



Arming the patient's immune system to fight cancer

4Q and full year 2017 presentation

15 February 2018



Important notice and disclaimer

This report contains certain forward-looking statements based on uncertainty, since they relate to events and depend on circumstances that will occur in future and which, by their nature, will have an impact on the results of operations and the financial condition of Targovax. Such forward-looking statements reflect the current views of Targovax and are based on the information currently available to the company. Targovax cannot give any assurance as to the correctness of such statements.

There are a number of factors that could cause actual results and developments to differ materially from those expressed or implied in these forward-looking statements. These factors include, among other things, risks or uncertainties associated with the success of future clinical trials; risks relating to personal injury or death in connection with clinical trials or following commercialization of the company's products, and liability in connection therewith; risks relating to the company's freedom to operate (competitors patents) in respect of the products it develops; risks of non-approval of patents not yet granted and the company's ability to adequately protect its intellectual property and know-how; risks relating to obtaining regulatory approval and other regulatory risks relating to the development and future commercialization of the company's products; risks that research and development will not yield new products that achieve commercial success; risks relating to the company's ability to successfully commercialize and gain market acceptance for Targovax's products; risks relating to the future development of the pricing environment and/or regulations for pharmaceutical products; risks relating to the company's ability to secure additional financing in the future, which may not be available on favorable terms or at all; risks relating to currency fluctuations; risks relating to the company's ability to retain key personnel; and risks relating to the impact of competition.



Agenda

- 4Q 2017 Highlights
- Financials 4Q and full year 2017
- ONCOS program update
- TG program update
- 2017 review & 2018 outlook



Highlights from the fourth quarter 2017

Clinical data

- One-year survival rate, immune activation, and safety data for 2nd cohort in the **TG01 phase I/II trial** in resected pancreatic cancer
- The two combination trials with ONCOS-102 in melanoma and mesothelioma both passed their initial, planned safety reviews
- TG02 passed the initial safety review in colorectal cancer, and also reported immune activation data in the first treated patients

Patents

 Targovax was granted a composition-of-matter patent in the US for TG02

Corporate

- Patrick Vink was appointed as the new chairman of the Board
- TRVX included in the OSEBX index from 1 December 2017



Highlights from the fourth quarter 2017 – post-period

Clinical data

- ONCOS-102 generated immune activation in 4/4 patients treated in the checkpoint inhibitor refractory melanoma trial
- The safety lead-in cohort of ONCOS-102 in mesothelioma was completed without any concerns
- ONCOS-102 generated immune activation in the first treated patients in the mesothelioma trial

Corporate

Dr. Michael Bogenstätter was appointed Chief Business
 Officer of Targovax, and started in January 2018



Agenda

4Q 2017 Highlights

Financials - 4Q and full year 2017

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Targovax has a sound financial position, with cash to complete the planned clinical program well into 2019

Operations			
Cash end of Q4	NOK 262m	USD 32m	Des 31 st 2017
Net cash flow	NOK -24m	USD -3m	Total Q4
Annual run rate	NOK 110m	USD 14m	Last four quarters

The share	OSE: TRVX			
Market Cap	NOK 900m	USD ~110m	At share price NOK ~17	
Daily turnover	NOK 4m	USD 0.5m	Rolling 6 month avg.	
Analyst coverage	DNB, ABG Sundal Collier, Arctic, Redeye, Norske Aksjeanalyser, Edison			



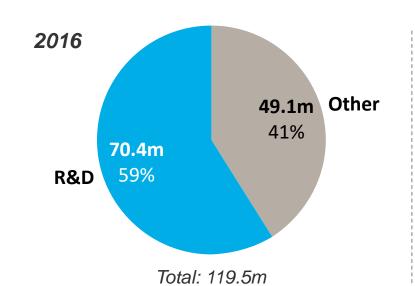
Total operating expenses for 2017 totaled 120m NOK, with a cash position of 262m NOK at end of Q4

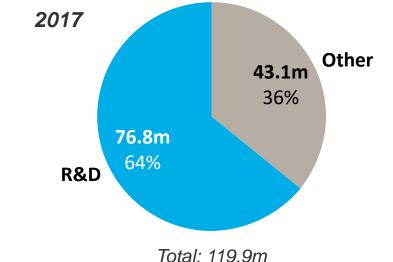
	4Q16	4Q17	2016	2017
Total revenue	0	0	0	0
External R&D expenses	-12	-12	-45	-46
Payroll and related expenses	-13	-13	-49	-48
Other operating expenses	-6	-7	-25	-26
Total operating expenses	-31	-32	-120	-120
Operating loss	-31	-32	-120	-120
Net financial items	-1	-0	-3	-2
Loss before income tax	-32	-33	-123	-122
Net change in cash	-21	-24	-2	90
Net cash EOP	172	262	172	262



Relative spend on R&D grew by 5% in 2017, driven by increased clinical activity

R&D spend (NOK million)



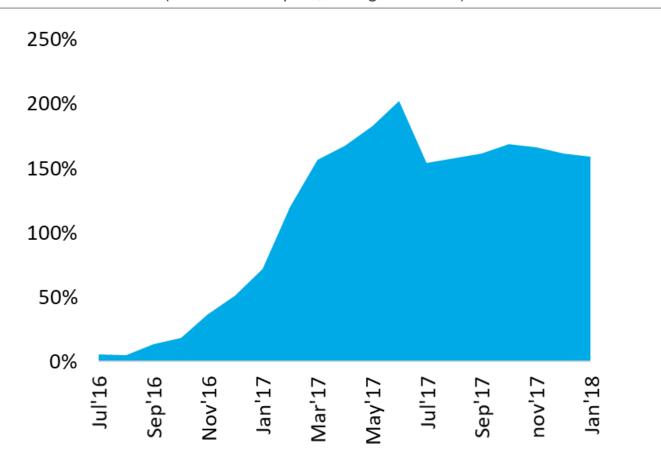


(Amounts in NOK thousands)	FY 2016		FY	FY 2017	
	Total	of which R&D	Total	of which R&D	
External R&D expenses	45.0	45.0	45.6	45.6	
Payroll and related expenses	49.2	24.4	48.3	30.0	
Other operating expenses	25.3	1.0	26.1	1.2	
Total	119.5	70.4	119.9	76.8	



Targovax is listed on the Oslo Stock Exchange, and included in the OSEBX index as of December 2017

TRVX share turnover (% of share capital, rolling 12 month)



- NOK ~900 m market cap
- NOK 4m NOK avg. daily turnover in last 6 months
- NOK 223m total turnover in 4Q
- 216k shares avg. daily volume in 4Q
- >4,100 owners
- 52.6m shares* (57.4 fully diluted)



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11

Targovax has two immuno-oncology programs in clinical development

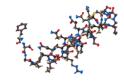
ONCOSOncolytic virus

- Genetically designed adenovirus
- Makes cancer antigens visible to immune system
- Induces T-cells specific to patients' tumor



TG
RAS neoantigen vaccine

- Cocktail of synthetic peptides
- Mimics cancer causing RAS neoantigens
- Induces T-cells specific to RAS mutations





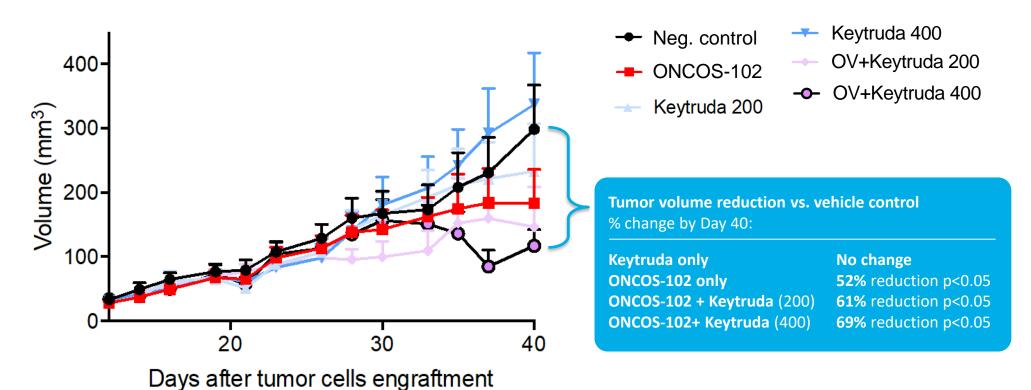
ONCOS clinical program overview

Combination with PD-1 Melanoma CPI in refractory patients Phase I Proof-of-concept 12 patients Memorial Sloan Kettering Combination with chemo Mesothelioma Randomized trial Phase I/II - controlled 30 patients Ultra-orphan indication Compassionate **Initial Phase I trial** use program Solid tumors **Finland** 7 indications 115 patients Ovarian / colorectal Collaboration with Ludwig & CRI Phase I/II - controlled Combination with Medimmune's 78 patients durvalumab (Imfinzi™) Prostate Partnered with SOTIO Phase I Combination with DC therapy 10 patients

13

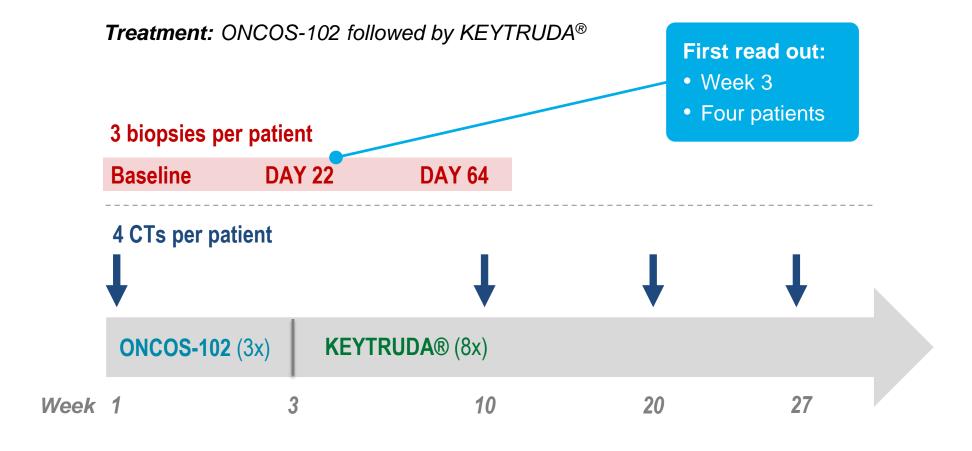
70% reduction in tumor volume with CPI combination in mouse melanoma model

Effect of ONCOS-102 and Keytruda in humanized mouse melanoma model, change in tumor volume





ONCOS-102 trial in CPI refractory melanoma





Melanoma trial: ONCOS-102 is well-tolerated, and induces immune activation in the first four patients

Safety Innate immune activation **Adaptive immune** activation

- ✓ First safety review completed with no safety concerns
- ✓ ONCOS-102 first time in melanoma treatment
- ✓ Systemic increase of several pro-inflammatory cytokines (4/4 patients)
- ✓ Increase in the relative level of cytotoxic CD8+ T cells (4/4 patients)
- ✓ Increase in PD-1 expression on CD8+ T cells (4/4 patients)

- **Clinical efficacy**
- First data expected in 2H 2018

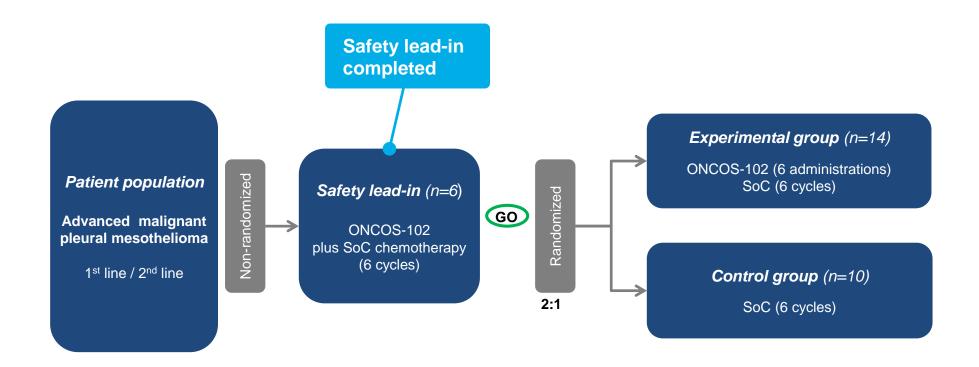


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17

ONCOS-102 in Mesothelioma – Phase I/II study design





Mesothelioma trial: ONCOS-102 passed the safety lead-in, and the randomized phase II part now open for recruitment

Safety

Innate immune activation

Adaptive immune activation

Clinical efficacy

- ▼ Phase Ib safety lead-in cohort of six patients completed with no safety concerns
- ✓ ONCOS-102 first time in combination with chemotherapy
- ✓ Systemic increase of several pro-inflammatory cytokines (3/3 patients with available samples)

✓ Increase in infiltration of cytotoxic CD8+ T cells into lesions (2/2 patients with pre- and post-biopsies available)

First data expected in 1H 2018



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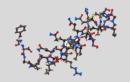
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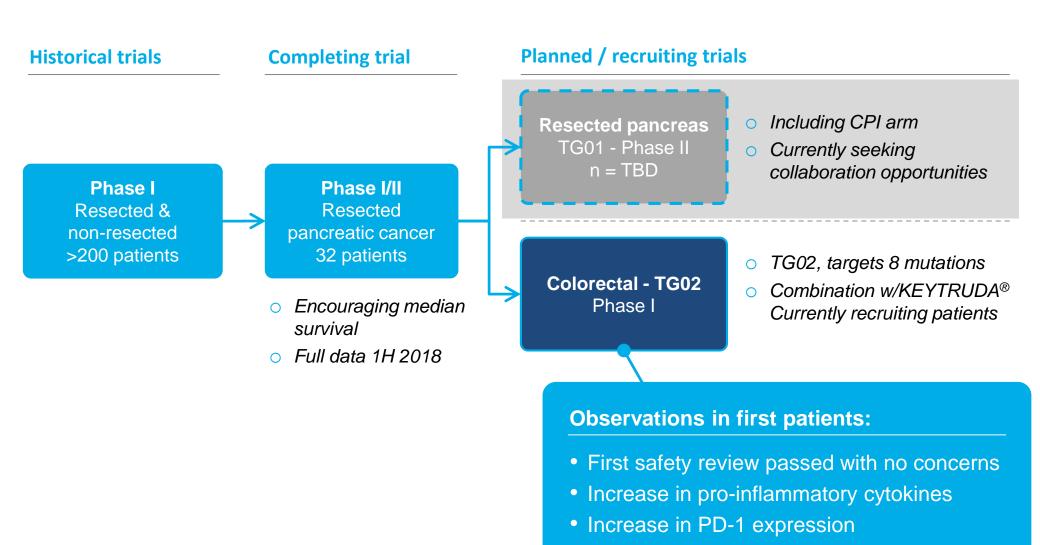
TG RAS neoantigen vaccine

- Cocktail of synthetic peptides
- Mimics cancer causing RAS neoantigens
- Induces T-cells specific to RAS mutations

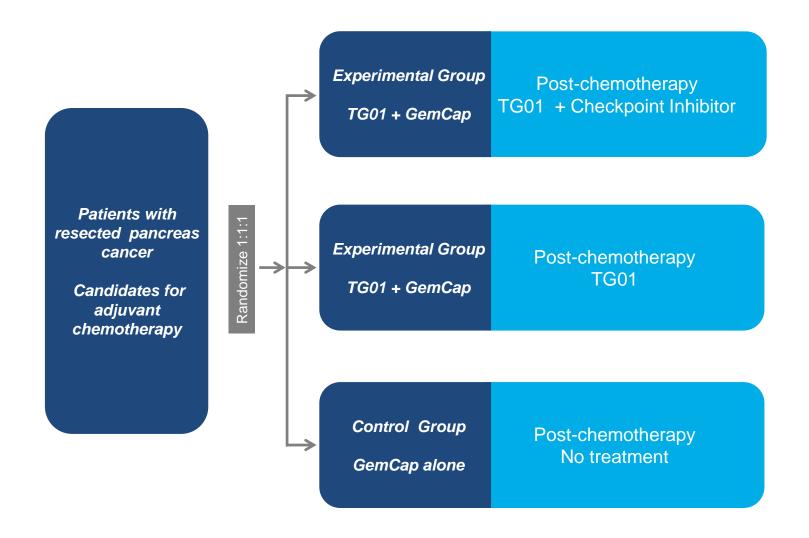




Clinical development overview for the TG program



Planned TG01 phase II in resected pancreatic cancer





Resected pancreatic cancer is the lead indication, but all RAS mutated cancers are potential TG targets



2 Colorectal cancer

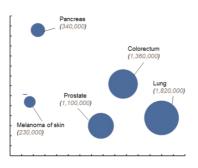












- TG01 lead indication
- Completing phase I/II
- Planning phase II
- 40.000 patients per year

- TG02 lead indication
- Phase I trial recruiting
- 50% RAS mutated
- Up to 500.000 patients per year

- TG02 potential future indication
- 30% RAS mutated
- Up to 500.000 patients per year

- TG02 + TG03 ultimate long-term potential
- 30% of all cancers
- Up to 30% of all cancer patients



Source: Global data, Riva et al. Plos One 2017

Estimated total addressable patient number with RAS mutations in US, EU and China

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Highlights from 2017

Clinical data

- TG01 showed encouraging survival and immune activation data in resected pancreatic cancer, which was presented at ASCO 2017
- ONCOS-102 passed the initial safety review in the two first combination trials in melanoma (CPI) and mesothelioma (chemo)

Patents

 The TG IP position was significantly strengthened by the grant of method-of-use and composition-of-matter patents in the US

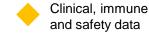
Corporate

- Oslo Stock Exchange and included in the OSEBX index
- Targovax raised 200m in a private placement, securing financing of the ongoing clinical program well into 2019
- The management team was strengthened by the addition of Dr. Erik D Wiklund as CFO and Dr. Michael Bogenstätter as CBO
- Patrick Vink was appointed as the new chairman of the Board



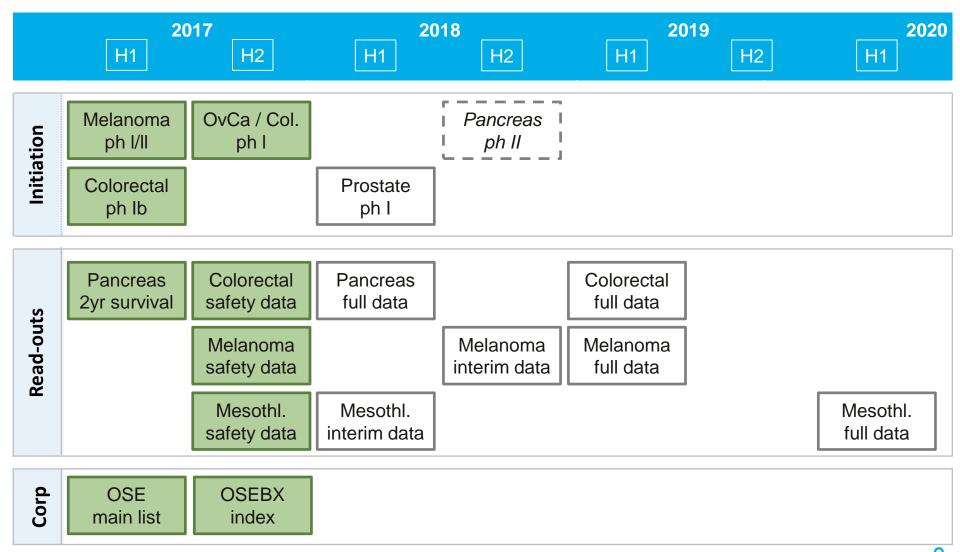
Overview of Targovax' full clinical program







News flow – Multiple near-term value inflection points



Arming the patient's immune system to fight cancer

Broad clinical program



- ✓ Six shots on goal
- ✓ Several upcoming data points

ONCOS



- ✓ Demonstrated ability to increase T-cell count
- ✓ Potential to make CPIs effective in more indications

TG



- Unique approach for targeting RAS mutations
- ✓ Potential to benefit up to 1/3 of all cancer patients

