

ORYZON provides a corporate update in the context of COVID-19

MADRID, SPAIN and CAMBRIDGE, MA, UNITED STATES, April 17th, 2020 - Oryzon Genomics, S.A. (ISIN Code: ES0167733015, ORY), a public clinical-stage biopharmaceutical company leveraging epigenetics to develop therapies in diseases with strong unmet medical need, announces today that, since the emergence of the COVID-19 pandemic, the company has been implementing and executing contingency plans necessary to ensure the health and safety of its employees, partners and clinical trial participants. These plans have been formulated to enable business continuity, and allow the company to fulfill its responsibility to continue developing experimental drugs for serious illnesses with unmet medical needs.

Oryzon has implemented a set of measures aligned with these principles and has carefully followed the recommendations enacted by the authorities. Among them, it has implemented a comprehensive work-from-home policy. Those employees involved on critical research and development and laboratory activities follow reinforced measures for prevention and disinfection, including safety guidelines to reduce close interactions and limiting the number of people required on-site, and the implementation of flexible schedule of work- hours.

Oryzon has not canceled or postponed recruitment in ongoing clinical trials, but the operations of ongoing clinical trials have been adapted following the instructions of the regulatory agencies (FDA, EMA, AEMPS) in order to protect the health of participating patients, their families and healthcare professionals, and to preserve the integrity of the trial data as much as possible. Given the advanced age and vulnerability of patients participating in ALICE (elderly leukemia patients who are not eligible for conventional chemotherapy) and in REIMAGINE-AD, ETHERAL-EU and ETHERAL-US (elderly Alzheimer's patients), the company has reduced the requirement for hospital visits by these patients, being replaced when possible by remote monitoring. It is foreseeable that, in some cases, there is a risk that certain data may be incomplete due to visits or evaluations not carried out. The recruitment in CLEPSIDRA trial (Phase IIa clinical trial investigating iadademstat in small cell lung cancer patients) has been finalized.

Also, given the current general situation of lock-down, Oryzon has decided to postpone the activation of its Phase IIb trial in agitation-aggression in patients with borderline-personality disorder (PORTICO trial) for a few months. The company is continually evaluating the situation and will report promptly when additional information becomes available.

The company believes it has sufficient manufactured drug material to supply its ongoing clinical studies. Preparations for new clinical trials in CNS and cancer continue.

Finally, the Company has also implemented the necessary actions in order to guarantee the fulfillment of its normal financial obligations. Cash, cash equivalents and marketable securities totaled \$39.6 million as of December 31, 2019.

About Oryzon

Founded in 2000 in Barcelona, Spain, Oryzon (ISIN Code: ES0167733015) is a clinical stage biopharmaceutical company considered as the European champion in Epigenetics. Oryzon has one of the strongest portfolios in the field. Oryzon's LSD1 program has rendered two compounds, vafidemstat and iadademstat, in clinical trials. In addition, Oryzon has ongoing programs for developing inhibitors against other epigenetic targets. Oryzon has a strong technological platform for biomarker identification and performs biomarker and target validation for a variety of malignant and neurological diseases. Oryzon has offices in Spain and the United States. For more information, visit www.oryzon.com

FORWARD-LOOKING STATEMENTS

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