



We want to save patients with severe cancer and autoimmune diseases

Göran Forsberg, CEO
May 2022

NASDAQ STOCKHOLM MAIN LIST (CANTA.ST)

Safe Harbor Statement



Statements in the Investor Presentation, including those regarding the possible or assumed future or other performance of the Company or its industry or other trend projections, constitute forward-looking statements. By their nature, forward-looking statements involve known and unknown risks, uncertainties, assumptions and other factors as they relate to events and depend on circumstances that will or may occur in the future, whether or not outside the control of the Company. No assurance is given that such forward-looking statements will prove to be correct. Prospective investors should not place undue reliance on forward-looking statements. They speak only as at the date of this Investor Presentation and the Company undertakes no obligation to update these forward-looking statements. Past performance does not guarantee or predict future performance. Moreover, the Company undertakes no obligation to review, update or confirm expectations or estimates or to release any revisions to any forward-looking statements to reflect events that occur or circumstances that arise in relation to the content of the Investor Presentation.

Cantargia – The IL1RAP company



FIRST IN CLASS ANTIBODY THERAPIES AGAINST NOVEL IL1RAP TARGET

- Five Phase I/II trials, with positive interim data in pancreatic cancer and NSCLC
- Differentiated by broad MOA and unique binding properties
- Synergistic with established therapies



PLATFORM WITH BROAD POTENTIAL TO ADDRESS HIGH UNMET NEEDS

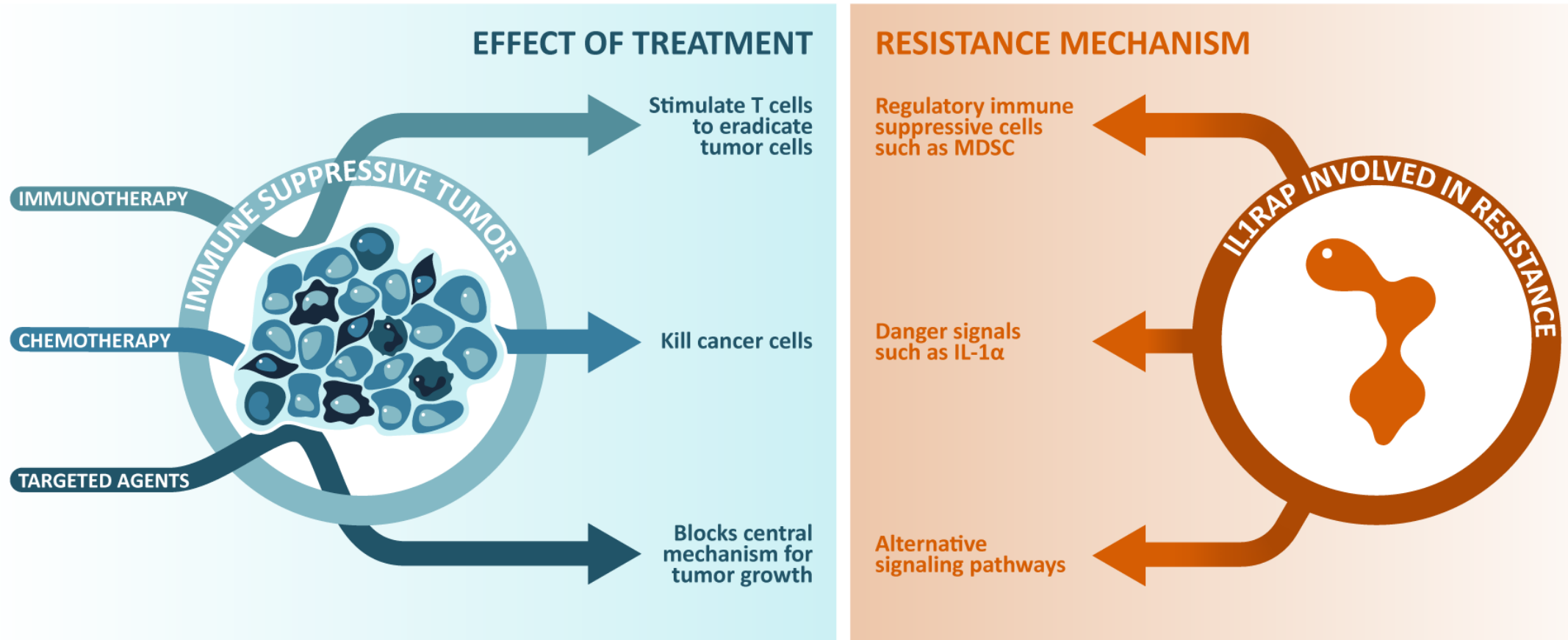
- Target IL1RAP found on most solid tumor forms and leukemias
- IL1RAP signalling key in large number of inflammatory diseases beyond oncology
- Robust patent portfolio on antibody target in oncology (to 2032) and lead asset (to 2035)



INGREDIENTS FOR SUCCESS

- Solid cash position (559 MSEK, 59 MUSD end Q4 2021)
- Clear development plan with multiple upcoming catalysts
- Strong management team with experience in bringing products through development to market

Cantargia - strategy to improve current cancer therapies



IL1RAP: A NOVEL TARGET WITH SEVERAL OPPORTUNITIES

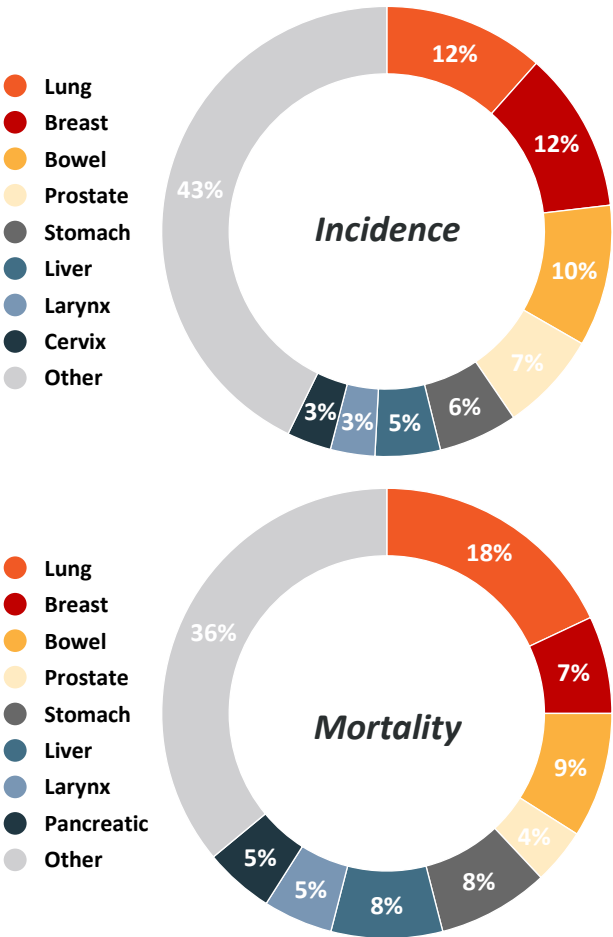
Cantargia – Save lives and create value through IL1RAP

Project	Disease	Type of treatment	Discovery phase	Preclinical phase	Clinical phase I	Clinical phase II	Clinical phase III	Commercial phase
CAN04 Nadunolimab	PDAC	1 st line	<i>Gemcitabine/nab-paclitaxel</i>					
			<i>FOLFIRINOX</i>					
	NSCLC	1 st line	<i>Cisplatin/gemcitabine</i>					
		2 nd /3 rd line	<i>Docetaxel</i>					
	Non-squamous NSCLC	1 st /2 nd line	<i>Carboplatin/pemetrexed</i>					
		1 st line	<i>Pembro/carboplatin/pemetrexed</i>					
	TNBC	1 st /2 nd line	<i>Carboplatin/gemcitabine</i>					
	Biliary tract cancer	1 st line	<i>Cisplatin/gemcitabine</i>					
CAN10	Colon cancer	3 rd line	<i>FOLFOX</i>					
			<i>Pembro</i>					
CANxx	Solid tumors	ICI combo						
CAN10	Myocarditis; Systemic sclerosis							
CANxx	New opportunities within IL1RAP platform							

- Potentially more effective treatment against novel target in clinically validated pathway
- First in class platform technology against novel target
- Well-financed to build a broad, diversified pipeline
- Right team and clear plan to position our projects and maximize value

PDAC – pancreatic cancer; NSCLC – non-small cell lung cancer; TNBC – triple negative breast cancer; ICI – immune checkpoint inhibitor; Pembro – pembrolizumab

Cantargia addresses NSCLC & PDAC



Non-Small Cell Lung Cancer



2.1m

Annual global incidences

12%

Fraction of cancer incidence

1.8m

Annual global mortalities

18%


Fraction of cancer mortality

19%

Five-year survival rate

Treatment: Surgery, Radiation, Chemotherapy, Immunotherapy

Pancreatic cancer



0.5m

Annual global incidences

3%

Fraction of cancer incidence

0.4m

Annual global mortalities

5%

Fraction of cancer mortality

9%

Five-year survival rate

Treatment: Chemotherapy, Surgery, Radiation

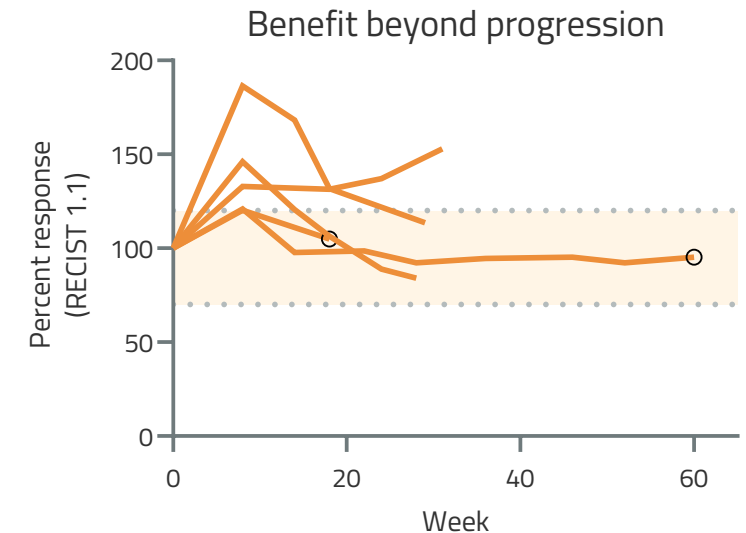
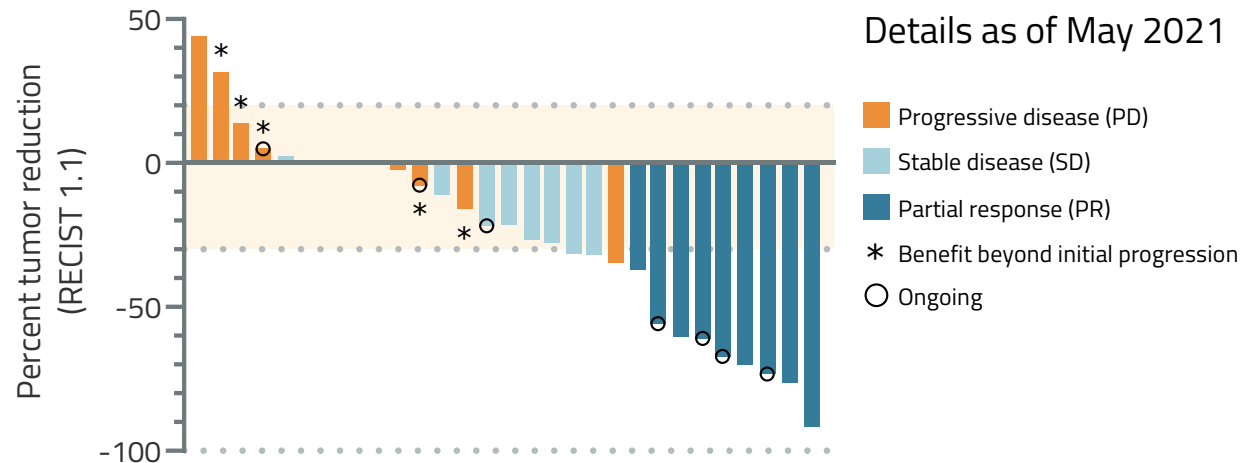
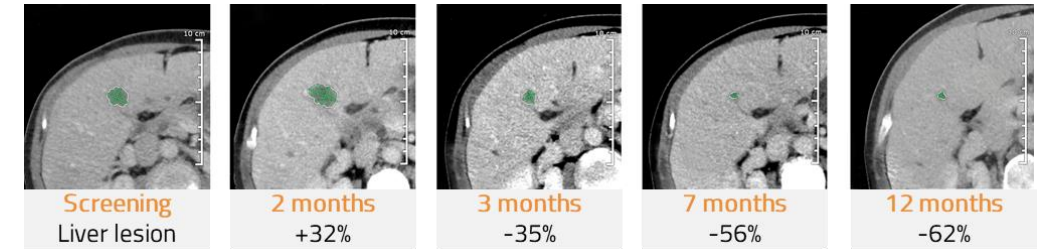
SIGNIFICANT UNMET NEEDS IN LUNG AND PANCREATIC CANCER

Positive interim data in pancreatic cancer

Nadunolimab combination with Gem/Abraxane in 1st line (Dec 2021), n=33:

- 27% response rate with durable responses, two patients still on treatment
- Pseudoprogression-like response in 5 (15%) patients predict long PFS
- Promising PFS (7.2 mo) and OS (12.7 mo, 64 % events)

UPDATE: Results on 73 pts to be presented at ASCO 2022



PFS AND OS LONGER THAN EXPECTED GIVEN HISTORICAL CONTROL

Advancing PDAC development to phase 2/3

PanCAN's Precision PromiseSM adaptive clinical trial platform designed together with the FDA

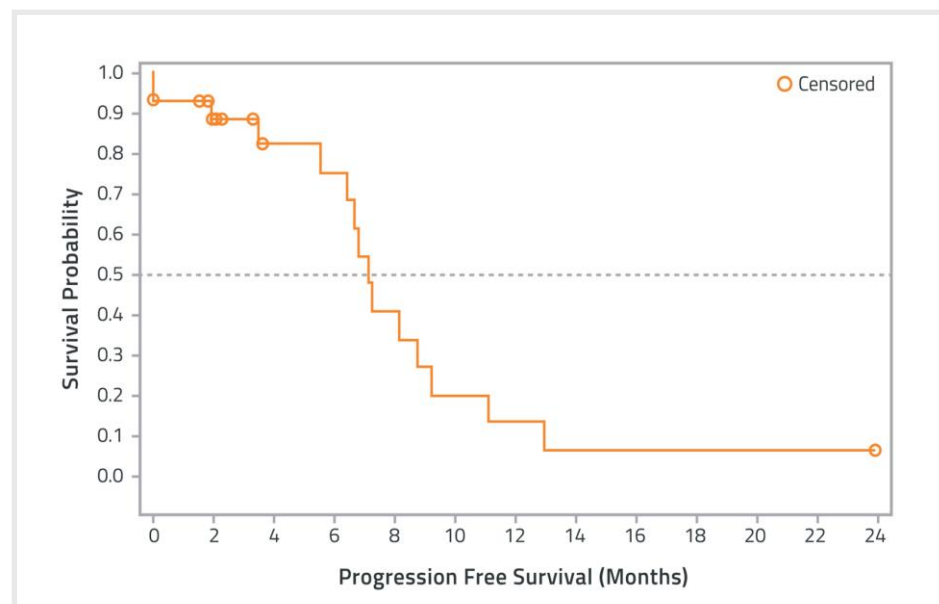
Nadunolimab selected for inclusion

- 15 leading US clinical centers - additional sites planned
- Patients randomized to receive nadunolimab with gemcitabine and nab-paclitaxel, or chemotherapy alone
- Bayesian design, successful completion of a 100-patient adaptively randomized Stage 1 may be followed by a 75-patient fixed-randomized Stage 2
- Trial results for nadunolimab arm expected 2027 or earlier
- Additional meetings with regulatory authorities to take place; pre-IND planned for submission to the US FDA in Q2 2022
- Cantargia funds nadunolimab arm and responsible for drug supply

ADVANCING WITH PANCAN FURTHER VALIDATES NADUNOLIMAB IN PDAC

Combination strategy in NSCLC – Promising efficacy

	Total NSCLC (27 pts)	Historical control ^{1,2}	Non-squamous NSCLC (15 pts)	Historical control ³	Squamous NSCLC (11 pts)	Historical control ⁴
ORR	48%	22-28%	53%	19%	36%	38%
PFS	7.2 mo	5.1 mo	NR	-	NR	-
Ongoing treatment	11 pts (41%)	-	6 pts (40%)	-	5 pts (45%)	-



Nadunolimab combination with Gem/Cis in 1st line:

- 13 of 27 evaluable patients with non-squamous non-small cell lung cancer (NSCLC) showed objective response including 1 complete response (ORR 48% vs historical control data of 22-28%)
- No major side effects observed except those from chemotherapy or nadunolimab alone. *Neutropenia frequency higher than expected from chemo (but can be treated with dose reductions or G-CSF)*
- Trial expanding - 40 additional patients with non-squamous NSCLC

STRONG INTERIM RESULTS, UPDATE AT ASCO 2022

¹ Schiller et al, N Engl J Med 2002

³ Gandhi et al, N Engl J Med 2018

² Scagliotti et al, J Clin Oncol 2008

⁴ Paz-Ares et al, N Engl J Med 2018

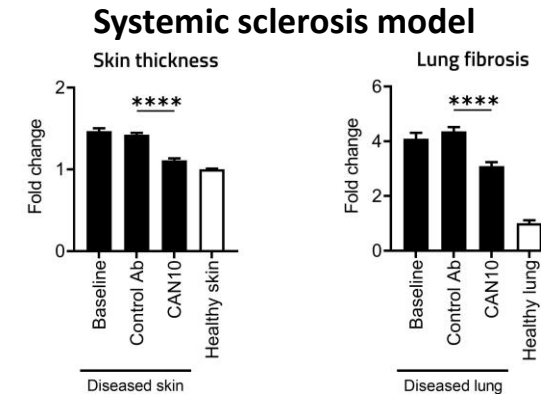
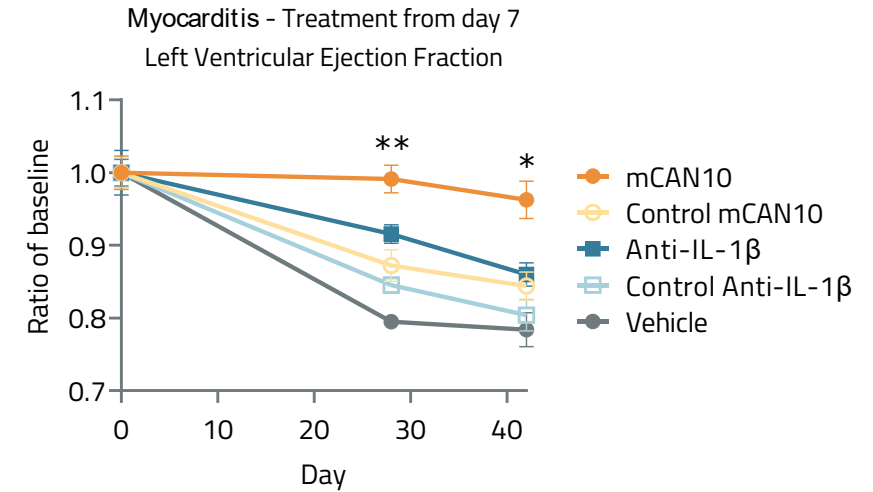
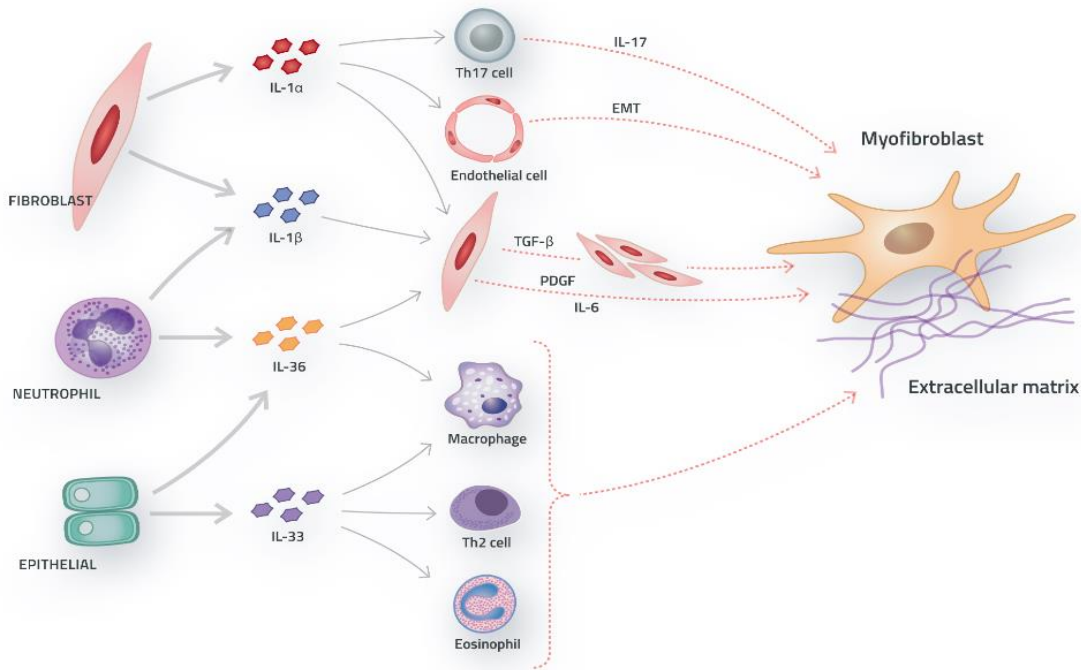
CIRIFOUR – Broadening into IO combinations

- First arm (15 pts): Combination with pembrolizumab in patients no longer responding to PD-(L)1 therapy (NSCLC, HNSCC, malignant melanoma and bladder cancer)
- Very good safety, only one treatment related grade 3 AE (febrile neutropenia); 5 pts on treatment (2 >31 weeks; 2 >49 weeks); data update (incl. efficacy) at ASCO 2022
- Second arm (up to 24 pat): Combination with 1st line pembrolizumab and carboplatin/pemetrexed in non-squamous NSCLC starting Q2 2022
- Primary endpoint safety, secondary endpoints include biomarkers and efficacy



**TRIAL DESIGNED TO ADVANCE NADUNOLIMAB OUTSIDE CHEMOTHERAPY COMBINATIONS
IMPORTANT STEP FOR COMBINATION WITH IO AND CHEMOTHERAPY**

CAN10 – Unique properties in preclinical disease models



CLINICAL TRIAL STRATEGY UNDER DESIGN TO VALIDATE PRECLINICAL RESULTS

Several upcoming value inflection points

Newsflow over next 6-9 months

Nadunolimab (CAN04)

- ASCO: New results for PDAC, NSCLC and Keytruda combination (CANFOUR, CIRIFOUR)
- Phase 2/3 Precision Promise (PDAC)
- New preclinical and translational results
- New clinical trials (Interim results, safety)
 - CAPAFOUR PDAC FOLFIRINOX
 - CESTAFOUR Basket trial (NSCLC, CRC, BTC)
 - TRIFOUR TNBC

CAN10

- Preclinical progress
- Development milestones
- ...and initiation of clinical trial early 2023



SIGNIFICANT DATA TO SECURE NEWSFLOW

Cantargia – The IL1RAP company



FIRST IN CLASS ANTIBODY THERAPIES AGAINST NOVEL IL1RAP TARGET

- Five Phase I/II trials, with positive interim data in pancreatic cancer and NSCLC
- Differentiated by broad MOA and unique binding properties
- Synergistic with established therapies



PLATFORM WITH BROAD POTENTIAL TO ADDRESS HIGH UNMET NEEDS

- Target IL1RAP found on most solid tumor forms and leukemias
- IL1RAP signalling key in large number of inflammatory diseases beyond oncology
- Robust patent portfolio on antibody target in oncology (to 2032) and lead asset (to 2035)



INGREDIENTS FOR SUCCESS

- Solid cash position (559 MSEK, 59 MUSD end Q4 2021)
- Clear development plan with multiple upcoming catalysts
- Strong management team with experience in bringing products through development to market