Newron reports amendment of terms of up to EUR 38 million financing agreement to support advancement of Phase 3 evenamide program

Total proceeds expected to fund global ENIGMA-TRS studies and extend cash runway beyond pivotal 12-and 26-week results, supporting development of evenamide as the potential first add-on therapy for treatment-resistant schizophrenia

Milan, Italy, November 27 , 2026, 07:00 CET – Newron Pharmaceuticals S.p.A. ("Newron", or the "Company") (SIX: NWRN, XETRA: NP5), a biopharmaceutical company focused on the development of novel therapies for patients with diseases of the central and peripheral nervous system, today announced that it has agreed with the investor group participating in Newron's up to EUR 38 million financing in February 2026 on an amendment of certain terms related to the third tranche of the financing agreement, corresponding to gross proceeds of EUR 5.5 million.

Under the amendment, the investor group is now expected to subscribe to the third tranche no later than December 31, 2026, strengthening the Company's financial position as it advances the ENIGMA-TRS Phase 3 program. The shares will be issued at a subscription price to be calculated pursuant to an agreed formula.

The investor group is expected to subscribe to an additional number of newly issued shares for further proceeds of EUR 12 million upon disclosure of results from the ENIGMA-TRS studies, conditional upon such results being positive.

About Newron Pharmaceuticals

Newron (SIX: NWRN, XETRA: NP5) is a biopharmaceutical company focused on the development of innovative therapies for patients with diseases of the central and peripheral nervous system. Headquartered in Bresso near Milan, Italy, the Company has a strong track record of advancing neuroscience-based treatments from discovery to market. Newron's lead compound, evenamide, is a first-in-class glutamate modulator and has the potential to be the first add-on therapy for treatment-resistant schizophrenia (TRS) and for poorly responding patients with schizophrenia. Evenamide is currently developed in the global pivotal ENIGMA-TRS Phase 3 development program. Clinical trial results to date demonstrate the benefits of this drug candidate in the TRS as well as poorly responding patient population, with significant improvements across key efficacy measures increasing over time, as well as a favorable safety profile, which is uncommon for available antipsychotic medications. Newron has signed development and commercialization agreements for evenamide with EA Pharma (a subsidiary of Eisai) for Japan and other Asian territories, as well as Myung In Pharm for South Korea. Newron's first marketed product, Xadago®/safinamide has received marketing authorization for the treatment of Parkinson's disease in the European Union, Switzerland, the UK, the USA, Australia, Canada, Latin America, Israel, the United Arab Emirates, Japan and South Korea. The product is commercialized by Newron's partner Zambon, with Supernus Pharmaceuticals holding marketing rights in the U.S., and Meiji Seika responsible for development and commercialization in Japan and other key Asian territories. For more information, please visit: www.newron.com and connect with us on LinkedIn.

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