## PRESS RELEASE

# Relief Reports that NeuroRx Announced that RLF-100™ (Aviptadil) has Successfully Demonstrated Ten-Day Accelerated Recovery from Respiratory Failure among Critically Ill Patients with Covid-19 Treated with High Flow Nasal Oxygen at 28 Day Interim Endpoint

**Geneva, Switzerland, February 24, 2021 –** RELIEF THERAPEUTICS Holding AG (SIX: RLF, OTCQB: RLFTF) ("**Relief**" or the "**Company**"), a biopharmaceutical company with its lead compound RLF-100™ (aviptadil) in advanced clinical development, announces that NeuroRx, Inc. has reported data from the Phase 2b/3 trial of RLF-100™ for the treatment of Respiratory Failure in Critical COVID-19 patients. NeuroRx is solely responsible for clinical development and regulatory submissions related to RLF-100™ in the U.S.

The relevant NeuroRx press release can be found here.

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### ABOUT RELIEF

Relief focuses primarily on clinical-stage programs based on molecules of natural origin (peptides and proteins) with a history of clinical testing and use in human patients or a strong scientific rationale. Currently, Relief is concentrating its efforts on developing new treatments for respiratory disease indications. Its lead drug candidate RLF-100™ (aviptadil) is being investigated in two placebo-controlled U.S. late-stage clinical trials in respiratory deficiency due to COVID-19. Relief holds a patent issued in the United States and various other countries covering potential formulations of RLF-100™.

RELIEF THERAPEUTICS Holding AG is listed on the SIX Swiss Exchange under the symbol RLF and quoted in the U.S. on OTCQB under the symbol RLFTF.

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