

### 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 nonapproval 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' 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.



### TARGOVAX AT A GLANCE

Lead product ONCOS-102 directed to the \$20+ billion market for checkpoint inhibitors

Best-in-class results of ONCOS-102 & checkpoint combo in melanoma

targovax

**Encouraging results of ONCOS-102 & chemotherapy combo in mesothelioma** 

Powerful immune activation supporting IO-combinations

Strong patent position & pipeline with multiple additional value-creating opportunities



## SOLID CLINICAL AND PRECLINICAL PIPELINE

Product candidate	Preclinical	Phase 1	Phase 2	Collaborator	Next expected event
	Mesothelioma Combination w/ pemetrexed/cisplatin			MERCK	<b>1H 2021</b> Survival updates Define next steps
ONCOS-102	<b>Melanoma</b> Combination w/Keytruda				<b>1H 2021</b> Define next steps
	Colorectal cancer Combination w/Imfinzi			AstraZeneca CANCER RESEARCH INSTITUTE	Update by collaborator
	Prostate cancer Combination w/DCvac			Sotio	Update by collaborator
ONCOS-200 series	Next Gen viruses			leidos	Updates at conferences
Novel mutRAS concepts				VALO THERAPEUTICS  OBLIQUE THERAPEUTICS	



### LARGE MARKET POTENTIAL FOR IMMUNE ACTIVATORS

CPIs are revolutionizing cancer therapy...

...but only a minority of patients respond...

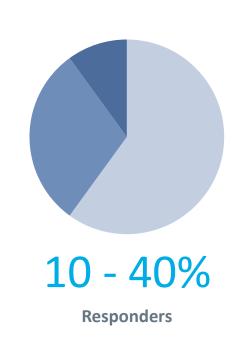
...leading to a high medical need for immune activators

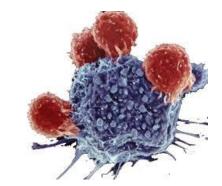
\$20+ bn

Global Cl I Illancet

44%

Patients eligible for CPI<sup>2</sup>:





## ONCOS-102 DRIVES A STRONG IMMUNE RESPONSE TRIGGERING ANTI-TUMOR IMMUNITY

Virus injection Immune activation **Anti-tumor** T-cell generation immunity T-cell tumor infiltration Intra-tumoral or intra-Oncolysis of tumor cells Antigen processing stimulated by GM-CSF peritoneal injection Inflammatory response by Tumor cell killing Tumor cell infection TLR-9 and other pathways T-cell activation in Synergy with lymph nodes Armed with immunecheckpoint inhibitors Tumor antigen release activating transgene

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ONCOS-102	<b>Melanoma</b> Combination w/Keytruda				



### ONCOS-102 IN ANTI-PD1 REFRACTORY MELANOMA

#### **Patient population**

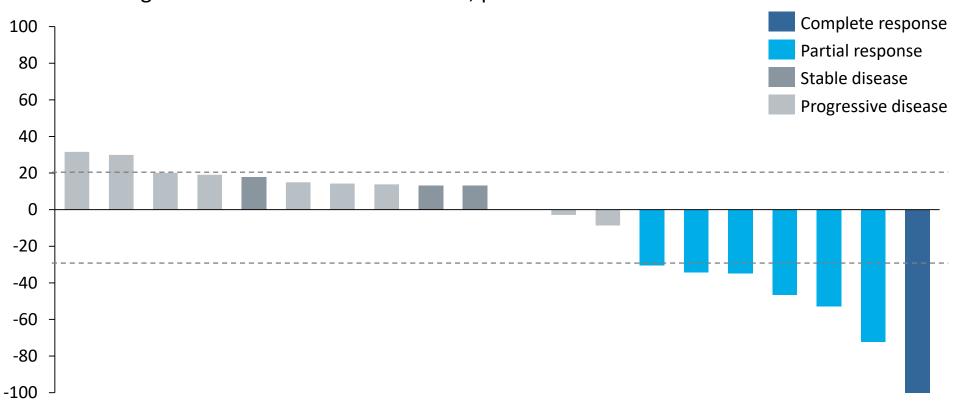
- Advanced, unresectable melanoma
- O Disease **progression** despite prior treatment with anti-PD1
- O Poor prognosis, with **few treatment alternatives**
- 20 patients, 11 stage III and 9 stage IV

#### **Treatment regime**

- Part 1: 3 ONCOS-102 injections followed by 5 months of Keytruda
- Part 2: 12 ONCOS-102 injections; priming and concomitant use
  - More advanced patients in part 2

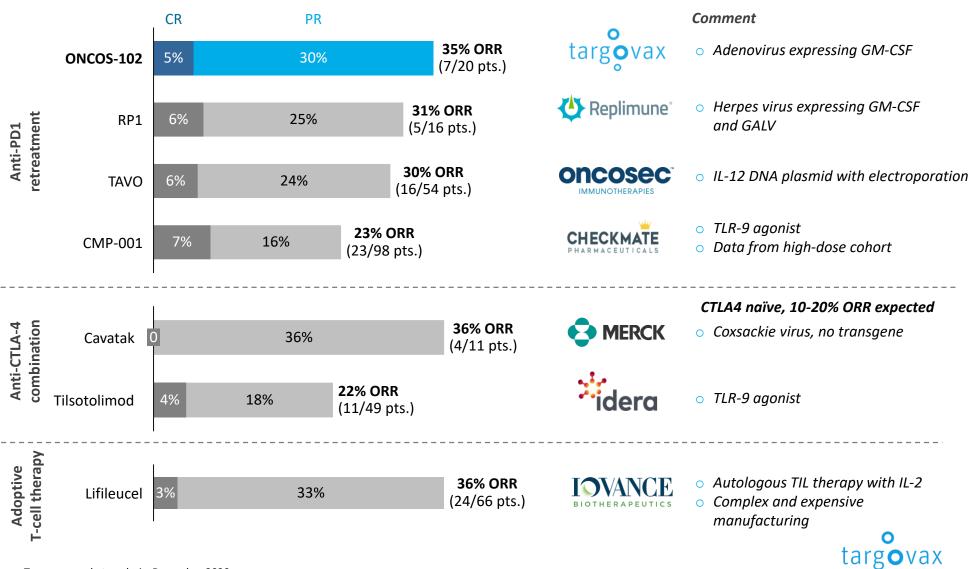
### **BEST-IN-CLASS RESPONSE RATE WITH ORR OF 35%**

Best change in tumor burden from baseline, percent



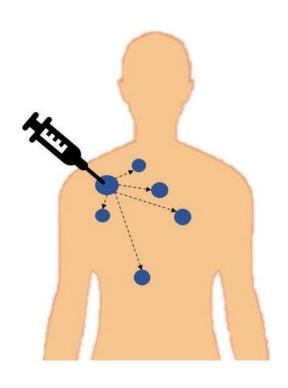


## ONCOS-102 EFFICACY IS COMPETITIVE TO LEADING DRUG CANDIDATES IN ANTI-PD1 REFRACTORY MELANOMA



## MULTIPLE EXAMPLES OF SYSTEMIC (ABSCOPAL) EFFECT

TWO PATIENTS WHERE A NON-INJECTED LESION COMPLETELY DISAPPEARED



- Threshold: tumor reduction of 30%¹ or more in a lesion
- Examples of complete remission of noninjected lesion seen in two patients
- Findings are based on early data assessment, systemic effects will be further assessed



## SUMMARY: EXCELLENT OUTCOME SUPPORT CONTINUED DEVELOPMENT IN ANTI-PD1 REFRACTORY MELANOMA



#### **Excellent safety profile confirmed**

ONCOS-102 and Keytruda combination is well-tolerated



#### **Excellent clinical outcome**

- Confirmed ONCOS-102 ability to reactivate CPI refractory tumors
- 35% ORR: Tumor responses were observed in 7 out of 20 evaluable patients
- Systemic effect: Tumor regression in non-injected lesions observed in multiple patients, including two lesions that regressed completely



#### **Next steps**

- Analyze more immunological data
- Planning for a confirmatory melanoma trial in combination with anti-PD1 checkpoint inhibitor

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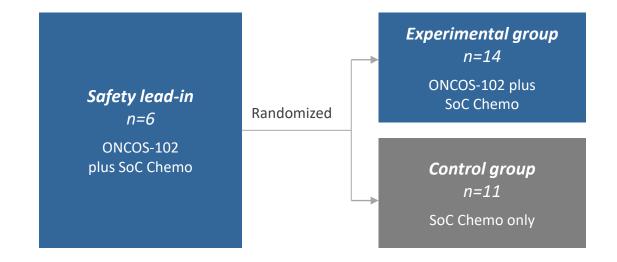


#### ADVANCED MALIGNANT PLEURAL MESOTHELIOMA

## PHASE I/II TRIAL IN COMBINATION WITH CHEMO

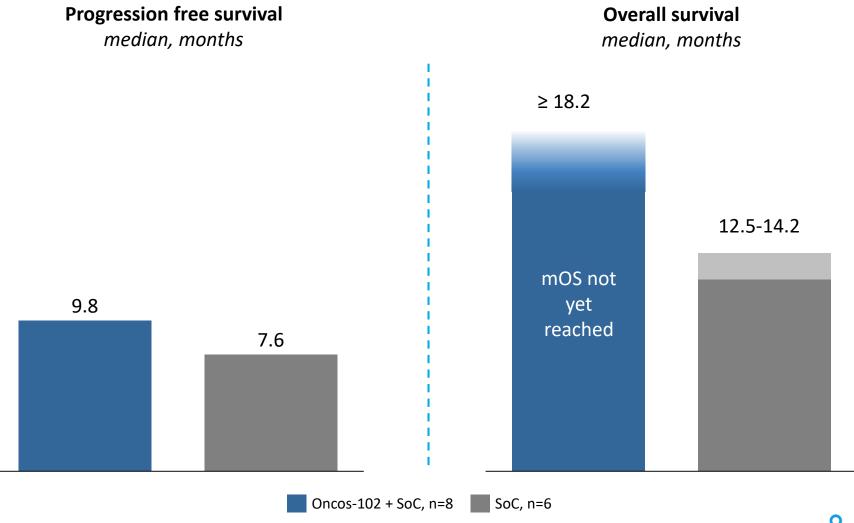
#### **Trial design**

- First and second (or later) line
- Standard of Care (SoC)
   Chemo: Pemetrexed and cisplatin, 6 cycles
- ONCOS-102: 6 intra-tumoral injections



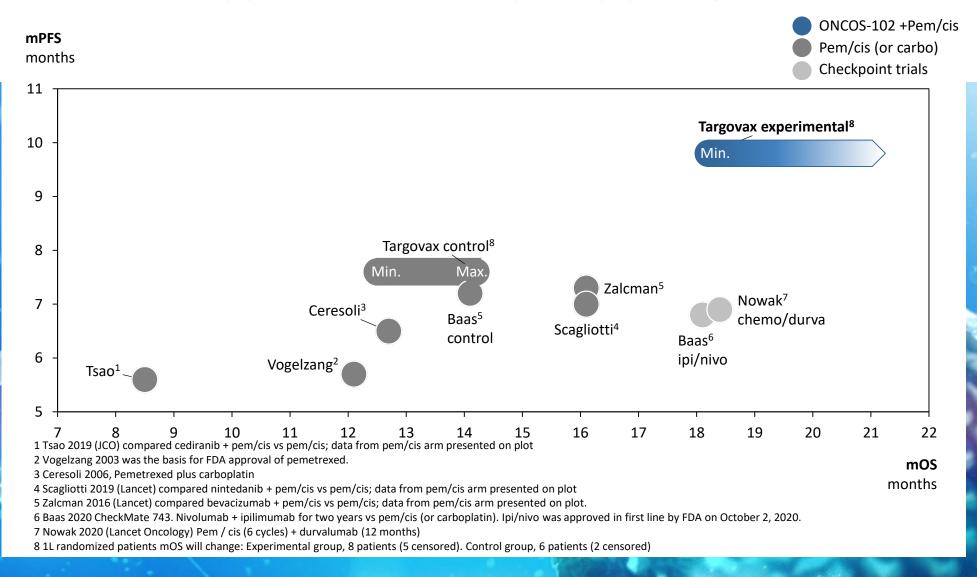
#### ONCOS-102 MESOTHELIOMA PHASE 1/2 COMBINATION WITH SoC CHEMO

## **ENCOURAGING CLINICAL OUTCOMES IN 1ST LINE**





## FIRST LINE DATA ARE MATURING AND ALREADY COMPETITIVE - MOS WILL BE 18.2 MONTHS OR MORE



## FUNDED WELL BEYOND IMPORTANT VALUE INFLECTION POINTS

#### The company

Cash at end of 3Q

**78 / 8** 

NOK million USD million

Raised NOK 75m in Oct 2020

Net cash flow - total 3Q

-24/-2.5

NOK million USD million

Market cap

800 / 90

NOK million USD million

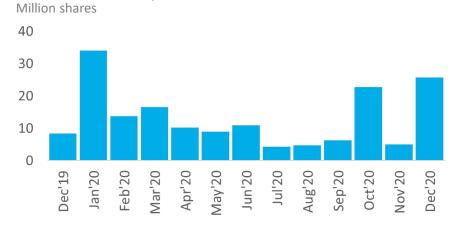
Analyst coverage

Carnegie, DNB, H.C. Wainwright

#### **Share liquidity**

+200% of shares traded last 12 months

Share turnover per month<sup>1</sup>



Daily value traded Average last 12 months

4.4 / 0.52

NOK million USD million



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## Save the date

# Capital Markets Day 18 February 2021

