AngioDynamics Receives 510(k) Clearance for AlphaVac Mechanical Thrombectomy System

LATHAM, N.Y.– (BUSINESS WIRE) – Jun. 8, 2021 – AngioDynamics, Inc. (NASDAQ: ANGO), a leading provider of innovative, minimally invasive medical devices for vascular access, peripheral vascular disease, and oncology, today announced that it has received 510(k) clearance from the United States Food and Drug Administration (FDA) for the AlphaVac Mechanical Thrombectomy System, an off-circuit, multi-purpose mechanical aspiration thrombectomy device for the non-surgical removal of thrombi or emboli from the vasculature.

AngioDynamics continues to anticipate the commercial release of the AlphaVac System in the second half of the calendar year 2021.

The Company will report financial results for the fourth quarter and fiscal year 2021 before the market opens on Tuesday, July 13, 2021. The Company's management will host a conference call at 8:00 a.m. ET to discuss the results.

Management will also host a virtual Investor and Technology Day following its earnings conference call at 9:30 a.m. ET that same day.

Further details, including registration, webcast, and dial-in information, will be provided at a later date.

About AngioDynamics, Inc.

AngioDynamics is a leading and transformative medical technology company focused on restoring healthy blood flow in the body's vascular system, expanding cancer treatment options, and improving quality of life for patients.

The Company's innovative technologies and devices are chosen by talented physicians in fast-growing healthcare markets to treat unmet patient needs. For more information, visit www.angiodynamics.com.