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PRESS RELEASE

GenSight Biologics announces the availability of a prospectus in connection with the listing of new shares on Euronext Paris

Paris, France, November 22, 2023, 7.30 am CET – GenSight Biologics (Euronext: SIGHT, ISIN: FR0013183985, PEA-PME eligible) (the "**Company**"), a biopharma Company focused on developing and commercializing innovative gene therapies for retinal neurodegenerative diseases and central nervous system disorders, announced today the availability of a listing prospectus approved by the French *Autorité des Marchés Financiers* (the "**AMF**") under number 23-481, on November 21, 2023, in the context of the admission to trading on the regulated market of Euronext Paris of new shares of the Company that will be or may be issued upon (i) a capital increase without shareholders' preferential right reserved to a category of persons satisfying determined characteristics for an amount of €4,399,686.28, (ii) the automatic conversion of 60 convertible bonds subscribed by Sofinnova Crossover I SLP, by Invus Public Equities LP and by UPMC Enterprises issued on August 4, 2023, and (iii) the potential conversion of 120 convertible bonds subscribed by CVI Investments, Inc. on December 28, 2022.

The listing prospectus approved by the AMF under number 23-481 comprises:

- the 2022 universal registration document filed by the Company with the AMF on May 10, 2023 under number D.23-0406;
- the amendment to the 2022 universal registration document filed by the Company with the AMF on November 21, 2023 under number D.23-0406-A01;
- a securities note; and
- the summary of the prospectus (included in the securities note).

These documents are available free of charge on the website of the Company (https://www.gensight-biologics.com) and of the AMF (https://www.amf-france.org).

Contacts

GenSight Biologics
Chief Financial Officer
Thomas Gidoin
tgidoin@gensight-biologics.com

LifeSci Advisors
Investor Relations
Guillaume van Renterghem
gvanrenterghem@lifesciadvisors.com
+41 (0)76 735 01 31

About GenSight Biologics

GenSight Biologics S.A. is a clinical-stage biopharma company focused on developing and commercializing innovative gene therapies for retinal neurodegenerative diseases and central nervous system disorders. GenSight Biologics' pipeline leverages two core technology platforms, the Mitochondrial Targeting Sequence (MTS) and optogenetics, to help preserve or restore vision in patients suffering from blinding retinal diseases. GenSight Biologics' lead product candidate, LUMEVOQ® (GS010; lenadogene nolparvovec), is an investigational compound and has not been registered in any country at this stage, developed for the treatment of Leber Hereditary Optic Neuropathy (LHON), a rare mitochondrial disease affecting primarily teens and young adults that leads to irreversible blindness. Using its gene therapy-based approach, GenSight Biologics' product candidates are designed to be administered in a single treatment to each eye by intravitreal injection to offer patients a sustainable functional visual recovery.

Disclaimer

This announcement is an advertisement and not a prospectus within the meaning of the Prospectus Regulation. Any decision to purchase shares must be made solely on the basis of publicly available information on the Company.

No communication and no information in respect of the offering by the Company of its shares may be distributed to the public in any jurisdiction where registration or approval is required. No steps have been taken or will be taken in any jurisdiction where such steps would be required. The offering or subscription of shares may be subject to specific legal or regulatory restrictions in certain jurisdictions.

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