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CORRECTION
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Corrigendum: Expanding the Clinico-Genetic Spectrum of Myofibrillar Myopathy: Experience From a Chinese Neuromuscular Center

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A Corrigendum on

Expanding the Clinico-Genetic Spectrum of Myofibrillar Myopathy: Experience From a Chinese Neuromuscular Center

by Luo, Y.-B., Peng, Y., Lu, Y., Li, Q., Duan, H., Bi, F., et al. (2020). Front. Neurol. 11:1014. doi: 10.3389/fneur.2020.01014

In the original article, there was a mistake in **Table 1** as published. For patients 16 and 17, the protein alteration resulting from the c. 107545delG mutation should be p.Ala35849Glnfs*16. The corrected **Table 1** appears below.

The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

TABLE 1 | Genetics of the present MFM patient cohort.

Patient no.	Gene	Chromosome	Exon	Transcript no.	Nucleotide	Protein	Reference
1	*DES*	2	7	NM_1927	c.1256C>T	p.Pro419Leu	None
2	*DES*	2	7	NM_1927	c.1256C>T	p.Pro419Leu	None
3	*DES*	2	7	NM_1927	c.1256C>T	p.Pro419Leu	None
4	*DES*	2	7	NM_1927	c.1256C>T	p.Pro419Leu	None
5	*DES*	2	6	NM_1927	c.1096_1098delACA	p.Asn366del	(14)
6	*DES*	2	6	NM_1927	c.1096_1098delACA	p.Asn366del	(14)
7	*DES*	2	6	NM_1927	c.1096_1098delACA	p.Asn366del	(14)
8	*DES*	2	6	NM_1927	c.1076_1077ins GGCCAGTGG	p.Glu359delins GluAlaSerGly	None
9	*BAG3*	10	3	NM_004281	c.626C>T	p.Pro209Leu	(15)
10	*BAG3*	10	3	NM_004281	c.626C>T	p.Pro209Leu	(15)
11	*FLNC*	7	36	NM_001458	c.6004+3G>A	splicing	None
12	*FLNC*	7	33	NM_001458	c.5468C>T	P.Thr1823Met	None
13	*FHL1*	X	5	NM_001159702	c.386G>A	p.Cys129Tyr	None
14	*TTN*	2	344	NM_001267550	c.95134T>C	p.Cys31712Arg	(16–22)
15	*TTN*	2	344	NM_001267550	c.95185T>C	p.Trp31729Arg	(23)
16	*TTN*	2	69	NM_001267550	c. 19993G>T	p.Glu6665X	None
			363	NM_001267550	c. 107545delG	p.Ala35849Glnfs*16	None
17	*TTN*	2	69	NM_001267550	c. 19993G>T	p.Glu6665X	None
			363	NM_001267550	c. 107545delG	p.Ala35849Glnfs*16	None
18	None	-	-	-	-	-	-