

FOR PATIENTS WITH POMPE DISEASE

# KEEP ACTIVE POSSIBLE

Initiating MYOZYME® upon diagnosis of Pompe disease increases acid  $\alpha$ -glucosidase (GAA) activity—to stabilize respiratory function and improve and maintain mobility so patients can stay active and preserve their quality of life<sup>1</sup>



All photos are actor portrayals, not actual patients.

**Pompe disease calls for an immediate response: treat with MYOZYME.<sup>2,3</sup>**

MYOZYME is indicated for long-term enzyme replacement therapy (ERT) in adults and pediatric patients of all ages with a confirmed diagnosis of Pompe disease (acid  $\alpha$ -glucosidase deficiency).<sup>1</sup>

 **Myozyme**<sup>®</sup>  
(alglucosidase alfa)

# MYOZYME®: approved long-term treatment for Pompe disease that addresses the underlying cause to keep patients going<sup>1</sup>



## In adult patients with LOPD, MYOZYME:

- **Helped to retain breathing function**, stabilizing pulmonary function and preserving ventilatory status in most patients<sup>20,25</sup>
- **Maintained independent walking**, sustaining long-term mobility in adult patients with LOPD<sup>20</sup>



## In pediatric patients with LOPD, MYOZYME:

- **Improved respiratory health**, stabilizing or improving pulmonary function, and reducing the need for ventilatory support<sup>24,27</sup>
- **Enabled continued participation in activities of daily living**, increasing or stabilizing motor function, allowing patients to maintain mobility<sup>22,24,26,27</sup>



## In patients with IOPD, MYOZYME:

- **Increased survival and achievement of motor function milestones**, prolonging life, increasing ventilator-free survival, and stabilizing cardiac health and motor function<sup>1,15,29-31</sup>



Please review Product Information before prescribing.  
Please Scan the QR code to access Full SMPC.



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**Myozyme®**  
(alglucosidase alfa)