

Transgene announces the resume of trading of its shares on Euronext Paris as of the opening of the markets

Strasbourg, France, 27 November 2025, 00:15 a.m. CET – **Transgene (Euronext Paris: TNG)** had requested the suspension of trading of its shares (ISIN: FR0005175080) on Euronext Paris **as from Wednesday, 26 November 2025**, before market opening, pending the publication of the results of its capital increase.

Following the announcement of the results of its capital increase on 26 November 2025, trading of its shares on Euronext Paris will resume on Thursday, 27 November 2025, at market opening.

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About Transgene

Transgene (Euronext: TNG) is a biotechnology company focused on designing and developing targeted immunotherapies for the treatment of cancer. The Company's clinical-stage programs consist of a portfolio of viral vector-based immunotherapeutics. TG4050, the first individualized therapeutic vaccine based on the *myvac*[®] platform is the Company's lead asset, with demonstrated proof of principle in patients in the adjuvant treatment of head and neck cancers. The Company has other viral vector-based assets, including BT-001, an oncolytic virus based on the Invir.IO[®] viral backbone, which is in clinical development. The Company also conducts innovative discovery and preclinical work, aimed at developing novel viral vector-based modalities.

With Transgene's *myvac*[®] platform, therapeutic vaccination enters the field of precision medicine with a novel immunotherapy that is fully tailored to each individual. The *myvac*[®] approach allows the generation of a virus-based immunotherapy that encodes patient-specific mutations, identified and selected through advanced Artificial Intelligence technologies.

With its proprietary platform Invir.IO[®], Transgene is building on its viral vector engineering expertise to design a new generation of multifunctional oncolytic viruses.

Additional information about Transgene is available at: www.transgene.com

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This press release contains forward-looking statements, which are subject to numerous risks and uncertainties, which could cause actual results to differ materially from those anticipated. The occurrence of any of these risks could have a significant negative outcome for the Company's activities, perspectives, financial situation, results, regulatory authorities' agreement with development phases, and development. The Company's ability to commercialize its products depends on but is not limited to the following factors: positive pre-clinical data may not be predictive of human clinical results, the success of clinical studies, the ability to obtain financing and/or partnerships for product manufacturing, development and commercialization, and marketing approval by government regulatory authorities. For a discussion of risks and uncertainties which could cause the Company's actual results, financial condition, performance or achievements to differ from those contained in the forward-looking statements, please refer to the Risk Factors ("Facteurs de Risque") section of the Universal Registration Document, available on the AMF website (<http://www.amf-france.org>) or on Transgene's website (www.transgene.com). Forward-looking statements speak only as of the date on which they are made, and Transgene undertakes no obligation to update these forward-looking statements, even if new information becomes available in the future.

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France

In France, the offer of Transgene shares described in this press release will be made in the context of (i) an offer reserved to specified categories of beneficiaries, pursuant to article L. 225-138 of the French Commercial Code, (ii) a public offering primarily intended to retail investors through the PrimaryBid platform, pursuant to article L. 225-136 of the French Commercial Code, and (iii) an offer reserved to TSGH, pursuant to article L. 225-138 of the French Commercial Code.

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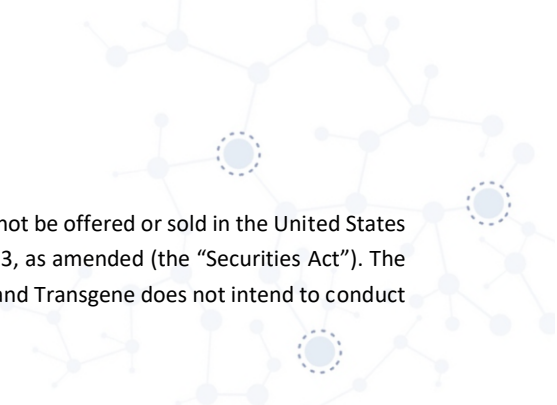
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MIFID

MIFID II Product Governance/Target Market: solely for the purposes of the requirements of article 9.8 of the EU Delegated Directive 2017/593 relating to the product approval process, the target market assessment in respect of the shares of Transgene has led to the conclusion in relation to the type of clients criteria only that: (i) the type of clients to whom the shares are targeted is eligible counterparties and professional clients and retail clients, each as defined in Directive 2014/65/EU, as amended (“**MiFID II**”); and (ii) all channels for distribution of the shares of Transgene to eligible counterparties and professional clients and retail clients are appropriate. Any person subsequently offering, selling or recommending the shares of Transgene (a “distributor”) should take into consideration the type of client assessment; however, a distributor subject to MiFID II is responsible for undertaking its own target market assessment in respect of the shares of Transgene and determining appropriate distribution channels.

General

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