Press release
February 4, 2015
Gothenburg

XVIVO Perfusion breaks new ground in Germany and Texas

The first XPS™ in Europe is going to be delivered to Germany and the first XPS™ is delivered to the densely populated Texas in the USA. Installation and training will take place during the month of February in both these clinics.

XVIVO Perfusion's XPS™ has been approved by FDA in the USA for the evaluation of marginal lungs before lung transplantation and has received a CE mark in Europe. XPS™ is used with good clinical results at leading clinics in the USA and there is already one at a clinic in Asia. In Europe, EVLP using STEEN Solution™ was previously done manually, in other words without XPS™. Access to XPS™ enables EVLP to be standardized and simplified. There is now great interest in XPS™ in both the USA and in Europe. Further university hospitals in Europe and the USA plan to start using EVLP (lung evaluation) combined with XPS™ and STEEN Solution™.

"The first XPS™ for Europe is an important milestone for XVIVO Perfusion and we are very pleased that Europe is now part of the expansion of EVLP using XPS™ and STEEN Solution™. This is a big step forward in being able to save more patients with terminal lung disease and give them a new life," says Magnus Nilsson, CEO of XVIVO Perfusion.

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XVIVO Perfusion AB is a medical technology company which develops solutions and systems for assessing and preserving organs outside the body and for selecting usable organs and maintaining them in optimal condition pending transplantation.
The company is headquartered in Gothenburg, Sweden, and has one office in the USA. The XVIVO share is listed on NASDAQ OMX First North and has the ticker symbol XVIVO. More information can be found on the website www.xvivoperfusion.com. The Certified Adviser is Redeye, www.redeye.se.

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