s41431-018-0231-2**

Journal Article | Published

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Author	Marsh, Ashley P ; <b>Novarino, Gaia</b> <sup>IST Austria</sup> ; Lockhart, Paul L ; Leventer, Richard J
Department	<b>Novarino Group</b>
Abstract	Clinical Utility Gene Card. 1. Name of Disease (Synonyms): Pontocerebellar hypoplasia type 9 (PCH9) and spastic paraplegia-63 (SPG63). 2. OMIM# of the Disease: 615809 and 615686. 3. Name of the Analysed Genes or DNA/Chromosome Segments: AMPD2 at 1p13.3. 4. OMIM# of the Gene(s): 102771.
Publishing Year	<b>2018</b>
Date Published	2018-08-08
Journal Title	European Journal of Human Genetics
Acknowledgement	FP7 (Grant Agreement Number 261469)
IST-REX-ID	<b>50</b>

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