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

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): **April 23, 2018**

TRACON Pharmaceuticals, Inc.		
(Exact name of registrant as specified in its charter)		
Delaware	**001-36818**	**34-2037594**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

4350 La Jolla Village Drive, Suite 800 San Diego, California	**92122**
(Address of principal executive offices)	(Zip Code)

Registrant's telephone number, including area code: (858) 550-0780

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter). Emerging growth company ☒

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☒

Item 7.01 Regulation FD Disclosure.

Charles P. Theuer, M.D., Ph.D., and other executive officers will be presenting the information attached as Exhibit 99.1 to this Current Report on Form 8-K at various upcoming meetings beginning April 23, 2018.

By furnishing this information, TRACON makes no admission as to the materiality of any information in this report and the exhibit hereto. The information contained in this report and the exhibit hereto is intended to be considered in the context of TRACON's filings with the Securities and Exchange Commission and other public announcements that TRACON makes, by press release or otherwise, from time to time. TRACON undertakes no duty or obligation to publicly update or revise the information contained in this report or the exhibit hereto, although it may do so from time to time as its management believes is appropriate. Any such updating may be made through the filing of other reports or documents with the Securities and Exchange Commission, through press releases or through other public disclosure.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	**Description**
99.1	Corporate Presentation, dated April 2018

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 hereunto duly authorized.

TRACON Pharmaceuticals, Inc.

Dated: April 23, 2018

By: /s/ Charles P. Theuer
Charles P. Theuer, M.D., Ph.D.
President and Chief Executive Officer

Exhibit 99.1

TRACON PHARMACEUTICALS
April 2018

NASDAQ: TCON

Forward-Looking Statements

This presentation contains statements that are, or may be deemed to be, "forward-looking statements." In some cases these forward-looking statements can be identified by the use of forward-looking terminology, including the terms "believes," "estimates," "anticipates," "expects," "plans," "intends," "may," "could," "might," "will," "should," "approximately," "potential," or, in each case, their negatives or other variations thereon or comparable terminology, although not all forward-looking statements contain these words. These statements relate to future events or our future financial performance or condition, business strategy, current and prospective product candidates, planned clinical trials and preclinical activities, product approvals, research and development costs, current and prospective collaborations, timing and likelihood of success of development activities and business strategies, plans and objectives of management for future operations, and future results of anticipated product development efforts, including potential benefits derived therefrom. These statements involve substantial known and unknown risks, uncertainties and other important factors that may cause our actual results, levels of activity, performance or achievements to differ materially from those expressed or implied by these forward-looking statements. These risks, uncertainties and other factors include, but are not limited to, risks associated with conducting clinical trials, whether any of our product candidates will be shown to be safe and effective, our ability to finance continued operations, our reliance on third parties for various aspects of our business, competition in our target markets, our ability to protect our intellectual property, and other risks and uncertainties described in our filings with the Securities and Exchange Commission, including under the heading "Risk Factors". In light of the significant uncertainties in our forward-looking statements, you should not place undue reliance on these statements or regard these statements as a representation or warranty by us or any other person that we will achieve our objectives and plans in any specified time frame, or at all. The forward-looking statements contained in this presentation represent our estimates and assumptions only as of the date of this presentation and, except as required by law, we undertake no obligation to update or revise publicly any forward-looking statements, whether as a result of new information, future events or otherwise after the date of this presentation.

This presentation also contains estimates, projections and other information concerning our industry, our business, and the markets for our drug candidates, as well as data regarding market research, estimates and forecasts prepared by our management. Information that is based on estimates, forecasts, projections, market research or similar methodologies is inherently subject to uncertainties and actual events or circumstances may differ materially from events and circumstances reflected in this information.

Investment Highlights

Late Stage Pipeline with Multiple Near Term Readouts for Lead Asset

- ***Oncology Phase 3*** → TAPPAS trial under SPA in orphan indication; multiple ongoing Phase 2 trials
- ***Ophthalmology Phase 2*** → AVANTE randomized wet AMD trial
- ***Immune Oncology Phase 1*** → Lung cancer trial combined with Opdivo

Significant Commercial Opportunity

- Opportunity to replicate revenue of the standard of care VEGF inhibitor companion therapies that generate > $18B in annual revenue in oncology and eye disease

Strategic Partnerships

- Lead Asset in Oncology: Regional partnership with **Ambrx**
- Lead Asset in Ophthalmology: Global partnership with **Santen**
- Asset in Prostate Cancer: Global partnership with **Janssen**

Proprietary Product Development Platform

- Advances TRACON's proprietary pipeline
- Basis for partnership with Janssen with potential to generate significant long term revenue

Capital Efficient Product Development Platform

Internal product development platform allows TRACON to conduct clinical trials without a CRO

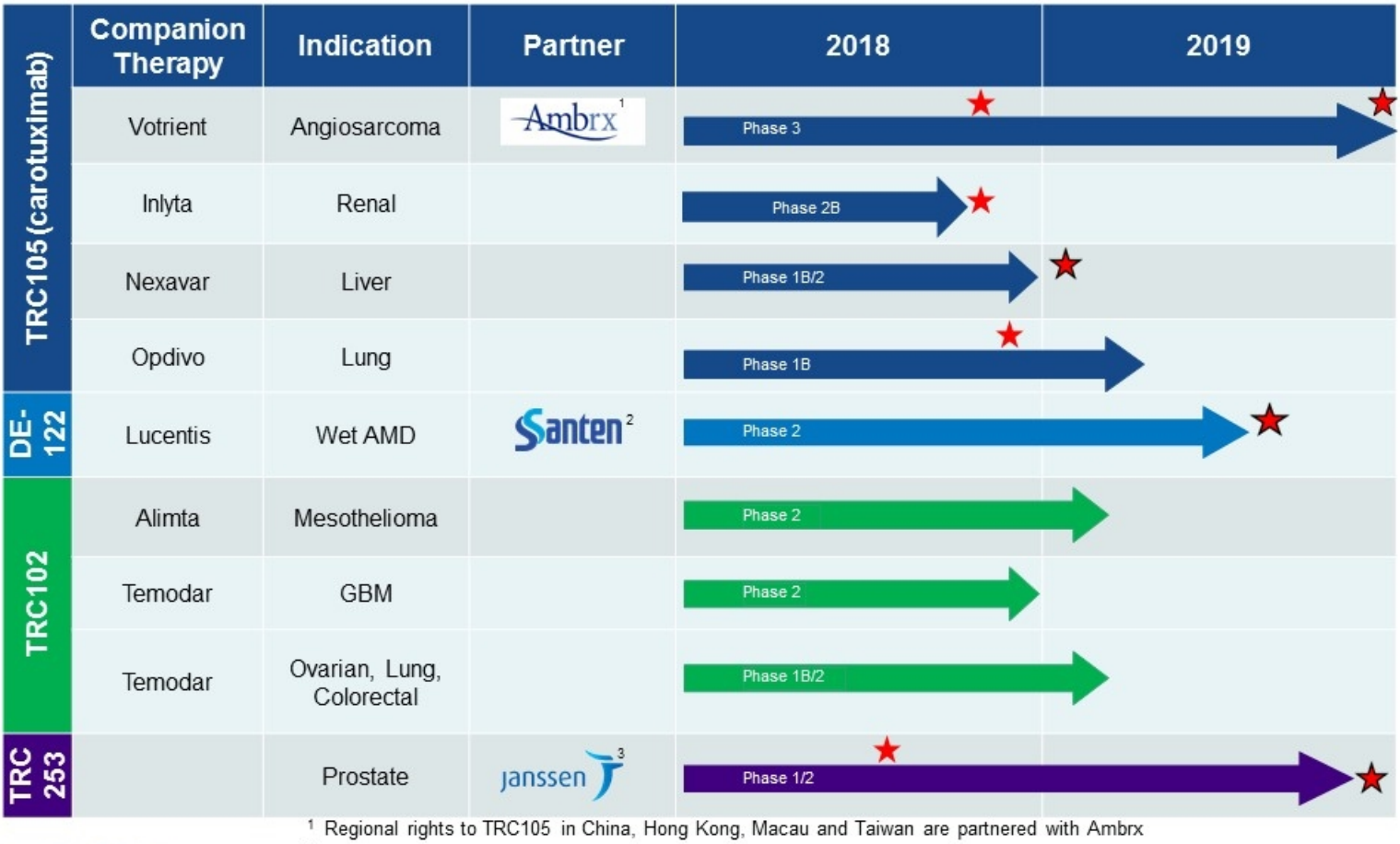
- Opportunity to achieve shorter timelines at lower cost

Management team with comprehensive CMC, Regulatory and QA expertise

- Development of multiple products through launch

Opportunity to expand the portfolio through in-licensing programs at no up-front cost (e.g., Janssen transaction)

Multiple Expected Near-Term Value Inflection Points



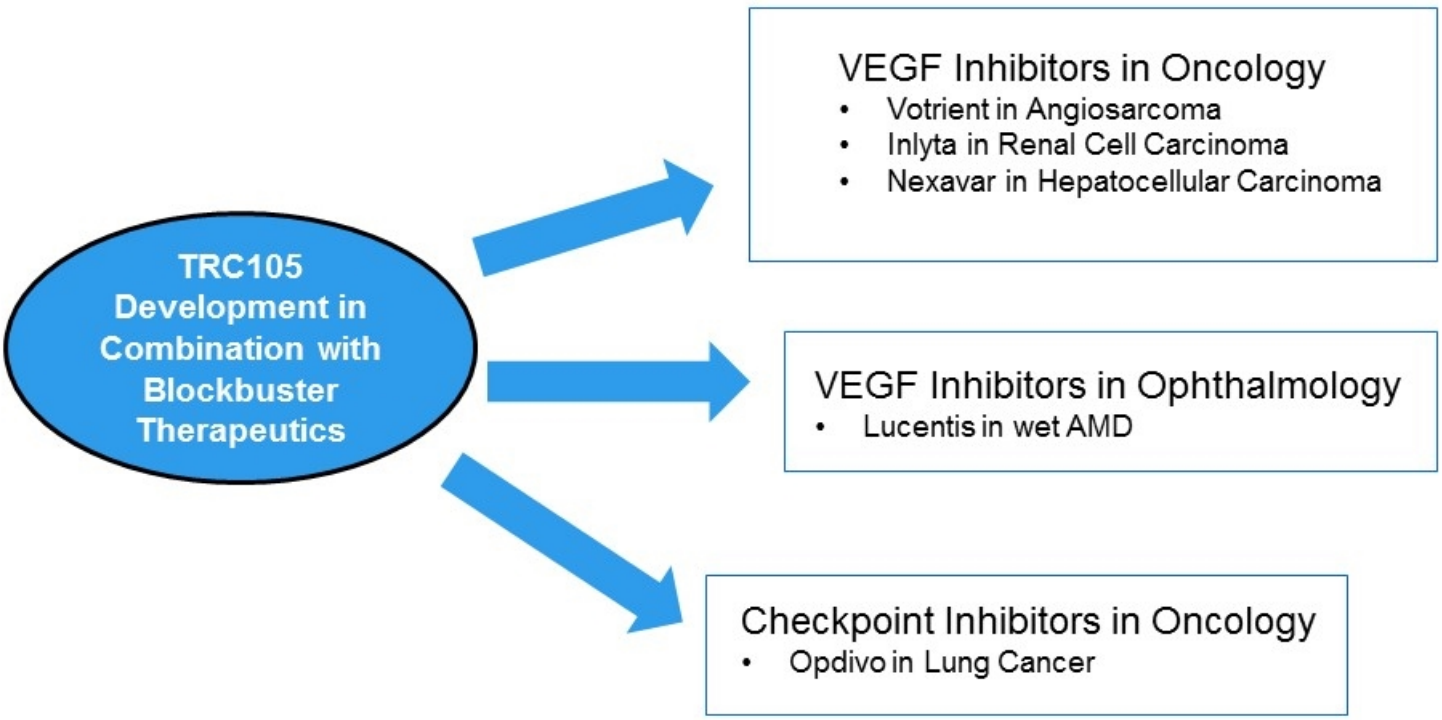
[1] Regional rights to TRC105 in China, Hong Kong, Macau and Taiwan are partnered with Ambrx

[2] Global Rights to DE-122 are partnered with Santen Pharmaceutical Co., Ltd. (Santen)

[3] Janssen Pharmaceutica N.V. (Janssen) has a buyback option to TRC253

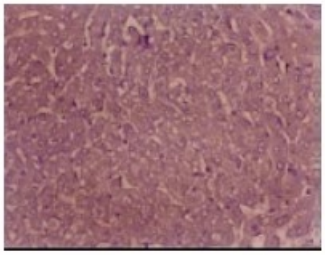
TRACON PHARMA

Lead Asset Development Strategy

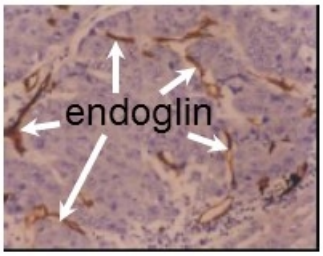


TRC105 Target:
Endoglin is an Essential Non-VEGF Angiogenic Target

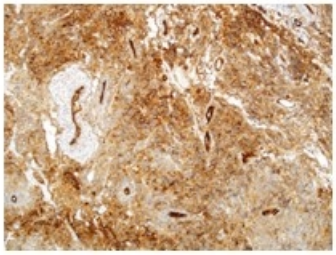
- Expressed on proliferating blood vessels in cancer and AMD
  - Essential for angiogenesis
  - Unfavorable prognostic marker
  - Up-regulated following VEGF inhibition
- Attenuated expression (Osler-Weber-Rendu syndrome) associated with improved cancer survival
- Genetic knockdown reverses resistance to VEGF inhibition
- Targeting VEGF and endoglin concurrently improves antitumor effects
- Targeting endoglin on myeloid derived suppressor cells (MDSCs) potentiates PD-1/PD-L1 inhibition in preclinical models



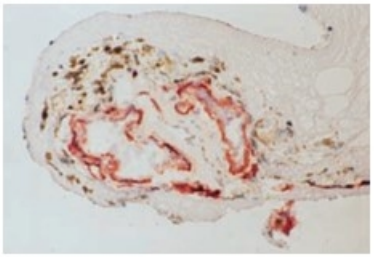
Normal Human Liver



Human Liver Cancer



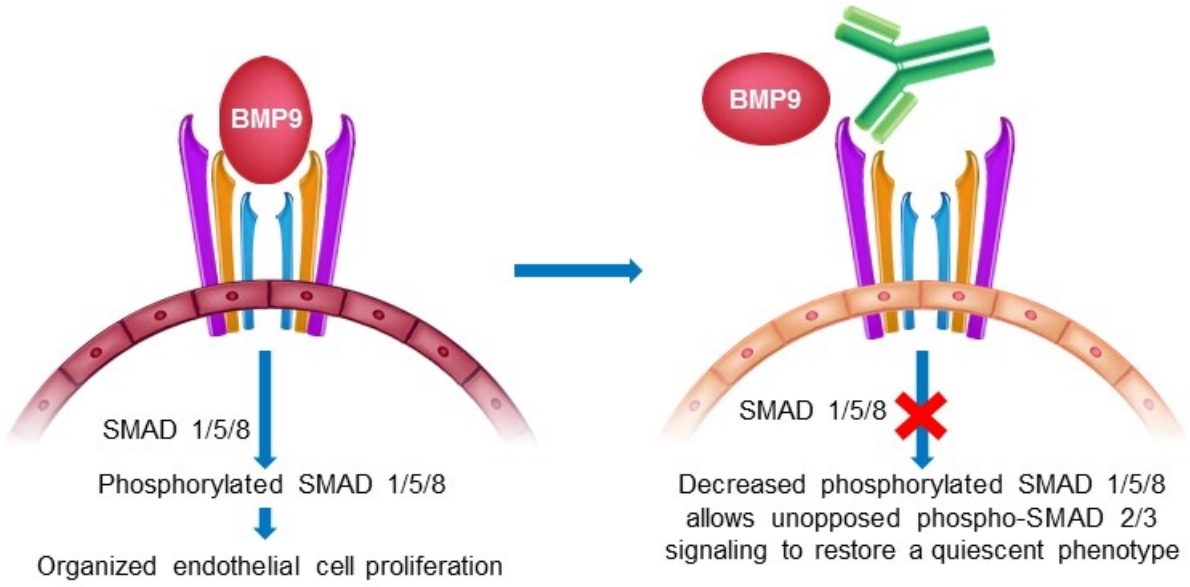
Angiosarcoma



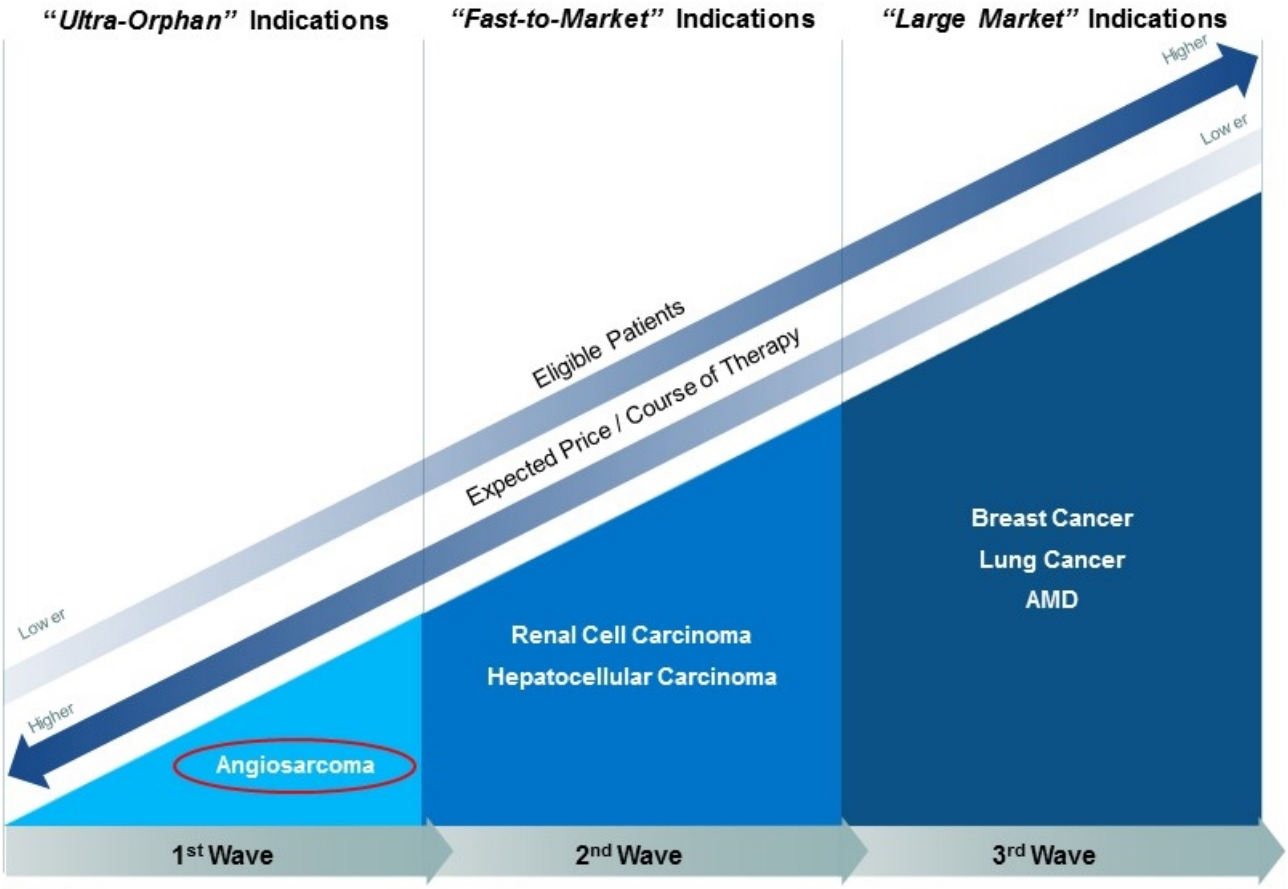
Human AMD Membrane

TRC105: Our Lead Endoglin Antibody

- TRC105 binds a precise endoglin epitope to inhibit BMP binding and angiogenesis
- TRC105 also potently mediates antibody-dependent cell mediated cytotoxicity (ADCC)



TRC105 Tiered Clinical Development



TRACON PHARMA

Figure for illustrative purposes only.

Lead Indication: Angiosarcoma

- ***Orphan indication***: ~ 600 cases annually in the US and 1,200 in Europe; greater incidence in Asia[1]
- ***High Unmet Need***: 5-year survival rate < 12% compared to 5-year survival rate of ~ 56% for all soft tissue sarcoma[2]
  - Treatment with chemotherapy (taxanes or doxorubicin) in the front line setting is associated with PFS of ~ 5 months and OS < 1 year[3]
  - Treatment with VEGF inhibitors in the second line setting is associated with PFS of 1.8 - 3.8 months and OS < 1 year
- ***Two subtypes***: About 50% of patients present with a primary cutaneous lesion
- ***Market size***: Estimated at >$150M in US/EU assuming premium pricing similar to the price of oncology therapeutics approved in other orphan indications[4]

[1]Suveillance, Epidemiology, and End Results Program, NCI, www.seer.cancer.gov; RARECARE database, www.rarecare.eu
[2]www.cancerresearchuk.org
[3]Penel et al, JCO 2008; Italiano et al, Cancer 2012
[4]TRACON estimate

TRACON PHARMA

Profile of Unmet Need in Initial Pivotal Indication: VEGF Inhibitors Have Limited Activity in Angiosarcoma

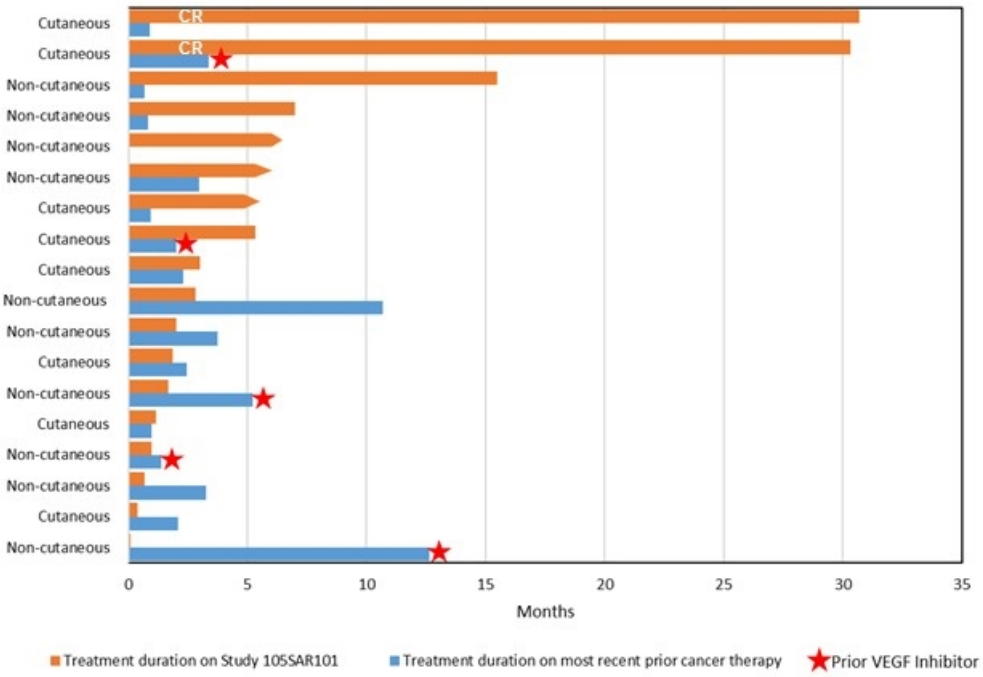
VEGF Inhibitor	Study	Patient Population	Activity
Votrient®[1]	*Retrospective analysis (CTOS 2016)*	*Angiosarcoma (n = 40)*	• *ORR = 20% (No CRs)* • *PFS = 3.0 months* • *OS = 9.9 months*
Votrient	Retrospective analysis (ASCO 2014)	Soft tissue sarcoma, including 6 angiosarcoma patients	• No CR's
Nexavar®	Single agent study (Maki 2009)	Angiosarcoma (n = 37)	• ORR = 14% (1/37 CR) • PFS = 3.8 months
Nexavar	Single agent study (French sarcoma group)	Angiosarcoma (n = 41)	<u>Cutaneous angiosarcoma</u> • ORR = 15% (2/26 CR) • PFS = 1.8 months <u>Visceral angiosarcoma</u> • ORR = 13% (No CRs) • PFS = 3.8 months
Avastin®	Single agent study (Agulnik 2013)	Angiosarcoma (n = 23)	• ORR = 9% (No CRs) • PFS = 3.0 months

[1] Votrient is the only VEGF inhibitor approved for the treatment of soft tissue sarcoma based on the superior PFS versus placebo (4.6 versus 1.6 months) in the Phase 3 PALETTE study.

TRC105 + Votrient is Active in Angiosarcoma

- PFS in 13 VEGF inhibitor-naïve patients of 7.8 months vs. 3 month PFS expected with Votrient
- Most VEGF inhibitor patients had superior time on treatment with TRF105 + Votrient compared to prior chemotherapy. US and EU regulators allowed enrollment of treatment naive angiosarcoma patients into the Phase 3 TAPPAS trial

Study Duration of 9 Angiosarcoma Patients Treated with TRC105 + Pazopanib in the Original Phase 1b/2 Trial and 9 Patients in the Expansion Cohort

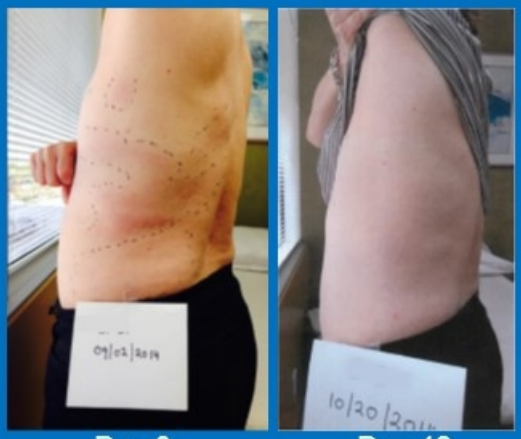


*Treatment duration is calculated from date of first dose to date of last dose
*Last response assessment used as date of progression for ongoing patients to calculate mPFS

Data as of November 2017

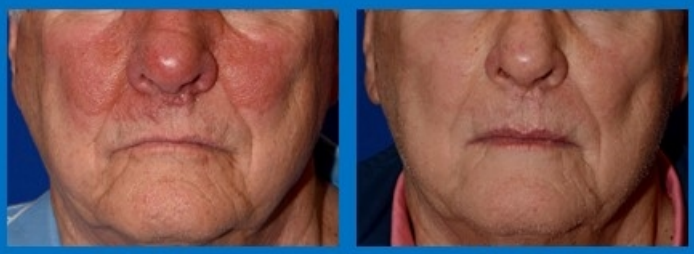
TRC105 + Votrient Phase 1b/2 Observations

Patient #1 off study (due to AE) after 30+ months with ongoing CR



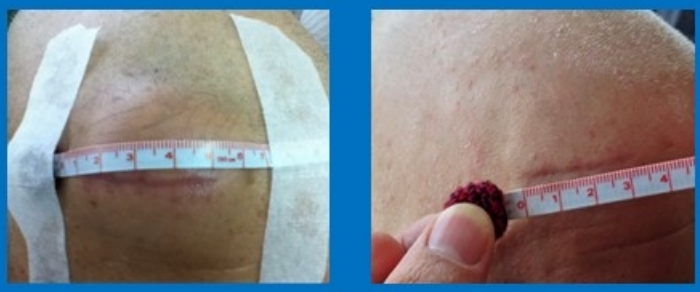
Day 0 Day 48

Patient #2 maintained a CR for 28+ months



Day 0 Day 37

Patient #3 remained on treatment for 16 months



Day 0 Day 84

Data as of November 2017

TRACON PHARMA

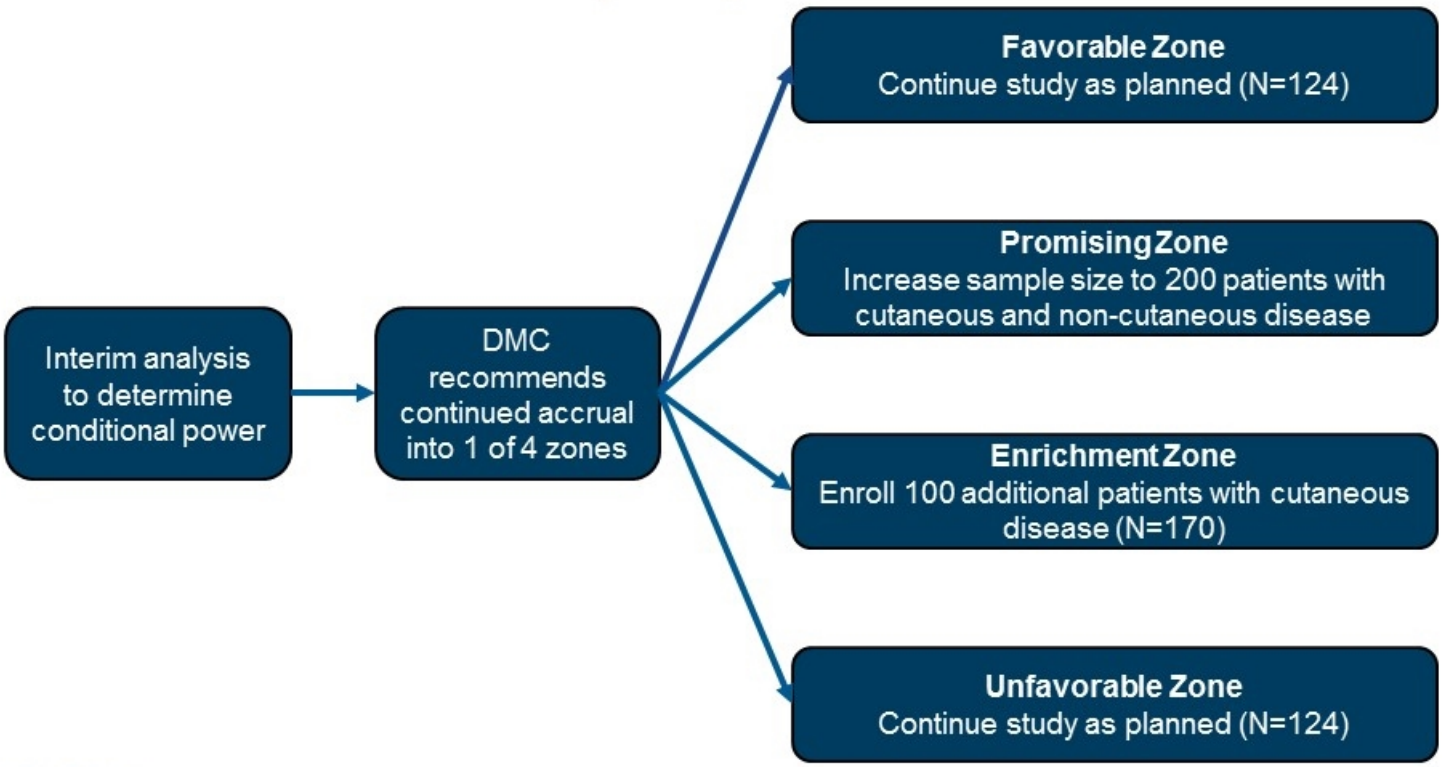
Phase 3 TAPPAS Trial in Angiosarcoma

- Primary Endpoint: PFS
- Independent blinded central review
- Key Secondary Endpoints: ORR, OS
- Key eligibility
  - Age ≥ 12
  - Unresectable angiosarcoma
  - Measurable disease by RECIST 1.1
  - No prior treatment with VEGF inhibitor
  - No more than 2 prior lines of treatment
  - ECOG PS 0-1
- Strata
  - Cutaneous vs Non-cutaneous
  - Prior chemotherapy: 0 vs 1 or 2
- N=124-200 (Adaptive design)

TRACON PHARMA

Phase 3 TAPPAS Trial in Angiosarcoma

Adaptive design recognized as Most Innovative Clinical Trial of 2017. Allows for sample size re-estimation or enrichment of cutaneous disease at the time of the interim analysis expected in 2H 2018.

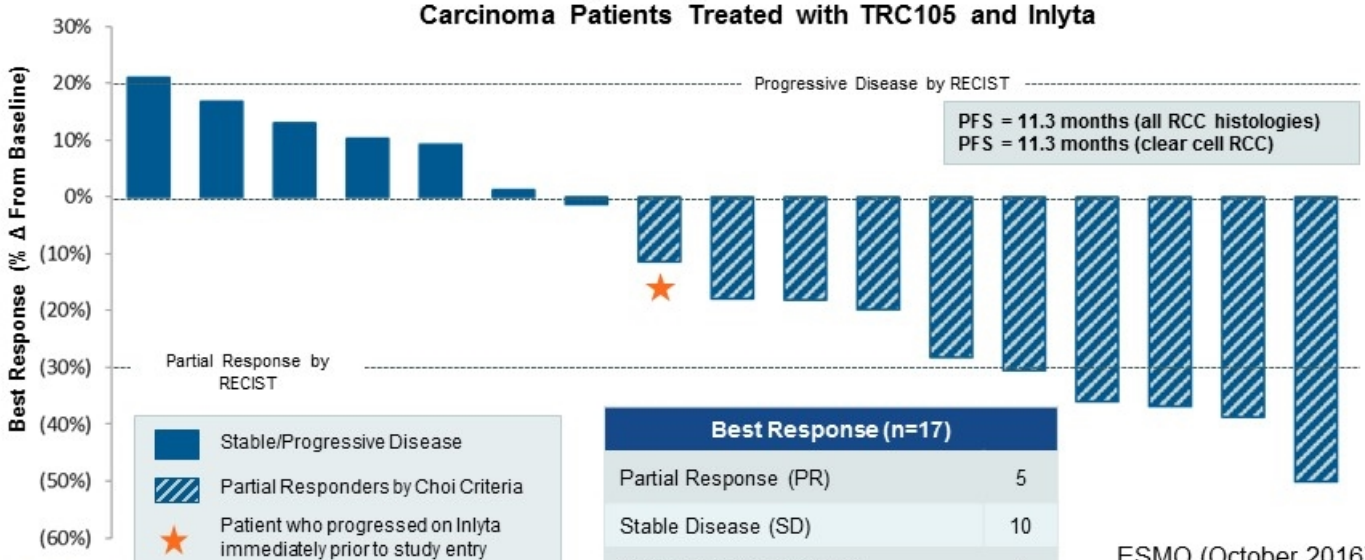


TRACON PHARMA

TRC105 + Inlyta® in Renal Cell Carcinoma

- 18 patients treated in a Phase 1b clinical trial who failed at least one VEGF inhibitor
- Partial response rate by RECIST of 29% (4 of which were in the fourth line setting)
  - Exceeded partial response rate of Inlyta following VEGFR TKI treatment in the Inlyta Phase 3 AXIS trial of 11%
- Median PFS in clear cell RCC of 11.3 months
  - Exceeded PFS of Inlyta following VEGFR TKI treatment in the Inlyta Phase 3 AXIS trial of 4.8 months
- Exploratory analysis indicated two biomarkers (baseline TGF-β R3 and osteopontin) correlated with activity

Maximum Percentage Change in Target Lesion Size in Renal Cell Carcinoma Patients Treated with TRC105 and Inlyta



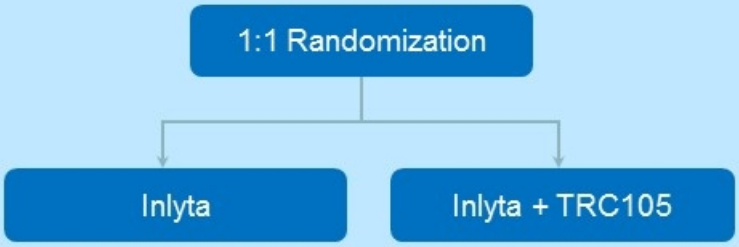
Best Response (n=17)	
Partial Response (PR)	5
Stable Disease (SD)	10
Progressive Disease (PD)	2

ESMO (October 2016)

TRACON PHARMA

Phase 2 TRAXAR Trial in Renal Cell Carcinoma

- Primary Endpoint: PFS by RECIST 1.1
- Independent blinded central review
- Key eligibility
  - Advanced or metastatic clear cell RCC
  - Measurable disease by RECIST 1.1
  - Progression following 1 prior VEGF inhibitor
  - 1 prior mTOR inhibitor allowed
  - 1 prior immunotherapy allowed

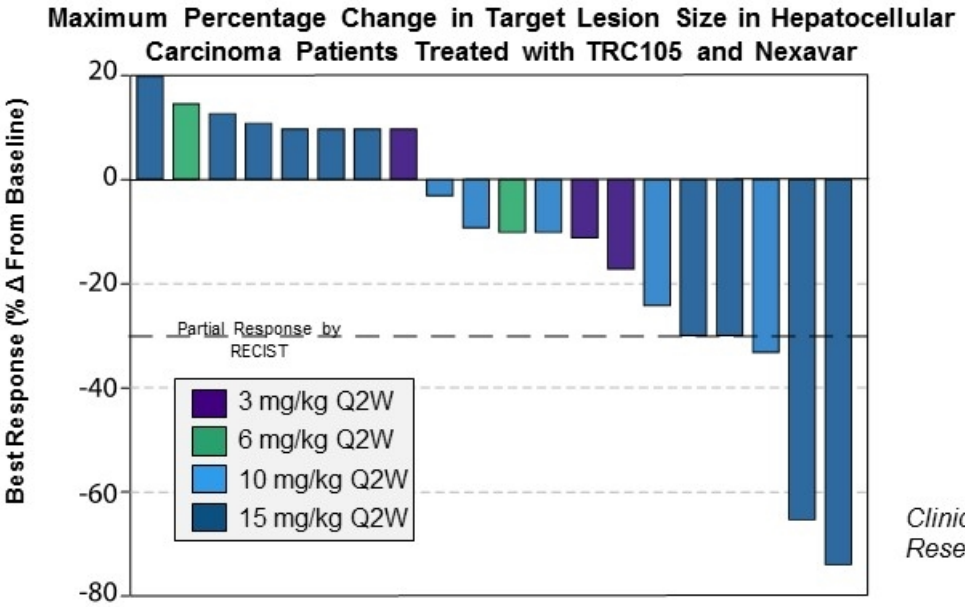


Fully enrolled (N=150) event driven trial with data expected mid 2018

TRACON PHARMA

TRC105 + Nexavar in Hepatocellular Carcinoma

- NCI Phase 1/2 study published in *Clinical Cancer Research* - partial response rate by RECIST of 25% across 4 dose levels; partial response rate of 33% for patients treated at two highest dose levels (10 or 15 mg/kg TRC105)
  - Exceed partial response rate of Nexavar in Phase 3 pivotal studies of 2 - 3%
  - Median OS of 15.5 months exceeded median OS of Nexavar in its pivotal Phase 3 of 10.7 months
- Multicenter Phase 2 trial in up to 33 patients is enrolling to confirm response rate
  - Interim data presented at GI ASCO (January 2018): partial responses in 2 of first 8 evaluable patients
  - Full data expected at GI ASCO January 2019
- Late stage development in hepatocellular cancer to be led by Ambrx in China



Clinical Cancer Research (May 2017)

TRACON PHARMA

TRC105 + Opdivo® in Lung Cancer

- Endoglin is a TGF-β co-receptor expressed on fibroblasts and myeloid derived suppressor cells (MDSCs), cell types not addressed by checkpoint inhibition
  - TGF-β signaling implicated as a primary means of tumor immune evasion that complements checkpoint inhibition and tumor mutational burden
- TRC105 potentiates the activity of PD-1 inhibition in syngeneic mouse tumor models
  - Oral presentations from Leiden University researchers announced at International Microenvironment Cancer Society meeting in June
- TRC105 is being studied with Opdivo in second line non-small cell lung cancer in a Phase 1 trial
  - Opdivo single agent response rate in this setting is 20%[1]
  - Correlation between response and MDSC tumor content will be assessed

TRACON PHARMA

[1]Opdivo® package insert

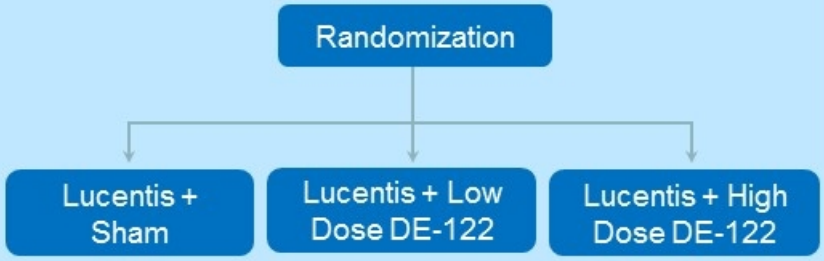
Development in AMD Partnered with Santen

- Global ophthalmology company with $1.8 billion in annual revenue leads global development and commercialization for DE-122 (ophthalmic formulation of TRC105) in wet AMD and other eye diseases
- Deal terms
  - $20 million received thus far
  - Santen pays all development costs
  - Up to $145 million in additional milestone payments
  - Royalties in the high single digits to low teens
- Failed Phase 2 and 3 studies from Ophthotech and Regeneron leave substantial opportunity for a superior mechanism of action to build on $8B VEGF inhibitor market in wet AMD; regulatory path well defined; substantial commercial opportunity

Clinical Development of DE-122 in wet AMD

- Phase 1/2 PAVE trial results presented February 10, 2018 at the Angiogenesis, Exudation and Degeneration meeting at Bascom Palmer Eye Institute
  - Safe with no serious adverse events
  - 8 out of 12 subjects demonstrated bioactivity: improved macular edema or visual acuity
- Phase 2 AVANTE randomized controlled trial is enrolling - data expected 2019

- Primary Endpoint: Best Corrected Visual Acuity following six monthly intravitreal injections
- Double masked
- N=51



TRC102: Reversing Resistance to Chemotherapy

- Small molecule designed to reverse resistance to chemotherapy and complement PARP inhibitors
- Inhibits base excision repair, a dominant pathway of DNA repair that allows for resistance to alkylating chemotherapy (e.g., Temodar®) and antimetabolite chemotherapy (e.g., Alimta®)
- Clinical development funded by NCI

Combination	Well Tolerated	Signs of Activity in Phase 1b/2	Ongoing Development
TRC102 + Alimta (Published in *Investigational New Drugs*, 2012)	√	Stable disease in some patients with squamous cell lung cancer, a tumor type where Alimta is inactive	Phase 2 trial with Alimta in mesothelioma
TRC102 + Fludara (Published in *Oncotarget*, 2017)	√	Partial response and stable disease in some patients previously treated with Fludara	
TRC102 + Temodar (Presented at ASCO 2017)	√	Partial responses in some patients with lung, KRAS+ colorectal and ovarian cancer; induced biomarkers of DNA damage Rad51, pNbs1, and/or γ-H2AX	Phase 2 expansion cohorts added in lung, colorectal, and ovarian cancer; Phase 2 trial with Temodar in glioblastoma

Deal with Janssen

- TRC253 and TRC694 in-licensed from Janssen at no cost
  - TRC253 is an antagonist of the F877L and other AR mutations that are resistance mechanisms for Xtandi® and apalutamide
  - TRC694 is a selective inhibitor of NF-kB-inducing kinase (NIK)
- TRACON was chosen because of our efficient product development platform
- $5M equity investment made by JJDC

TRC253

- Janssen has rights to re-acquire TRC253 following Phase 1/2 for $45M
  - Additional potential milestones of $137.5M and low single digit royalty
- If kept by TRACON, we would owe regulatory and commercial milestones of up to $45M and a low single digit royalty to Janssen

TRC694

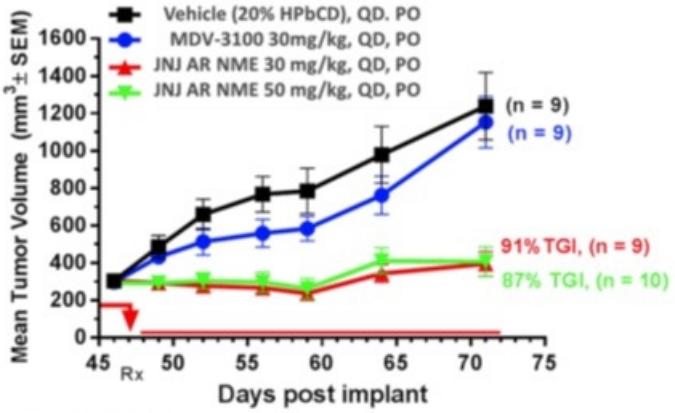
- Janssen has a right of first negotiation for TRC694 following Phase 1
- TRACON will owe development and regulatory milestones of up to $60M and low single digit royalty

TRACON PHARMA

TRC253: Novel Androgen Receptor (AR) Mutant Inhibitor

- Designed to treat AR resistant prostate cancer
  - Occurs in ~10% of mCRPC cases
- Active against wild-type AR and many clinically relevant ligand binding domain mutations
- Clear path to POC data in targeted population using a companion diagnostic
- Phase 1/2 trial enrolling

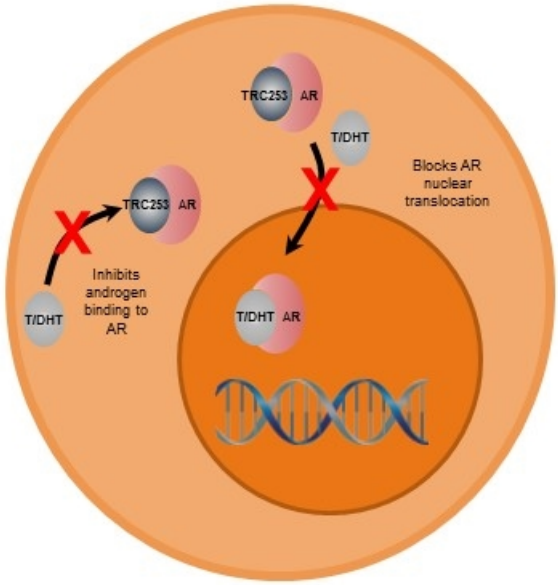
AR F877L-driven xenograft model



TRACON PHARMA

Hickson, I. AACR 2016 Annual Meeting.
Joseph, JD, et al. Cancer Discovery 2013.

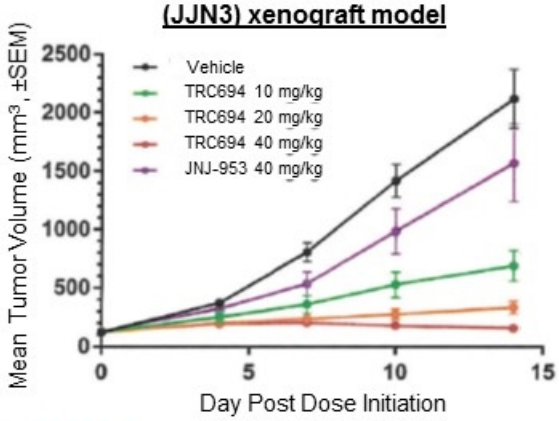
Multiple Mechanisms of Action



TRC694: Novel NF-kB Inducing Kinase (NIK) Inhibitor

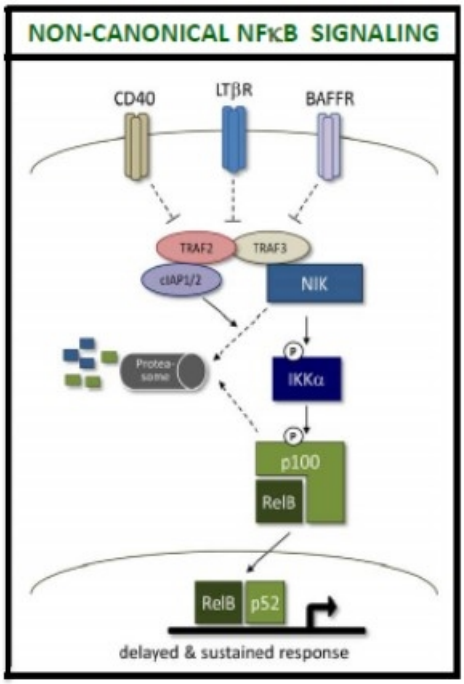
- NIK pathway is dysregulated in hematologic malignancies
  - Multiple myeloma (~12-20% of cases), mantle cell lymphoma (~17%), diffuse large B-cell lymphoma (~9-15%), CLL (~4% at diagnosis, higher later)
- Clear path to POC data in targeted population using a precision medicine approach

NIK-translocated myeloma cell line (JJN3) xenograft model



TRACON PHARMA

NIK is Critical for Non-Canonical NFκB Activation



Krappmann & Vincendeau, 2016

Ongoing Business Development Strategy

- Leverage the TRACON Product Development Platform to develop promising oncology assets while diversifying the TRACON portfolio
  - Transactions similar to the Janssen transaction where TRACON develops asset(s) to certain value inflection points in return for substantial economics and/or downstream commercial rights
  - For companies with little or no development infrastructure in the US, conduct proof-of-concept clinical trials in the US in exchange for substantial economics and/or product rights in the US

Expected Milestones Across All Programs

Milestone	Expected Timing
Initial Response Data from TRC105 Phase 2 multicenter trial in HCC	1H 2018 ✓
Present DE-122 Phase 1/2 PAVE trial data in wet AMD (Santen)	1H 2018 ✓
Present preclinical data from TRC105 + checkpoint inhibitor combination studies	mid 2018
Complete dose escalation in TRC253 Phase 1/2 trial in prostate cancer	mid 2018
Top-line data from TRC105 Phase 2 TRAXAR trial in RCC	2H 2018
Interim Analysis from TRC105 Phase 3 pivotal TAPPAS trial in angiosarcoma	2H 2018
Present data from TRC105 + Opdivo Phase 1 trial	2H 2018
Response data from TRC105 Phase 2 multicenter trial in HCC	1H 2019
File IND and initiate Phase 1 trial of TRC694	1H 2019
DE-122 Phase 2 randomized AVANTE trial data in wet AMD	2H 2019
Janssen opt-in decision to reacquire TRC253 for $45M + expenses following completion of TRC253 Phase 1/2 trial in prostate cancer	2H 2019
Top-line data from TRC105 Phase 3 pivotal TAPPAS trial in angiosarcoma	2H 2019

Financial Overview (as of December 31, 2017)

Ticker	TCON (NASDAQ)
Cash, Cash Equivalents and Short-term Investments	$34.5 million*
Debt – Outstanding Principal	$8.0 million
Common Shares O/S	17.7 million*
Covering Analysts	Jim Birchenough (Wells Fargo) Bert Hazlett (BTIG) Chad Messer (Needham) Maury Raycroft (Jefferies) Alex Schwartz (Stifel)

*Does not include proceeds or share issuances from $38.7 million March 2018 PIPE transaction

Investment Highlights

Late Stage Pipeline with Multiple Near Term Readouts for Lead Asset

- ***Oncology Phase 3*** → TAPPAS trial under SPA in orphan indication; multiple ongoing Phase 2 trials
- ***Ophthalmology Phase 2*** → AVANTE randomized wet AMD trial
- ***Immune Oncology Phase 1*** → Lung cancer trial combined with Opdivo

Significant Commercial Opportunity

- Opportunity to replicate revenue of the standard of care VEGF inhibitor companion therapies that generate > $18B in annual revenue in oncology and eye disease

Strategic Partnerships

- Lead Asset in Oncology: Regional partnership with **Ambrx**
- Lead Asset in Ophthalmology: Global partnership with **Santen**
- Asset in Prostate Cancer: Global partnership with **Janssen**

Proprietary Product Development Platform

- Advances TRACON's proprietary pipeline
- Basis for partnership with Janssen with potential to generate significant long term revenue

TRACON PHARMACEUTICALS
April 2018

NASDAQ: TCON

Backup

Complementing VEGF Inhibition Represents a Substantial Potential Commercial Opportunity for TRC105

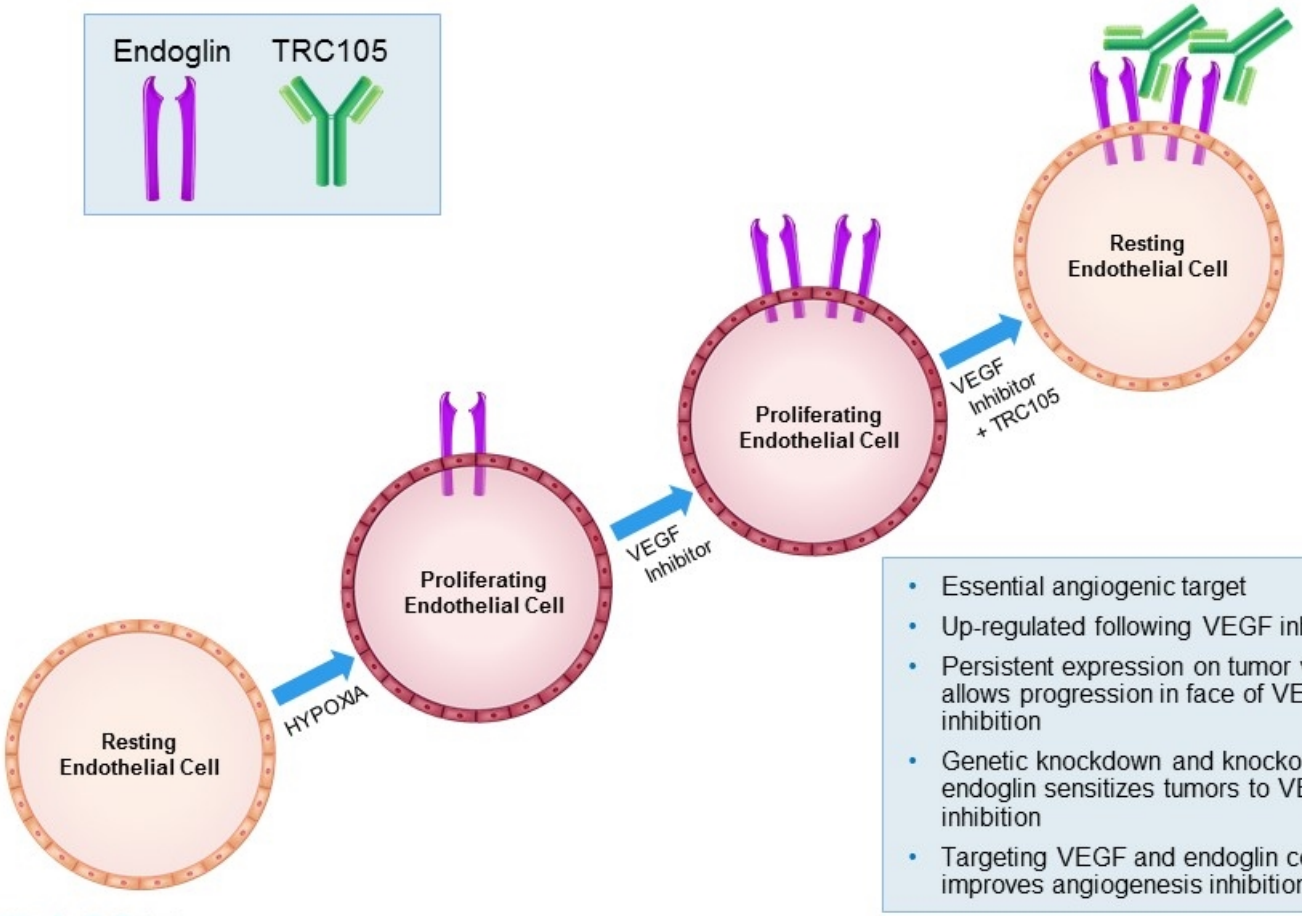
Indication	Approved VEGF Inhibitors	2017 VEGF Inhibitor Revenue[1]
2nd Line Renal Cell Carcinoma	Inlyta	$401 million
1st Line Hepatocellular Carcinoma	Nexavar	$1.0 billion[2]
2nd Line Soft Tissue Sarcoma	Votrient	~$150 million[3]
Colorectal Cancer, Lung Cancer	Avastin, Cyramza, Zaltrap, Stivarga	>$5 billion[4]
Wet AMD	Eylea Lucentis	$5.2 billion $3.2 billion

Substantial opportunity to build upon multiple established VEGF inhibitor franchises by improving patient outcomes through improved inhibition of angiogenesis

1 Company reports, SEC filings, DataMonitor.
2 Nexavar is approved in HCC, RCC and thyroid cancer. The majority of Nexavar's sales are in HCC.
3 Votrient is approved in both HCC and advanced STS. Estimated sales for Votrient in STS (based on 2014 total sales less DataMonitor estimates in RCC).
4 Based on Company estimates of sales by indication for Avastin and Cyramza.

TRACON PHARMA

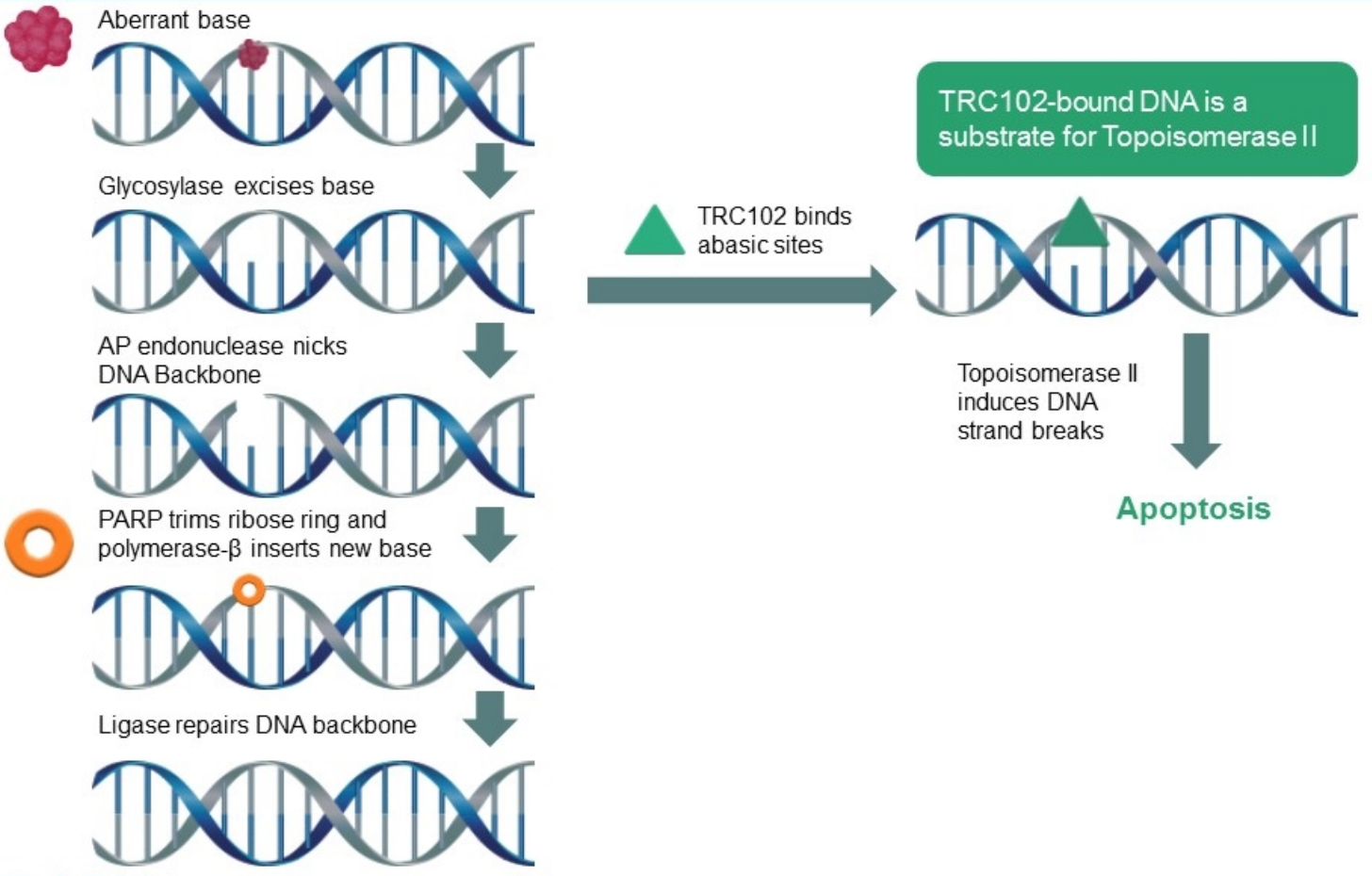
Targeting Endoglin Complements VEGF Inhibition



- Essential angiogenic target
- Up-regulated following VEGF inhibition
- Persistent expression on tumor vessels allows progression in face of VEGF inhibition
- Genetic knockdown and knockout of endoglin sensitizes tumors to VEGF inhibition
- Targeting VEGF and endoglin concurrently improves angiogenesis inhibition

TRACON PHARMA

TRC102: Reversing Resistance to Chemotherapy



TRACON PHARMA