

Company Announcement no. 2/2015

To: NASDAQ OMX Copenhagen A/S

Hørsholm, Denmark, 14 January 2015

### **U.S. District Court Sets Briefing Schedule in Veloxis Litigation**

Veloxis Pharmaceuticals A/S (OMX: VELO) (Veloxis) today confirmed that the U.S. District Court for the District of Columbia has set a briefing schedule that enables complete briefing in its pending case by March 3, 2015. Veloxis is seeking an order requiring FDA to grant final approval to Envarsus® XR. While Veloxis currently expects the matter to be resolved by mid-year 2015, it is possible that delays may occur. There can be no assurance that Veloxis will be successful in its action against FDA.

#### **For more information, please contact:**

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#### **About Envarsus®**

Tacrolimus is a leading immunosuppression drug used for the prevention of transplant allograft rejection after organ transplantation. Envarsus® (tacrolimus prolonged-release tablets) has received marketing authorization in the EU for prophylaxis of organ rejection in kidney and liver transplant recipients. In the US, Envarsus®, known as Envarsus® XR (tacrolimus extended-release tablets), has received Tentative Approval as a once-daily tablet version of tacrolimus for prophylaxis of organ rejection in kidney transplant patients in combination with other immunosuppressants. Envarsus® XR has received orphan drug designation in the U.S. Veloxis plans to commercialize Envarsus® XR in the US through its own sales force and in the EU through its partnership with Chiesi Farmaceutici SpA.

#### **About Veloxis Pharmaceuticals**

Based in Hørsholm, Denmark, with an office in New Jersey, Veloxis Pharmaceuticals A/S, or Veloxis, is a specialty pharmaceutical company. Veloxis' unique, patented delivery technology, MeltDose®, is designed to enhance the absorption and bioavailability of select orally administered drugs. Veloxis is listed on the NASDAQ OMX Copenhagen under the trading symbol OMX: VELO.

For further information, please visit [www.veloxis.com](http://www.veloxis.com).