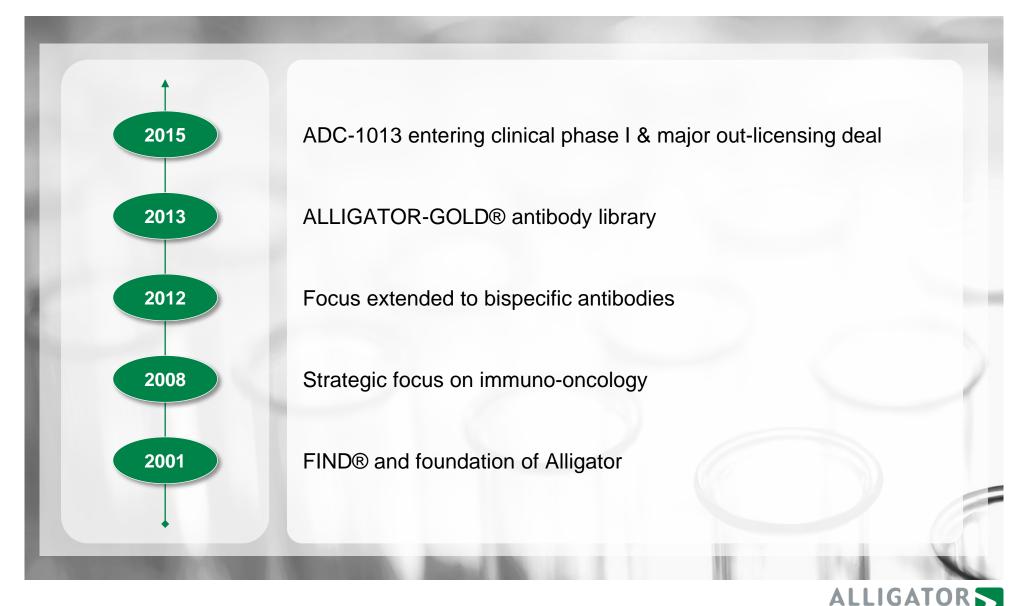


Alligator develops agonistic antibodies for tumor directed immunotherapy

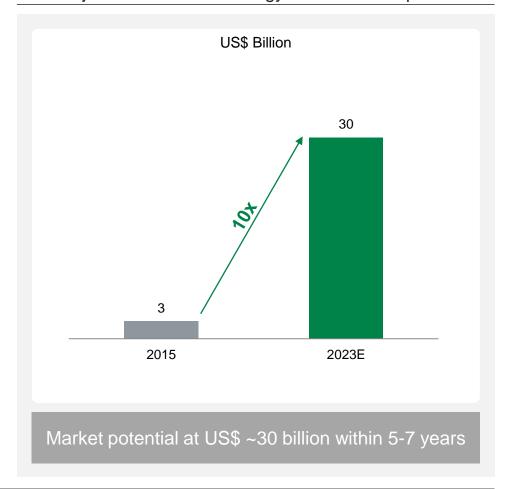


Rapid development within the field of immuno-oncology

Sales of existing immuno-oncology treatments

US\$ Million 2 634 566 21.5% 35.8% 1 369 960 706 1 126 42.8% 360 2011 2012 2013 2014 2015 ■ Yervoy® Opdivo® ■ Keytruda® Treatment costs at ~150,000 annually per patient

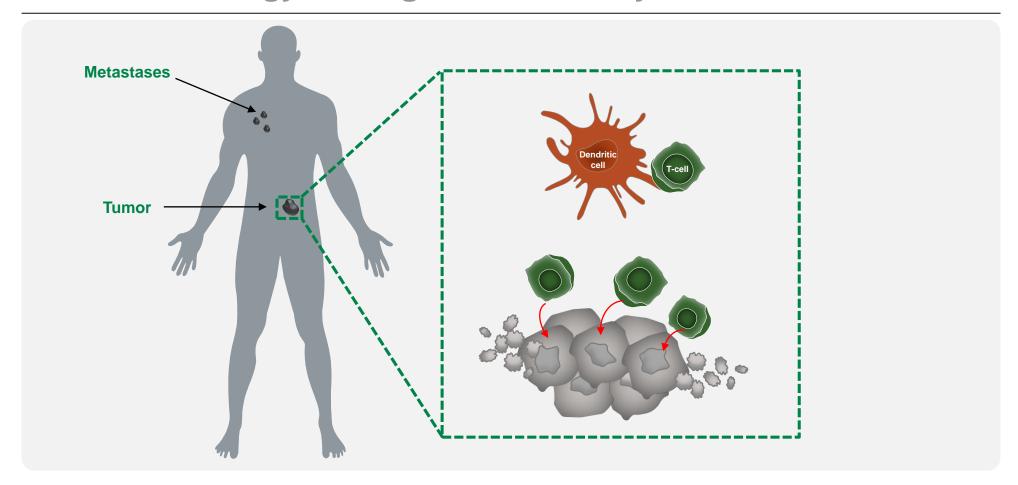
Projected immuno-oncology market development



The I/O market holds the largest upside potential within the global pharmaceutical market



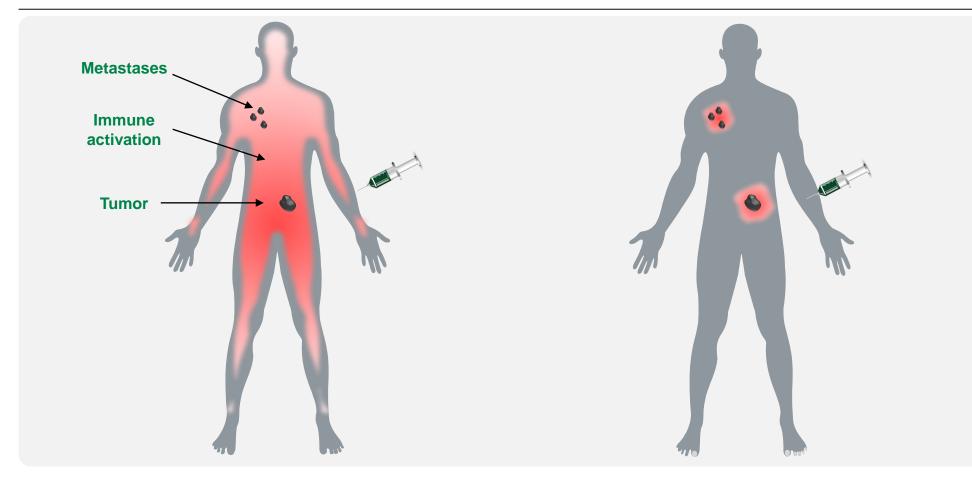
Immuno-oncology – using the immune system to kill cancer



Cancer immunotherapy switches the balance from immunosuppression to immune activation, resulting in immune-mediated tumor eradication



Tumor-directed immunotherapy



SYSTEMIC IMMUNO-ACTIVATION

General immune activation with severe toxicity

TUMOR-DIRECTED IMMUNO-ACTIVATION

Tumor-selective immune activation with less toxicity



Alligator drug development pipeline

RESEARCH	THE SEMIONE DEVELOR MENT	PHASE I	PHASE II
ADC-1013* (CD40)			
ATOR-1015 (OX40/CTLA-4)			
ATOR-1016 (TNFR-SF/TAA			
(TNFR-SF)			
(TNFR-SF/ND)			

TNFR-SF: Tumor Necrosis Factor Receptor-Superfamily

TAA: Tumor-Associated Antigen

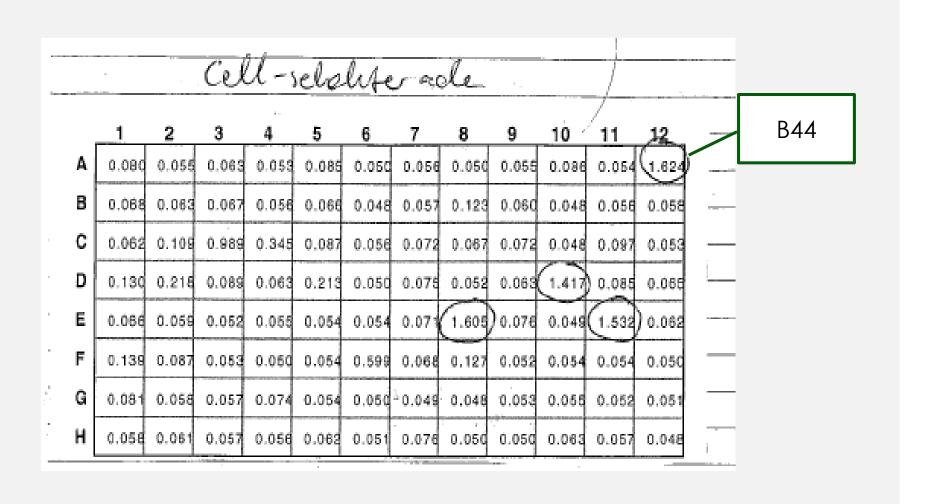
ND: Not Disclosed

*Partnered with Janssen Biotech Inc., developed as JNJ-64457107

All product candidates suitable for combination therapy with other I-O drugs, e.g. anti-PD-1 and anti-PD-L1



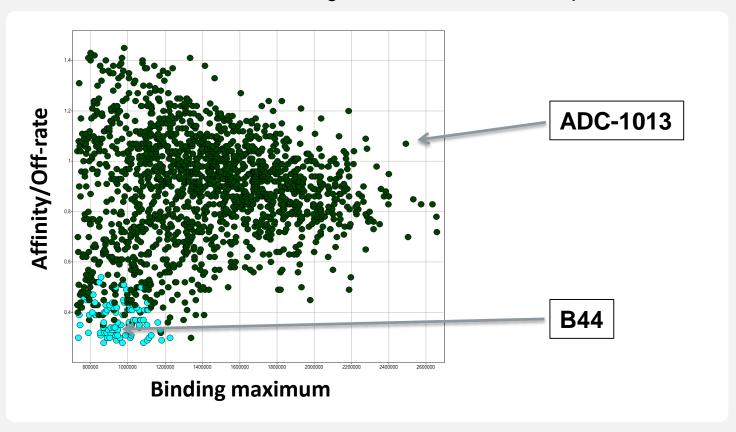
ADC-1013 – First experiment in 1999





Lead Optimization of CD40 Antibody

- Affinity maturation using FIND®. Affinity improved from 1 to 0.01 nM.
- Increased tumor retention through elevation of isoelectric point

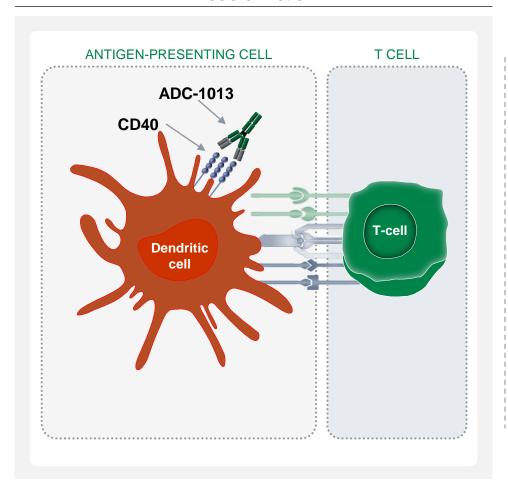


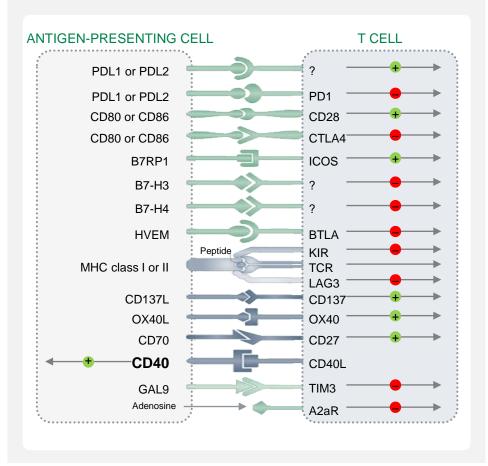


CD40 is a key immuno-oncology target

Mode of Action

Co-stimulating receptors





CD40 is a promising target for combination with T-cell activating antibodies such as PD-1



ADC-1013 clinical trial



- → 40 patients with solid tumors
- → 5 clinical sites in UK, DK and SE



Dosing & administration

- •FiH, first dose April 2015
- Dose escalation
- •Intra-tumoral & intravenous

Primary endpoint

Safety and tolerability

Secondary endpoints

- ■PK & PD
- Immunogenicity
- Clinical efficacy



Partnership with Janssen



Description of agreement

- Exclusive world-wide license
- Alligator continues as sponsor for the ongoing Phase I clinical trial
- Additional Phase I study initiated by Janssen
- All development costs covered by Janssen

Royalty / Milestone potential

- Up-front payment plus additional milestones of up to US\$ 700 million
- Tiered high single digit to low double digit royalties on all future sales



ADC-1013 out-licensing – success factors

- Strong scientific rationale, internal expertise and scientific collaborations
- Well positioned: potential for first in class and best in class
- Extensive pre-clinical data package
 - Data package included all data likely to be requested, including strong benchmark data
- Solid intellectual property
- High quality development, no shortcuts
- Timing and guts

