

**Company announcement**

No. 20/2022

Inside information

**Orphazyme A/S in restructuring**

Ole Maaløes Vej 3

DK-2200 Copenhagen N

[www.orphazyme.com](http://www.orphazyme.com)

Company Registration No. 32266355

## Statutory restructuring plan adopted by creditors

**Copenhagen, Denmark, April 7, 2022** – Orphazyme A/S in restructuring (ORPHA.CO) (“Orphazyme” or the “Company”), a late-stage biopharmaceutical company, announces that today an in-court meeting was held where the statutory restructuring plan as proposed and published by the Company through company announcement no. 15/2022 on March 31, 2022 and company announcement no. 17/2022 on April 4, 2022, was adopted by the Company’s creditors in accordance with section 11 e of the Danish Insolvency Act. No amendments to the restructuring proposal were made at the meeting.

Following the adoption of the statutory restructuring plan, the Company will continue its restructuring proceedings in accordance with the restructuring plan.

**For additional information, please contact****Orphazyme A/S in restructuring**

Anders Vadsholt, Chief Executive Officer and Chief Financial Officer

+45 2898 9055

John Sommer Schmidt, Restructuring Administrator

+45 8620 7500

**About Orphazyme**

Orphazyme is a late-stage biopharmaceutical company developing arimoclomol for Niemann-Pick disease type C (NPC). Orphazyme is headquartered in Denmark. Orphazyme’s shares are listed on Nasdaq Copenhagen (ORPHA).

**About arimoclomol**

Arimoclomol is an investigational drug candidate that amplifies the production of heat shock proteins (HSPs). HSPs can rescue defective misfolded proteins and improve the function of lysosomes. Arimoclomol is administered orally, and has now been studied in 10 Phase 1, four Phase 2, and three pivotal Phase 2/3 trials. Arimoclomol has received Orphan Drug Designation (ODD) for NPC in the US and EU. Arimoclomol has received Fast-Track Designation (FTD), Breakthrough Therapy Designation (BTD), and Rare Pediatric Disease Designation (RPDD) from the U.S. Food and Drug Administration (FDA) for NPC. On June 17, 2021, Orphazyme received a Complete Response Letter from the FDA regarding its New Drug Application for arimoclomol for the treatment of NPC. The Company plans to request a Type C Meeting with the FDA in Q2 2022.

**Forward-looking statement**

This company announcement may contain certain forward-looking statements under the U.S. Private Securities Litigation Reform Act of 1995 and otherwise, including forward-looking statements about the company’s restructuring proceedings. Although the Company believes its expectations are based on reasonable assumptions, all statements other than statements of historical fact included in this company announcement about future events are subject to (i) change without notice and (ii) factors beyond the Company’s control, including pursuant to regulatory or judicial intervention. Except as required by law, the Company assumes no obligation to update these forward-looking statements publicly, or to update the reasons actual results could differ materially from those anticipated in the forward-looking statements, even if new information becomes available in the future.