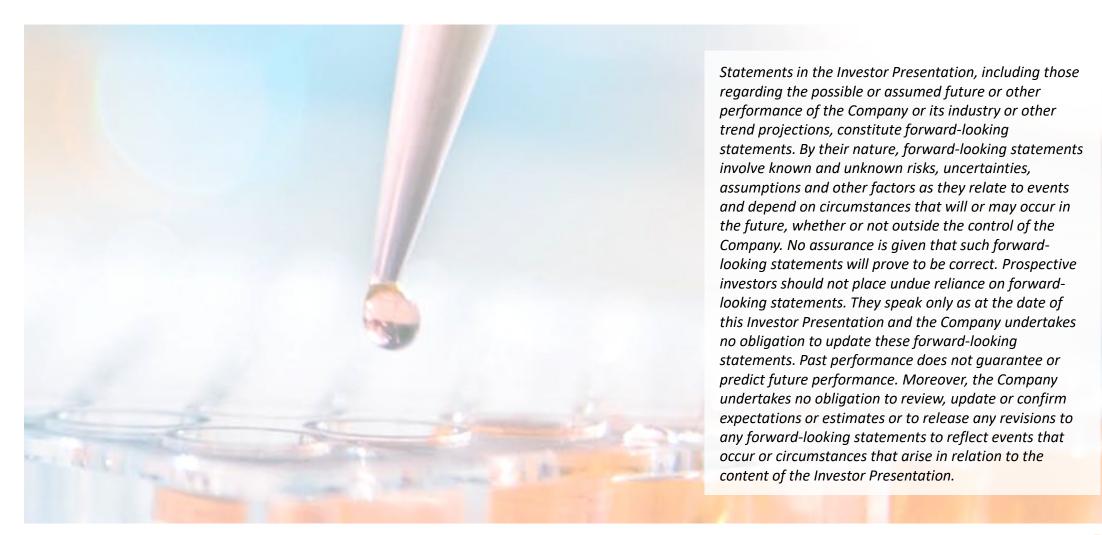


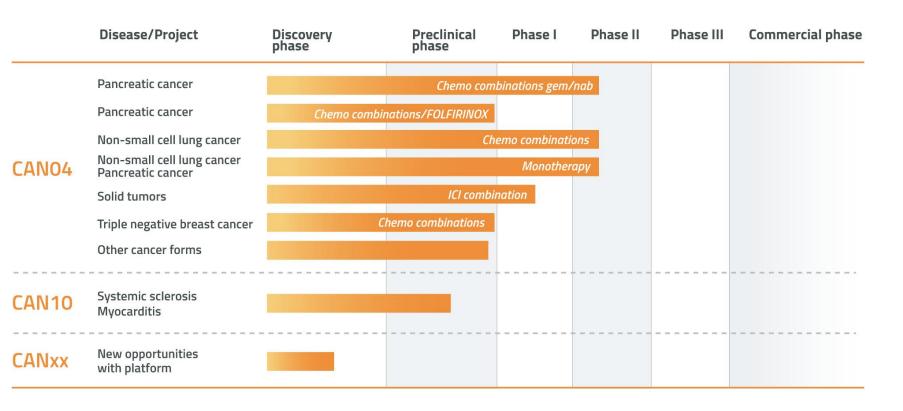
Safe Harbour Statement







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

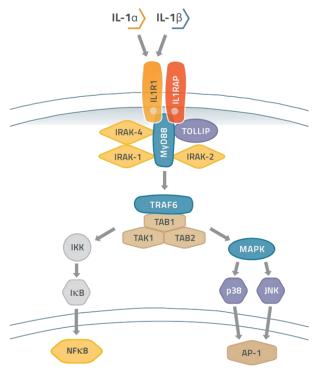
- Market cap: SEK 2.6bn (USD ~310m) (21 May-21)
- Cash: SEK 903m (USD 108m) (31 Dec-20)

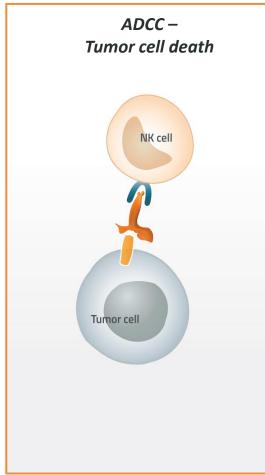
Current owners (31 Mar 2021)				
Swedbank Robur Funds	9.7%			
4th AP fund	7.7%			
Alecta	6.8%			
1st AP fund	6.3%			
Six Sis AG	5.5%			
Avanza Pension	3.9%			
Handelsbanken fonder	3.1%			
Sunstone LSV	3.0%			
SEB AB, Luxemburg	2.7%			
Morgan Stanley	2.0%			

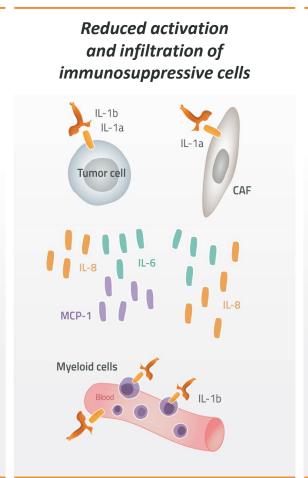


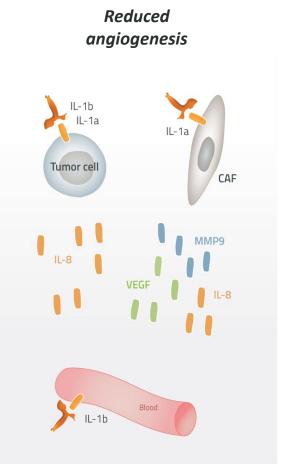


CAN04 – Mechanism of action





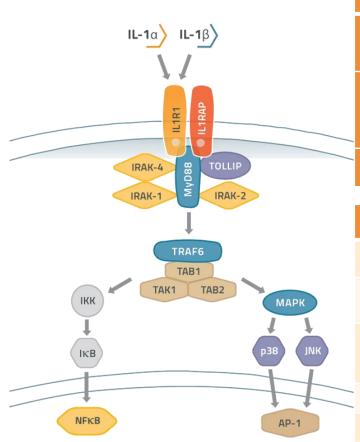




CAN04 BLOCKS BOTH FORMS OF IL-1 AND CAN ERADICATE CELLS MEDIATING THE EFFECTS OF IL-1



CAN04 – Differentiated and superior MOA

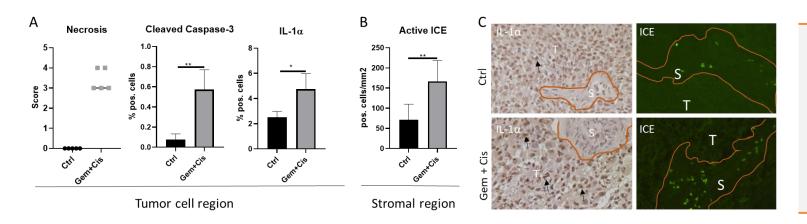


Cancer context	IL-1α	IL-1β	comment
Localization	Cellbound and solubleCancer cells and stroma	• Soluble	 IL-1α trigger and IL-1β enhance inflammation Often work in pair
Function	 Stimulates inflammation - IL1R1 - IL-1, IL1R1 and IL1RAP in complex Note: Significant differences in an 	 No known difference 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	

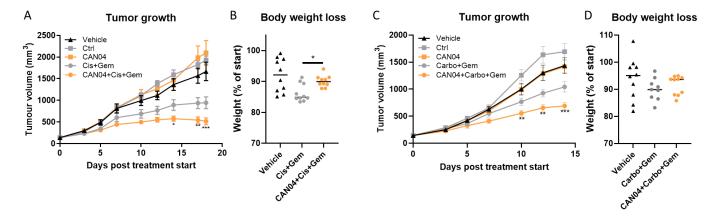
Company	Compound	IL-1α	IL-1β	ADCC	Indication/dev phase
Cantargia	CAN04	++	++	++	Pancreatic cancer, NSCLC phase IIa
Xbiotech/ Janssen	Xilonix XB2001	++	-	+	Autoimmunity, dermatologyPancreatic cancer, phase I
Novartis	Canakinumab Gevokizumab	-	++	-	Autoimmunity, registeredNSCLC, phase IIICancer comb, phase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



Targeting IL1RAP allows unique synergistic effects with chemotherapy (AACR 2020)



- Upregulation of both forms of IL-1 in PDX-model as response to Gem/Cis
- IL-1α (DAMP) on cancer cells trigger inflammasome activation in tumor microenvironment (e.g. IL-1β)



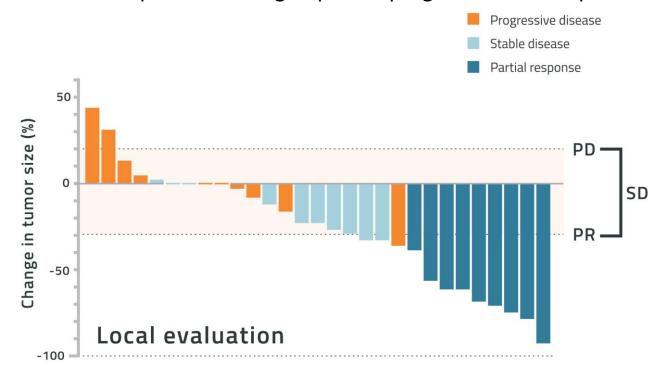
- CAN04 increases efficacy of Pt based chemotherapy regimes
- CAN04 counteracts weight loss after chemotherapy

SYNERGY WITH CHEMOTHERAPY IN LINE WITH CURRENT DEVELOPMENT STRATEGY

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.

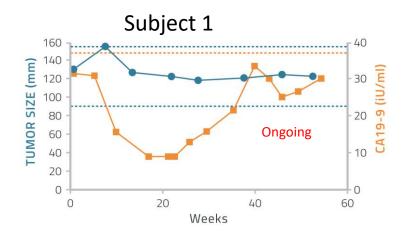


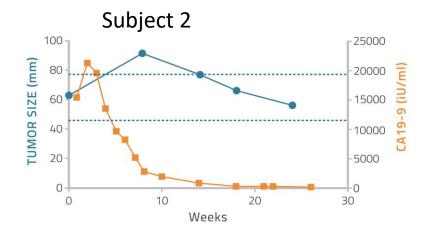
- → CAN04 in combination with gem/abraxane in 1st line
- → 27% confirmed responses, 15% pseudoprogression
- → Median duration of response 6.8 months
- → No major side effects observed except those from chemotherapy or CAN04 alone. Neutropenia frequency higher than expected from chemo (treated with dose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

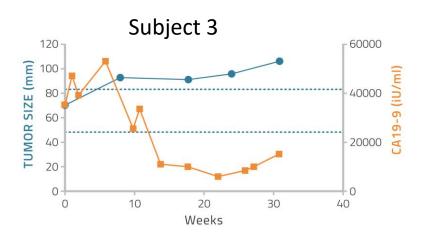


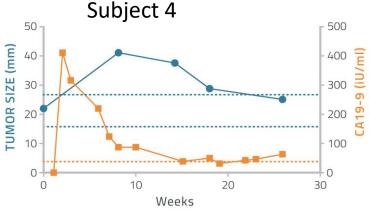
Patients with Pseudoprogression-like response

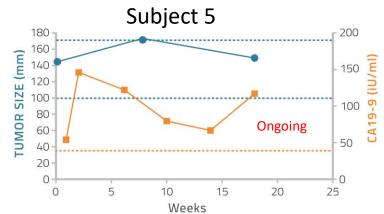
- All presented PD at 1st CT scan evaluation (8 weeks)
- All showed concomitant reduction of CA19-9







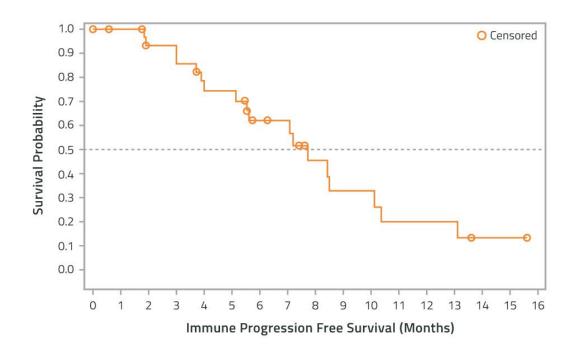


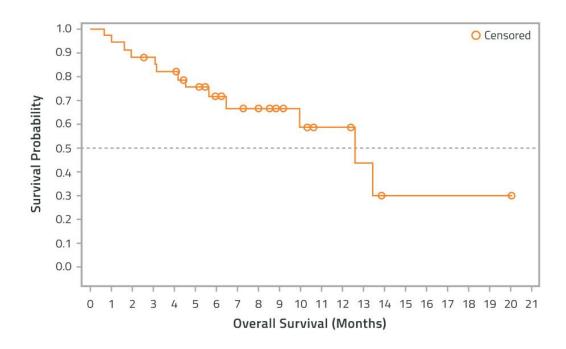


antarqia

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



CANO4/GN in PDAC safety summary and benchmark

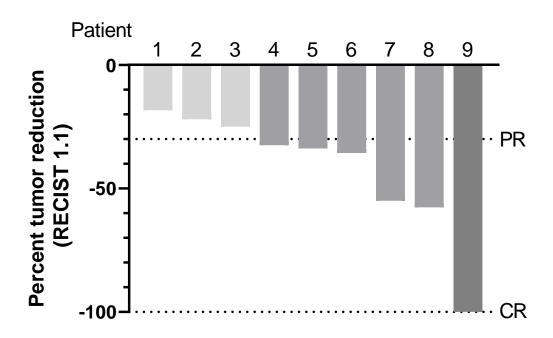
Grade 3 or higher AEs	Gem/Abraxane (von Hoff) N=421	CANFOUR CAN04/GN N=36	FOLFIRINOX (Conroy 2011) N=171
Neutropenia	38%	67%	46%
Febrile neutropenia	3%	17%	5%
Thrombocytopenia	13%	19%	9%
Anemia	13%	14%	8%
Fatigue	17%	6%	24%
Peripheral neuropathy	17%	0%	9%
Diarrhea	6%	3%	13%
Elevated ALT	ND	3%	7%
IRR	ND	3%	ND

The beneficial effect in fatigue and chemotherapy-induced neuropathy² (nabpaclitaxel or oxaliplatin) can be mediated by IL-1 blockade.

- G-CSF not used proactively/prophylactically in this trial. In later trials, G-CSF counteracts neutropenia.
- Median duration of treatment 4.8 months (reference 3.9 months)
- Most common reasons for termination: gastrointestinal events or general health deterioration



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)

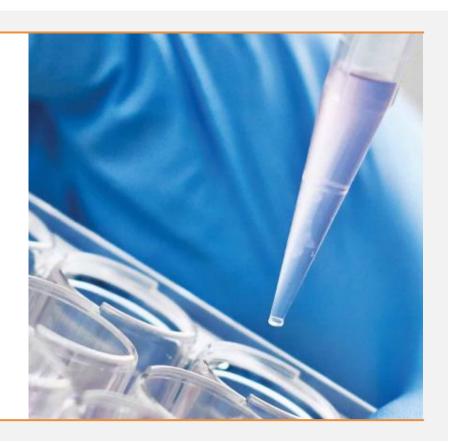


POSITIVE INTERIM DATA, RECRUITMENT CONTINUE FOR PRIMARY ANALYSIS BROADENING OF NSCLC DEVELOPMENT INTO ADDITIONAL MARKET SEGMENTS



CIRIFOUR Phase I clinical trial

- → First patient started 2020, results expected H2 2021
- → Combination with checkpoint inhibitor in patients no longer responding to PD1/PDL-1 therapy
- → Primary endpoint safety, secondary endpoints include biomarkers and efficacy
- Indications include NSCLC, HNSCC, malignant melanoma and bladder cancer (18 patients)
- → Strong US centers, Coord investigator Prof Roger Cohen, UPenn
- https://clinicaltrials.gov/ct2/show/NCT04452214



TRIAL DESIGNED TO ADVANCE CAN04 OUTSIDE CHEMOTHERAPY COMBINATIONS IMPORTANT STEP FOR COMBINING CAN04 WITH IO AND CHEMOTHERAPY



Nadunolimab clinical development status

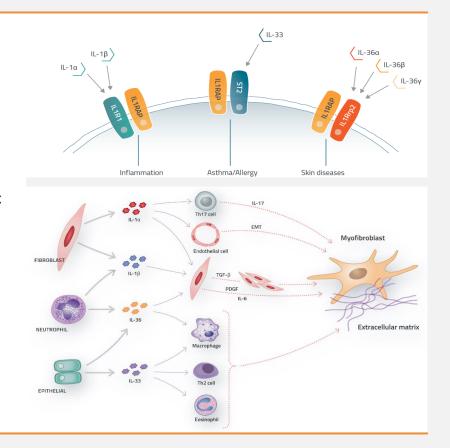
Study	Indication	CAN04 combination	Status	Planned milestone(s)
CANFOUR	NSCLC	Gemcitabine/cisplatin	Recruitment ongoing,	Results planned for Q3 2021
CANFOUR	PDAC	Gemcitabine/nab- paclitaxel	Extension phase ongoing. (Dosing schedule, lower doses, G-CSF)	Main study results presented 20 May LPI extension 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





CAN10 – New development project

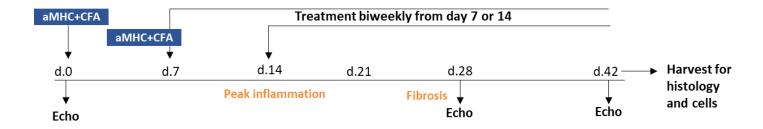
- → IL1RAP binding antibody potently blocking IL-1, IL-33 and IL-36
- Unique anti-inflammatory activity observed in different mouse models (myocarditis, psoriasis, inflammation)
- → Development focusing on unmet medical need in systemic sclerosis and myocarditis. Disease selection in collaboration with experts based on scientific rational, medical need, development opportunity and competition
- → Clinical trials start early 2022

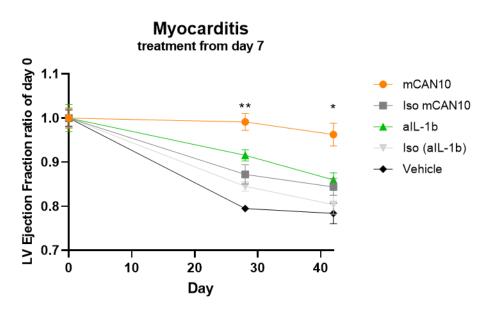


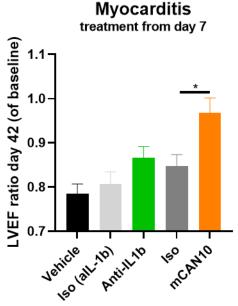
UNIQUE OPPORTUNITY FOR CAN10 IDENTIFIED IN LIFE-THREATENING DISEASES



mCAN10 improves heart function in experimental autoimmune myocarditis











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



Cantargia highlights



UNIQUE IMMUNOTHERAPY ANTIBODY CAN04 IN PHASE IIA CLINICAL DEVELOPMENT

- First in class antibody with broader MOA than competitors
- Positive interim data set and further clinical milestones during 2021



VISION OF BECOMING AN IMPORTANT PART IN FUTURE CANCER TREATMENTS

Combination therapy strategy based on synergies with established therapies



PLATFORM WITH MANY POTENTIAL THERAPEUTIC AREAS

Cancer and large number of autoimmune/inflammatory diseases



HIGHLY RELEVANT RESEARCH WITHIN CLINICALLY VALIDATED MECHANISMS

Focus on opportunities with major unmet medical need



ROBUST PATENT PORTFOLIO - GRANTED IP FOR THERAPEUTIC TARGET IL1RAP AND CAN04

Global patent families – antibody target in oncology (2032) and CAN04 (2035)



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

- Market cap: SEK 2.6bn (USD ~310m) (21 May-21)
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