UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
of the Securities Exchange Act of 1934

For the month of June 2021
Commission File Number: 001-37643

PURPLE BIOTECH LTD.
(Translation of registrant's name into English)

4 Oppenheimer Street, Science Park, Rehovot 7670104, Israel
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the Registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ☐

Indicate by check mark if the Registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ☐

On June 4, 2021, Purple Biotech Ltd. (the "Company" or the "Registrant") issued a press release, "**Purple Biotech Presents New Clinical Data from NT219 at the 2021 ASCO Annual Meeting**", which is attached hereto as Exhibit 99.1. The Company also announced that it has made available an updated Company Presentation on its website. A copy of the updated Company Presentation is attached hereto as Exhibit 99.2 and may be viewed at the Company's website at www.purple-biotech.com.

Exhibits

99.1	Press Release
99.2	Purple Bio Company Presentation June 2021

Incorporation by Reference

This Form 6-K, including all exhibits attached hereto, is hereby incorporated by reference into each of the Registrant's Registration Statement on Form S-8 filed with the Securities and Exchange Commission on May 20, 2016 (Registration file number 333-211478), the Registrant's Registration Statement on Form S-8 filed with the Securities and Exchange Commission on June 6, 2017 (Registration file number 333-218538), the Registrant's Registration Statement on Form F-3, as amended, originally filed with the Securities and Exchange Commission on July 16, 2018 (Registration file number 333-226195), the Registrant's Registration Statement on Form S-8 filed with the Securities and Exchange Commission on March 28, 2019 (Registration file number 333-230584), the Registrant's Registration Statement on Form F-3 filed with the Securities and Exchange Commission on September 16, 2019 (Registration file number 333-233795), the Registrant's Registration Statement on Form F-3 filed with the Securities and Exchange Commission on December 2, 2019 (Registration file number 333-235327), the Registrant's Registration Statement on Form F-3 filed with the Securities and Exchange Commission on May 13, 2020 (Registration file number 333- 238229), the Registrant's Registration Statement on Form S-8 filed with the Securities and Exchange Commission on May 28, 2020 (Registration file number 333-238481) and each of the Registrant's Registration Statements on Form F-3 filed with the Securities and Exchange Commission on July 10, 2020 (Registration file numbers 333-239807 and 333-233793), to be a part thereof from the date on which this report is submitted, to the extent not superseded by documents or reports subsequently filed or furnished.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

June 4, 2021

PURPLE BIOTECH LTD.

By: */s/ Isaac Israel*
Isaac Israel
Chief Executive Officer

Exhibit 99.1

Purple Biotech Presents New Clinical Data from NT219 at the 2021 ASCO Annual Meeting

NT219 was Well-tolerated with Minimal Adverse Events in Initial Clinical Data from Ongoing Phase 1/2 Clinical Trial in Adults with Advanced Solid Tumors

Partial Response Observed in a Patient with Refractory Gastroesophageal Junction Cancer

REHOVOT, Israel, June 4, 2021 - Purple Biotech Ltd. ("Purple Biotech", or the "Company") (NASDAQ/TASE: PPBT), a clinical-stage company developing first-in-class, effective and durable therapies by overcoming tumor immune evasion and drug resistance, announced today the presentation of new data from the first dose level cohort of its ongoing Phase 1/2 clinical trial of NT219, at the 2021 ASCO Annual Meeting, being held virtually June 4-8, 2021. The Phase 1/2 study is evaluating NT219 as monotherapy for the treatment of solid tumors, in addition to a subsequent dose escalation of NT219 in combination with cetuximab, an epithelial growth factor receptor (EGFR) blocking monoclonal antibody, for the treatment of recurrent and/or metastatic solid tumors and squamous cell carcinoma of the head and neck cancer.

As of the cutoff date (April 25th, 2021), six patients have been enrolled into the study, including three subjects with advanced solid tumors in the first cohort receiving 3 mg/kg of NT219 as a single agent, and three subjects in the second cohort receiving 6mg/kg of NT219 as monotherapy.

Initial results from the first dose level cohort revealed NT219 was well-tolerated with minimal adverse events. In addition, a partial response was observed in a patient with refractory gastroesophageal junction cancer, previously treated with four prior lines of therapies. For this patient, who has been treated for 22 weeks, a complete remission was seen at the largest target lesion and at one non-target lesion, while stable disease was observed at the other non-target lesion.

"We are encouraged by these initial safety and efficacy results from this first-in-human study of NT219," said Alberto Bessudo, M.D., a medical oncologist and hematologist at California Cancer Associates for Research & Excellence, who presented the data at ASCO. "This study is especially compelling because NT219 uniquely targets the IRS protein to degradation by utilizing a covalent, irreversible inhibition strategy. Based on the preclinical results observed to date, by targeting the IRS1/2 and STAT3 pathways NT219 has the potential to significantly shrink tumors, prevent and reverse tumor resistance when administered as a monotherapy, as well as in combination with existing oncology therapies."

"Our innovative approach to overcoming tumor immune evasion and drug resistance is focused on interactions within the microenvironment of the tumor, not just targeting the tumor as an isolated feature," said Bertrand C. Liang, M.D., Ph.D., Chief Medical Officer of Purple Biotech. "It has been demonstrated clinically that IRS and STAT3 proteins are central in defining tumor responsiveness to therapy. Indeed, STAT3 has been seen to play a role in treatment response, and IRS is a novel target with recent promising findings and interest in oncology. While still early in the trial, we are very excited by these initial clinical results and expect to report additional data from higher dose levels of this study in the second half of this year."

NT219 is a dual inhibitor, novel small molecule that simultaneously targets IRS1/2 and STAT3, known important oncogenic drivers and major drug resistance pathways in hard-to-treat cancers. The primary objectives of the open-label Phase 1/2 trial are to evaluate safety, assess pharmacokinetics, identify the recommended dose to be studied in the Phase 2 portion, and establish preliminary efficacy of NT219. The Phase 1 portion of the study will encompass a dose escalation evaluation of NT219 monotherapy administered weekly in patients with refractory advanced solid tumors. Upon reaching the third dose level of NT219, a second cohort of patients, with recurrent or metastatic squamous cell carcinoma of the head and neck or colorectal adenocarcinoma, will be administered weekly with NT219, and dose escalated, in combination with cetuximab.

About Purple Biotech

Purple Biotech Ltd. is a clinical-stage company developing first-in-class therapies by overcoming tumor immune evasion and drug resistance. The Company's oncology pipeline includes NT219 and CM24. NT219 is a dual inhibitor, novel small molecule that simultaneously targets IRS1/2 and STAT3. The Company is currently advancing NT219 as a monotherapy treatment of solid tumors, followed by a dose escalation of NT219 in combination with cetuximab for the treatment of recurrent and/or metastatic squamous cell carcinoma of the head and neck cancer (SCCHN) or colorectal adenocarcinoma in a phase 1/2 study, and an expansion phase of NT219 at its recommended phase 2 level in combination with cetuximab in patients with recurrent and/or metastatic squamous cell carcinoma of the head and neck. CM24 is a humanized monoclonal antibody that blocks CEACAM1, an immune checkpoint protein that supports tumor immune evasion and survival through multiple pathways. The Company is advancing CM24 as a combination therapy with anti-PD-1 checkpoint inhibitors in selected cancer indications in a phase 1b study followed by a phase 2 for the treatment of non-small cell lung cancer and pancreatic cancer. The Company has entered into a clinical collaboration agreement, as amended, with Bristol Myers Squibb for the planned phase 1/2 clinical trials to evaluate the combination of CM24 with the PD-1 inhibitor nivolumab (Opdivo®) in patients with non-small cell lung cancer and in combination with nivolumab in addition to nab-paclitaxel (ABRAXANE®) in patients with pancreatic cancer. The Company is also the owner of Consensi®, an FDA-approved fixed-dose combination of celecoxib and amlodipine besylate, for the simultaneous treatment of osteoarthritis pain and hypertension that was approved by the FDA for marketing in the U.S. Consensi® is being sold in the U.S. by Burke Therapeutics, the marketing partner of the Company's U.S. distributor, Coeptis Pharmaceuticals. The Company has also partnered to commercialize Consensi in China and South Korea. The Company's corporate headquarters are located in Rehovot, Israel. For more information, please visit https://www.purple-biotech.com.

Forward-Looking Statements and Safe Harbor Statement

Certain statements in this press release that are forward-looking and not statements of historical fact are forward looking statements within the meaning of the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Such forward-looking statements include, but are not limited to, statements that are not statements of historical fact, and may be identified by words such as "believe", "expect", "intend", "plan", "may", "should", "could", "might", "seek", "target", "will", "project", "forecast", "continue" or "anticipate" or their negatives or variations of these words or other comparable words or by the fact that these statements do not relate strictly to historical matters. You should not place undue reliance on these forward-looking statements, which are not guarantees of future performance. Forward-looking statements reflect our current views, expectations, beliefs or intentions with respect to future events, and are subject to a number of assumptions, involve known and unknown risks, many of which are beyond our control, as well as uncertainties and other factors that may cause our actual results, performance or achievements to be significantly different from any future results, performance or achievements expressed or implied by the forward-looking statements. Important factors that could cause or contribute to such differences include, among others, risks relating to: the plans, strategies and objectives of management for future operations; product development for NT219 and CM24; the process by which early stage therapeutic candidates such as NT219 and CM24 could potentially lead to an approved drug product is long and subject to highly significant risks, particularly with respect to a joint development collaboration; the fact that drug development and commercialization involves a lengthy and expensive process with uncertain outcomes; our ability to successfully develop and commercialize our pharmaceutical products; the expense, length, progress and results of any clinical trials; the impact of any changes in regulation and legislation that could affect the pharmaceutical industry; the difficulty in receiving the regulatory approvals necessary in order to commercialize our products; the difficulty of predicting actions of the U.S. Food and Drug Administration or any other applicable regulator of pharmaceutical products; the regulatory environment and changes in the health policies and regimes in the countries in which we operate; the uncertainty surrounding the actual market reception to our pharmaceutical products once cleared for marketing in a particular market; the introduction of competing products; patents obtained by competitors; dependence on the effectiveness of our patents and other protections for innovative products; our ability to obtain, maintain and defend issued patents; the commencement of any patent interference or infringement action against our patents, and our ability to prevail, obtain a favorable decision or recover damages in any such action; and the exposure to litigation, including patent litigation, and/or regulatory actions, and other factors that are discussed in our Annual Report on Form 20-F for the year ended December 31, 2020 and in our other filings with the U.S. Securities and Exchange Commission ("SEC"), including our cautionary discussion of risks and uncertainties under "Risk Factors" in our Registration Statements and Annual Reports. These are factors that we believe could cause our actual results to differ materially from expected results. Other factors besides those we have listed could also adversely affect us. Any forward-looking statement in this press release speaks only as of the date which it is made. We disclaim any intention or obligation to publicly update or revise any forward-looking statement or other information contained herein, whether as a result of new information, future events or otherwise, except as required by applicable law. You are advised, however, to consult any additional disclosures we make in our reports to the SEC, which are available on the SEC's website, https://www.sec.gov.

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Exhibit 99.2


PURPLE
BIOTECH
CORPORATE PRESENTATION
NASDAQ/TASE: PPBT
June 2021

Forward-looking Statements and Safe Harbor

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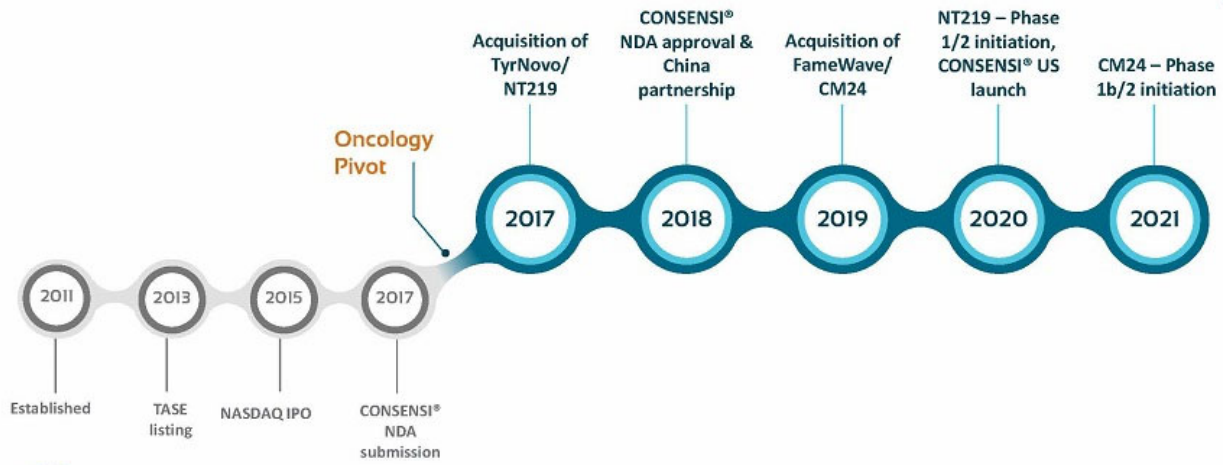
Our Transformation into Oncology



Business Highlights

CM24 - First-in-class α-CEACAM1 mAb, validating clinical collaboration with Bristol Myers Squibb

NT219 - First-in-class, small molecule, dual inhibitor of IRS 1/2 and STAT3

H2:21 - Two phase 1 study readouts

Strong balance sheet and cash position

Purple Biotech (NASDAQ/TASE: PPBT)

ADSs outstanding: 17.5M

$61M cash as of December 31st, 2020

Cash runway through 2024

Advancing Clinical-stage Novel Oncology Therapies

Program	Indication	Preclinical	Phase 1	Phase 2	Phase 3	Partner	Value Drivers
CM24 (CEACAM-1)	Solid tumors (monotherapy) (Completed) Solid tumors (combination with nivolumab) NSCLC (combination with nivolumab) Pancreatic Cancer (combination with nivolumab and nab-paclitaxale)					Bristol Myers Squibb	Expansion arms on RP2D: Initiation Q4:21 **Topline data: H2:21**
NT219 (IRS1/2 & STAT3)	Solid tumors (monotherapy) R/M SCCHN & CRC (dose escalation with cetuximab); R/M SCCHN (expansion - combination with cetuximab on RP2D)						Expansion arms on RP2D: Initiation Q4:21 **Topline data: H2:21**

Expansion | Done / Ongoing | Planned study

Multiple data read-outs expected in the next 12 months

Experienced Leadership

Isaac Israel
Chief Executive Officer
Former CEO of BeeContact Ltd. (TASE:BCNT). NextGen Biomed (TASE: NXGN)

Eric K. Rowinsky, MD
Chairman of the Board
Former CMO at ImClone, Stemline, Board member at Biogen Inc.

Gil Efron
Deputy CEO and Chief Financial Officer
Former Deputy CEO & CFO at Kamada (NASDAQ:KMDA)

Hadas Reuveni, Ph.D
Vice President, R&D
Formerly at Keryx (NASDAQ:KERX)

Bertrand Liang, MD, Ph.D, MBA/AMP, FAAN
Chief Medical Officer
Formerly at Biogen Idec, Amgen, NCI

Michael Schickler, Ph.D
Head of Clinical & Regulatory Affairs
Formerly at Hoffmann-La Roche, CEO at CureTech

Advancing First-in-Class Oncology Therapies

CM24 - an α-CEACAM1 mAb

CEACAM1* Plays a Key Role in Cancer Biology

01 | ADHESION

Horst, 2011

Oncogene

*"CEACAM1 creates a pro-angiogenic tumor microenvironment that **supports tumor vessel maturation**"*

Ferri, 2020

The Journal of Immunology

*"Neutrophil extracellular trap-associated CEACAM1 as a putative therapeutic target to **prevent metastatic progression** of colon carcinoma"*

02 | IMMUNE CELLS/ IMMUNE EXCLUSION

Tsuzuki, 2020

*"**Immune-checkpoint molecules** on regulatory T-cells as a potential therapeutic target in head and neck squamous cell cancers"*

Tsang, 2020

Cancer Biotherapy and Radiopharmaceuticals

"[Blockade] ***enhances natural killer cell cytotoxicity*** *against tumor cells through blockade of the inhibitory CEACAM1 / CEACAM5 immune checkpoint pathway"*

03 | IMMUNO-ONCOLOGY

Blumberg, 2015

nature

*"CEACAM1 **regulates TIM-3**-mediated tolerance and exhaustion"*

Shively, 2013

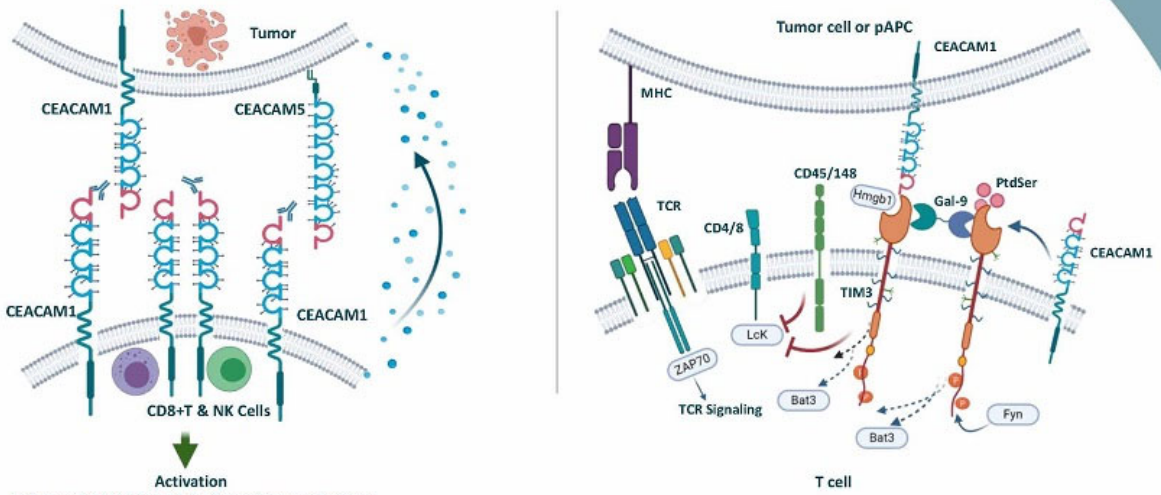
Experimental Cell Research

*"CEACAM1 **regulates Fas-mediated apoptosis** in Jurkat T-cells via its interaction with β-catenin"*

PURPLE

*Carcinoembryonic Antigen Cell Adhesion Molecule

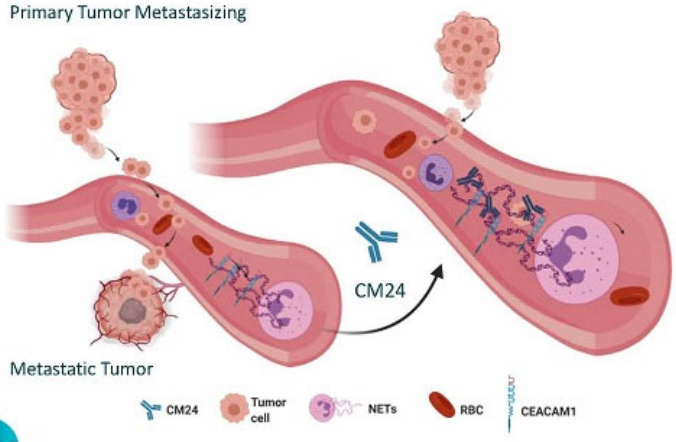
CM24 MOA | Immuno-oncology

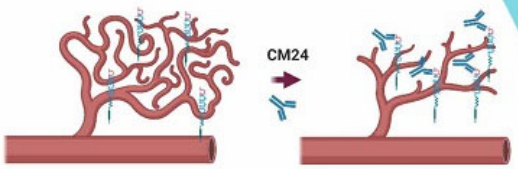


Markel et al, J Immunol 2002, 2006; Immunology, 2008; Cancer Immunol Immunother 2010; Ortenberg et al, Mol Cancer Ther 2012; Zhou, 2009; Li, 2013; Huang, 2015; Acharya N, et al. J Immunotherapy Canc 8:e911-22, 2020.

CM24 MOA | Adhesion

Neutrophil extracellular trap-associated CEACAM1 as a putative therapeutic target to prevent metastatic progression:



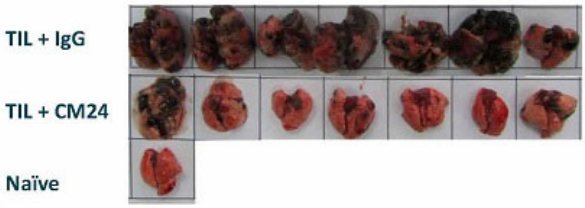


CEACAM1 creates a pro-angiogenic tumor microenvironment that supports tumor vessel maturation. CM24 is proposed to attenuate/block tumor growth and metastatic processes.

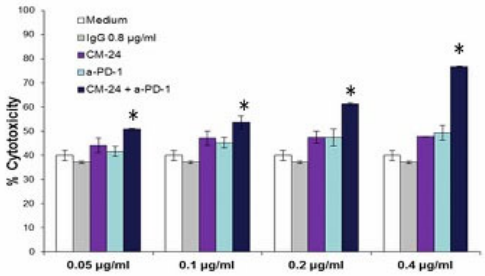
Rayes RF, et al. Neutrophil Extracellular Trap-Associated CEACAM1 as a Putative Therapeutic Target to Prevent Metastatic Progression of Colon Carcinoma. J Immunol. 2020; Gerstel, D. et al. CEACAM1 creates a pro-angiogenic tumor microenvironment that supports tumor vessel maturation. Oncogene 30, 4275–4288 (2011)

Anti-cancer Effect Following Treatment

Preclinical Data With CM24 + TIL and CM24 + α-PD1



- Xenograft, lung lesion, Mel 526 (IV)
- Tumor burden was monitored 26 days post last CM24 treatment



Combination index (CI) = 0.15

$$CI = \frac{(D)_1}{(D_x)_1} + \frac{(D)_2}{(D_x)_2} < 1 \rightarrow \text{synergy}$$

Significant benefits as both single agent and in combination with α-PD-1

CM24 Phase 1 Monotherapy Trial

UCLA | Yale-New Haven Hospital

- Open-label, multi-dose escalation monotherapy study to assess safety and tolerability
- Heavily pre-treated patients with a median of 4 prior regimens (range 2-8) Overall, treatment was well tolerated
- Most of the responding patients were treated with the highest dose levels of 3mg/kg & 10mg/kg
- PK analysis revealed non-linearity, modeling recommended to continue administration of higher doses to reach saturation

27 patients:

Colorectal	11
Melanoma	7
Ovarian	4
Gastric	3
NSCLC	2

**24 patients evaluated*

	# pts
0.01mg/kg	1
0.03mg/kg	1
0.1 mg/kg	3
0.3 mg/kg	3
1.0 mg/kg	3
3.0 mg/kg	9
10.0 mg/kg	7

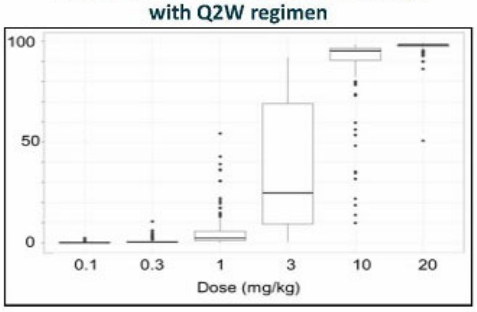
PURPLE

No DLTs up to 10 mg/kg	No discontinuation of study drug due to an AE	No drug related mortalities	33.3% SD (RECIST)

PK/PD Modeling Provides Dosage & Schedule Guidance

- Consistent with observed PK showing high clearance at doses <10 mg/kg, plot shows low TMDD saturation at low doses
- 10 mg/kg has a broad range of saturation, and >10 mg/kg, Q2W dose is needed for saturation across population
- Nivolumab administered Q2W or Q4W, representing good clinical and commercial fit for CM24

Simulated TMDD[1] saturation at Ctrough with Q2W regimen



The Phase 1b/2a study will continue escalating the CM24 dose above 10mg/kg q2wk, in combination with nivolumab

[1]Target-mediated drug disposition. [2] OPDIVO® is a registered trademark of Bristol-Myers Squibb. Ctrough is the drug concentration reached by CM24 before the next dose is administered

Large Market Opportunity in NSCLC & Pancreatic Cancer

- NSCLC accounts for ~200K new cases/year in the US; with a 5-year relative survival rate of 23%[2]
- Immunotherapy is now recommended as first line therapy in all patients with NSCLC and no driver mutations[3]
- 5-year overall survival rates with chemotherapy in 2L is 2.6% and with I/O Opdivo® is 13.4%[4]

- Pancreatic Cancer accounts for ~60K new cases/year in the US; with a 5-year relative survival rate of 10%[2]
- I/O approaches have been limited to patients with high microsatellite instability (MSI-H) or high tumor mutational burden (TMB-H)
- 5-year overall survival rates with chemotherapy in 2L is 3%[2]

Combining nivolumab with CM24 in a clinical collaboration with

Bristol Myers Squibb™

- CEACAM1 expression correlates with poor prognosis in NSCLC and Pancreatic cancer[1]
- Preclinical data support significant synergy
- Receptor saturation with CM24 is better achieved with a Q2W regimen, aligning with the nivolumab regimen

1 Dango et al, Lung Cancer 2008; 60:426 & Calinescu et al, Journal of Immunology Research 2018: 7169081.
2 American Cancer Society, Cancer Facts & Figures 2019, and the ACS website, https://seer.cancer.gov/statfacts/html/pancreas.html
3 Economopoulou P, Mountzios G. The emerging treatment landscape of advanced non-small cell lung cancer. Ann Transl Med. 2018;6(8):138. doi:10.21037/atm.2017.11.07
4 Gettinger S, et al "Five-year outcomes from the randomized, phase III trials CheckMate 017/057: Nivolumab vs docetaxel in previously treated NSCLC" WCLC 2019; Abstract OA14.04.

CM24 Phase 1/2 Combination Study Design (NCT04731467)

A Phase 1/2 open label multi center study of CM24 in combination with:

- Nivolumab in selected cancer indications (Phase 1) and NSCLC (Phase 2)
- Nivolumab and nab-paclitaxel in pancreatic cancer (Phase 2)

Primary endpoints:

Phase 1: Evaluate the safety, PK and the MTD/RP2 dose

Phase 2: Evaluate preliminary efficacy in 2nd line NSCLC and pancreatic cancer

Measurement of CEACAM1 based bio-marker.

Exploring further studies in other tumor types as well as monotherapy

2021	2022	2023-24
Dose Escalation	**Expansions**	
Doses: 10, 15, 20mg/kg q2wk + nivolumab (480mg q4w) 3+3 design $9 \leq n \leq 18$ Indications: NSCLC, Pancreatic, Ovarian, CRC, Melanoma, Papillary Thyroid Carcinoma	CM24 (@RP2 dose) + nivolumab (480mg) q4w I/O refractory NSCLC; 2nd line n=13+14 (Simon 2 Stage Design) CM24 (@RP2 dose) + nivolumab (480mg) q4w + nab-paclitaxel Locally advanced, unresectable pancreatic cancer; 2nd line n=13+14 (Simon 2 Stage Design)	Clinical collaboration with: Bristol Myers Squibb™

PURPLE

Advancing First-in-Class Oncology Therapies

NT219 – A Small Molecule Dual Inhibitor of IRS 1/2 and STAT3

NT219 - Dual Inhibitor of IRS1/2 & STAT3

IRS1/2

- Scaffold proteins, mediating mitogenic, metastatic, angiogenic and anti-apoptotic signals from IGF1R, IR, IL4R and other oncogenes, overexpressed in multiple tumor types
- Regulates major survival pathways such as the PI3K/AKT, MEK/ERK and WNT/β-catenin
- Activated as a feedback response to anti-cancer therapies

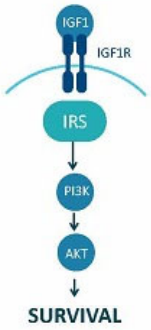
STAT3

- Well-established transcription factor associated with the tumorigenic phenotype
- Provides a crucial axis to support cell proliferation and survival
- Active in tumor JAK/STAT3 and TGF-β resistance mechanisms

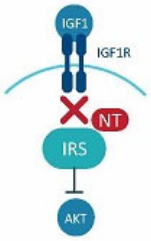
[1]Hodas Reuveni et al.; Cancer Res 2013;73:4383-4394. 2013 ; [2]Machado-Neto, et al. Clinics (Sao Paulo, Brazil) vol. 73,suppl 1 e566s. 11 Oct. 2018, doi:10.6061/clinics/2018/e566s
[3]Naokazu Ibuki1,2, Mazyar Ghaffari1,3, Hodas Reuveni4 et al. DOI: 10.1158/1535-7163.MCT-13-0842 Published December 2014; [4]Ramplas T, Favlochia R, Stebbing J, Giamas G. 2016 May 19;35(20):2562-4. doi: 10.1038/onc.2015.392. Epub 2015 Oct 19. PMID: 26477311
[5]Yashner-Abramsonet al., Oncogene. 2016 May 19;35(20):2675-80. doi: 10.1038/onc.2015.229. Epub 2015 Jun 29. PMID: 26119932, [6]Sanchez-Lopez E,. Oncogene. 2016 May 19;35(20):2634-44. doi: 10.1038/onc.2015.326. Epub 2015 Sep 14. PMID: 26364612; PMCID: PMC4791217.
[7]Zhao C, et al. Trends Pharmacol Sci. 2016 Jun;37(1):47-61. doi: 10.1016/j.tips.2015.10.001. Epub 2015 Nov 12. PMID: 26576830, [8] Johnson, Daniel E et al. "Targeting the IL-6/JAK/STAT3 signaling axis in cancer." Nature reviews. Clinical oncology vol. 15,4 (2018): 234-248. doi:10.1038/nrclinonc.2018.8

Novel MOA: IRS Degradation By NT219

Blocking IGF1R-AKT Pathway[1]

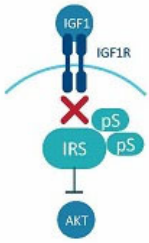


1 Binding to IRS



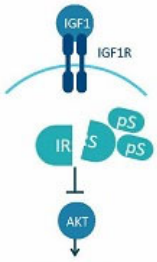
Covalent binding to IRS1/2 leads to the dissociation of IRS1/2 from IGF1R

2 Ser-phosphorylation



Serine phosphorylation prevents re-binding of IRS1/2 to the receptor

3 Degradation



APOPTOSIS

IRS1/2 is degraded by the proteasome

[1]*Reuveni et al. Cancer Res 2013*

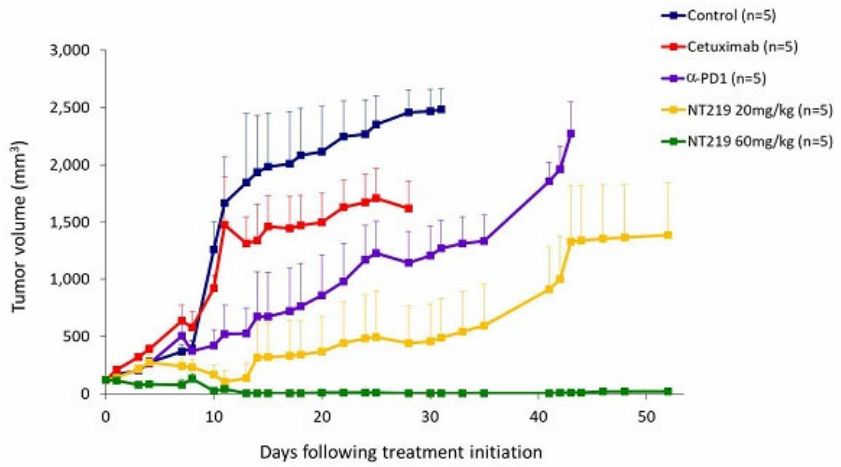
NT219 Efficacy as Monotherapy

Animal model

Head & Neck Cancer (SCC-9) NSG™, PBMCs-injected[1]

Drugs

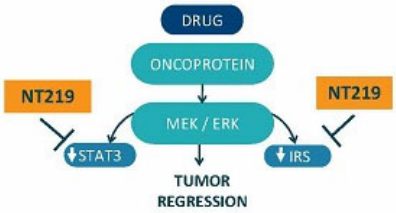
α-PD1
Cetuximab (Erbitux®)
NT219 20mg/kg
NT219 60mg/kg



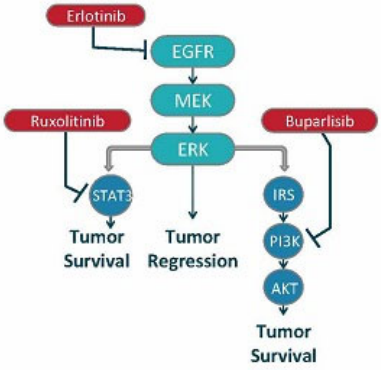
*1 NSG mice were injected SC with SCC-9 cells. PBMCs (18*10⁶ cells per mouse) administered 4 weeks prior to first treatment. NT219, α-PD1, and cetuximab were administered IV (NT219) and IP (α-PD1 and cetuximab) twice a week for 4 weeks. Graph reflects relevant data adapted from 2020 Multidisciplinary Head and Neck Cancers Symposium Poster presentation*

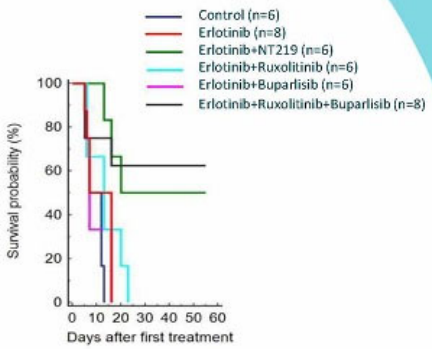
STAT3 and IRS are Essential in Therapeutic Resistance

Blocking survival pathways



Proof of Concept: PDX model of Head and Neck Cancer





By blocking both STAT3 and IRS pathways, NT219 prevents tumor resistance and re-sensitizes tumors to anti-cancer therapies

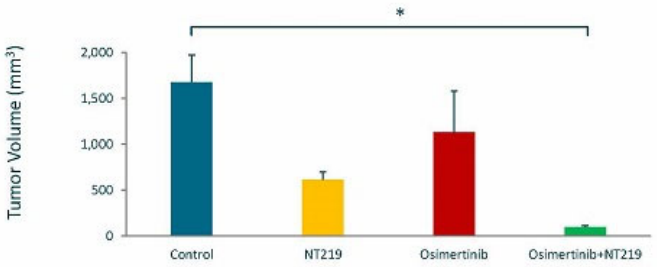
NT219 + Targeted Therapies Established Efficacy in PDX Models

NSCLC

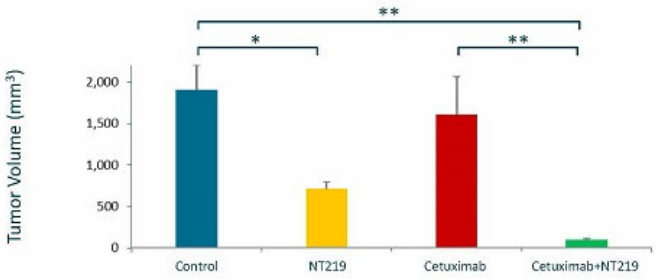
Exon 19 deletion EGFR and T790M, biopsy of bone marrow metastasis, patient previously progressed on afatinib and osimertinib

R/M SCCHN

metastasis, patient previously progressed on chemoradiation, several chemotherapies, and pembrolizumab



Osimertinib 5 mg/kg, NT219 65 mg/kg, mean tumor volume at the end point, 3 mice/group;



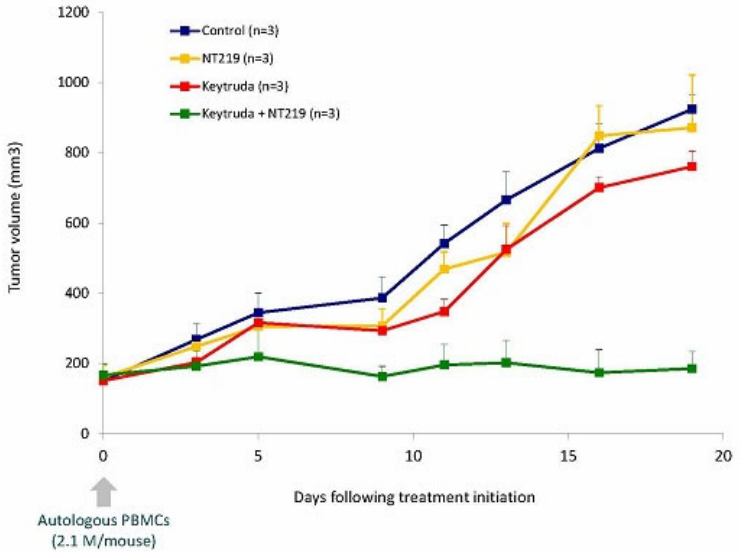
Treatments on days 0, 3 and 10, cetuximab - 1mg/mouse, 3 mice/group; PBMCs (1.4M cells/mouse) were injected on day 621

*** p<0.01, * p<0.02 based on one-way ANOVA with post hoc Tukey's HSD test*

NT219 + α-PD1 Re-sensitizes to Refractory α-PD1 Tumors

PDX Model
Humanized PDX of Esophagus Cancer (refractory to pembrolizumab)

Drug
Pembrolizumab (Keytruda®)



** Double autologous model - Tumors & PBMCs are from the same patient (#RA236) | Keytruda - 6mg/kg IP, NT219 - 60mg/kg IV*

First Market Opportunity

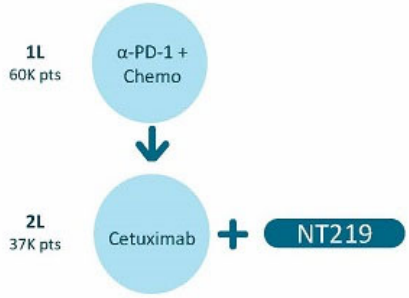
Recurrent or Metastatic Squamous Cell Carcinoma of Head and Neck (SCCHN)

Targeting the unmet medical need

- 1L Standard of care has shifted from chemotherapy towards immuno-oncology + chemotherapy
- < 20% of R/M SCCHN patients respond to α-PD1s
- 175k new cases/year are expected by 2024

Rationale for combining Cetuximab + NT219

- EGFR and PD(L)-1 are the only clinically validated targets in SCCHN
- < 15% of R/M SCCHN patients respond to Cetuximab
- Cetuximab inhibits EGFR signaling and promotes ADCC in EGFR expressing tumors
- STAT3 and IRS-to-AKT activation contributes to resistance to cetuximab in SCCHN



NT219 + Cetuximab has the potential to become an attractive 2nd line therapy

Global Data 2018: Head and Neck Squamous Cell Carcinoma: Opportunity Analysis and Forecasts to 2026 ; Internal best current estimates of patient numbers based on external research, 5 major global territories

NT219 Monotherapy and Combination Phase 1/2 Study Design (NCT04474470)

Study title

A phase 1/2 study with open-label, dose escalation phase followed by single-arm expansion to assess the safety, tolerability, PK, PD and efficacy of NT219, alone in adults with recurrent or metastatic solid tumors and in combination with Erbitux® (cetuximab) in head and neck cancer

Endpoints

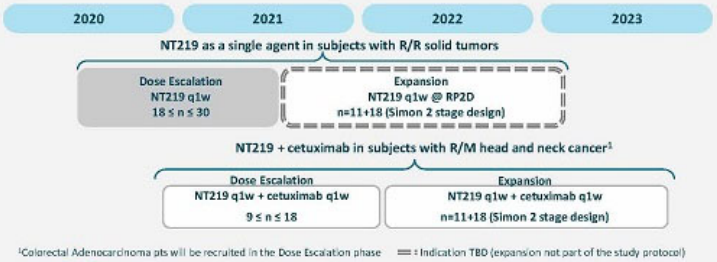
Primary endpoints:

Safety, pharmacokinetics and to determine the MTD

Secondary endpoints:

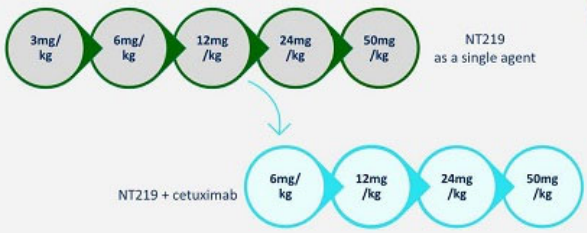
Obtain preliminary efficacy data

Study Design



[1]Colorectal Adenocarcinoma pts will be recruited in the Dose Escalation phase = : Indication TBD (expansion not part of the study protocol)

Dose Escalation Design



Interim Analysis – SAFETY and RESPONSE

3 mg/kg Dose Level as a Single Agent

- NT219 has been found to be well-tolerated with minimal adverse events
- In a patient with refractory GE junction disease, NT219 administration was associated with a partial response (PR):
  - ✓ Complete remission at the largest target lesion and one non-target central mesenteric lymph node
  - ✓ Stable Disease at the other non-target lesion pre-carinal lymph node

Response Analysis

	Cancer Type	Prior Lines of Therapy	Treatment Duration(Weeks)	Best Response*	
NT219 3mg/kg	Pancreatic Cancer	3	8	PD	
	GE Junction Cancer	4	22	PR	Target lesion: **Absent** Non target lesion 1: **Absent** Non target lesion 2: **Stable**
	Breast Cancer	11	8	PD	

*Interim data

Business Highlights

CM24 - First-in-class α-CEACAM1 mAb, validating clinical collaboration with Bristol Myers Squibb

NT219 - First-in-class, small molecule, dual inhibitor of IRS 1/2 and STAT3

H2:21 - Two phase 1 study readouts

Strong balance sheet and cash position

Purple Biotech (NASDAQ/TASE: PPBT)

ADSs outstanding: 17.5M

$61M cash as of December 31st, 2020

Cash runway through 2024

We are committed

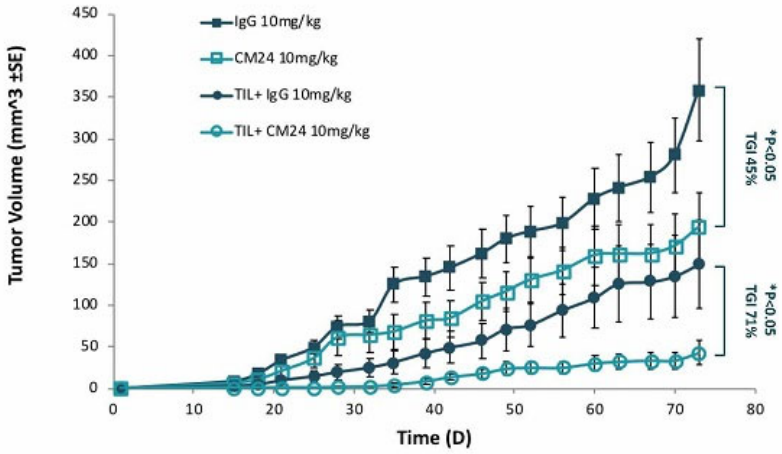
to providing cancer patients with first-in-class therapies to OVERCOME tumor drug resistance, ENHANCE treatment response and SLOW tumor progression


PURPLE
BIOTECH
THANK YOU
Contact Us:
ir@purple-biotech.com

Appendix A - CM24

Inhibition of Melanoma Growth Following CM24 and CM24 + TIL Treatment

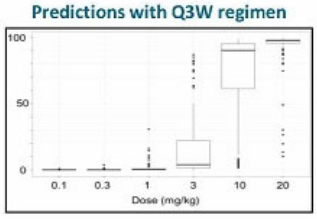
CM24 activity is Demonstrated as Single Agent and in Combination with TILS



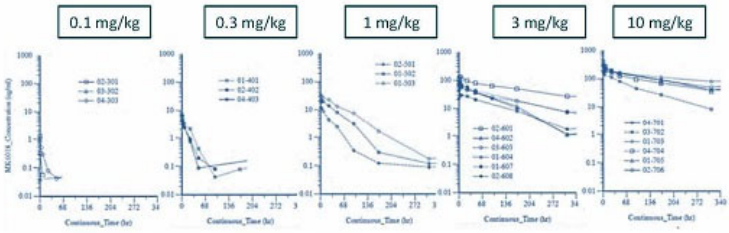
PHASE 1 PK DATA

Target saturation was not reached with doses up to 10mg/kg

Predictions with Q3W regimen



- Modeling with increased interval between doses demonstrates lower TMDD at 10mg/kg dose
- With Q3W, 20mg/kg dosing is not sufficient for saturation across population



Slower clearance with increasing dose

Higher half-life with increasing dose

Appendix B - NT219

Selected Publications

UC San Diego
SCHOOL of MEDICINE

Michael Karin

Oncogene (2016) 35, 2634–2644
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www.nature.com/onc

ORIGINAL ARTICLE

Targeting colorectal cancer via its microenvironment by inhibiting IGF-1 receptor-insulin receptor substrate and STAT3 signaling

E Sanchez-Lopez[1], E Flashner-Abramson[2], S Shalapour[1], Z Zhong[1], K Taniguchi[1,3], A Levitzki[2] and M Karin[1]

האוניברסיטה העברית בירושלים
THE HEBREW UNIVERSITY OF JERUSALEM

Alexander Levitzki

SHORT COMMUNICATION

Targeting melanoma with NT157 by blocking Stat3 and IGF1R signaling

THE UNIVERSITY OF TEXAS
MD Anderson Cancer Center

Menashe Bar-Eli

Published OnlineFirst May 7, 2013; DOI: 10.1158/0008-5472.CAN-12-3385

Therapeutics, Targets, and Chemical Biology

Cancer Research

Therapeutic Destruction of Insulin Receptor Substrates for Cancer Treatment

Hadas Reuveni[1,2,*], Efrat Flashner-Abramson[2], Lilach Steiner[1,2], Kirill Makedonski[1,2], Renduo Song[3], Alexei Shir[1], Meenhard Herlyn[4], Menashe Bar-Eli[3], and Alexander Levitzki[2,*]

UBC THE UNIVERSITY OF BRITISH COLUMBIA

Michael Cox

Published OnlineFirst September 29, 2014; DOI: 10.1158/1535-7163.MCT-13-0842

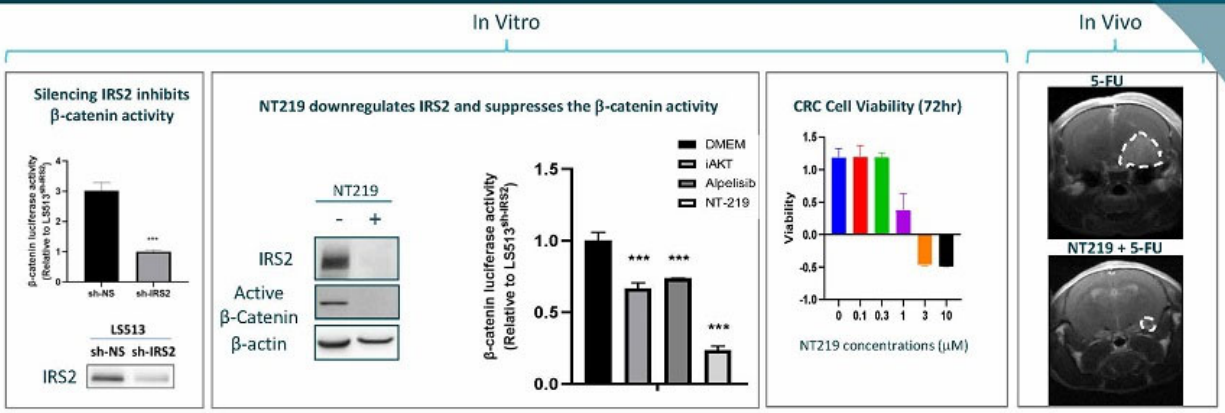
Small Molecule Therapeutics

Molecular Cancer Therapeutics

The Tyrphostin NT157 Suppresses Insulin Receptor Substrates and Augments Therapeutic Response of Prostate Cancer

Naokazu Ibuki[1,2], Mazyar Ghaffari[1,2], Hadas Reuveni[3,4], Mitali Pandey[1], Ladan Fazli[1], Haruhito Azuma[2], Martin E. Gleave[1,5], Alexander Levitzki[6], and Michael E. Cox[1,5]

NT219 | Suppresses β-Catenin activity in CRC Cells and Inhibited CRC Brain Metastasis



Colon cancer LS-513 cells overexpressing IRS2 demonstrate enhanced β-catenin activity.

Targeted inhibition of IRS2 by NT219 or IRS2-SH RNA, suppresses the increased β-catenin activity and inhibit LS-513 cell viability.

Combination of 5-FU and NT219 significantly inhibited the growth of CRC tumors in brain, using intracranial model and extended mice survival.

NT219 | Pancreatic Cancer in Combination with Gemcitabine

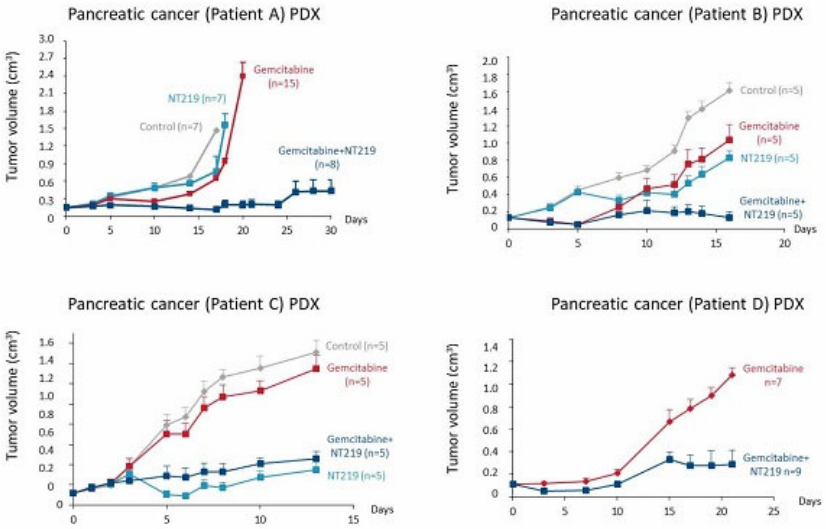
PDX model

Pancreatic Cancer

Drug

Gemcitabine (Gemzar®)

Highly effective anti cancer activity exhibited by NT219 in combination with Gemcitabine

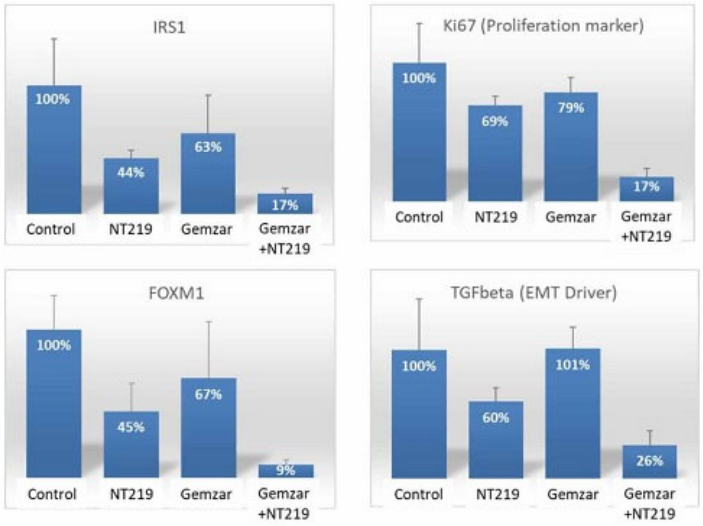


RNA Sequencing | Analysis of Tumors Following Treatment

PDX model
Pancreatic Cancer

Drug
Gemcitabine (Gemzar®)

Reduced expression of IRS1, Ki67, FOXM1 & TGFb is exhibited by pancreatic cancer treated with NT219 alone and in combination with gemcitabine



NT219 – DEMOGRAPHICS & SAFETY
3 mg/kg Dose Level as a Single Agent

Patients Demographics

Demographics of Patients treated with NT219 3mg/kg (n=3)	
Median age (range)	74 (69-79)
Male/Female, n (%)	2(66.6%)/1(33.3%)
Race	
White n (%)	3 (100%)
Prior Lines of Therapy	
3 n (%)	1 (33.3%)
4, n (%)	1 (33.3%)
11, n (%)	1 (33.3%)
Diagnosis, n	
Pancreatic Cancer	1
Gastroesophageal Junction Cancer	1
Breast Cancer	1
ECOG, n (%)	
1	3 (100%)
Median time from initial Diagnosis, Months (range)	62 (22-90)

Summary of Adverse Events

Adverse Event Description	Grade 1 n (#Events)	Grade 2 n (#Events)	Grade 3 n (#Events)
Abdominal Pain	1(2)	1(1)	
Alanine Aminotransferase Increased	1(1)		
Alkaline Phosphatase Increased			1(1)*
Aspartate Aminotransferase Increased	1(1)		
Back Pain		1(1)	
Belching	2(2)		
Cellulitis Left Foot		1(1)	
Dyspnea	2(2)		
Fatigue	1(1)		
Nausea	1(1)	1(1)	
Rash Pustular		1(1)	
Retching	1(2)		
Toxic Encephalopathy			1(1)**

*Transient- G2 after 2 weeks, **Transient- less than 24h

Presented at the ASCO annual meeting June 2021

Appendix C - CONSENSI®

CONSENSI® | From IND to the U.S. Market

CONSENSI® is the only NSAID whose labeling indicates reduction of blood pressure and consequent risk reduction of heart attack, stroke and death

Full U.S. Prescribing Information is available at http://www.consensi.com

Fixed dose combination of

Celecoxib, a COX-2 selective NSAID *(the active ingredient in Pfizer's Celebrex®)*

+

Amlodipine a blood pressure-lowering agent (a calcium channel blocker) *(the active ingredient in Pfizer's Norvasc®)*

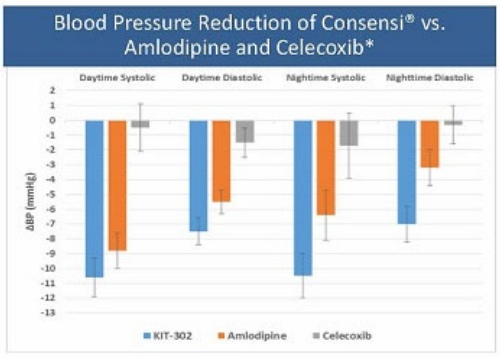
Launched in the USA- Coeptis Pharmaceuticals

Partnered in China- CSBio

Partnered in S. Korea- Kuhnil Pharmaceutical

Purple Biotech's clinical, regulatory and medical teams developed CONSENSI® internally from IND, through successful Phase III clinical trials, to FDA approval

CONSENSI® Phase III Trial Results



Consensi® demonstrated <u>even better</u> BP reduction than same amount of amlodipine given without celecoxib

* Error bars – standard error of mean

- Primary efficacy endpoint was successfully achieved (P=0.001)
- Demonstrated 2.5x better blood pressure reduction than FDA requirement (50% of amlodipine arm)
- Demonstrated consistent reduction in all measures of blood pressure
- Observed beneficial renal functions:

Measure	Consensi®	Amlodipine
Creatinine plasma level reduction	-3.22 µmol/L	-2.55 µmol/L
Peripheral edema (% patients)	8.2%	15.6%

- Additional Phase III/IV clinical trial to scientifically validate the renal benefits (not required for NDA submission) was completed. Topline results were announced in October, 2017

Purple Biotech Commercial Drug: CONSENSI®

CONSENSI® is the only NSAID whose labeling indicates reduction of blood pressure and consequent risk reduction of heart attack, stroke and death

Full U.S. Prescribing Information is available at: : http://www.consensi.com

Fixed dose combination of
Celecoxib,
a COX-2 selective NSAID
(the active ingredient in Pfizer's Celebrex®)

+

Amlodipine,
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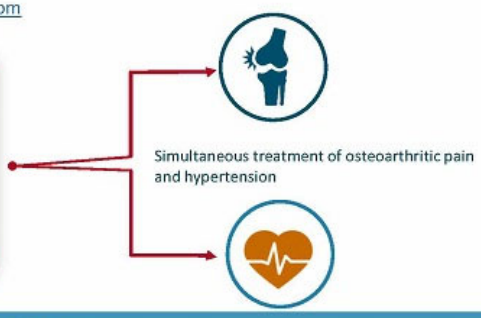


Simultaneous treatment of osteoarthritic pain and hypertension

- Approved for marketing by U.S. FDA on May 31, 2018
- Clinical data showed Consensi™ was more effective at lowering blood pressure than amlodipine alone
- Clinical data also demonstrated beneficial renal function measures
- Formulated with 200 mg celecoxib and three different dosages (2.5, 5, 10 mg) of amlodipine
- Manufactured by Dexcel Pharma – Israel's largest private pharmaceutical company

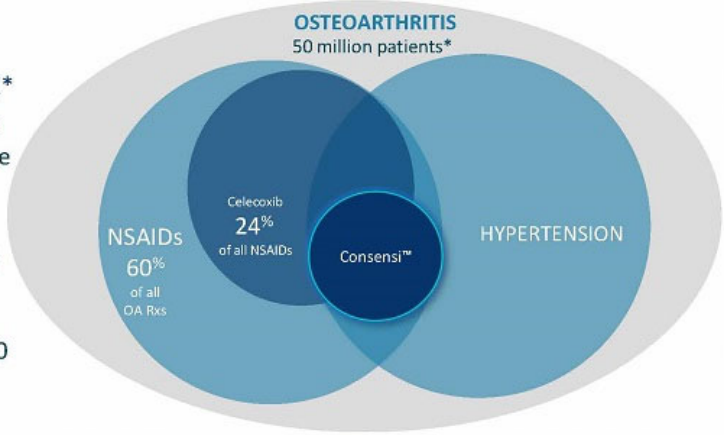
**Celebrex® is a registered trademark of G.D. Searle LLC (a subsidiary of Pfizer Inc.).*
Norvasc® is a registered trademark of Pfizer Inc.

CONSENSI® U.S. Target Markets

CONSENSI® targets osteoarthritis (OA) patients currently treated with NSAIDs (celecoxib as well as others) who also suffer from existing or newly diagnosed hypertension

ARTHRITIS PREVALENCE*

- More than 50 million adults in the U.S. have doctor-diagnosed osteoarthritis
- 67 million people are expected to have doctor-diagnosed osteoarthritis by 2030



HYPERTENSION PREVALENCE**

- 29% of U.S. adults older than 18
- 65% of U.S. adults older than 60

COMORBIDITIES

- 44% of adults with high blood pressure have osteoarthritis**

PURPLE

** Arthritis Foundation: http://www.arthritis.org/ ** Hypertension Among Adults in the United States: National Health and Nutrition Examination Survey, 2011–2012*