

## STENTYS First Half 2014 Revenues Rise 18%

**PRINCETON, N.J. and PARIS – July 24, 2014 – STENTYS (FR0010949404 – STNT), a medical technology company commercializing in Europe the world's first and only Self-Apposing® stent to treat acute myocardial infarction (AMI), today announces revenues for the second quarter and first half of 2014, as well as cash position at June 30, 2014.**

- **Half-year and quarterly revenues\***

€ thousands	Half year – 6 months			Quarterly – 3 months		
	H1 2014	H1 2013	Var. (%)	Q2 2014	Q2 2013	Var. (%)
<b>Revenues</b>	<b>1,934.0</b>	1,638.4	+18.0%	<b>1,080.0</b>	913.4	+18.2%

\* Audited data

Revenues increased 18.2% to €1,080 thousand in the second quarter of 2014 compared with €913 thousand in the same quarter of 2013.

For the first half of 2014, sales rose 18.0% to €1,934 thousand from €1,638.4 thousand for the same period in 2013. This growth was generated exclusively with the Company's current range of bare-metal and drug-eluting Self-Apposing® stents and further illustrates the popularity of STENTYS' technology among cardiologists in Europe and internationally.

- **A robust financial position**

At June 30, 2014, STENTYS had cash and equivalents totaling €23 million. The Company believes its current cash balance is sufficient to continue its marketing and clinical development.

**Gonzague Issenmann, Chief Executive Officer and co-founder of STENTYS, commented:** "Following a good first half, revenues should continue growing at a steady rate during the second half of the year, primarily as a result of the commercial development in recently opened regions (South America, Asia) and the CE marking of our Sirolimus-eluting stent which is expected before the end of the year."

- **Upcoming financial publication**

STENTYS expects to publish its financial results for the first half of 2014 on August 28, 2014.

### About the STENTYS Self-Apposing® Stent

The STENTYS Self-Apposing® Stent addresses the stent-sizing dilemma that cardiologists are confronted with when treating heart attack patients or patients with atypical artery anatomy. Its flexible, self expanding design takes the shape of the patient's unique vessel anatomy and apposes to the irregular contours of a blood vessel, in particular after an AMI as the vessel dilates and the clot dissolves. It reduces the risk of malapposition and complications associated with conventional stents in this setting. The STENTYS Self-Apposing Stent has been marketed in Europe since receiving CE Mark in 2010. The STENTYS Sirolimus-eluting stent should receive the CE Mark during the second half of 2014.

### About STENTYS

STENTYS is developing and commercializing innovative solutions for the treatment of patients with acute myocardial infarction (AMI, or heart attack) and complex coronary artery disease. STENTYS' Self-Apposing® Stents are designed to adapt to vessels with ambiguous or fluctuating diameters, particularly in the post-infarction phase, in order to prevent the malapposition problems associated with conventional stents. In the APPOSITION III clinical trial, STENTYS stents demonstrated a very low one year mortality rate among 1,000 high-risk AMI patients when compared to recent studies with conventional stents. **More information is available at [www.stentys.com](http://www.stentys.com).**

This press release contains forward-looking statements about the Company's business. Such forward looking statements 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, including with respect to a pre-market approval for the Company's BMS, 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 2011 Registration Document (*document de référence*) filed with the French *Autorité des Marchés Financiers* on August 27, 2013 under number R.13-040 as such section may be updated from time to time.

**STENTYS**

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