

Data updated 1 November 2016

**CNNM4 mutations, Jalili syndrome type, and statistics (including cases with unreported mutations): statistics**

Go to:	<a href="#">Mutations</a>		<a href="#">References</a>			<a href="#">Statistics</a>		
Origin	Total Cases Reported	Cases Examined	Families	Sibships	M	F	Gender n/a	JS Type
Kosovo*	2	2	1	1	2	0		A
Kosovo*	2	2	1	1	1	1		A
Kosovo*	1	1	1	1	0	1		A
Kosovo*	2	2	1	1	2	0		A
Kosovo*	2	2	1	1	0	2		A
Kosovo*	1	1	1	1	1	1		A
Total per column	10	10	6	6	6	5	0	
Kosovan mutations	10	10	6	6	6	5	0	

[References](#)

s on sibships and patients.\*

Reference (abbreviated) - See last column

Parry et al (2009)<sup>2</sup>, Michaeladis et al<sup>3</sup>

Polok et al (2009)<sup>4</sup> (Family A)

Zobor et al (2012)<sup>6</sup>

Luder et al (2013)<sup>8</sup>

Gerth-Kahlert et al (2013)<sup>10</sup>

Kiessling F et al (2016)<sup>16</sup>

including unreported mutations