



GFB

Gruppo Familiari Beta-sarcoglicanopatie

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NEWSLETTER N.4

A MILESTONE IN THE HISTORY OF DUCHENNE DYSTROPHY AND GENE THERAPY

The first gene therapy treatment for Duchenne Dystrophy in children in the 4-5 age group was approved by the US FDA on June 22. More information at the link: www.DMDtreatmentoption.com

BREAKING NEWS

Sarepta Therapeutics Announces
U.S. FDA Approval of Therapy to
Treat Duchenne Muscular Dystrophy



THE PATIENT IRANIAN GROUP IS BORN

The GFB group of Persian-speaking patients is born!

The group consists of patients residing in Iran and is coordinated by Shaghayegh Parsanejad, GFB interpreter.

Coordinator: Shaghayegh Parsanejad

Join the Whatsapp group by writing to Shagha: +39 389 613 6092

Instagram: [@gfbricercadm](https://www.instagram.com/gfbricercadm)

E-mail: segreteria@beta-sarcoglicanopatie.it

SUPPORTERS 2022

In 2022 GFB received these contributions:

240 €	Donazione NN per la ricerca scientifica da Roma
24.000 €	Contributo NN per la ricerca scientifica
2.219 €	Donazione NN per progetto ausili da Roma
50 €	Donazione NN per la ricerca scientifica da Treviso
450 €	Contributo NN per la ricerca scientifica
12.689 €	Contributo 5x1000 anno 2021
420 €	Contributo NN per la ricerca scientifica
40 €	Contributo NN per la ricerca scientifica da Salerno
3.000 €	Contributo NN per la ricerca scientifica da Lecco
300 €	Contributo NN per la ricerca scientifica
100 €	Contributo NN per la ricerca scientifica
43.508 €	TOTALE 2022