

Erratum to: The Face of Lysosomal Storage Disorders in India: A Need for Early Diagnosis

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The original article contained mistake in Table 2. Corrected table is provided below.

Table 2 List of mutations detected in patients with LSD presenting to authors' centre

Serial number	Diagnosis	Mutation
1-7.	Gaucher disease (7)*	Homozygous <i>L444P</i>
8. ^a	Gaucher disease (1)	Homozygous <i>G383D</i>
9. ^a	Gaucher disease (1)	Homozygous <i>R359Q</i>
10. ^a	Gaucher disease (1)	Homozygous <i>S356F</i>
11.	Gaucher disease (1)	Homozygous <i>S125R</i>
12.	Gaucher disease (1)	<i>Homozygous F213I(c754A)</i>
13.	Gaucher disease (1)	Homozygous <i>R48W</i>
14.	Gaucher disease (1)	Rec Ex2 (c.44 T>C+46A>G+ IVS2+ 1g>a): R170C (c.625 C>T, exon 6)
15.	Gaucher disease (1)	<i>L444P/A456P/R496C/55 bpdel</i>
16.	Gaucher disease (1)	<i>L444P, R463C</i>
17.	Niemann-Pick disease (1)	Homozygous <i>R543X</i>
18.	Pompe disease (1)	<i>c.1003G>A (p.G335R)</i>
19.	Maroteaux-Lamy syndrome (MPS VI) (1)	Homozygous <i>W450C</i>
20.	Metachromatic leukodystrophy (1)	Compound heterozygous for <i>G34E</i> and <i>P136L</i>
21. ^b	Farber Disease (1)	Homozygous <i>IVS6+4A>G</i>

Figures in parentheses indicate the absolute number of patients diagnosed with the respective mutation

^a Published: Reference number 10

^b Published: Reference number 9

The online version of the original article can be found at <http://dx.doi.org/10.1007/s12098-014-1628-8>.

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