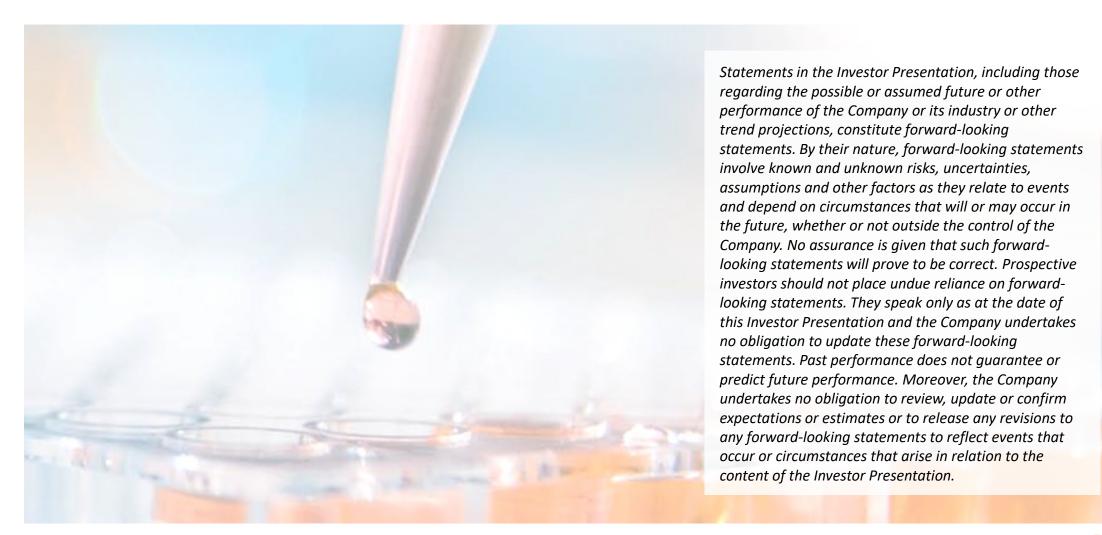


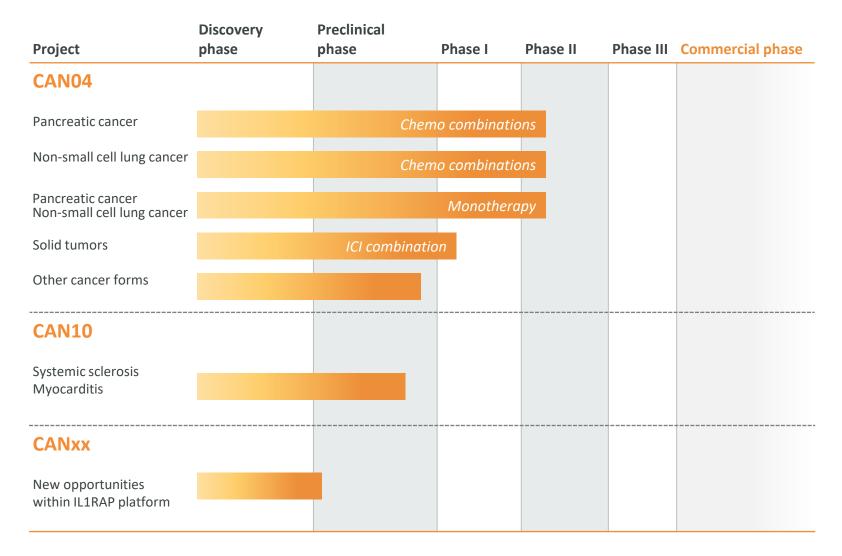
### Safe Harbour Statement







# 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

- Positive interim data set response rates higher than historical control
- First in class antibody with broader MOA than competitors



#### 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 3.2bn (USD ~380m) (Apr 20-21)
- Cash: SEK 903m (USD 108m) (31 Dec-20)

Current owners (30 Dec 2020)				
Swedbank Robur Funds	9.7%			
4th AP fund	7.7%			
Alecta	6.6%			
1st AP fund	6.3%			
Öhman Bank S.A.	5.3%			
Handelsbanken fonder	3.8%			
Avanza Pension	3.8%			
Sunstone LSV	3.5%			
Morgan Stanley	2.0%			
JP Morgan	1.8%			

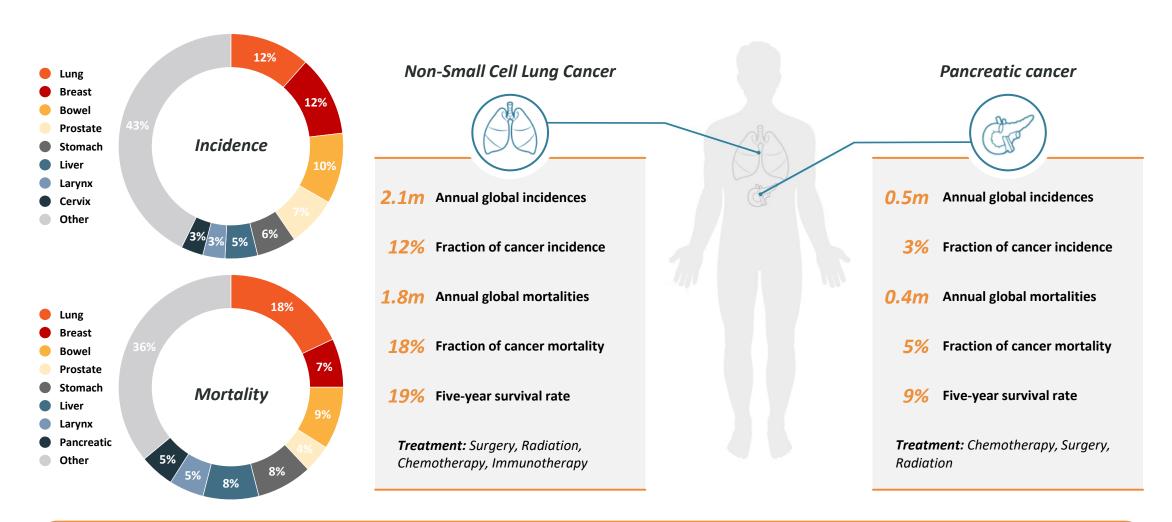
Others



49.5%



### Cantargia addresses NSCLC & PDAC



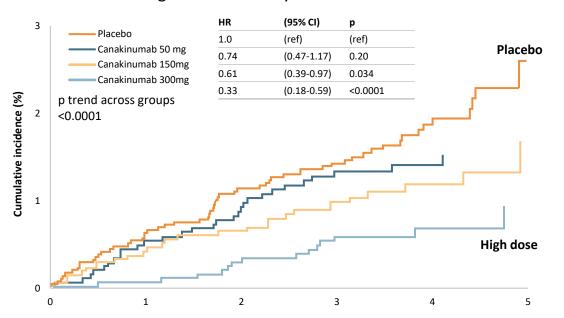
SIGNIFICANT UNMET NEEDS IN LUNG AND PANCREATIC CANCER



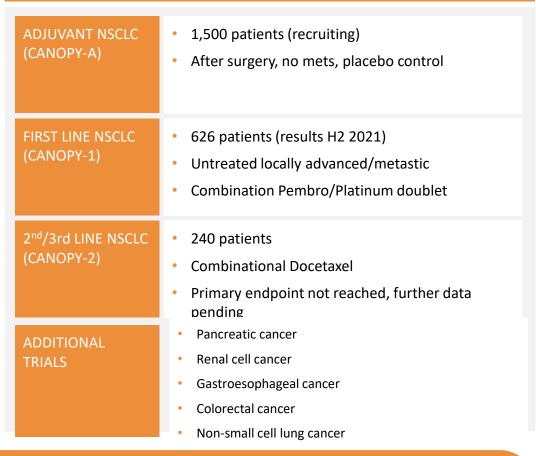
# Inflammatory IL-1 pathway validated as target

#### Cantos trial (n=10,061)

- Canakinumab (Novartis)
- Reduced lung cancer incidence by 67% and death by 77%
- Reduced non-lung cancer death by 37%



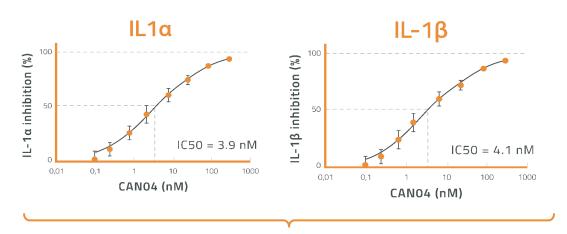
#### **Canakinumab phase III trials**



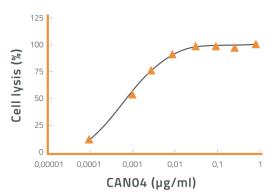
antarqia

ROLE OF IL-1B IN EARLY STAGE CANCER
TWO PHASE III TRIALS AS WELL AS PHASE I-II TRIALS ONGOING
CAN04 PROVIDE MORE POTENT BLOCKADE OF PATHWAY

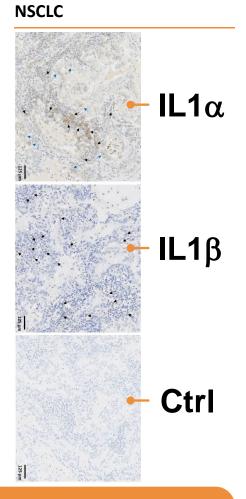
### CAN04 – Dual anticancer mechanism



Inhibition of IL-1 signaling



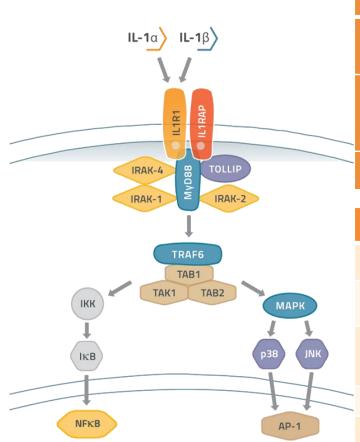
ADCC enhanced antibody



CAN04 BLOCKS BOTH FORMS OF IL-1 AND STIMULATES DIRECT KILLING OF CANCER CELLS



# CAN04 – Differentiated and superior MOA



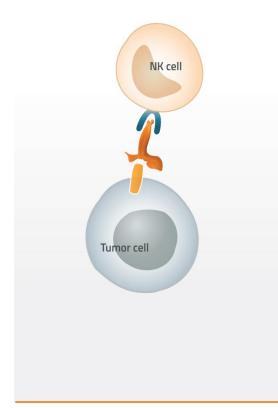
Cancer context	IL-1α	IL-1β	comment
Localization	<ul><li>Cellbound and soluble</li><li>Cancer cells and stroma</li></ul>	Soluble	<ul> <li>IL-1α trigger and IL-1β enhance inflammation</li> <li>Often work in pair</li> </ul>
Function	<ul> <li>Stimulates inflammation - IL1R1 -</li> <li>IL-1, IL1R1 and IL1RAP in complex</li> <li>Note: Significant differences in an</li> </ul>	<ul> <li>No known difference in signal induced by the 2 forms</li> </ul>	
Clinical data from blockade	<ul> <li>Signal of benefit in CRC and NSCLC</li> </ul>	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	++	-	+	<ul><li>Autoimmunity, dermatology</li><li>Pancreatic cancer, phase I</li></ul>
Novartis	Canakinumab Gevokizumab	-	++	-	<ul><li>Autoimmunity, registered</li><li>NSCLC, phase III</li><li>Cancer comb, phase II</li></ul>
Flame Biosci.	FL-101	-	++	_	• NSCLC
Buzzard	Isunakinra	++	++	-	Cancer phase I
SOBI	Kineret	++	++	_	• Autoimmunity, reg
Regeneron/ Kiniksa	Rilonacept	++	++	-	<ul><li>Autoimmunity, reg</li><li>Pericarditis</li></ul>
R-Pharm	RPH-104	+	++	-	Pericarditis, inflammatory disease

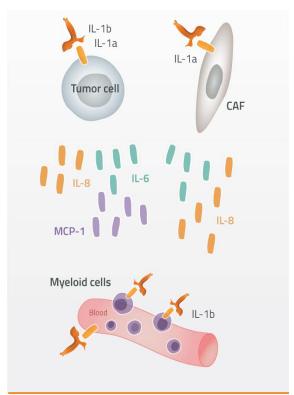


### CAN04 – Mechanism of action

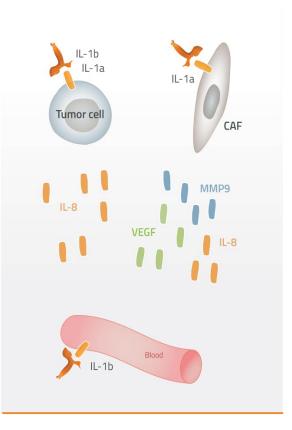
ADCC – Tumor cell death



Reduced activation and infiltration of immunosuppressive cells



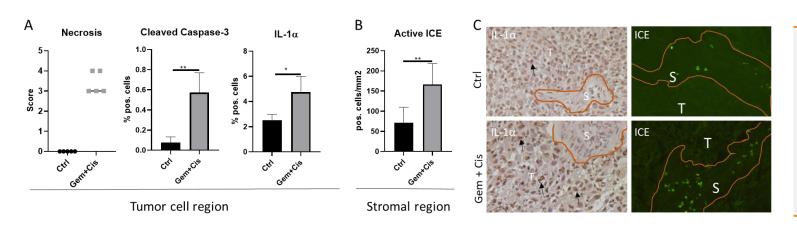
### Reduced angiogenesis



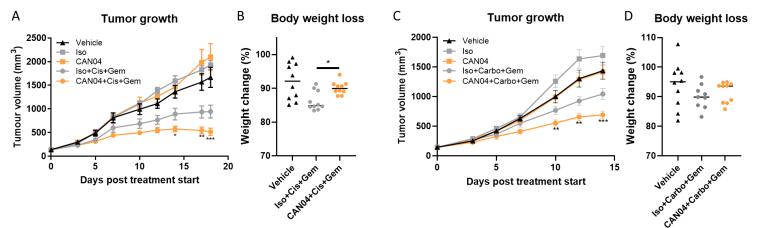
CANO4 BLOCKS BOTH FORMS OF IL-1 AND CAN ERADICATE CELLS MEDIATING THE EFFECTS OF IL-1



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



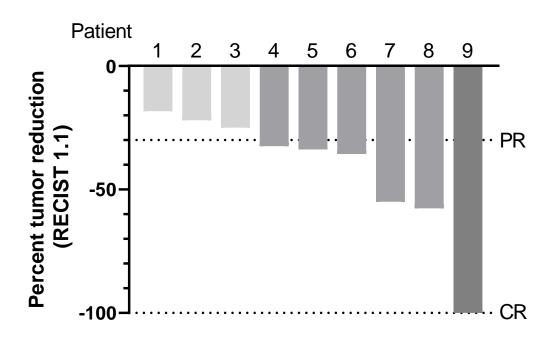
- Upregulation of both forms of IL-1 in PDX-model as response to Gem/Cis
- IL-1α (DAMP) on cancer cells trigger inflammasome activation in tumor microenvironment (e.g. IL-1β)



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

SYNERGY WITH CHEMOTHERAPY IN LINE WITH CURRENT DEVELOPMENT STRATEGY

### Tumor shrinkage – NSCLC combination



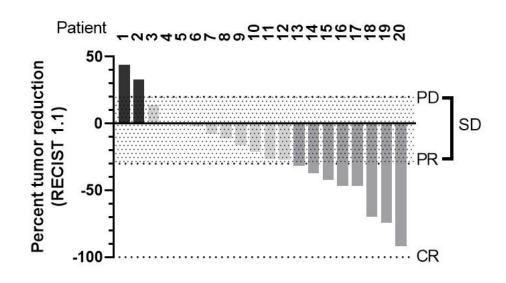
- → CAN04 in combination with gem/cis in 1<sup>st</sup> line chemotherapy
- → 6 of 9 evaluable patients with metastatic non-small cell lung cancer (NSCLC) showed objective response including 1 complete response (67% vs historical control data 22–28%)
- → The complete response has lasted more than 1 year
- 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

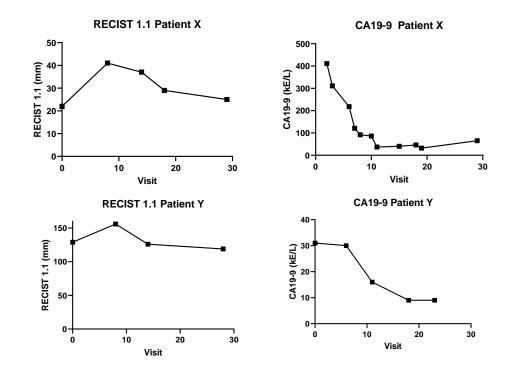


### Positive interim data – pancreatic cancer



- → CAN04 combination with gemcitabine/abraxane in 1<sup>st</sup> line PDAC
- → 8 out of 20 evaluable patients with metastatic PDAC showed response (40% vs historical control data 23%). Two responses durable for 12 months
- → No major side effects were observed apart from those expected with chemotherapy or CANO4 alone. Neutropenia frequency higher than expected from chemo (treated with dose reductions/GCSF), fatigue and neuropathy lower than expected

FULLY RECRUITED -31 PATIENTS FOR PRIMARY ANALYSIS
ONGOING EXTENSION PHASE 20–40 PATIENTS TO STUDY DOSE/RESPONSE
PREPARATIONS FOR LATE STAGE DEVELOPMENT INITIATED



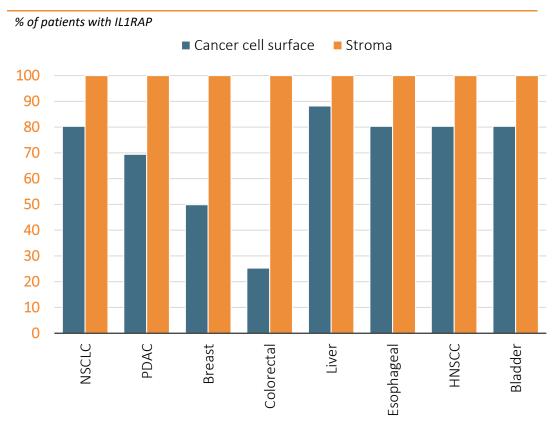
- Two patients had tumor shrinkage of 39% and 24% after initial PD.
   Both recorded as SD
- CA19-9 decreased by 92% and 71%. CA19-9 is a biomarker for tumor burden

NOTABLE RESPONSE PROFILE DEVIATING FROM CHEMO ALONE SUGGEST ADDITIONAL EFFECT FROM CAN04



# IL1RAP in several cancer with high medical need

#### **IL1RAP**





- Discovery of IL1RAP on cancer cells
- Antibodies against IL1RAP antitumor effects
- IP on antibody therapy against IL1RAP

### PRIMARY INDICATIONS

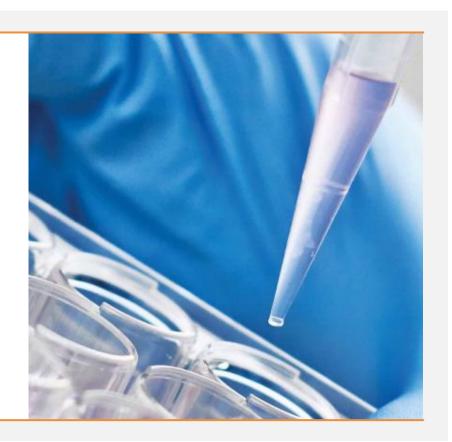
- Non-small cell lung cancer NSCLC
- Pancreatic cancer PDAC
- Biomarker studies ongoing, identify patients most likely to respond
- Opportunity to expand development in additional cancer forms with high unmet medical need

CANO4 DEVELOPMENT CAN BE EXPANDED TO ADDITIONAL INDICATIONS ONWARDS



### US Phase I clinical trial

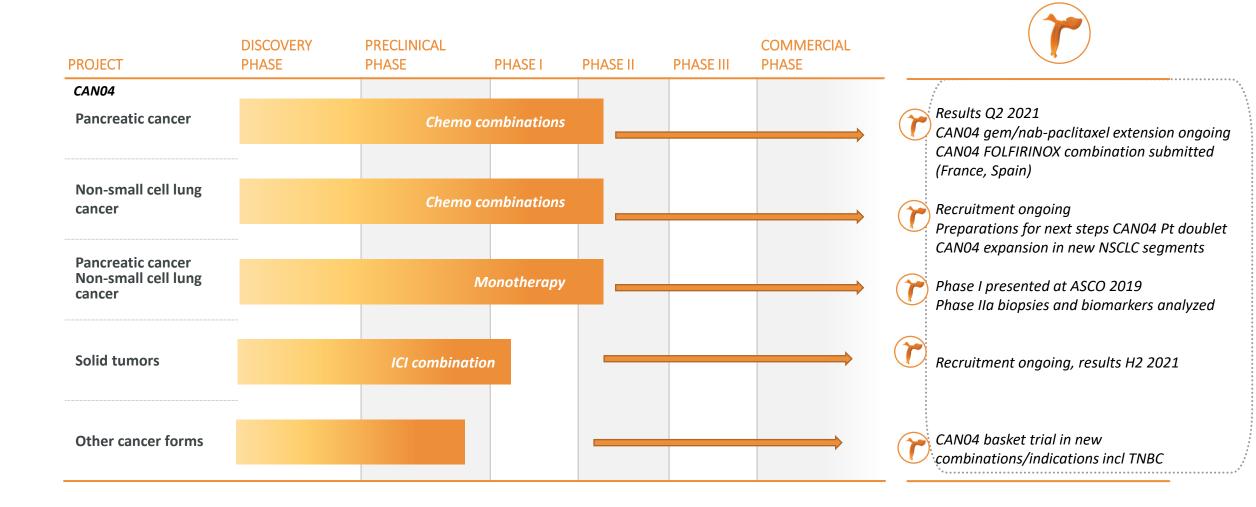
- → First patient started 2020
- → Combination with checkpoint inhibitor in patients no longer responding to PD1/PDL-1 therapy
- → Primary endpoint safety, secondary endpoints include biomarkers and efficacy
- → Indications include NSCLC, HNSCC, malignant melanoma and bladder cancer (18 patients)
- → 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



### CAN04— Broadening development

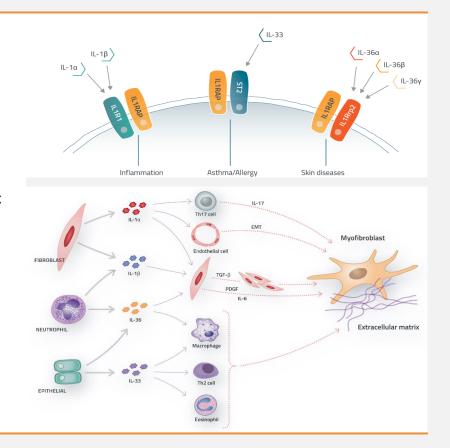






## CAN10 – New development project

- → 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

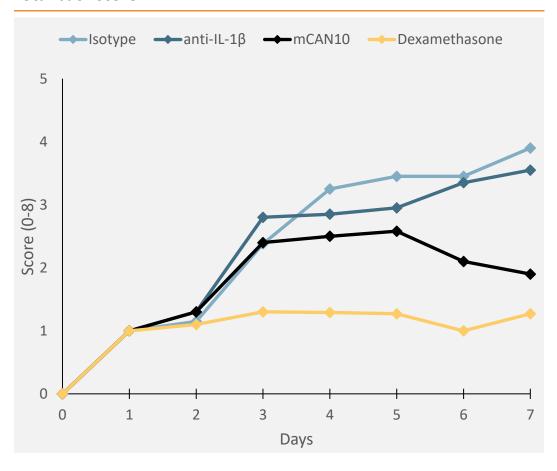


UNIQUE OPPORTUNITY FOR CAN10 IDENTIFIED IN LIFE-THREATENING DISEASES



### CAN10 counteract inflammation in disease model

#### **Total back score**



- Mechanistic proof of concept for IL1RAP blockade in inflammatory driven psoriasis model
- → Effect not dependent on IL-1β blockade
- → Effects in myocarditis model to be presented at conference during Q2 2021 − CAN10 superior to IL-1 blockade.
- → Well tolerated as single dose up to 50 mg/kg (tox model)

CAN10 HAS UNIQUE ANTI-INFLAMMATORY PROPERTIES





## Cantargia has several near-term value inflection points

#### 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 and new indications like TNBC

#### CAN10

- → Preclinical progress
- → Development milestones



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 CANO4

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



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