

## **2019 Financial Calendar**

PARIS – December 10, 2018 – 5.45 pm (CET) - STENTYS (FR0010949404 — STNT), French group specialized in medical technologies for interventional cardiology, today announced its preliminary financial calendar for 2019.

Event	Date *
Full-Year 2018 Revenues	Thursday, January 10, 2019
Full-Year 2018 Results	Thursday, March 28, 2019
Q1 2019 Revenues	Thursday, April 11, 2019
Shareholders' General Meeting	Thursday, May 9, 2019
Q2 2019 Revenues	Thursday, July 11, 2019
Half-Year 2019 Results	Thursday, September 26, 2019
Q3 2019 Revenues	Thursday, October 10, 2019

Financial year end on 31 December

\* Subject to modification. Press releases are published after financial markets close.

## **About STENTYS**

The STENTYS group develops and markets minimally-invasive cardiovascular solutions for the needs of interventional cardiology. Its extensive range of innovative products, including drug-eluting stents, coronary and drug-eluting balloons as well as cardiovascular accessories, is marketed in over 60 countries. Thanks to its flagship product, Xposition S, the self-apposing stent that adapts to vessels with variable diameters and enables the treatment of complex arterial disorders, and to its portfolio of balloons and accessories, STENTYS covers all coronary indications.

Additional information is available at www.stentys.com

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STENTYS is listed on Compartment C of Euronext Paris ISIN: FR0010949404 – Ticker: STNT





## **Forward-looking Statements**

This press release contains forward-looking statements about the Company that are based on numerous assumptions regarding the Company's present and future business strategies and the environment in which it will operate in the future which may not be accurate. Such forward-looking statements involve known and unknown risks which may cause the Company's actual results, performance or achievements to differ materially from any future results, performance or achievements expressed or implied by such forward-looking statements. Such factors include, among others, risks associated with the development and commercialization of the Company's products, market acceptance of the Company's products, its ability to manage growth, the competitive environment in relation to its business area and markets, its ability to enforce and protect its patents and proprietary rights, uncertainties related to the U.S. FDA approval process, slower than expected rates of patient recruitment for clinical trials, the outcome of clinical trials, and other factors, including those described in the Section 4 "Risk Factors" of the Company's 2016 Registration Document (*document de référence*) filed with the French *Autorité des Marchés Financiers* (AMF) on November 29, 2017 under number D.17-1084.

