

We want to save patients with severe cancer and autoimmune diseases Clinical investigations with our lead antibody CANO4 to our proprietary target

## Göran Forsberg, CEO

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## 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. 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 – Opportunity to save lives and create value

Project	Discovery phase	Preclinical phase		Phase I	Phase I Phase II		Phase III		Commercial phase			
CANO4												
Non-small cell lung cancer		(	Themo	combination	15							
Pancreatic cancer		combination	15									
Non-small cell lung cancer Pancreatic cancer				Monotherap	y							
Solid tumors		ICI combinat	ion									
Other cancer forms												
CAN10												
Systemic sclerosis Myocarditis												
CANxx												
New opportunities with platform												

- ightarrow Potentially more effective treatment against novel target in clinically validated pathway
- ightarrow Right team and clear plan to position our projects and maximize value
- ightarrow First in class platform technology against novel target



## Cantargia at a glance



### Unique immunotherapy antibody CAN04 in phase IIa clinical development

- Positive interim data set with response rates higher than historic data
- Further phase II milestones during 2020



### Platform with many potential therapeutic areas

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



### Vision of becoming an important part in future cancer treatments

Combination therapy strategy based on synergies with established therapies



### 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's main list >6,000 shareholders and long term investors

- Market cap: SEK 2.6bn (USD ~300m) (Aug 31, 2020)
- Cash : SEK 458 MSEK (USD 53m) (30 Jun 2020)

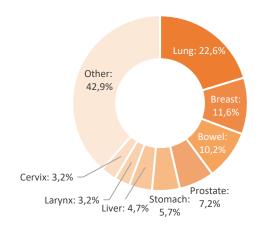
### Current owners (30 June 2020)

4th AP fund	7.8%
Swedbank Robur Funds	7.4%
Alecta	6.6%
1st AP fund	6.3%
Sunstone	5.2%
Öhman Bank S.A.	4.9%
Avanza Pension	4.4%
SEB S.A. (Nordic Cross)	3.0%
Morgan Stanley (HBM)	2.4%
Handelsbanken fonder	2.3%
Others	49.7%



## Cantargia addresses a huge market

Incidence, Globally 2018 Type of cancer:

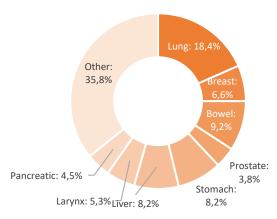


## Incidence, Globally 2018

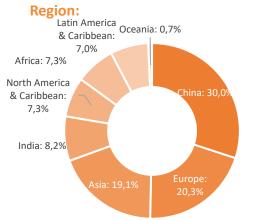


### Mortality, Globally 2018

**Type of cancer:** 



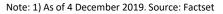
### Mortality, Globally 2018



	Lung cancer	Pancreatic cancer				
Incidence 2018 (globally)	2,093,876	458,918				
Fraction of cancer incidence	13.0%	2.9%				
Mortality 2018	1,761,007	432,242				
Fraction of cancer mortality	19.9%	4.9%				
5 year survival	18.6%	8.5%				
Treatment	Surgery, Radiation, Chemotherapy, Immunotherapy	Chemotherapy, Surgery, Radiation				

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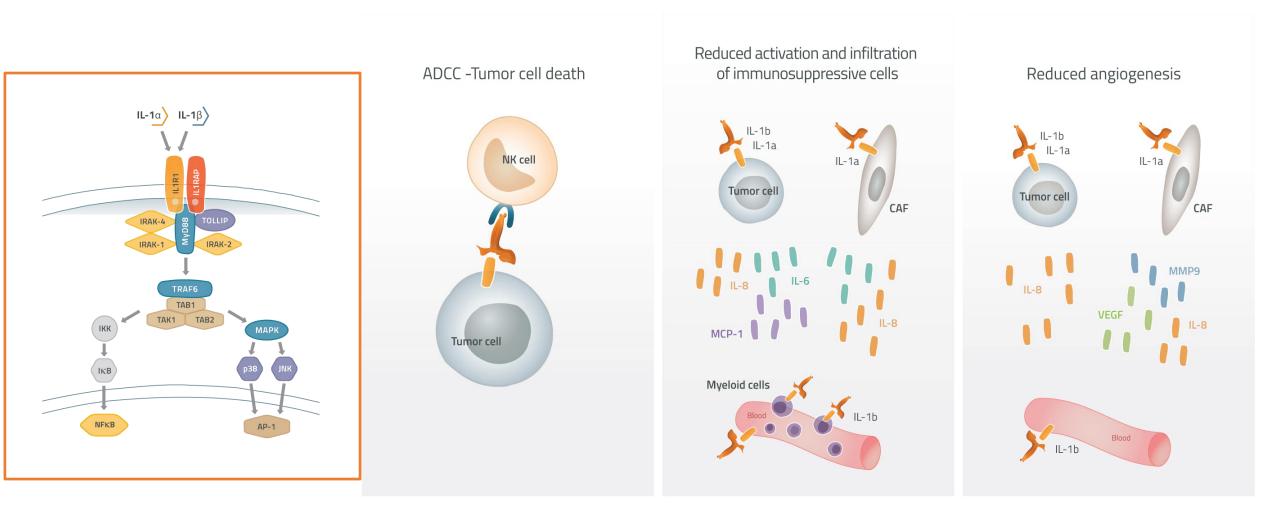
### Significant unmet needs in lung and pancreatic cancer



## Lead antibody CAN04



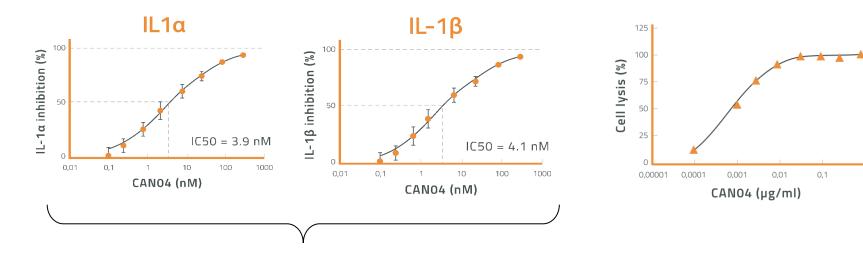
## CAN04 – Mechanism of action





Note: ADCC = Antibody-Dependent Cellular Cytotoxicity. CAF = Cancer-Associated Fibroblast. NK = Natural Killer. MCP = Monocyte Chemoattractant Protein. MMP = Matrix Metallopeptidase. IL = Interleukin. VEGF = Vascular Endothelial Growth Factor

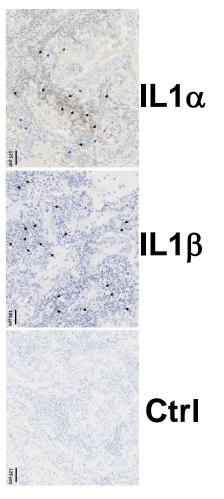
## CAN04 – Dual anticancer mechanism



• Inhibition of IL-1 signaling:

• ADCC enhanced antibody





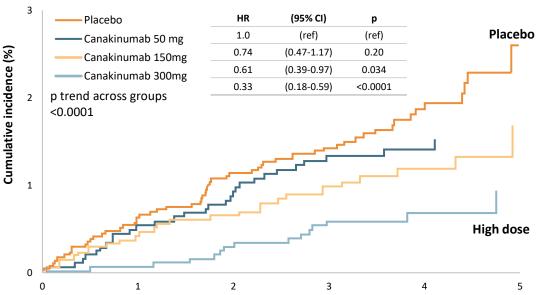
**CAN04** blocks tumor inflammation and stimulates direct killing of cancer cells

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# Validating study – Counteracting tumor inflammation

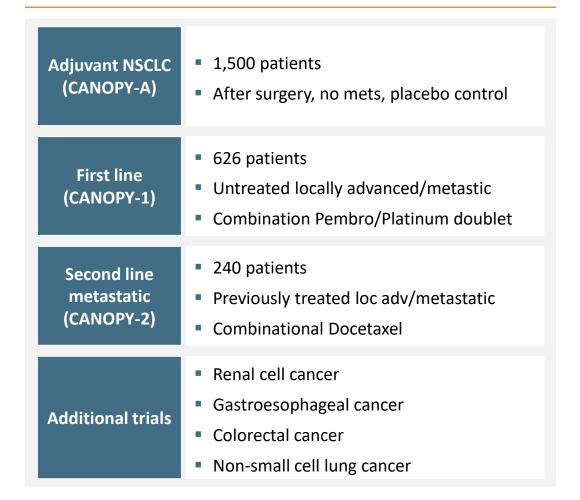
### CANTOS trial (n=10,061)

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



- $ightarrow\,$  Clinical validation of IL-1 pathway
- $\rightarrow$  Dose/response
- $ightarrow\,$  Cantargia's CAN04 has broader MOA

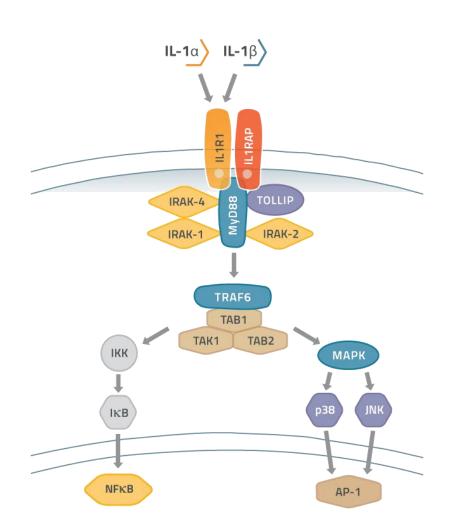
### Canakinumab phase III trials



CANTOS data support CAN04 as well as broader IL1RAP platform activities



## CAN04 – Superior MoA against other IL-1 blocking approaches



Company	Compound	IL-1α	IL-1β	ADCC	Indication/dev phase		
Cantargia	CAN04	++	++	++	Pancreatic cancer, NSCLC     phase IIa		
Xbiotech/ Janssen	Xilonix	++	-	+	<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>		
Buzzard	Isunakinra	++	++	-	Cancer phase I		
SOBI	Kineret	++	++	-	Autoimmunity, reg		
Regeneron/ Kiniksa	Rilonacept	++	++	-	<ul><li>Autoimmunity, reg</li><li>Pericarditis</li></ul>		
Use of IL1RAP a hematological o Two families Valid until 20	ancers	Use of IL1RAP as target for solid tumors Valid until 2032 Granted (EPO*, Japan,			<ul> <li>The product candidate CAN04</li> <li>Valid until 2035</li> <li>Granted (EPO, USA, China)</li> </ul>		

USA, China)

\*divisional application opposed in Europe

 Granted (EPO, USA, Japan, China)

### Cantargia has strong IP and superior MoA in CAN04



# Positive phase IIa interim combination data

	Initiated	On therapy	Evaluable	CR/PR	SD	PD	NE
PDAC	10	7	7	4 <sup>1)</sup>		2 <sup>2)</sup>	1 <sup>2)</sup>
Historical				23%	27%	20%	30%
NSCLC	4	3	3	2 <sup>1)</sup>	1		
Historical				22-28%	18%	40%	<20%



"After I presented the CAN04 monotherapy data at ASCO 2019, the CANFOUR trial has advanced with the combination therapy. The initial results are very encouraging in non-small cell lung cancer (pretreated with checkpoint inhibitor) and pancreatic cancer and suggest that CAN04 could be a valuable contribution to improve the chemoterapy regimes in these diseases" *Prof Ahmad Awada, Institute Jules Bordet, Brussels, Belgium, Coordinating investigator CANFOUR-study* 

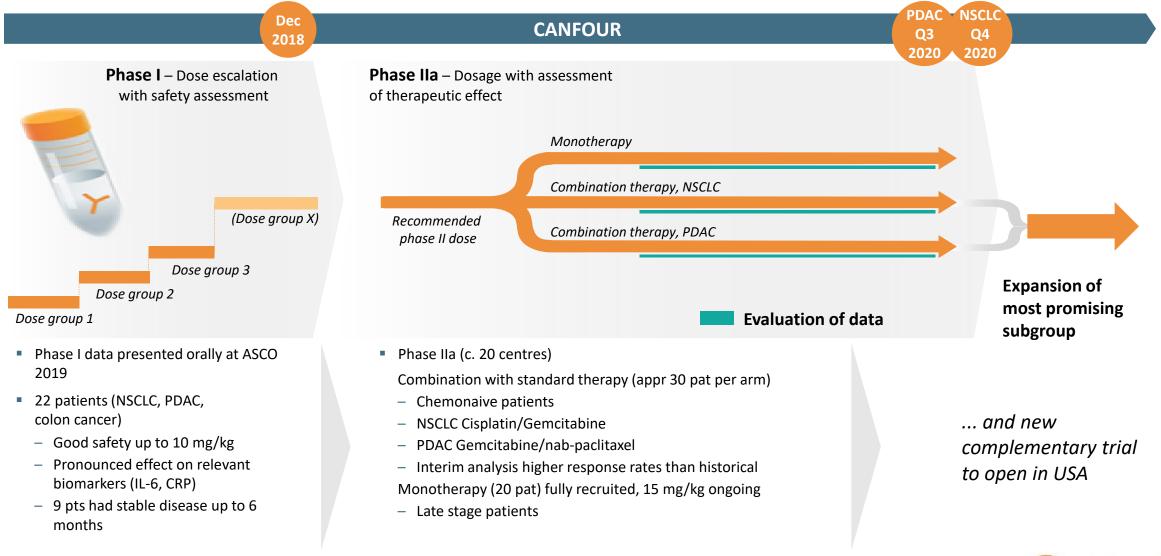
- → By adding CAN04 response rates are higher than historical data using these standard first line chemotherapies alone
- → 4 of 7 evaluable patients with metastatic pancreatic cancer (PDAC) showed objective response. 1 additional patient showed pseudoprogression. Pronounced effect of biomarker CA19-9
- → 2 of 3 evaluable patients with metastatic non-small cell lung cancer (NSCLC) showed objective response including 1 complete response
- → No major side effects were observed apart from those expected with chemotherapy or CAN04 alone

### Strong tumor shrinkage in majority of patients



Note: 1) All patients except 1 PDAC and 1 NSCLC have responses confirmed on second scan. 3 of 4 PDAC patients with objective response has a sustained decrease of >90 % of CA19-9. In NSCLC, 1 patient has a confirmed complete response (CR). 2) 1 patient has ongoing tumor shrinkage after initial progression and a strong reduction in CA19-9. 1 patient terminated after rapid clinical progression without CT-scan

## CAN04 – CANFOUR clinical trial

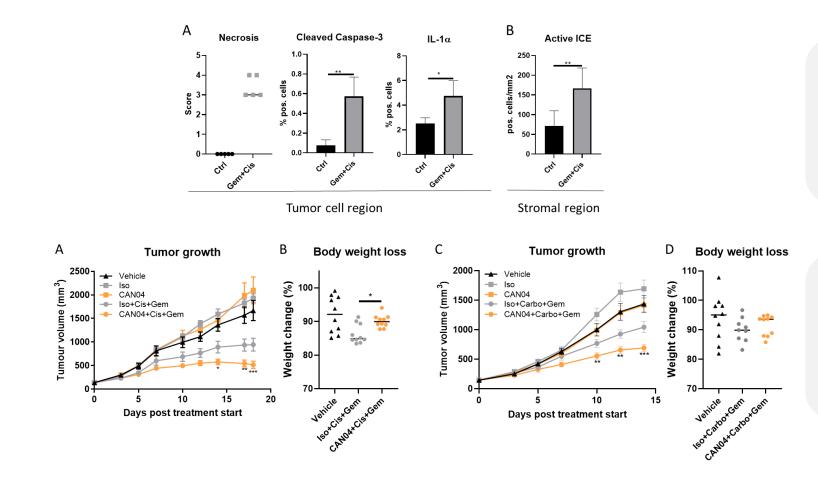


### Generation of data instrumental for next phase of development

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



→ Upregulation of both forms of IL-1 (IL-1a/ICE) as potential tumor defense to chemotherapy

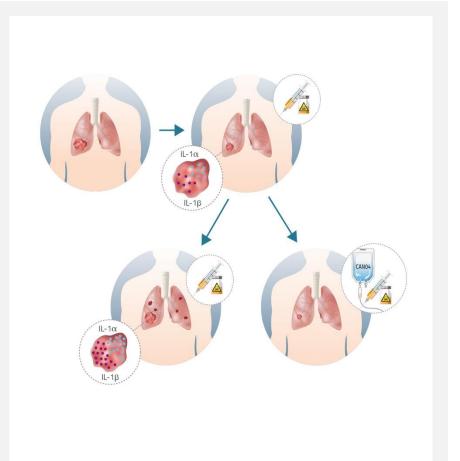
- → CAN04 increase efficacy of chemotherapy regimes
- $\rightarrow$  CAN04 counteract weight loss after chemotherapy

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Synergy with chemotherapy in line with current development strategy <sup>13</sup>

## **Chemotherapy resistance**

- $\rightarrow\,$  Most chemotherapies induce chemoresistance already after a few months of therapy
- $\rightarrow\,$  Chemotherapy upregulate both IL-1 $\alpha$  and IL-1 $\beta$
- $\rightarrow\,$  Blocking IL-1 signalling counteracts chemoresistance in preclinical models
- $\rightarrow\,$  High blood levels of inflammatory cytokines IL-1 and IL-6 leads to poor gemcitabine efficacy in patients
- ightarrow IL-1 mediated chemoresistance for several classes of chemotherapy
  - ightarrow Platinum based chemotherapy, 5FU, Gemcitabine



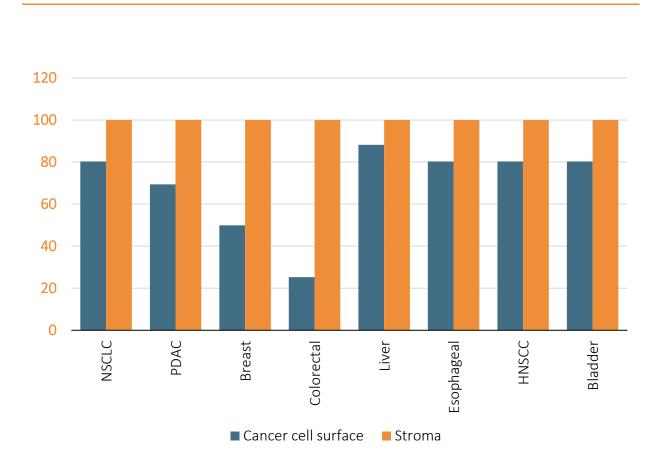
Several lines of evidence suggest CAN04 counteract chemoresistance



## CAN04 oncology expansion



# IL1RAP in several cancer with high medical need



IL1RAP



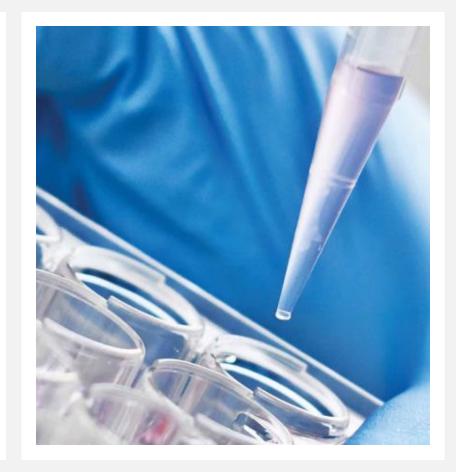
- $\rightarrow\,$  Biomarker studies ongoing, identify patients most likely to respond
- ightarrow Opportunity to expand development in additional cancer forms with high unmet medical need

### CAN04 development can be expanded to additional indications onwards <sup>16</sup>



# US phase I clinical trial

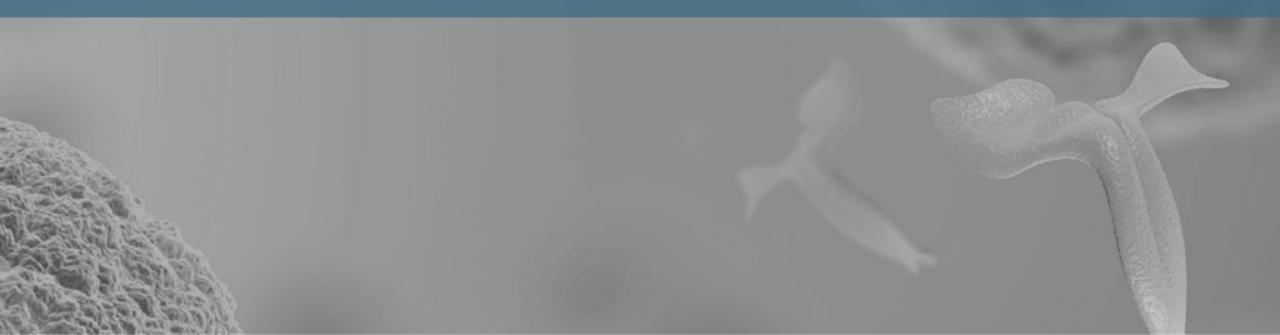
- ightarrow IND granted May 2020, FPI planned Q3 2020
- $\rightarrow$  Combination with checkpoint inhibitor in patients that no longer respond to PD1/PDL-1 therapy
- $\rightarrow\,$  Primary endpoint safety, secondary endpoints include biomarkers and efficacy
- → Indications include NSCLC, HNSCC, malignant melanoma and bladder cancer (18 patients)
- ightarrow Strong US centers, Coord investigator Prof Roger Cohen, UPenn
- $\rightarrow$  https://clinicaltrials.gov/ct2/show/NCT04452214



Trial designed to advance CAN04 outside chemotherapy combinations 17

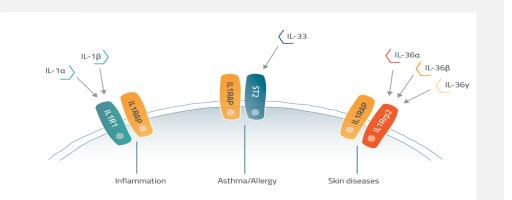


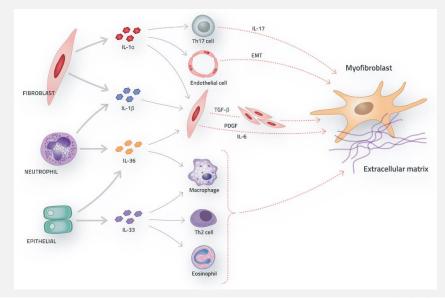
## Untapped possibilities in autoimmune diseases



## CAN10 – New development project

- $\rightarrow\,$  IL1RAP binding antibody potently blocking IL-1, IL-33 and IL-36
- $\rightarrow$  Unique anti-inflammatory activity observed in mouse model
- → 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
- ightarrow Clinical trials start early 2022





### Unique opportunity for CAN10 identified in life-threatening diseases



## Milestones and summary



# Significant value inflection points

## **Newsflow next 6-9 months**

### CAN04

- ightarrow FPI checkpoint combination clinical trial
- ightarrow LPI Phase IIa combination in PDAC and NSCLC
- ightarrow Phase IIa combination results PDAC and NSCLC
- ightarrow Next step combination therapy
- ightarrow Phase IIa biomarker/biopsy results
- ightarrow New clinical trial in disease/combination outside CANFOUR

### CAN10

- ightarrow Preclinical progress
- ightarrow Production development



## Significant data to secure newsflow



## Cantargia at a glance



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### Platform with many potential therapeutic areas

Cancer and large number of autoimmune/inflammatory diseases



### Vision of becoming an important part in future cancer treatments

Combination therapy strategy based on synergies with established therapies



### 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)



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