

ThromboGenics on Track to file Ocriplasmin by end of 2011

Retinal community showing increasing interest in ocriplasmin's clinical profile

Leuven, Belgium – 9 August, 2011 – ThromboGenics NV (Euronext Brussels: THR), a biopharmaceutical company focused on developing innovative ophthalmic medicines, today issued an update on its filing plans and pre-commercialization activities for its lead product ocriplasmin.

ThromboGenics is pleased to report that it is making good progress in preparing the dossiers that it plans to file with the EMA and FDA in order to gain marketing approval for ocriplasmin in Europe and the U.S. respectively. The Company expects to make the first ocriplasmin filing with the EMA in the next two months with the FDA filing taking place before year end. To-date ThromboGenics' discussions with both regulatory authorities have been encouraging and supportive.

In parallel with this regulatory activity, ThromboGenics has continued to invest in its pre-commercialization activities for ocriplasmin. A key element of these activities has been the presentation of the positive Phase III MIVI-TRUST trial data by leading retinal specialists at many of the most important ophthalmology conferences in both the U.S. and Europe. The retinal community's reaction to these presentations and to ThromboGenics' other medical education activities for ocriplasmin has also been very positive.

Patrik de Haes, the CEO of ThromboGenics said, "I am very pleased to report that we are making excellent progress towards our key goal of bringing ocriplasmin to market by the end of 2012. We plan to make our first regulatory filing in Europe in the next two months and we intend to file with the FDA before year end. Our continuing dialogue with the retinal community based on its positive Phase III results gives me great confidence that ocriplasmin, if approved, will rapidly become an attractive pharmacological treatment option for patients with important retinal disorders such as symptomatic vitreo-macular adhesion and macular hole."

- Ends -

For further information please contact:

ThromboGenics

Dr. Patrik De Haes, CEO

Tel: + 32 16 75 13 10

patrik.dehaes@thrombogenics.com

Chris Buyse, CFO

Tel: + 32 16 75 13 10

chris.buyse@thrombogenics.com

Citigate Dewe Rogerson

David Dible/ Nina Enegren/ Sita Shah

Tel: +44 (0) 207 638 95 71
sita.shah@citigatedr.co.uk**About ThromboGenics**

ThromboGenics is a biopharmaceutical company focused on the discovery and development of innovative medicines for the treatment of eye disease, vascular disease and cancer. The Company's lead product ocriplasmin (microplasmin) has successfully completed two Phase III clinical trials for the pharmacological treatment of symptomatic vitreomacular adhesion (sVMA). Ocriplasmin is also being evaluated in Phase II clinical development for additional vitreoretinal conditions. ThromboGenics is also developing novel antibody therapeutics in collaboration with BioInvent International. These include TB-402 (anti-Factor VIII), a long-acting anticoagulant in Phase II, and TB-403 (anti-PlGF) in Phase Ib/II for cancer in partnership with Roche.

ThromboGenics is headquartered in Leuven, Belgium. The Company is listed on NYSE Euronext Brussels under the symbol THR. More information is available at www.thrombogenics.com.

Important information about forward-looking statements

Certain statements in this press release may be considered "forward-looking". Such forward-looking statements are based on current expectations, and, accordingly, entail and are influenced by various risks and uncertainties. The Company therefore cannot provide any assurance that such forward-looking statements will materialize

and does not assume an obligation to update or revise any forward-looking statement, whether as a result of new information, future events or any other reason. Additional information concerning risks and uncertainties affecting the business and other factors that could cause actual results to differ materially from any forward-looking statement is contained in the Company's Annual Report.