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## PRESS RELEASE

**FOR IMMEDIATE RELEASE**

### **GENFIT ANNOUNCES TRADING RESUMPTION OF ITS SHARES AND OF ITS OCEANES ON EURONEXT PARIS AND EURONEXT ACCESS**

**Lille (France), Cambridge (Massachusetts, United States), March 27, 2019 – 3:00 pm CET –** GENFIT SA (Euronext: GNFT – ISIN: FR0004163111) (“**GENFIT**” or the “**Company**”), a French biopharmaceutical company focused on discovering and developing drug candidates and diagnostic solutions targeting liver diseases, in particular those of metabolic origin, and hepatobiliary diseases, announces today that it has requested that trading of its ordinary shares on the regulated market of Euronext in Paris and of its OCEANES on Euronext Access resumes as from 4:00 p.m. CET, today.

Trading of the ordinary shares and of the OCEANES of GENFIT was halted at the request of the Company on March 27, 2019 from 9:00 am CET in the context of its initial public offering on the Nasdaq Global Select Market, the terms of which have been previously announced today, in order to allow for the confirmation of allocations to investors and for the commencement of trading of the Company's American Depositary Shares (“ADSs”) on the Nasdaq Global Select Market.

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This press release does not constitute an offer to sell or the solicitation of an offer to buy securities, and shall not constitute an offer, solicitation or sale in any jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of that jurisdiction.

#### **ABOUT GENFIT**

GENFIT is a biopharmaceutical company focused on discovering and developing drug candidates and diagnostic solutions targeting liver diseases, in particular those of metabolic origin, and hepatobiliary diseases. GENFIT concentrates its R&D efforts in areas of high unmet medical needs corresponding to a lack of approved treatments. GENFIT's lead proprietary compound, elafibranor, is a drug candidate currently being evaluated in one of the most advanced Phase 3 studies worldwide (RESOLVE-IT) in NASH, considered by regulatory authorities as a medical emergency because it is silent, with potentially severe consequences, and with a prevalence on the rise. Elafibranor has also obtained positive preliminary results in a Phase 2 clinical trial in PBC, a severe chronic liver disease. As part of its comprehensive approach to clinical

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management of NASH patients, GENFIT is conducting an ambitious discovery and development program aimed at providing patients and physicians with a blood-based test for the diagnosis of NASH, i.e. non-invasive and easy-to-access. With facilities in Lille and Paris, France, and Cambridge, MA (USA), the Company has approximately 150 employees. GENFIT is a public company listed in compartment B of Euronext's regulated market in Paris (Euronext: GNFT - ISIN: FR0004163111).

### DISCLAIMER

This press release and the information contained herein do not constitute an offer to sell or a solicitation of an offer to buy or subscribe to shares in GENFIT in any country. This press release has been prepared in both French and English. In the event of any differences between the two texts, the French language version shall supersede.

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