Meet the Onxeo Team during the 2017 J.P. Morgan Healthcare Conference

- *January 9-12, 2017 in San Francisco, CA*

In 2016, Onxeo has achieved a number of significant milestones, positioning the Company for success in 2017 and beyond. With a robust pipeline focused on addressing unmet medical needs in the treatment of rare cancers, Onxeo is poised to become the leader in orphan oncology.

Onxeo has a promising therapeutic pipeline with multiple assets in various stages of development, including:

- AsiDNA™ – First-in-class DNA Repair Signal-Interfering Technology with compelling preclinical and clinical data – **expected to enter a Phase I clinical trial as a systemic monotherapy in 2017**
- Livatag® – Novel, nanoparticle formulation of doxorubicin currently in a Phase III study in hepatocellular carcinoma (primary liver cancer) – **preliminary data expected mid-2017**
- Beleodaq® (belinostat) – Commercialized as a second-line treatment for Peripheral T-Cell Lymphoma (PTCL) – **recently announced promising preclinical results from studies evaluating Beleodaq® in combination with checkpoint inhibitors as a potential treatment option for certain tumor types**

For additional information or to schedule a meeting, please contact Lee Roth of The Ruth Group at lroth@theruthgroup.com or 646-536-7012.

About Onxeo

Onxeo is a biopharmaceutical company specializing in the development of innovative drugs for the treatment of orphan diseases, in particular in oncology, driven by high therapeutic demand in one of the fastest growing segments of the pharmaceutical industry. Onxeo's objective is to become a major international player in the field of rare cancers. Its growth strategy is founded on the development of innovative, effective, and safe drugs based on breakthrough technologies that can make a real difference in the treatment of orphan oncology diseases and considerably improve the quality of life of patients affected by rare and aggressive cancers. Onxeo's comprehensive portfolio features a broad orphan oncology pipeline, with four independent programs in various stages of clinical development, including Onxeo's first approved orphan oncology drug, Beleodaq®. The Company is headquartered in Paris, France and has approximately 50 employees. Onxeo is listed on Euronext in Paris, France (Ticker: ONXEO, ISIN Code: FR0010095596) and Nasdaq Copenhagen, Denmark (Ticker: ONXEO).

Onxeo's orphan oncology products are:

- **Livatag®** (Doxorubicin Transdrug™): Currently being evaluated in a Phase III trial (ReLive) in patients with hepatocellular carcinoma (primary liver cancer); Livatag is also under exploratory preclinical development to assess interest of its combination with other anti cancer agents.

- **Beleodaq®** (belinostat): FDA-approved in the US in 2014 under the agency's accelerated approval program as a second-line treatment for patients with peripheral T-cell lymphoma (PTCL) and currently marketed by Onxeo's partner in the US, Spectrum Pharmaceuticals; belinostat in combination with other cancer agents is currently in development in first-line treatment for patients with PTCL (BelCHOP) and a oral formulation is under development.
- **AsiDNA**: first-in-class siDNA (signal-interfering DNA) which has successfully undergone a proof-of-concept Phase I trial in metastatic melanoma via local administration
- **Validive®** (Clonidine Lauriad®): Positive final results from a Phase II trial in head and neck cancer patients with severe oral mucositis

In addition, Onxeo has successfully developed and registered two non-cancer products, which are currently being commercialized in the U.S. and Europe.

Learn more by visiting www.onxeo.com.

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