

## ADC-1013 clinical Phase I data to be presented at ASCO

**Lund, Sweden, May 15, 2019 – Alligator Bioscience (Nasdaq Stockholm: ATORX)**, today announces that Janssen Biotech, Inc. (Janssen) will present data from a Phase I clinical study of ADC-1013 (JNJ-7107) at the upcoming ASCO Annual meeting held in Chicago on May 31-June 4. The study results show that side effects were generally mild and transient.

The primary aim of the study was safety and to establish a recommended dose for Phase II. The study is still ongoing and has enrolled a total of 95 patients. The maximum tolerated dose has not yet been reached. Doses up to 1200 µg/kg i.v without premedication, and up to 2000 µg/kg with premedication have been shown to be safe and tolerable. Early evidence of clinical activity in this study included a partial response (PR) in a patient with renal cell cancer and 10 patients with prolonged stable disease (SD) ≥6 months.

“I am glad to see the promising safety and tolerability profile of ADC-1013 in patients with cancer. This together with the early signs of clinical activity, gives me great confidence in this CD40 drug candidate”, said Per Norlén, CEO of Alligator Bioscience.

A poster (#171) with the title **“A Phase 1 Study to Assess Safety, Pharmacokinetics (PK) and Pharmacodynamics (PD) of JNJ-64457107, a CD40 Agonistic Monoclonal Antibody, in Patients (pts) with Advanced Solid Tumors”** will be showcased at ASCO on June 1, 2019 from 8 - 11 a.m. in the session Developmental Immunotherapy and Tumor Immunobiology.

The licensing agreement with Janssen encompasses milestone payments up to a potential total value of USD 695 million. Alligator is also eligible to receive tiered royalties on worldwide net sales upon successful launch and commercialization of ADC-1013.

**For further information, please contact:**

Cecilia Hofvander, Director IR & Communications

Phone +46 46 540 82 06

E-mail: [cecilia.hofvander@alligatorbioscience.com](mailto:cecilia.hofvander@alligatorbioscience.com)

*This information is such information as Alligator Bioscience AB (publ) is obliged to make public pursuant to the EU Market Abuse Regulation. This information was submitted for publication, through the agency of the contact person set out above, at 11:00 p.m. CEST (5:00 p.m. ET) on May 15, 2019.*

**About ADC-1013 (JNJ-7107)**

ADC-1013 is a drug candidate intended for immunotherapy of different types of cancer. Results from a previous clinical Phase I first-in-human study performed by Alligator (International Journal of Cancer, <https://doi.org/10.1002/ijc.32141>) showed that ADC-1013 was generally well tolerated. Preclinical data have previously shown that the ADC-1013 antibody effectively activates T cells, mediated through binding to the co-stimulatory receptor CD40 on dendritic cells. The increased T cell activation enables the immune system to attack the cancer.

**About Alligator Bioscience**

Alligator Bioscience AB is a clinical-stage biotechnology company developing tumor-directed immuno-oncology antibody drugs. Alligator's growing pipeline includes five lead clinical and preclinical drug candidates: ADC-1013, ATOR-1015, ATOR-1017, ALG.APV-527 and ATOR-1144. Alligator's shares are listed on Nasdaq Stockholm (ATORX). The Company is headquartered in Lund, Sweden, and has approximately 55 employees. For more information, please visit [www.alligatorbioscience.com](http://www.alligatorbioscience.com).

ADC-1013 (JNJ-7107) is licensed to Janssen for global development and commercialization.