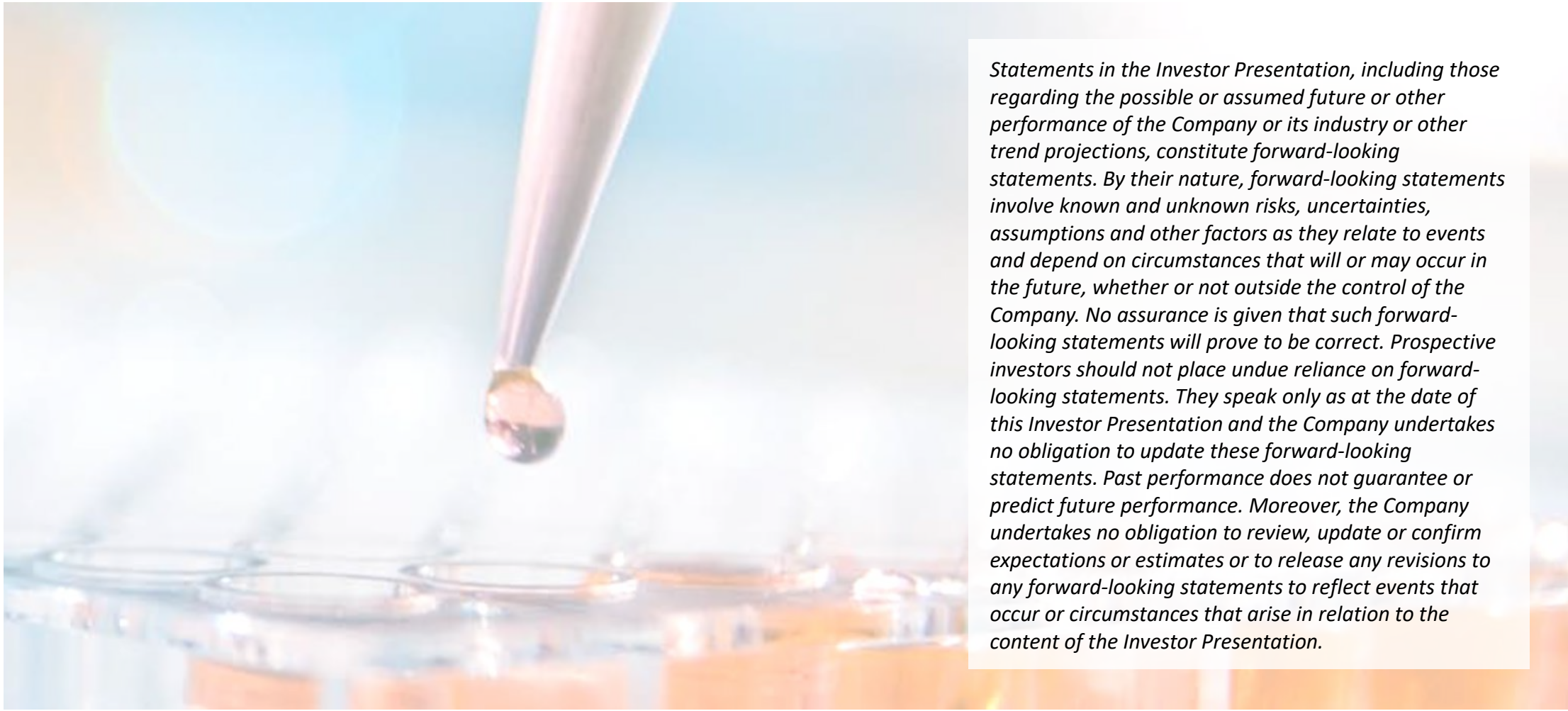
We want to save patients with severe cancer and autoimmune diseases

Clinical investigations with our lead antibody CAN04 to our proprietary target

Göran Forsberg, CEO

May 2021

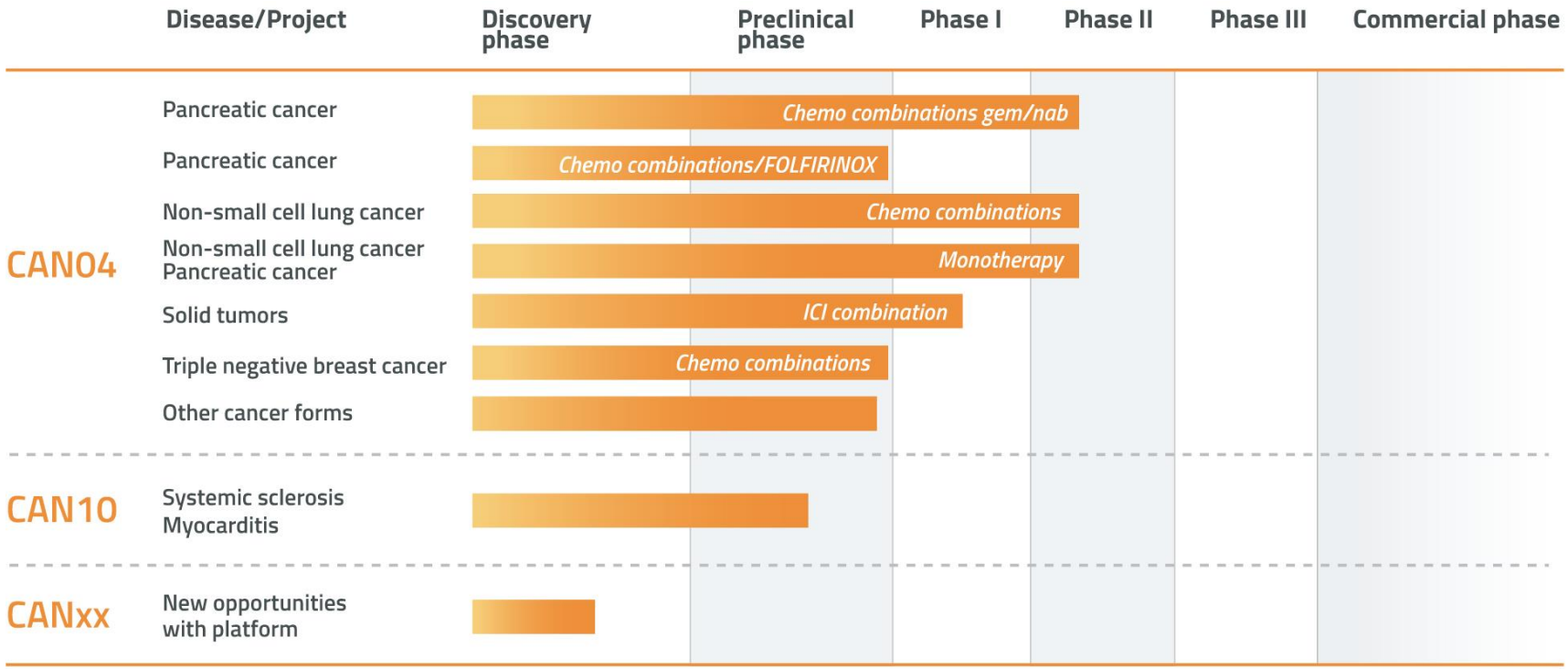
Safe Harbour Statement



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.

I. INTRODUCTION

Cantargia – Opportunity to save lives and create value



- 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

Cantargia highlights

UNIQUE IMMUNOTHERAPY ANTIBODY CAN04 IN PHASE IIA CLINICAL DEVELOPMENT

- First in class antibody with broader MOA than competitors
- Positive clinical interim data – PFS, durable responses and pseudoprogression

VISION OF BECOMING AN IMPORTANT PART IN FUTURE CANCER TREATMENTS

- Combination strategy based on synergies with established therapies

PLATFORM WITH MANY POTENTIAL THERAPEUTIC AREAS

- Target IL1RAP found on most solid tumor forms and leukemia
- IL1RAP signalling (IL-1, IL-33 and IL-36) in large number of diseases

HIGHLY RELEVANT RESEARCH WITHIN CLINICALLY VALIDATED MECHANISMS

- Focus on opportunities with major unmet medical need

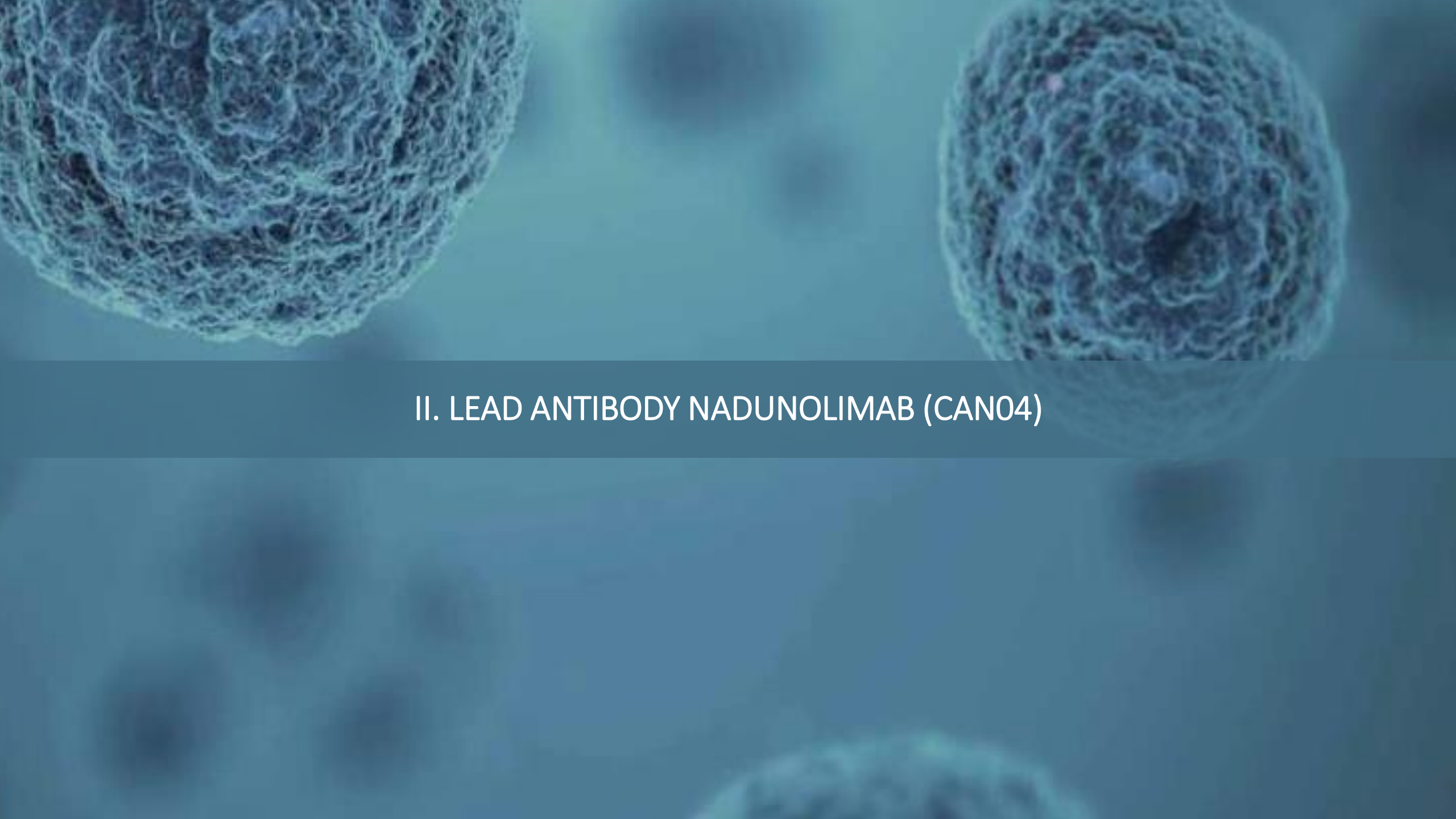
ROBUST PATENT PORTFOLIO

- Global patent families on IL1RAP as antibody target in oncology until 2032 and CAN04 until 2035

NASDAQ STOCKHOLM MAIN LIST ~10,000 SHAREHOLDERS AND LONG TERM INVESTORS

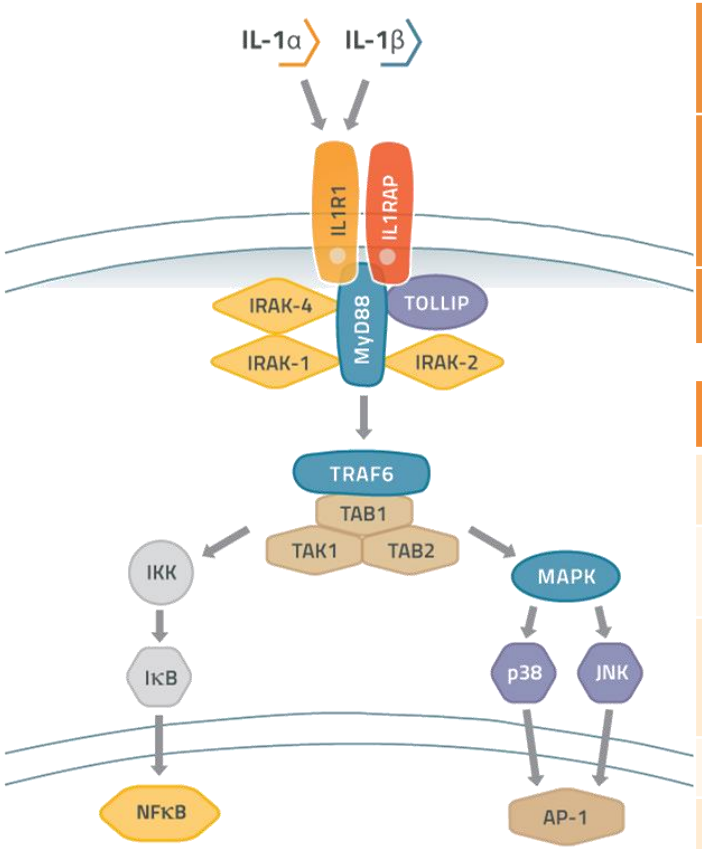
- Market cap: SEK 2.7bn (USD ~320m) (31 May-21)
- Cash: SEK 843m (USD 102m) (31 Mar-20)

Current owners (31 Mar 2021)	
Swedbank Robur Funds	9.7%
4th AP fund	7.7%
Alecta	6.8%
1st AP fund	6.3%
Six Sis AG	5.5%
Avanza Pension	3.9%
Handelsbanken fonder	3.1%
Sunstone LSV	3.0%
SEB AB, Luxemburg	2.7%
Morgan Stanley	2.0%



II. LEAD ANTIBODY NADUNOLIMAB (CAN04)

CAN04 – Differentiated and superior MOA



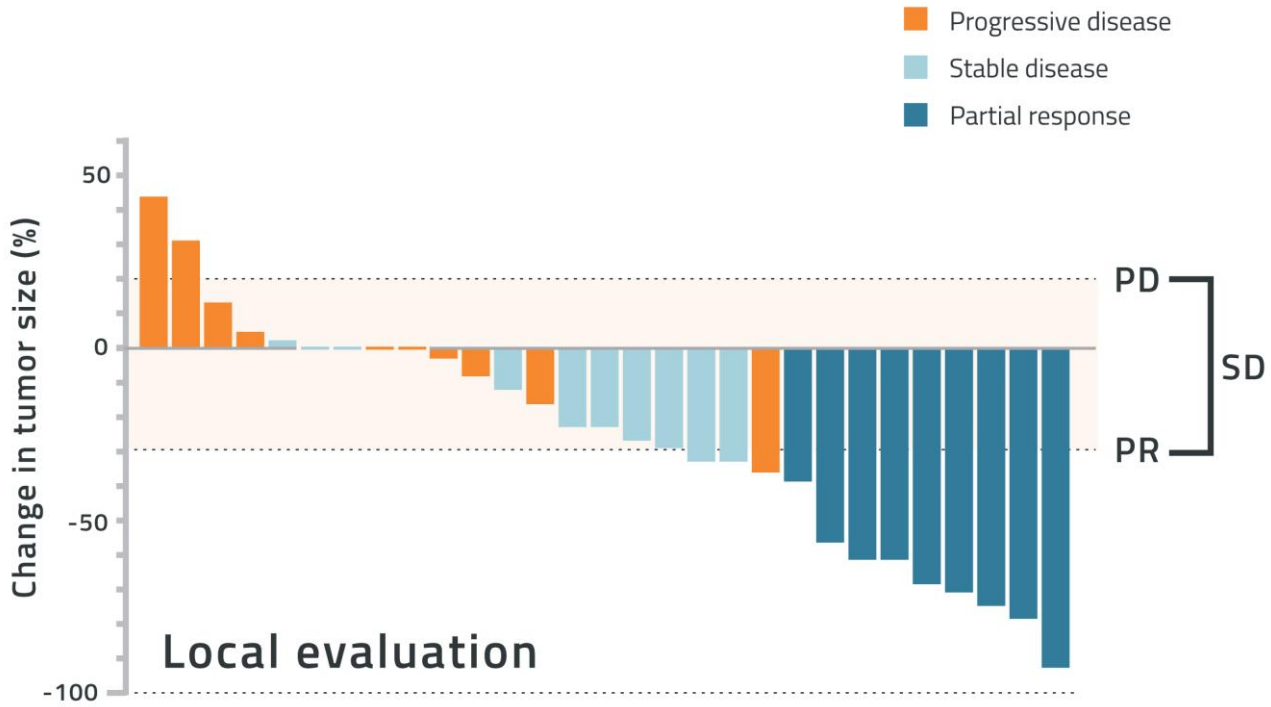
Cancer context	IL-1α	IL-1β	comment
Localization	• Cellbound and soluble • Cancer cells and stroma	• Soluble	• IL-1α trigger and IL-1β enhance inflammation • Often work in pair
Function	• Stimulates inflammation - IL1R1 -forming complex with IL1RAP • IL-1, IL1R1 and IL1RAP in complex - essential for signal • Note: Significant differences in amino acid sequence		• No known difference in signal induced by the 2 forms
Clinical data from blockade	• Signal of benefit in CRC and NSCLC	• CANTOS: reduce lung cancer incidence and death	

Company	Compound	IL-1α	IL-1β	ADCC	Indication/dev phase
Cantargia	CAN04	++	++	++	• Pancreatic cancer, NSCLC phase IIa
Xbiotech/ Janssen	Xilonix XB2001	++	–	+	• Autoimmunity, dermatology • Pancreatic cancer, phase I
Novartis	Canakinumab Gevokizumab	–	++	–	• Autoimmunity, registered • NSCLC, phase III • Cancer comb, phase II
Flame Biosci.	FL-101	–	++	–	• NSCLC
Buzzard	Isunakinra	++	++	–	• Cancer phase I
SOBI	Kineret	++	++	–	• Autoimmunity, reg
Regeneron/ Kiniksa	Rilonacept	++	++	–	• Autoimmunity, reg • Pericarditis
R-Pharm	RPH-104	+	++	–	• Pericarditis, inflammatory disease

Positive data in pancreatic cancer

Efficacy evaluation summary:

- Durable responses observed
- Promising PFS and OS
- Important finding of pseudoprogression-like response in 5 (15%) patients predicting long PFS.



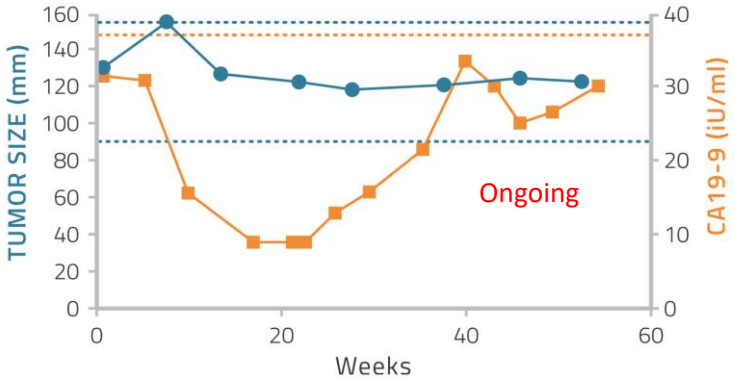
- CAN04 in combination with gem/abraxane in 1st line
- 27% confirmed responses, 15% pseudoprogression
- Median duration of response 6.8 months
- No major side effects observed except those from chemotherapy or CAN04 alone. *Neutropenia frequency higher than expected from chemo (treated with dose reductions/GCSF). Neuropathy and fatigue were less common*

EXTENSION PHASE TO OBTAIN MORE INFORMATION ON VARIOUS DOSE LEVELS ONGOING
DURABLE RESPONSES AND PSEUDPROGRESSION LEADS TO LONG PFS

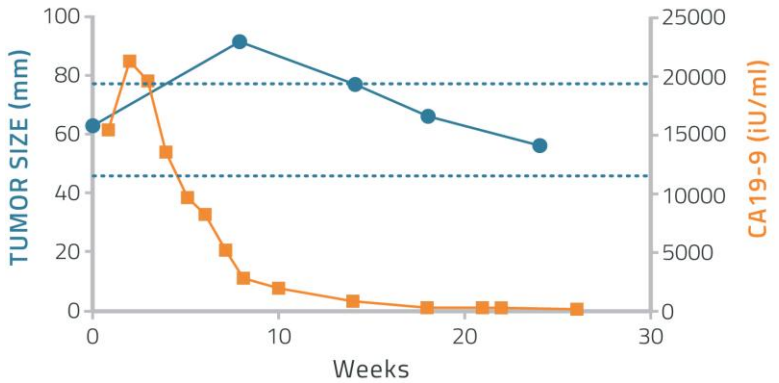
Patients with Pseudoprogression-like response

- All presented PD at 1st CT scan evaluation (8 weeks)
- All showed concomitant reduction of CA19-9

Subject 1



Subject 2



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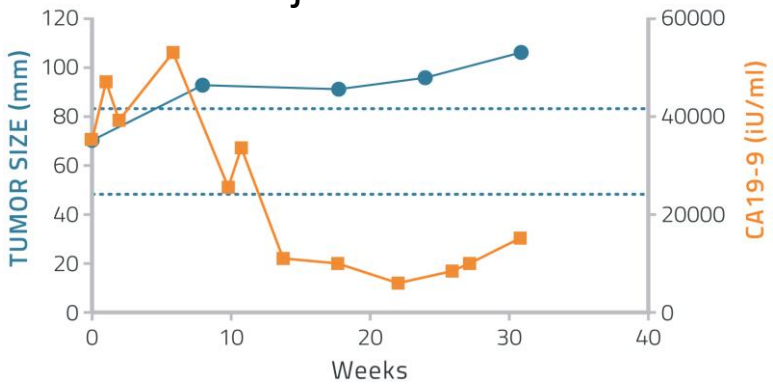

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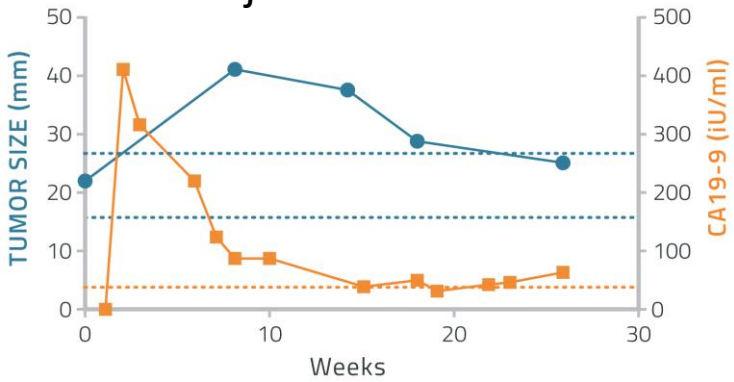

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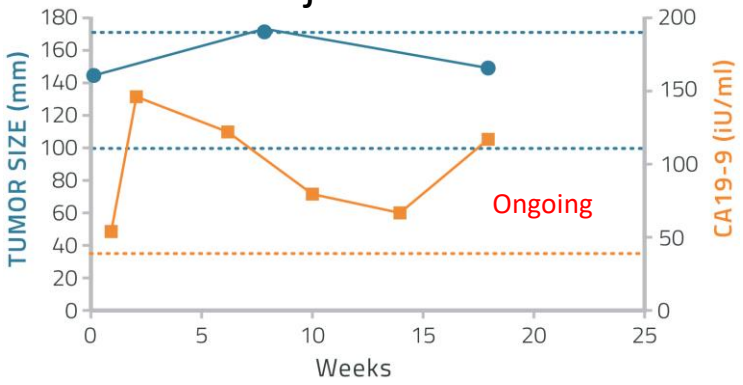
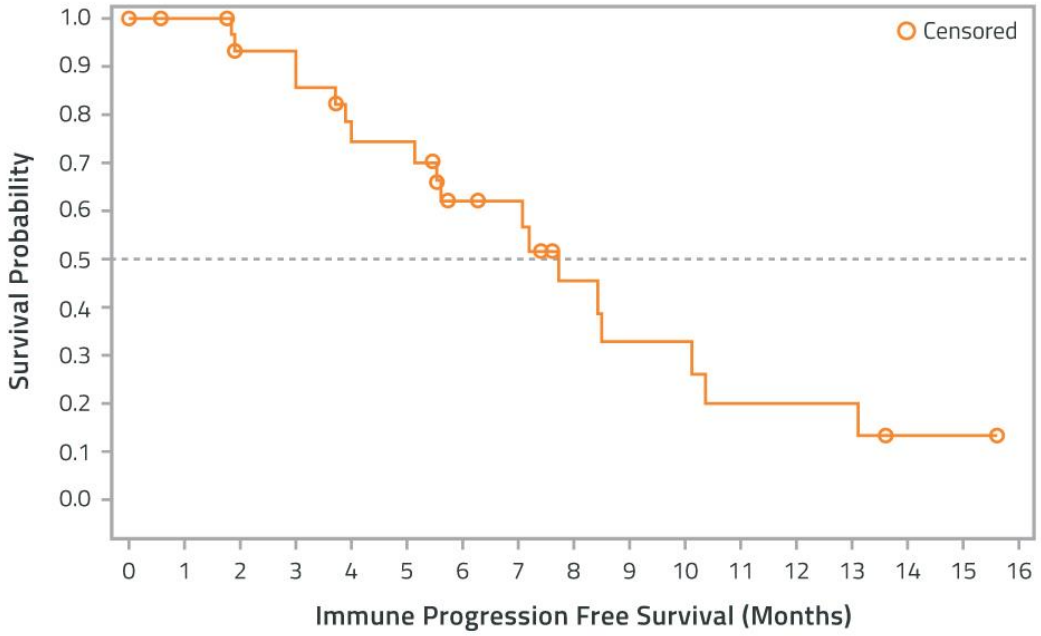
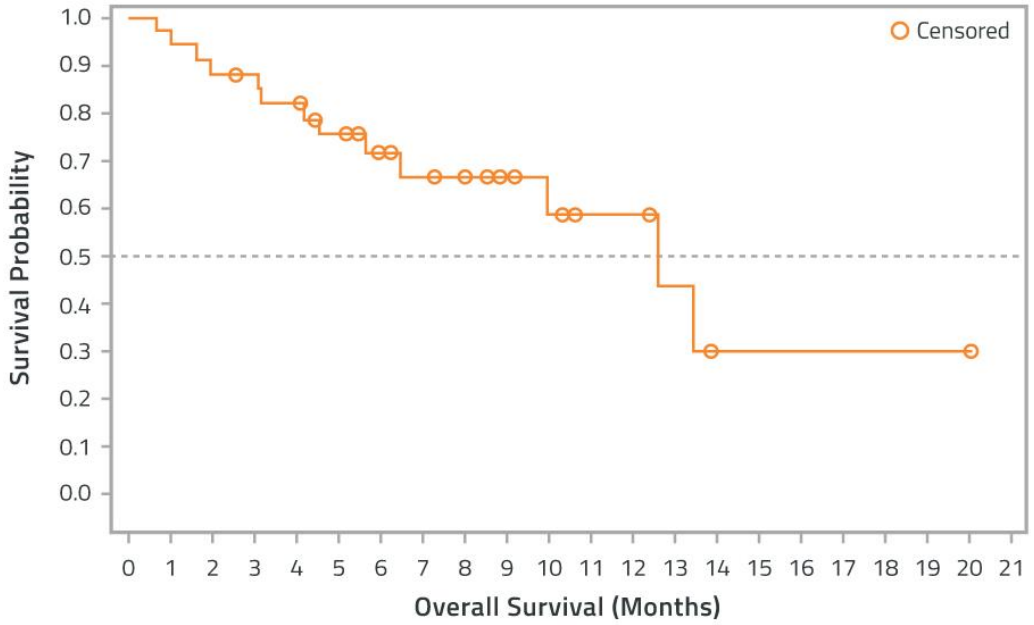


PSEUDOPROGRESSION VERY UNCOMMON IN PANCREATIC CANCER
INDICATE IMMUNE RELATED MECHANISM OF CAN04 LEADING TO LONG TERM BENEFIT

Progression Free Survival (iRECIST) and overall survival

- Median iPFS is 7.8 months (95% CI 5.2 to 10.2) with 55% of events.
- Median OS is 12.6 months (95% CI not estimable) with 42% of OS events.
  - Seven patients at cut-off are still receiving treatment.





OS and iPFS longer than expected from chemotherapy alone

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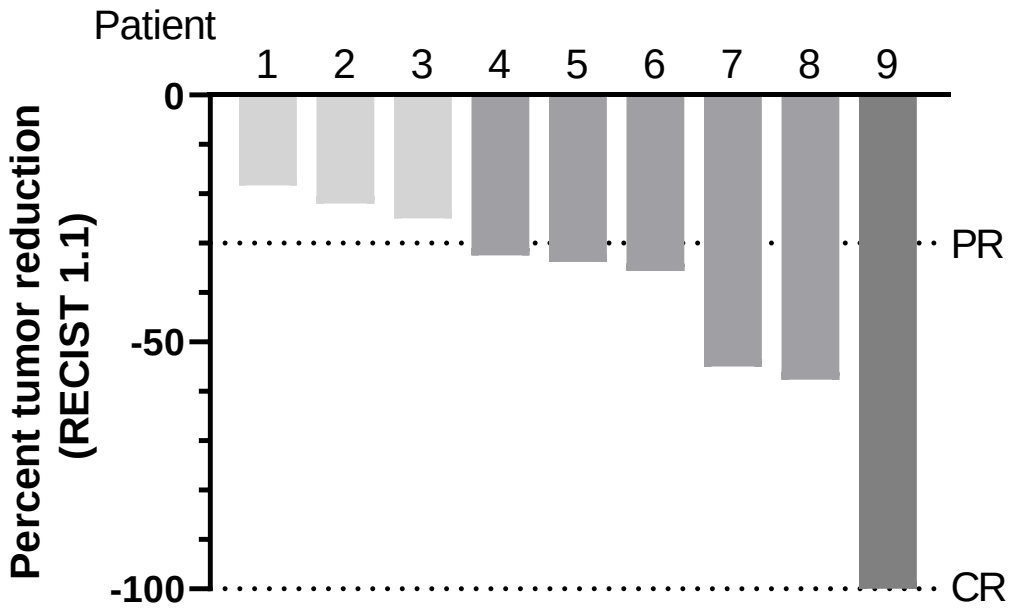
Tumor shrinkage – NSCLC combination



- CAN04 in combination with gem/cis in 1st line chemotherapy
- 6 of 9 evaluable patients with metastatic non-small cell lung cancer (NSCLC) showed objective response including 1 complete response (67% vs historical control data 22–28%)
- The complete response has lasted more than 1 year
- 5 patients were second line to pembrolizumab monotherapy, 4 patients first line
- No major side effects observed except those from chemotherapy or CAN04 alone. *Neutropenia frequency higher than expected from chemo (treated with dose reductions/GCSF)*

POSITIVE INTERIM DATA, RECRUITMENT CONTINUE FOR PRIMARY ANALYSIS
BROADENING OF NSCLC DEVELOPMENT INTO ADDITIONAL MARKET SEGMENTS

Note: CR Complete response, PR Partial response, SD Stable disease, PD progressive disease

Cantargia

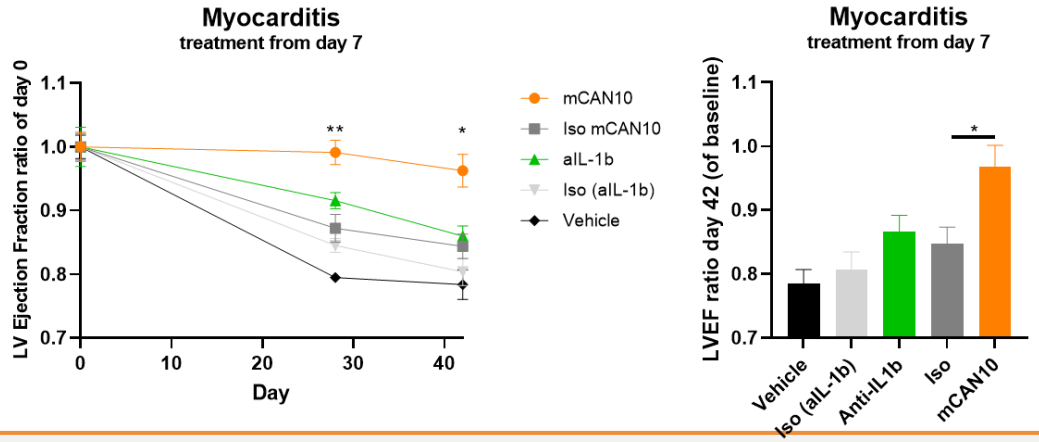
Nadunolimab clinical development status

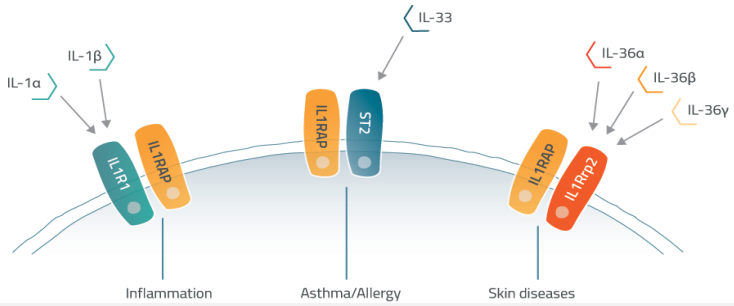
Study	Indication	CAN04 combination	Status	Planned milestone(s)
CANFOUR	NSCLC	Gemcitabine/cisplatin	Recruitment ongoing,	Results planned for Q3 2021
CANFOUR	PDAC	Gemcitabine/nab-paclitaxel	Extension phase ongoing. (Dosing schedule, lower doses, G-CSF)	Main study results presented 20 May LPI extension phase expected Q3 2021
CIRIFOUR	NSCLC, HNSCC, melanoma, bladder cancer	Pembrolizumab	Recruitment ongoing	LPI Q3 2021 Results H2 2021
-	PDAC	mFOLFIRINOX	Regulatory review ongoing	FPI Q2 2021
-	TNBC	Gemcitabine/carboplatin	Preparation together with GEICAM.	Submission Q2
-	Colorectal cancer	mFOLFOX	Preparation	Submission Q2
-	Biliary tract cancer	Gemcitabin/Cisplatin	Preparation	Submission Q2
-	NSCLC	Docetaxel	Preparation	Submission Q2

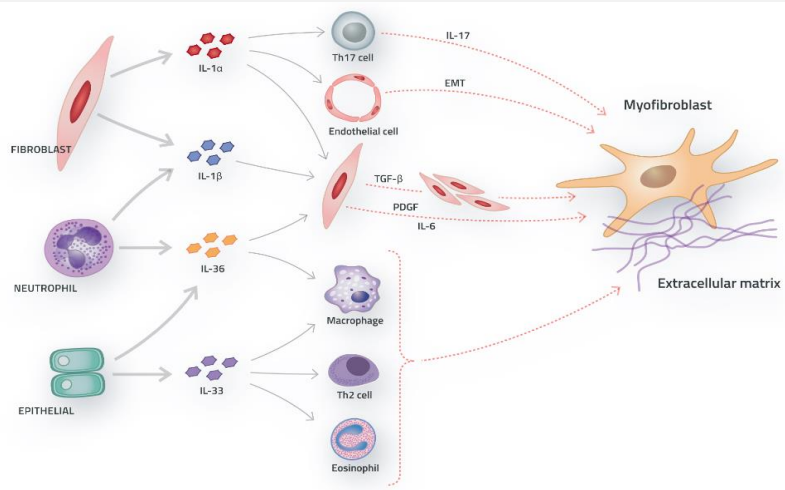
III. UNTAPPED POSSIBILITIES IN AUTOIMMUNE DISEASES

CAN10 – New development project

- IL1RAP binding antibody potently blocking IL-1, IL-33 and IL-36
- Development focusing on unmet medical need in systemic sclerosis and myocarditis. Disease selection in collaboration with experts based on scientific rational, medical need, development opportunity and competition
- Clinical trials start early 2022







UNIQUE OPPORTUNITY FOR CAN10 IDENTIFIED IN LIFE-THREATENING DISEASES

IV. MILESTONES AND SUMMARY

Cantargia reached several milestones and have several value inflection points in near future

Newsflow over next 6–9 months

CAN04

- New results PDAC, NSCLC and Keytruda combination
- Next steps combination therapy PDAC and NSCLC
- Phase IIa biomarker/biopsy results
- Start new clinical trials
  - FOLFIRINOX combination PDAC
  - Basket trial (NSCLC, CRC, BTC)
  - TNBC

CAN10

- Preclinical progress
- Development milestones
- …..and initiation of clinical trial early 2022

SIGNIFICANT DATA TO SECURE NEWSFLOW