

November 6, 2020

--Featuring a panel of distinguished guest speakers-

--Prelaunch activities for tenapanor well underway in anticipation of PDUFA goal date of April 29, 2021-

--Pipeline expansion with novel, first-in-class drug candidate for hyperkalemia advancing in the clinic-

FREMONT, Calit., Nov. 6, 2020. PRNewswire / - Andelyn, Inc. (Nasdar, ARDX), a biophamaceutical company locused on developing first-in-class medicines to improve treatment for people with kidney and cardiovascular diseases, today announced that it will hold a virtual Analyst Day on Thursday, November 12, 2020, from 9:30 a.m. to 12:00 p.m. ET. This year's event will focus on the company's launch and commercialization plans in anticipation of the potential approval of tempany for the control of sevent physichastes with chronic kidney desease (XKD) on dai/sis. Additionally, the company will review its pipeline therepeutic for hyperkalemia.



The management team will be joined by

German Hemandez, M.D., FASN, FACP, associate nephrologist at El Paso Kidney Specialists and clinical associate professor of medicine at Texas Tech University Health Sciences Center, will review lanapanor's unique profile.
Jennife Robinson, preaidant of Spherix Global Insights, will present data on the challenges of treating hyperphosphatemia in patients with CRD on dalysis.
Douglas Paul, PharmD, Ph.D., vereaident and patient at MME, will doubes value perceptions across payers, patients and healthcare professionals.

A live webcast of the event will be available on the Ardelyx website via the Events and Presentations page under the investor relations section at https://ir.ardelya

About Ardelys, inc. About Ardelys, inc. Baour Ardelys, is toosaed on developing invokive first-in-class medicines to enhance the lives of patients with kidney and cardiovascular diseases. Ardelys is advancing tenapanor, a novel product candidate to control serum phosphorus in adult patients with CKD on dalysis, for which the company's NDA is currently under review by the FDA, with a PDUFA goal date of April 29, 2021. Ardelys is also advancing Broting To a patients with advances of the protection of the patients with kidney and cardiovascular diseases. In addition, Ardelys, received FDA approval of IBSRELA[®] (tenapanor) on September 12, 2019. Ardelys has established agreements with Kyowa Kith in Japan, Fourn Pharma in China and Knight Therapeutics in Canada for the development and commercialization of temppanor in the respective territories.

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