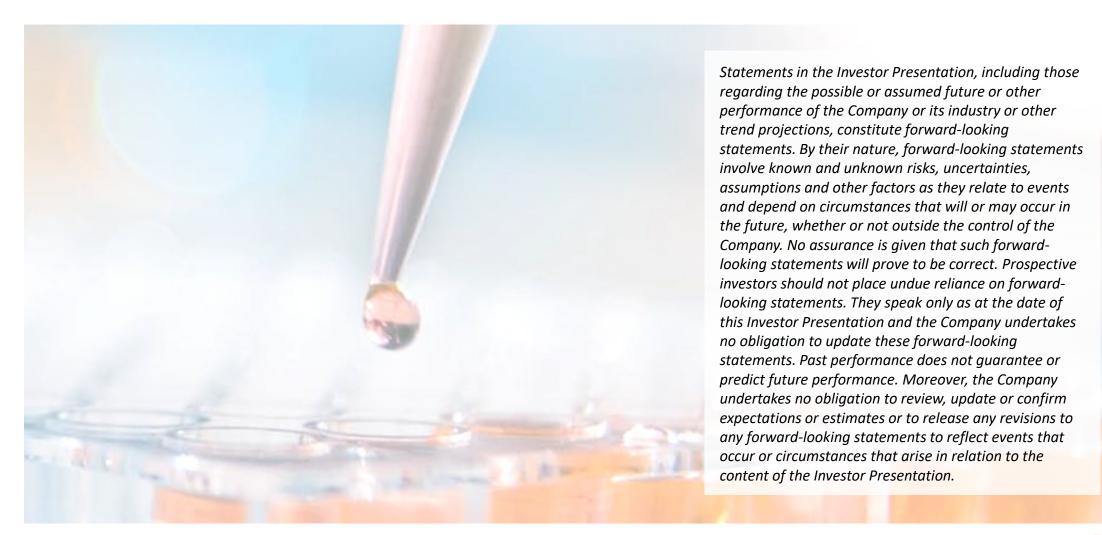


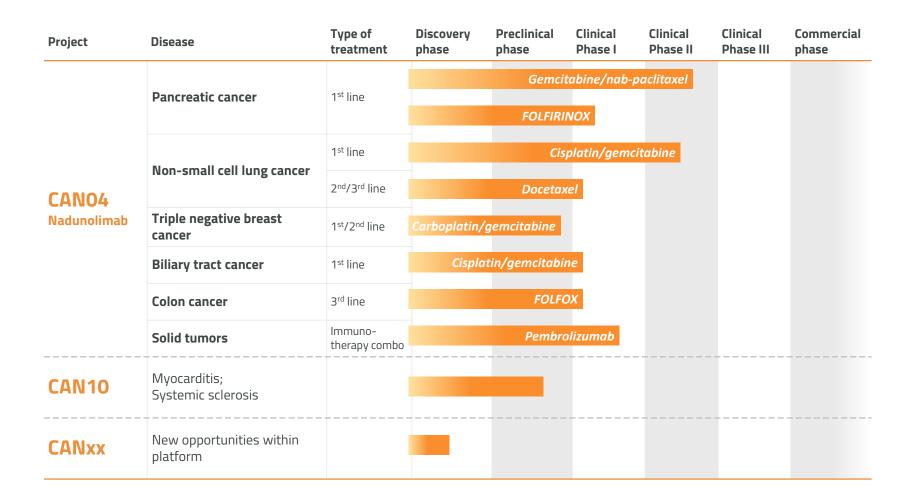
## Safe Harbour Statement







## 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 and further results during 2021



#### 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 ~12,000 SHAREHOLDERS AND LONG TERM INVESTORS

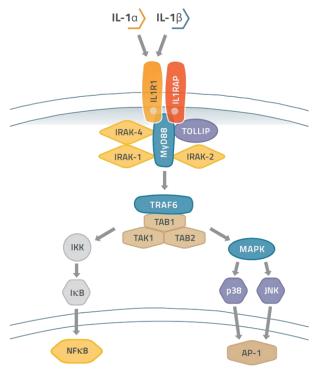
- Market cap: SEK 2.2bn (USD ~250m) (20 Sep-21)
- Cash: SEK 761m (USD 87m) (30 Jun-21)

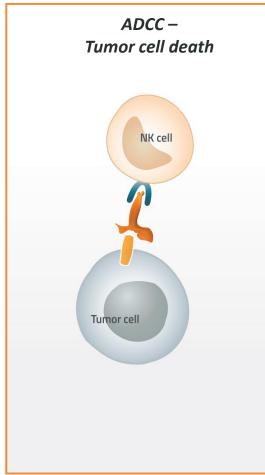
Current owners (30 Jun 2021)					
Swedbank Robur Funds	9.7%				
4th AP fund	8.7%				
Alecta	7.0%				
1st AP fund	6.3%				
Six Sis AG	5.7%				
Avanza Pension	4.4%				
SEB AB, Luxemburg	3.2%				
Sunstone LSV	3.0%				
Handelsbanken fonder	2.8%				
Unionen	2.0%				

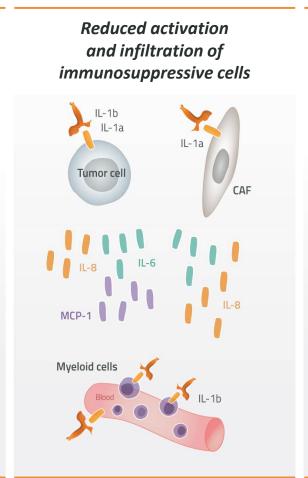


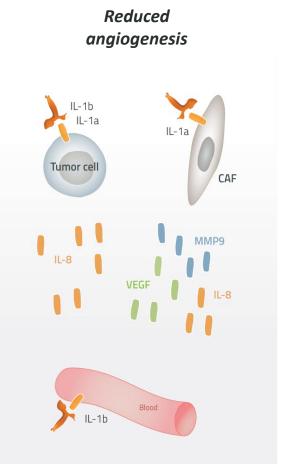


## CAN04 – Mechanism of action





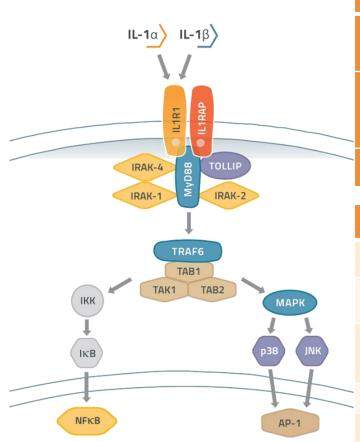




CAN04 BLOCKS BOTH FORMS OF IL-1 AND CAN ERADICATE CELLS MEDIATING THE EFFECTS OF IL-1



## CAN04 – Differentiated and superior MOA

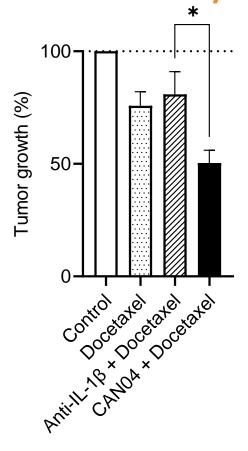


Cancer context	IL-1α	IL-1β	comment
Localization	<ul><li>Cellbound and soluble</li><li>Cancer cells and stroma</li></ul>	• Soluble	<ul> <li>IL-1α trigger and IL-1β enhance inflammation</li> <li>Often work in pair</li> </ul>
Function	<ul> <li>Stimulates inflammation - IL1R1 -</li> <li>IL-1, IL1R1 and IL1RAP 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



## CANO4 broad mechanism uniquely enhance docetaxel antitumor activity

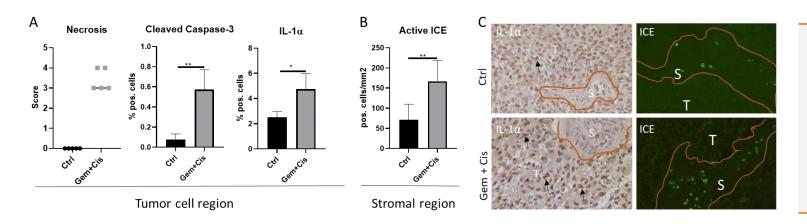


- → CAN04 in combination with docetaxel in MC38 syngeneic model
- → CAN04 increase efficacy of docetaxel
- $\rightarrow$  Control antibody blocking IL-1 $\beta$  did not have the same effect
- $\rightarrow$  In vitro experiment show docetaxel increase IL-1 $\alpha$  production
- → Highlight importance of blocking both forms of IL-1 to increase docetaxel efficacy
- → Clinical trial investigating CAN04 + docetaxel being initiated.

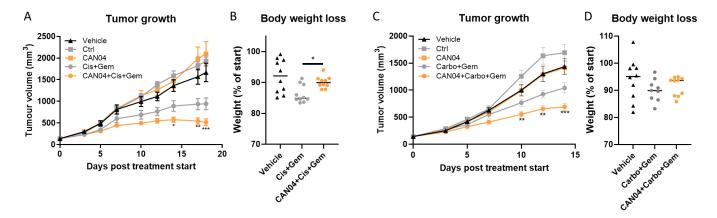
DIFFERENTIATING FROM IL-1B BLOCKADE, CAN04 INCREASE DOCETAXEL EFFICACY



# 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

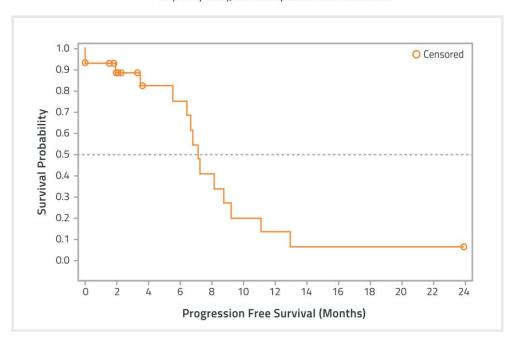
## Combination data in NSCLC show promising efficacy

#### Summary of key interim results

	Total NSCLC (27 pts)	Historical control <sup>1,2</sup>	Non-squamous NSCLC (15 pts)	Historical control <sup>3</sup>	Squamous NSCLC (11 pts)	Historical control <sup>4</sup>	PDAC (33 pts)	Historical control <sup>5</sup>
ORR	48%	22-28%	53%	19%	36%	38%	27%*	23%
PFS	7.2 mo	5.1 mo	NR**		NR**		7.8 mo	5.5 mo
Ongoing treatment	11 pts (41%)		6 pts (40%)		5 pts (45%)		7 pts (21%)	

<sup>\*15%</sup> additional patients benefit with a pseudoprogression-like response

<sup>\*\*</sup>NR (not reported); will be analyzed with more mature data



- → CAN04 in combination with gem/cis in 1<sup>st</sup> line chemotherapy
- → 13\* of 27 evaluable patients with non-sq non-small cell lung cancer (NSCLC) showed objective response including 1 complete response (48% vs historical control data 22-28%)
- → No major side effects observed except those from chemotherapy or CAN04 alone. Neutropenia frequency higher than expected from chemo (treated with dose reductions/GCSF)

## **DEVELOPMENT ADVANCING IN SEVERAL SEGMENTS OF NSCLC**



<sup>\*</sup>Incl 2 patients awaiting second conf scan

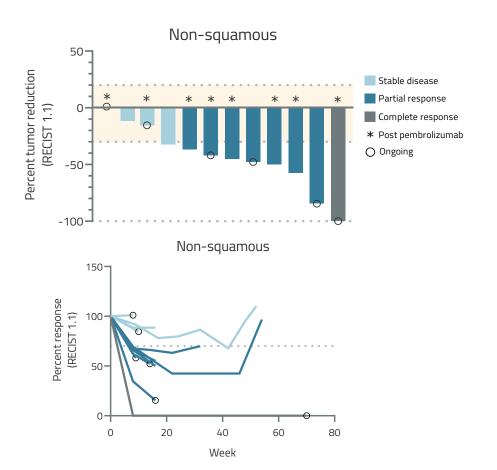
<sup>&</sup>lt;sup>1</sup> Schiller et al, N Engl J Med 2002; 346:92–98

<sup>&</sup>lt;sup>2</sup> Scagliotti et al, J Clin Oncol 2008; 26:3543–3551

<sup>&</sup>lt;sup>3</sup> Gandhi et al, N Engl J Med 2018; 378:2078-2092

<sup>&</sup>lt;sup>4</sup> Paz-Ares et al, N Engl J Med 2018; 379:2040-2051

## Strong signal in non-squamous NSCLC



- → CAN04 in combination with gem/cis in 1<sup>st</sup> line chemotherapy
- → 8 of 15 evaluable patients with non-sq non-small cell lung cancer (NSCLC) showed objective response including 1 complete response (53% vs historical control data 19%)
- → The complete response ongoing for >1.5 years
- → 8 patients were second line to pembrolizumab monotherapy, with 6 responses
- → No major side effects observed except those from chemotherapy or CAN04 alone. Neutropenia frequency higher than expected from chemo (treated with dose reductions/GCSF)



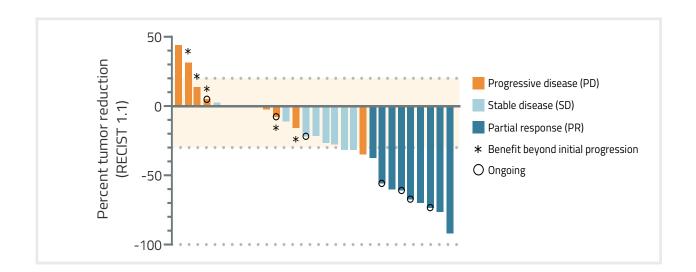
**DEVELOPMENT ADVANCING TOWARDS RANDOMIZED TRIAL END 2022** 

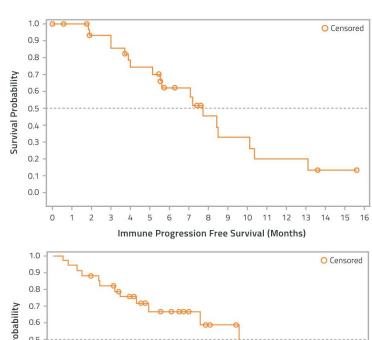


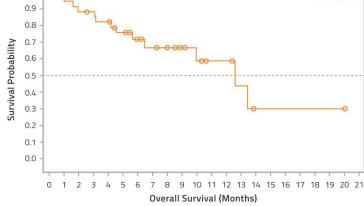
## Positive data in pancreatic cancer

CAN04 in combination with gem/abraxane in 1<sup>st</sup> line:

- Durable responses observed (median DOR 6.8 mo, 27% response rate)
- Important finding of pseudoprogression-like response in 5 (15%) patients predicting long PFS.
- Promising PFS (7.8 mo) and OS (12.6 mo, 42 % events), seven patients still on treatment

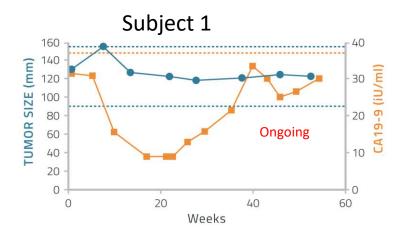


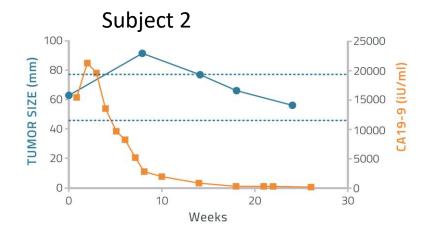


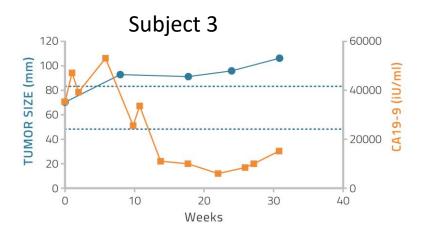


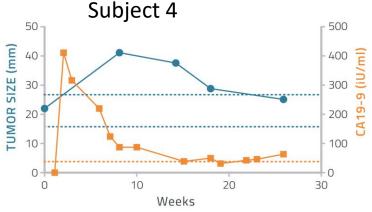
## Patients with Pseudoprogression-like response

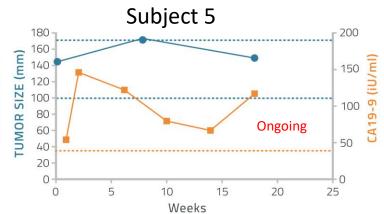
- All presented PD at 1<sup>st</sup> CT scan evaluation (8 weeks)
- All showed concomitant reduction of CA19-9











## 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 neuropathy<sup>2</sup> (nabpaclitaxel or oxaliplatin) can be mediated by IL-1 blockade.

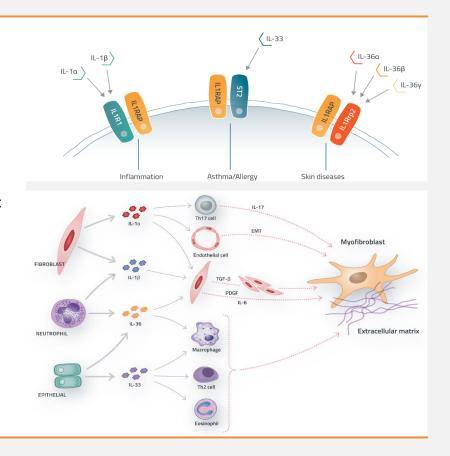
- G-CSF not used proactively/prophylactically in this trial. In later trials, G-CSF counteracts neutropenia.
- Median duration of treatment 4.8 months (reference 3.9 months)
- Most common reasons for termination: gastrointestinal events or general health deterioration





## CAN10 – New development project

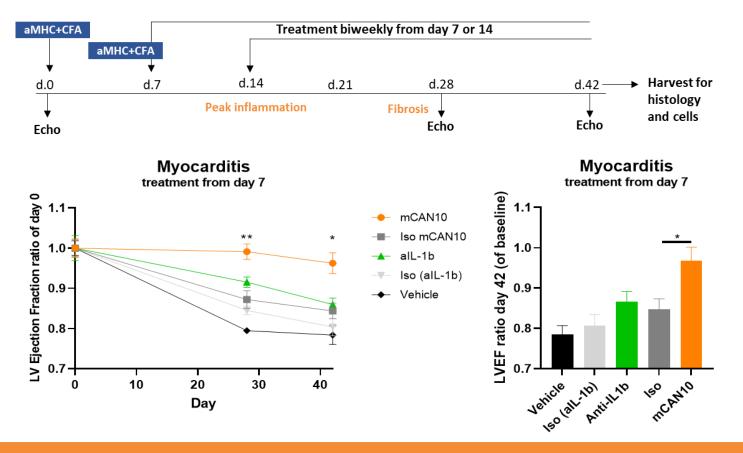
- → IL1RAP binding antibody potently blocking IL-1, IL-33 and IL-36
- Unique anti-inflammatory activity observed in different mouse models (myocarditis, psoriasis, inflammation)
- → Development focusing on unmet medical need in systemic sclerosis and myocarditis. Disease selection in collaboration with 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

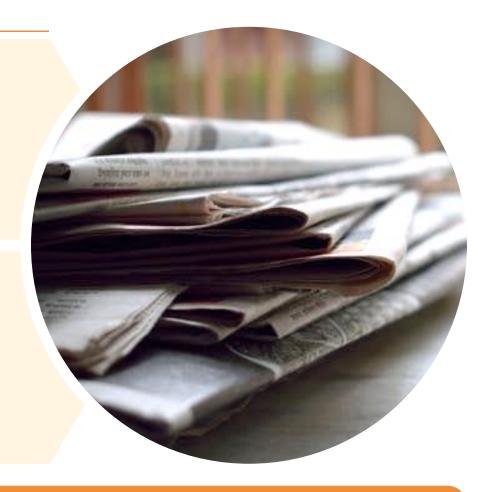
## **Newsflow over next 6 months**

## Nadunolimab (CAN04)

- → New results PDAC, NSCLC and Keytruda combination
- → Randomized trials PDAC and NSCLC
- → New preclinical and translational results
- → New clinical trials
  - CAPAFOUR FOLFIRINOX combination PDAC
  - CESTAFOUR Basket trial (NSCLC, CRC, BTC)
  - TRIFOUR TNBC

### CAN10

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



SIGNIFICANT DATA TO SECURE NEWSFLOW



## Cantargia highlights



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



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Combination therapy strategy based on synergies with established therapies



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Cancer and large number of autoimmune/inflammatory diseases



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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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