



# SCHOLAR ROCK

*Deep Insights  
Impactful Medicines*

June 2020

# Disclaimers

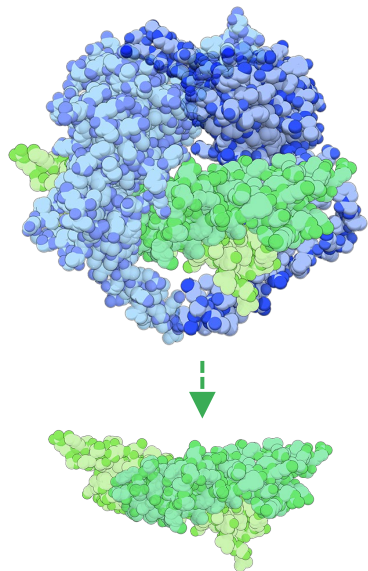
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Various statements in this presentation concerning Scholar Rock's future expectations, plans and prospects, including without limitation, Scholar Rock's expectations regarding its strategy, its product candidate selection and development timing, including timing for the initiation of and reporting results from its clinical trials for its product candidates, its disease indication selection and timing for such selection, the ability of SRK-015 to affect the treatment of patients suffering from Spinal Muscular Atrophy (SMA) either as a monotherapy or in conjunction with the current standard of care, the ability of SRK-181 to affect the treatment of cancer patients in a manner consistent with preclinical data constitute forward-looking statements for the purposes of the safe harbor provisions under The Private Securities Litigation Reform Act of 1995. The use of words such as "may," "might," "will," "should," "expect," "plan," "anticipate," "believe," "estimate," "target," "project," "intend," "future," "potential," or "continue," and other similar expressions are intended to identify such forward looking statements. Actual results may differ materially from those indicated by these forward-looking statements as a result of various important factors, including, without limitation, Scholar Rock's ability to provide the financial support and resources necessary to identify and develop multiple product candidates on the expected timeline, competition from others developing products for similar uses, the preliminary nature of interim clinical data, Scholar Rock's ability to obtain, maintain and protect its intellectual property, Scholar Rock's dependence on third parties for development and manufacture of product candidates including to supply any clinical trials, and Scholar Rock's ability to manage expenses and to obtain additional funding when needed to support its business activities and establish and maintain strategic business alliances and new business initiatives as well as those risks more fully discussed in the section entitled "Risk Factors" in the Quarterly Report on Form 10-Q for the quarter ended March 31, 2020, which is on file with the Securities and Exchange Commission, as well as discussions of potential risks, uncertainties, and other important factors in Scholar Rock's subsequent filings with the Securities and Exchange Commission. Any forward-looking statements represent Scholar Rock's views only as of today and should not be relied upon as representing its views as of any subsequent date. Scholar Rock explicitly disclaims any obligation to update any forward-looking statements unless required by law.

# Differentiated Approach to Highly Sought After Growth Factors Implicated in Devastating Diseases

## Scholar Rock's Target

### Growth Factor Precursor (Latent Form)



Targeting the activation of growth factor precursors to :

- Optimize potency and selectivity
- Localize effect

**Traditional Target**  
"Mature" Growth Factor

## Spinal Muscular Atrophy

**SRK-015:** Inhibitor of latent myostatin activation

**TOPAZ** Phase 2 trial ongoing

## Immuno-Oncology

**SRK-181:** Inhibitor of latent TGF $\beta$ 1 activation

**DRAGON** Phase 1 proof-of-concept trial ongoing

## Fibrosis

Inhibitors of latent TGF $\beta$  activation

**Advancing collaboration with Gilead towards product candidate selection**

*Growing preclinical and clinical progress using Scholar Rock's differentiated approach*

# 2020/2021: Transformative Years with Multiple Planned Clinical Read-Outs

1Q20 ✓ 4Q20 1H21 2021

## Initiated DRAGON SRK-181 Phase 1 POC trial

- Two-part trial with dose escalation (Part A) and dose expansion cohorts (Part B)
- Patients with locally advanced or metastatic solid tumors
- Exhibit primary resistance to anti-PD-(L)1 antibodies
- Evaluate multiple cancers, such as urothelial carcinoma, melanoma, and NSCLC, in combination with an approved anti-PD-(L)1 therapy

## Interim efficacy and safety results from SRK-015 TOPAZ trial

- Enrollment completed for all 3 cohorts in January<sup>†</sup>
- Analysis will be of patients who have progressed through at least 6 months of the treatment period

## Update on dose escalation from SRK-181 DRAGON Phase 1 POC trial

- Part A: SRK-181 as a single agent and in combination with anti-PD-(L)1 therapy

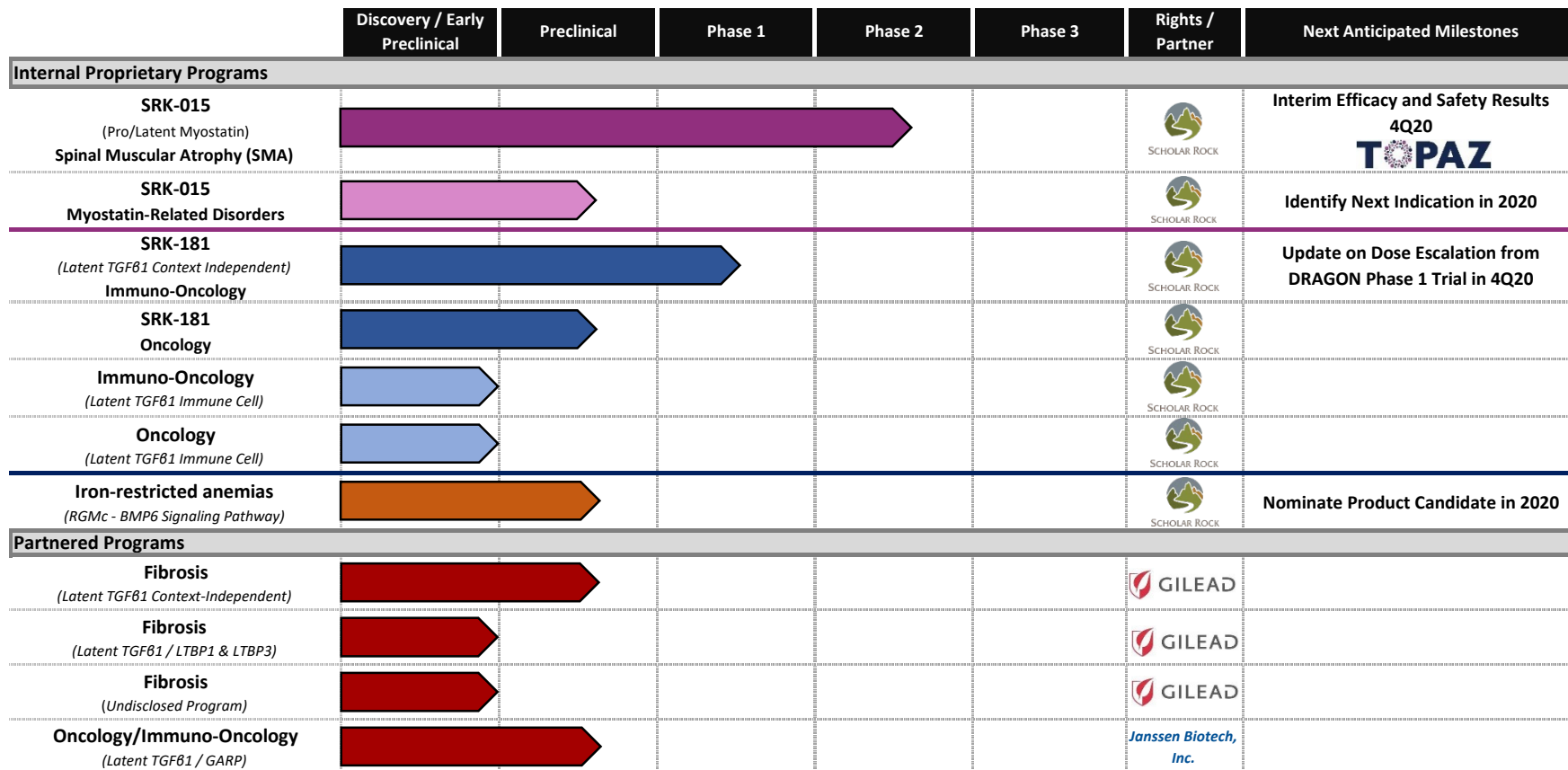
## Top-line 12-month results from SRK-015 TOPAZ trial

- Full 12-month treatment period
- Patients eligible to continue treatment for an additional 12 months

## Clinical response and safety data from SRK-181 Phase 1 POC trial

- Part A and Part B clinical data
- Multiple parallel cohorts to evaluate the anti-tumor activity of SRK-181 in combination with anti-PD-(L)1 therapy

# Building Differentiated Pipeline; Pairing Revolutionary Approach with Proven Modality



# **SRK-181: Potential Transformative Backbone for a New Era of Cancer Immunotherapy**



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# Human Tumor Analyses Reveal TGF $\beta$ as Key Determinant of Primary Resistance to Checkpoint Inhibitor (CPI) Therapies



## TGF $\beta$ 1 creates “immune-excluded” tumor microenvironment

Nature (on-line), Feb. 14, 2018

### TGF $\beta$ attenuates tumour response to PD-L1 blockade by contributing to exclusion of T cells

Sanjeev Mariathasan<sup>1\*</sup>, Shannon J. Turley<sup>1\*</sup>, Dorothee Nickles<sup>1\*</sup>, Alessandra Castiglioni<sup>1</sup>, Kobe Yuen<sup>1</sup>, Yulei Wang<sup>1</sup>, Edward E. Kadel III<sup>1</sup>, Hartmut Koeppen<sup>1</sup>, Jillian L. Astarita<sup>1</sup>, Rafael Cubas<sup>1</sup>, Suchit Jhunjhunwala<sup>1</sup>, Romain Banchereau<sup>1</sup>, Yagai Yang<sup>1</sup>, Yinghui Guan<sup>1</sup>, Cecile Chalouni<sup>1</sup>, James Ziai<sup>1</sup>, Yasin Şenbabaoğlu<sup>1</sup>, Stephen Santoro<sup>1</sup>, Daniel Sheinson<sup>1</sup>, Jeffrey Hung<sup>1</sup>, Jennifer M. Giltzane<sup>1</sup>, Andrew A. Pierce<sup>1</sup>, Kathryn Mesh<sup>1</sup>, Steve Lianoglou<sup>1</sup>, Johannes Riegler<sup>1</sup>, Richard A. D. Carano<sup>1</sup>, Pontus Eriksson<sup>2</sup>, Mattias Höglund<sup>2</sup>, Loan Somarriba<sup>3</sup>, Daniel L. Halligan<sup>3</sup>, Michiel S. van der Heijden<sup>4</sup>, Yohann Loriot<sup>5</sup>, Jonathan E. Rosenberg<sup>6</sup>, Lawrence Fong<sup>7</sup>, Ira Mellman<sup>1</sup>, Daniel S. Chen<sup>1</sup>, Marjorie Green<sup>1</sup>, Christina Derleth<sup>1</sup>, Gregg D. Fine<sup>1</sup>, Priti S. Hegde<sup>1</sup>, Richard Bourgon<sup>1</sup> & Thomas Powles<sup>8</sup>

Article

Cell

### Genomic and Transcriptomic Features of Response to Anti-PD-1 Therapy in Metastatic Melanoma

Authors

Willy Hugo, Jesse M. Zaretsky, Lu Sun, ...,  
Douglas B. Johnson, Antoni Ribas, Roger S. Lo

Volume 165, Issue 1, 24 March 2016, Pages 35-44

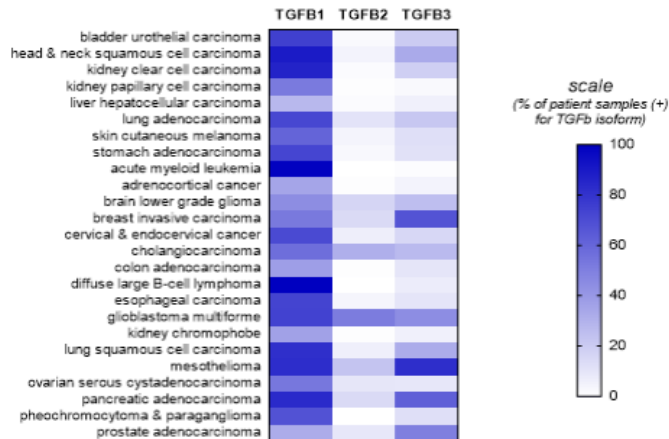
*Clinically-derived rationale points to significant opportunity to increase checkpoint therapy responses*

<sup>†</sup> Source: Carretero-Gonzalez et al. (2018) Oncotarget 9:8706-8715

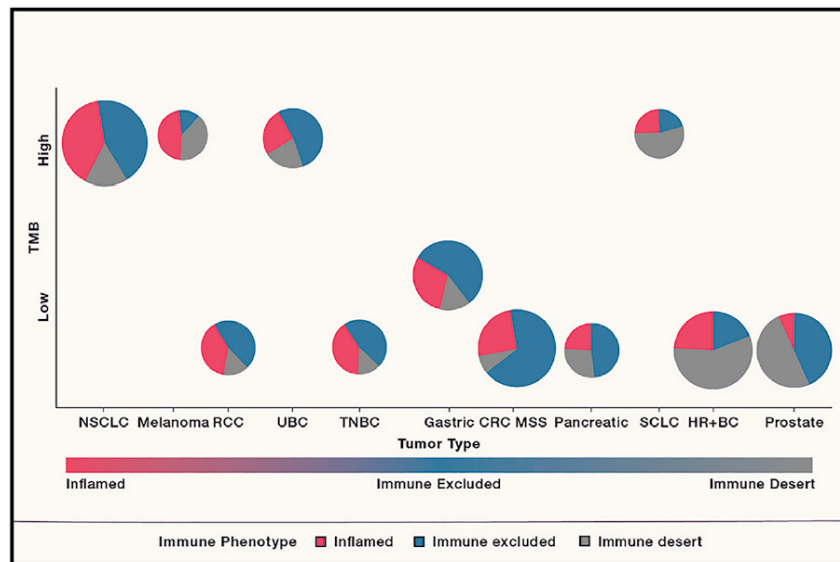
# Broad Potential for TGF $\beta$ Blockade Across Many Solid Tumors

- TGF $\beta$ 1 is most likely driver of TGF $\beta$  pathway signaling in human cancers
- TGF $\beta$ 1 expression correlates with TGF $\beta$  pathway activity in tumors

**Cancer Genome Atlas RNAseq analysis of >10,000 samples spanning 33 tumor types<sup>†</sup>**



## ***Substantial Proportion of Solid Tumors Exhibit Immune Exclusion<sup>††</sup>***

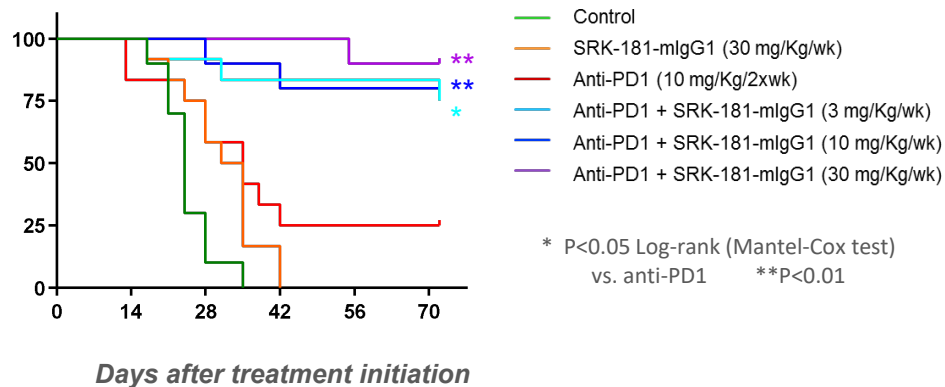
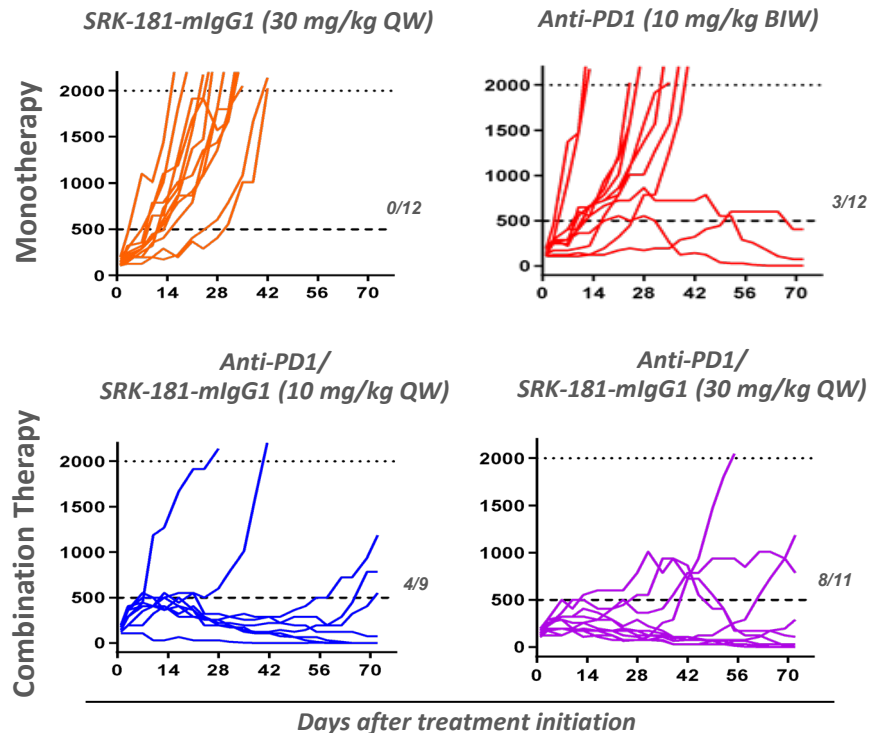


<sup>†</sup>Source: National Cancer Institute - Cancer Genome Atlas Program



# TGF $\beta$ 1 Blockade with SRK-181-mIgG1 Rendered Preclinical Tumor Models Susceptible to Anti-PD1 Therapy

## Cloudman S91 melanoma model: Combination treatment led to tumor regression and survival benefit



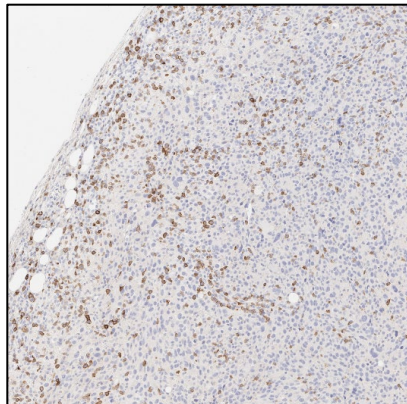
Similar results demonstrated in bladder model and breast cancer model (TGF $\beta$ 1/3 co-expressing)

### Selection Criteria for Mouse Tumor Models

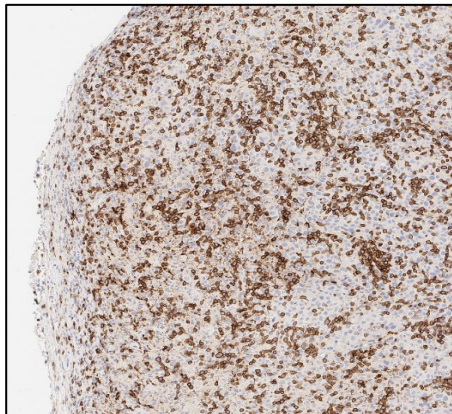
- Immune exclusion
- Minimal or no response to anti-PD-(L)1
- High TGF $\beta$ 1 over TGF $\beta$ 2/3 expression
- Evidence for TGF $\beta$  signaling

# SRK-181-mIgG1 Combination Therapy Enabled Infiltration and Expansion of CD8<sup>+</sup> T cells

*Anti-PD1*



*Anti-PD1/SRK-181-mIgG1 (10 mg/kg)  
led to increase in CD8<sup>+</sup> cells*



Turning “cold” tumors  
“hot,” and reduction in  
suppressive myeloid cells  
are consistent with  
significant anti-tumor  
responses

**In preclinical bladder cancer model, combination treatment with SRK-181-mIgG1/anti-PD1 led to:**

- Significant increase in effector T cells ( $p < 0.05$ )
  - Expansion of CD8<sup>+</sup> population to an average of 34% of the tumor's immune cells from a control average of 3.5%
- Significant decrease in intratumoral immunosuppressive myeloid cells ( $p < 0.05$ )
  - Reduction in TAM/MDSC population to 14% of the tumor's immune cells from a control average of 47%
  - Reduction in MDSC population to 1.4% from 11% of CD45<sup>+</sup> cells in the IgG control group

# TGFβ1 Isoform Specificity of SRK-181

## Improved Preclinical Toxicity Profile

Microscopic observations in heart	Control	LY2109761	PanTGFβAb	SRK-181			Legend
	vehicle iv, qwk x 4	300 mg/kg po, qd x 8	30 mg/kg iv, 1 dose	10 mg/kg iv, qwk x 4	30 mg/kg iv, qwk x 4	100 mg/kg iv, qwk x 4	
Valvulopathy							Unremarkable
Atrium - Mixed cell infiltrate							Minimal
Myocardium - Degeneration/necrosis							Slight
Myocardium - Hemorrhage							Moderate
Myocardium - Mixed cell infiltrate, base							
Coronary artery - Necrosis with inflammation							
Cardiomyocyte - Necrosis/inflammatory cell infiltrate							

### Repeat dose pilot toxicology study in adult female Sprague Dawley rats:

- Cardiac findings were exhibited in animals dosed with a pan-TGFβ antibody or LY2109761 (inhibitor of ALK5, common TGFβ receptor kinase) as expected based on published data†
- No cardiotoxicities (valvulopathy) were noted with SRK-181
  - NOAEL for SRK-181 was the highest dose evaluated of 100 mg/kg QW

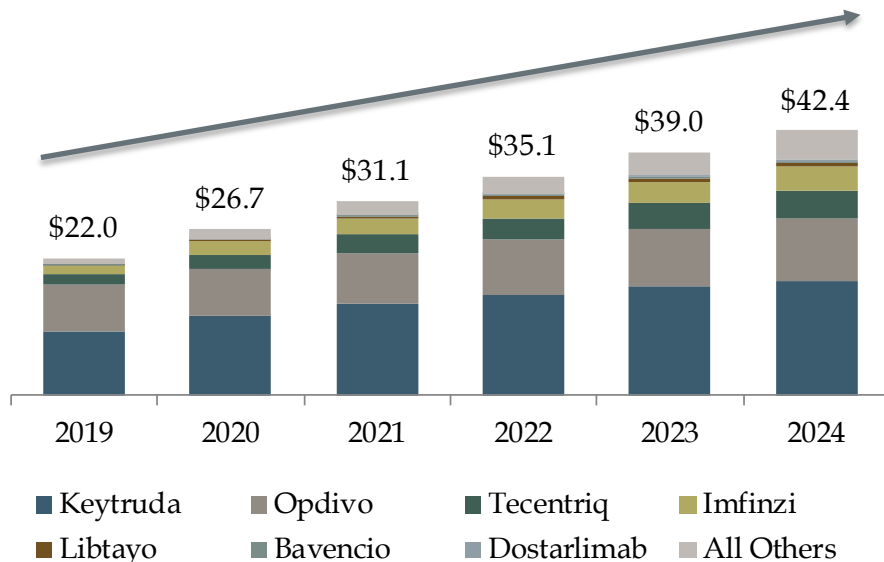
### 4-week GLP toxicology studies:

- Rats: NOAEL for SRK-181 was up to highest evaluated dose of 200 mg/kg QW
- Non-human primates: NOAEL for SRK-181 was up to highest evaluated dose of 300 mg/kg QW

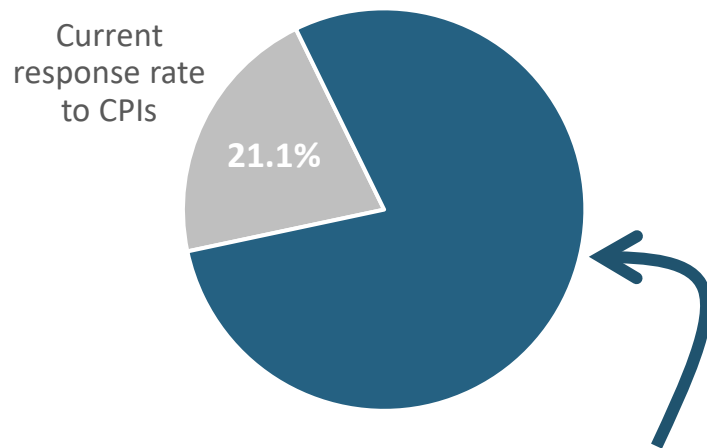
*Selectivity of SRK-181 offers potential to overcome toxicity and dose-limiting challenges of non-selective TGFβ pathway approaches*

# SRK-181 Has Potential to Increase Response to CPIs Through Clinically Relevant Resistance Pathway

Market for checkpoint inhibitor therapies expected to double over the next few years<sup>†</sup>...



...Yet Medical Need Not Addressed by Current Era of Immunotherapy<sup>††</sup>:



**SRK-181 has potential to substantially increase the addressable patient population for all checkpoint therapies**

# DRAGON Phase 1 POC Trial to Evaluate SRK-181's Ability to Overcome Primary Resistance to Checkpoint Inhibitors

*Update on Dose Escalation in 4Q20; Clinical Response and Safety Data in 2021*

## Part A

### **Part A1:**

- SRK-181 as a single agent
- Modified 3+3 dose escalation
- Assess SRK-181 dose range of 80-2400 mg (avg weight 80kg)

### **Part A2:**

- SRK-181 with approved anti-PD-(L)1
- 3+3 dose escalation
- Treat with same anti-PD-(L)1 as previously tried but did not experience a response

## Part B

- SRK-181 in combo with approved anti-PD-(L)1
- Multiple parallel cohorts – each will enroll up to 40 patients
- Target indications expected to include NSCLC, urothelial carcinoma, melanoma, amongst other solid tumor types
- Treat with same anti-PD-(L)1 as previously tried but did not experience a response
- Most recent dose of anti-PD-(L)1 therapy within six months of enrollment

### **SRK-181:**

*selective  
inhibitor of  
TGFβ1  
activation*

- Open-label, dose escalation, and dose expansion clinical trial
- Evaluate the efficacy, safety/tolerability, and PK/PD of SRK-181 in combination with approved anti-PD-(L)1 therapy
- Patients with locally advanced or metastatic solid tumors that exhibit primary resistance to anti-PD(L)1 therapy
- Lack of response characterized as stable or progressive disease following ≥3 cycles of anti-PD-(L)1 therapy either alone or in combination with chemotherapy

# SRK-015: Potential First Muscle-Directed Therapy for Spinal Muscular Atrophy



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# Significant Unmet Need Remains Despite Current Therapies

## Muscle-Directed Therapies Needed to Complement Disease-Stabilizing Benefits of SMN Upregulators\*

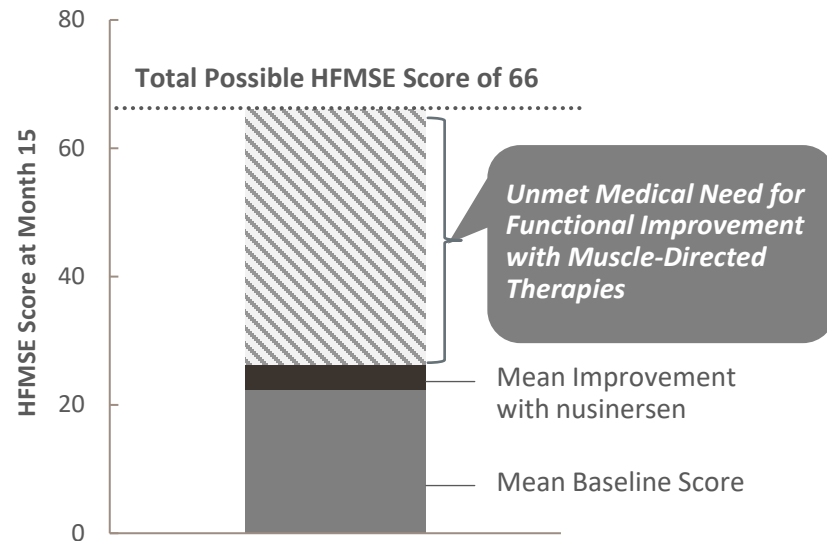
### SMN Upregulator Therapies

Address SMN deficiency to prevent further motor neuron deterioration

### Muscle-Directed Therapies (SRK-015)

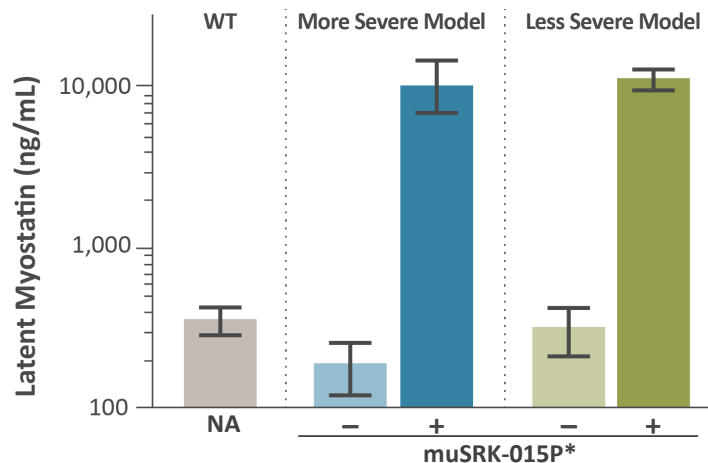
Act directly on muscle with aim to improve motor function

## Muscle Function in SMA (Human) *Hammersmith Functional Motor Scale Expanded (HFMSE)*

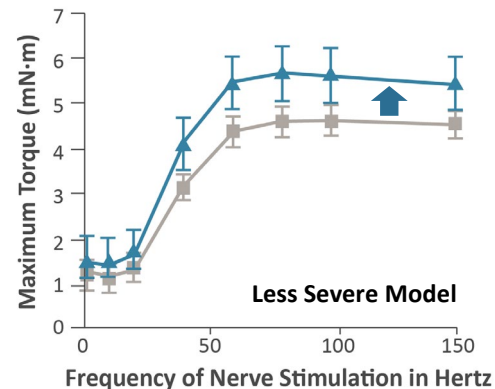
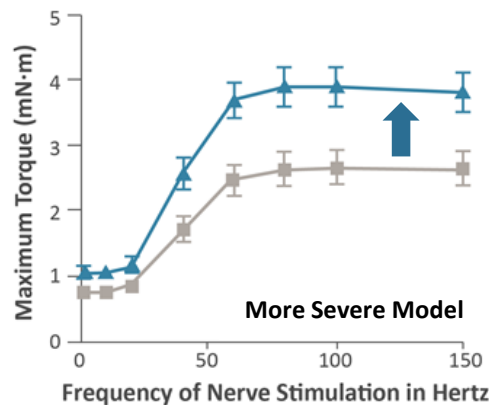


*Mean improvement in HFMSE score experienced by patients with later-onset SMA in the Phase 3 CHERISH clinical trial of nusinersen*

# Treatment of SMN $\Delta$ 7 Mouse Models Show Improvement in Muscle Strength



- Achieved multi-fold increase in serum latent myostatin levels indicating target engagement
- Confirms presence of target in disease setting
- Lower latent myostatin levels in the SMA group may be attributable to reduced overall muscle mass



SMN upregulator + muSRK-015P  
SMN upregulator only

## Treatment improved muscle strength:

- Maximal torque of the plantar flexor muscle group increased:
  - More severe model: 44%-51%
  - Less severe model: 20%-30%

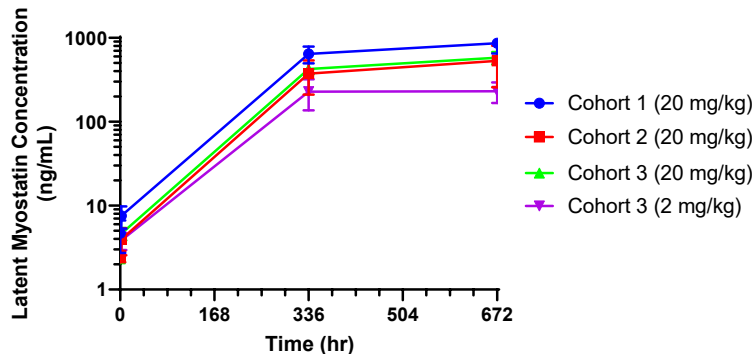
\*muSRK-015P is the parental clone of SRK-015 on a mouse IgG1 framework

Preclinical data published Nov 2018: Long, K., O'Shea, K., Khairallah, R., et al. Specific Inhibition of Myostatin Activation is Beneficial in Mouse Models of SMA Therapy. Human Molecular Genetics, ddy382 <https://scholarrock.com/platform/publications/>



# Preliminary TOPAZ Biomarker Data Provide First Demonstration of Target Engagement in Patients with SMA

## Latent Myostatin Change over Baseline in SRK-015 TOPAZ Trial



## Robust Target Engagement Observed

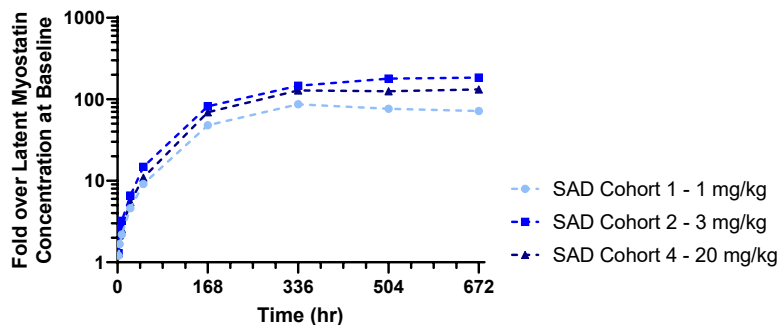
- ~100-fold increases in serum latent myostatin levels following single 20 mg/kg dose in all cohorts of TOPAZ

- Confirms presence of latent myostatin in patients with SMA

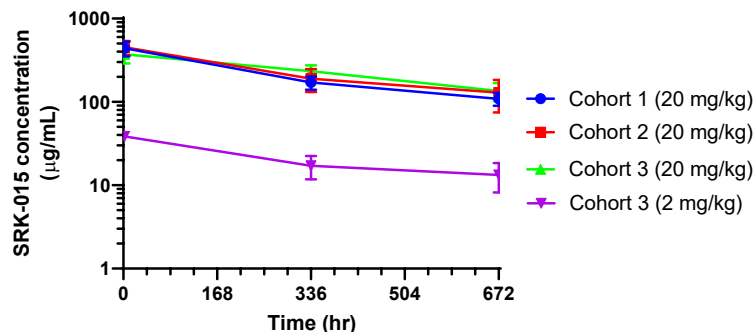
## Well-Behaved, Linear PK Profile

- Minimal variability across TOPAZ cohorts
- Dose proportional increase in serum drug exposure between low (2 mg/kg) and high (20 mg/kg) doses

## Latent Myostatin Change over Baseline in Phase 1 HV Trial



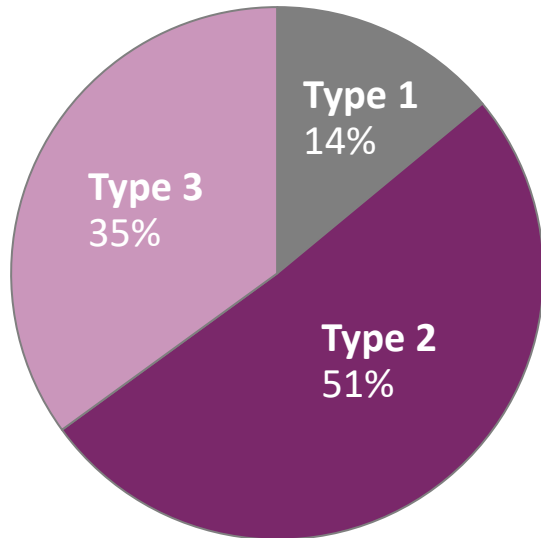
## Preliminary TOPAZ Phase 2 Pharmacokinetic (PK) Data



# SRK-015 Opportunity in Spinal Muscular Atrophy

## Overall Prevalence of 30,000-35,000 in U.S. and Europe

Relative Prevalence Among  
Patients Living With SMA



### **Type 1:**

- Infant-onset; often fatal

### **Type 2 and non-ambulatory type 3:**

- Later-onset but still early childhood
- Severe deficits in motor function

### **Ambulatory type 3:**

- Limited mobility and substantial morbidity

### **Type 4:**

- Population not well-characterized

### **Focus of TOPAZ Trial**

*Potential to use  
SRK-015 in conjunction  
with SMN upregulators*

*Potential to use SRK-015  
as monotherapy or in  
conjunction with SMN  
upregulators*

**...potential to expand more  
broadly in future trials**

# SRK-015 Phase 2 Trial Design



*Interim Efficacy and Safety Results 4Q20; Top-line 12-Month Data 1H21*

	Cohort 1	Cohort 2	Cohort 3
Design	<ul style="list-style-type: none"> <li>N= 23<sup>††</sup>; ages 5-21</li> <li>Open-label, single-arm</li> <li>20 mg/kg SRK-015 IV Q4W</li> <li>12-month treatment period</li> </ul>	<ul style="list-style-type: none"> <li>N= 15; ages 5-21</li> <li>Open-label, single-arm</li> <li>20 mg/kg SRK-015 IV Q4W</li> <li>12-month treatment period</li> </ul>	<ul style="list-style-type: none"> <li>N= 20; ages ≥2</li> <li>Double-blind, randomized (1:1) to 2 mg/kg or 20 mg/kg SRK-015 IV Q4W</li> <li>12-month treatment period</li> </ul>
Patients	<ul style="list-style-type: none"> <li>Ambulatory Type 3 SMA</li> <li>Receiving treatment with approved SMN upregulator or as monotherapy</li> </ul>	<ul style="list-style-type: none"> <li>Type 2 or non-ambulatory Type 3 SMA</li> <li>Receiving treatment with approved SMN upregulator</li> </ul>	<ul style="list-style-type: none"> <li>Type 2 SMA</li> <li>Initiated treatment with approved SMN upregulator before age 5</li> </ul>
Primary Objectives	<ul style="list-style-type: none"> <li>Safety</li> <li>Mean change from baseline in RHS</li> </ul>	<ul style="list-style-type: none"> <li>Safety</li> <li>Mean change from baseline in HFMSE</li> </ul>	<ul style="list-style-type: none"> <li>Safety</li> <li>Mean change from baseline in HFMSE</li> </ul>

*Each cohort evaluates a subpopulation that is sizable and has substantial unmet medical need*

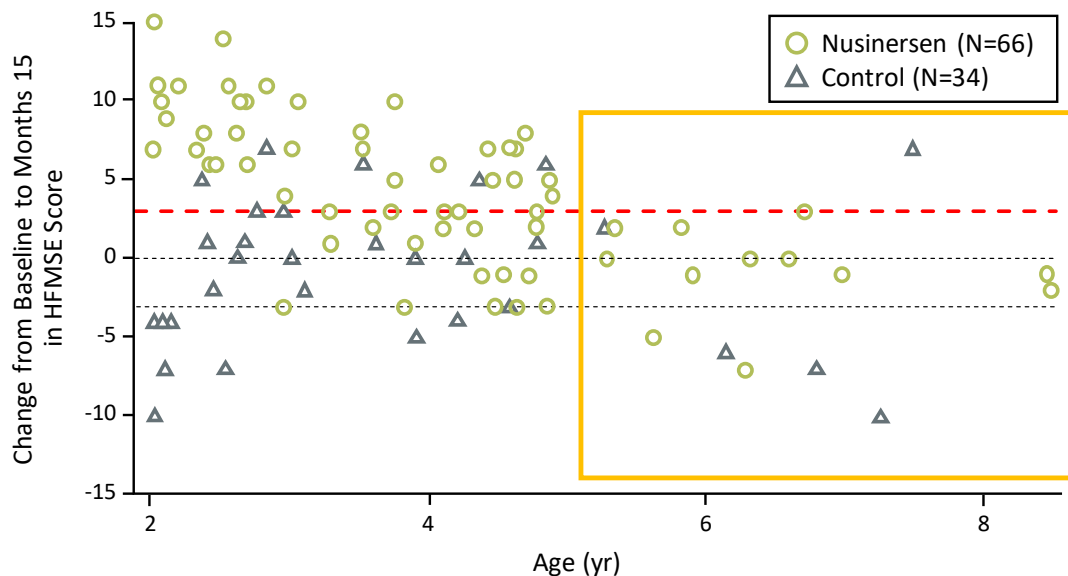
HFMSE – Hammersmith Functional Motor Scale Expanded; RHS – Revised Hammersmith Scale

<sup>†</sup>Source: Mercuri, E. et.al. Nusinersen versus Sham Control in Later-Onset Spinal Muscular Atrophy, *N Engl J Med* 2018; 378:625-635

<sup>††</sup> Baseline demographics presented as part of AAN virtual platform (May 2020) <https://scholarrock.com/platform/publications/>

# Later-Onset SMA: High Unmet Need to Improve Motor Function

## Nusinersen CHERISH Trial in Later-Onset SMA



*In patients with later-onset SMA who were age  $\geq 5$  at screening...*

- Primary benefit of nusinersen appeared to be stabilization of motor function (in HFMSE) rather than improvement from baseline
- Attainment of  $\geq 3$ -point increase was rare (<15% of patients) even with nusinersen treatment

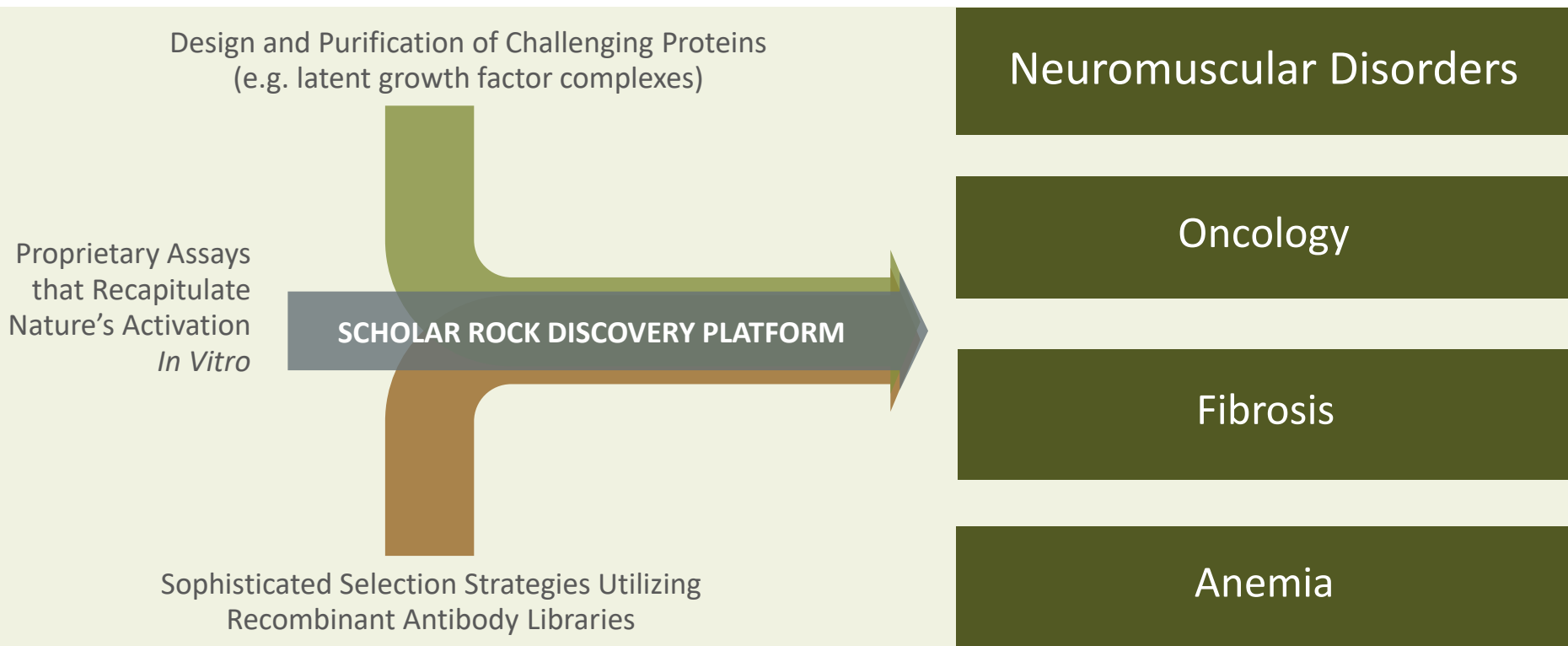
*3-point improvement in a patient is considered clinically meaningful and rare to observe*

# Building Differentiated Pipeline Through Highly Productive Platform



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# Proprietary Platform to Target Latent Growth Factor Activation



# Fibrosis Partnership with Gilead Advances with Achievement of First Milestone

## Scholar Rock's highly specific inhibitors of latent TGF $\beta$ activation:

- Aim to improve efficacy and tolerability compared to traditional non-selective approaches
- Demonstrated efficacy in *in vivo* preclinical models

Upfront cash and equity investment:  
**\$80 million\***

Additional milestones across 3 programs:  
**Up to \$1,425 million**

One-time preclinical milestone:  
**\$25 million**  
(achieved end of 2019)

High single- to low double-digit tiered royalties on net sales



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- Antibody discovery and preclinical research thru product candidate nomination
- Distinct antibodies
- Retains exclusive WW rights for oncology and cancer immunotherapy



GILEAD

- Upon option exercise, responsible for preclinical and clinical development and commercialization

*TGF $\beta$ -driven signaling broadly implicated as a central regulator of fibrosis*

# Upcoming Key R&D Milestones

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## Spinal Muscular Atrophy (SMA)

- Interim efficacy and safety results from SRK-015 TOPAZ Phase 2 trial in 4Q20\*
    - Patients from 3 cohorts progressed thru at least 6 months of the treatment period
  - Top-line results (12 months) from SRK-015 TOPAZ trial in 1H21\*
    - Patients are eligible to continue treatment for additional 12 months
  - Identification of second indication for SRK-015 in 2020
- 

## Oncology

- Update on dose escalation from SRK-181 DRAGON Phase 1 POC trial in 4Q20\*
    - SRK-181 as a single-agent as well as in combination with approved anti-PD-(L)1 therapy
  - Clinical response and safety data from SRK-181 Phase 1 POC trial in 2021\*
  - Continue to advance active discovery programs for context-dependent inhibition of TGFβ1
- 

## Fibrosis

- Continue to advance collaborative programs with Gilead towards product candidate selection
- 

## Anemias

- Nominate product candidate in RGMc program in 2020



# Appendix



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# Our Purpose

*Relentlessly focused on seeing new possibilities in validated biologies and antibody technologies to allow us to move with speed and urgency to deliver transformative medicines to patients with devastating diseases*



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## Revolutionary Approach

- First company to successfully target growth factor precursors
- Pursue high value targets proven challenging for traditional non-selective approaches
- Focus on biologically validated targets using proven modality (mAbs)
- Leverage protein science and antibody expertise to develop high-impact medicines for patients suffering neuromuscular disorders, cancer, fibrosis and anemia
- Develop broad IP portfolio covering compositions and methods

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## Differentiated Pipeline Portfolio

- Two product candidates, SRK-015 and SRK-181, in clinical development
- Multiple clinical read-outs offer near-term value inflection points
- Potential to expand pipeline with additional indications for each product candidate
- Strategic fibrosis collaboration with Gilead to develop potent and selective inhibitors of latent TGF $\beta$  activation
- Applying expertise across the TGF $\beta$  superfamily of growth factors and beyond

# Differentiated Approach with SRK-181

- *Fully human mAb*
- *Potent and selective inhibitor of latent TGFβ1 activation*
- *Minimal or no binding to latent TGFβ2 and TGFβ3 isoforms*
- *Designed for exquisite selectivity for TGFβ1 to avoid the cardiac tox often seen with traditional, less-selective approaches*



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## Strong preclinical data shows potential of SRK-181 in overcoming primary resistance to checkpoints

- ~80% of patients with metastatic solid tumors do not respond to CPIs<sup>†</sup>
- Strong human translational data and preclinical models implicate TGFβ as key culprit in primary resistance to CPIs<sup>††</sup>
- Exquisite selectivity of SRK-181 offers potential to reduce toxicity and avoid dose-limiting challenges
- Rationally designed preclinical studies demonstrate potential of SRK-181 in overcoming primary resistance

### Commenced dosing of DRAGON Phase 1 POC trial for SRK-181 in 2Q20

- Update on dose escalation in 4Q20
- Clinical response and safety data in 2021

## *Preclinical efficacy and safety results point toward a clinically feasible path forward*

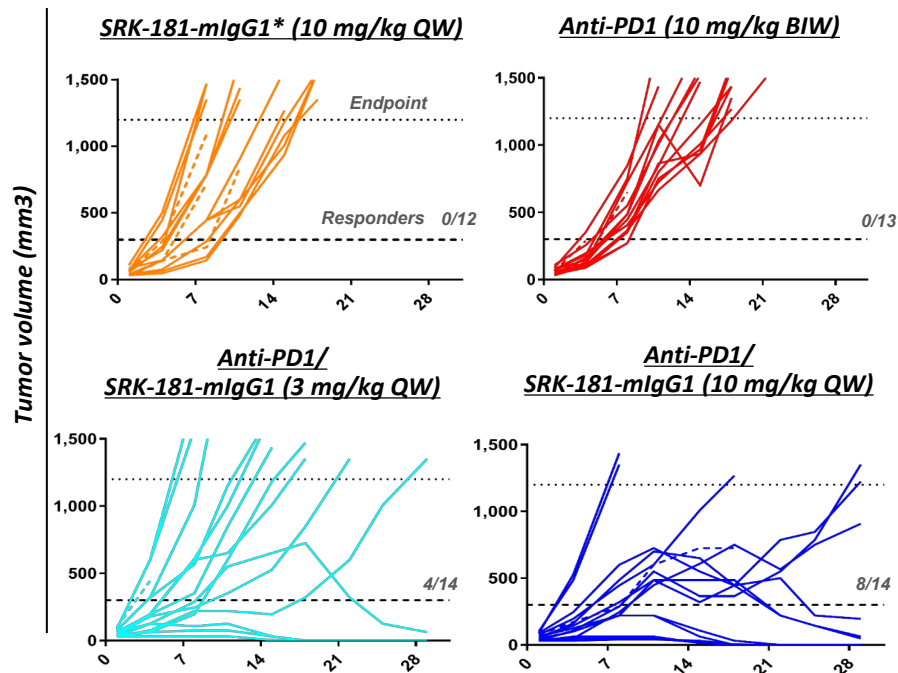
Refer to posters presented at SITC (Nov 2018) and AACR (April 2019) available at <https://scholarrock.com/platform/publications/>

<sup>†</sup>Source: Carretero-Gonzalez et al. (2018) Oncotarget 9:8706-8715

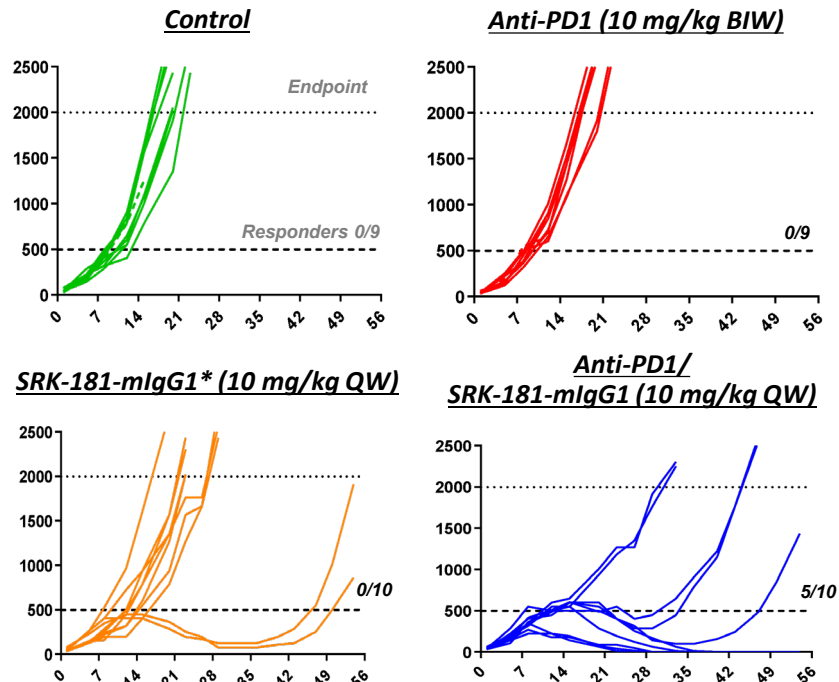
<sup>††</sup>Source: Mariathasan, Turley, et.al TGFβ attenuates tumour response to PD-L1 blockade by contributing to exclusion of T cells, Nature (online), Feb 2018

# TGF $\beta$ 1 Blockade with SRK-181-mIgG1 Rendered Preclinical Tumor Models Susceptible to Anti-PD1 Therapy

## Bladder Cancer



## Breast Cancer



Days after treatment initiation

## SRK-015: Selective Inhibitor of Latent Myostatin Activation

- *Fully human mAb*
- *Half-life of 23-33 days*
- *Orphan Drug Designation for SMA granted by FDA and EC*
- *Strong patent portfolio:*
  - *US Patent 9,758,576 covers mAbs that inhibit the activation of myostatin precursor (expiry in 2034)*
  - *US Patent 10,287,345 covers treatment methods for various myostatin-related conditions (expiry in 2037)*



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## Preclinical and clinical data provide strong rationale for developing in Spinal Muscular Atrophy (SMA)

- Need to improve motor function remains despite the availability of SMN upregulators for the treatment of SMA
- SMA disease features align well with attributes of myostatin biology
- Studies in SMN $\Delta$ 7 mouse models demonstrated substantial increases in muscle strength
- Emerging pharmacologic profile for SRK-015 supports chronic therapy:
  - Evaluating Q4W dosing regimen
  - Phase 1/Phase 2 prelim. PD data show robust target engagement
  - No clinically significant safety signals observed as of data cutoff in preliminary Phase 2 PK/PD analysis

### TOPAZ Phase 2 trial: interim efficacy and safety data in 4Q20

- Top-line 12-month analysis anticipated 1H21

## Therapeutic potential as muscle-directed therapy to complement any SMN upregulator

Source: Mercuri, E. et.al. Nusinersen versus Sham Control in Later-Onset Spinal Muscular Atrophy, *N Engl J Med* 2018; 378:625-635

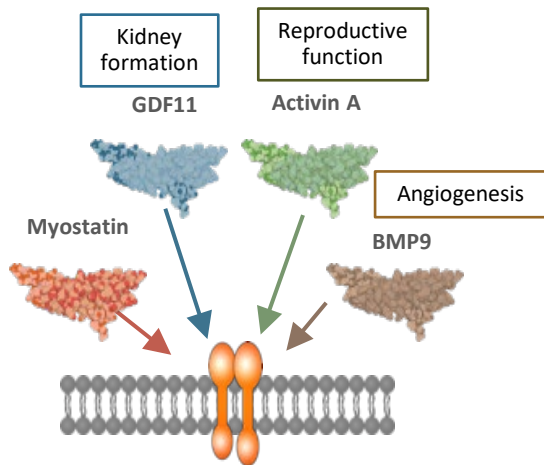
Refer to Phase 1 data poster at World Muscle Society (Oct 2019) at [www.scholarrock.com](http://www.scholarrock.com)

Refer to poster presentation at the MDA Clinical and Scientific Conference (March 2020) available at <https://scholarrock.com/platform/publications/>

# Traditional Approaches Can Raise Significant Safety Concerns

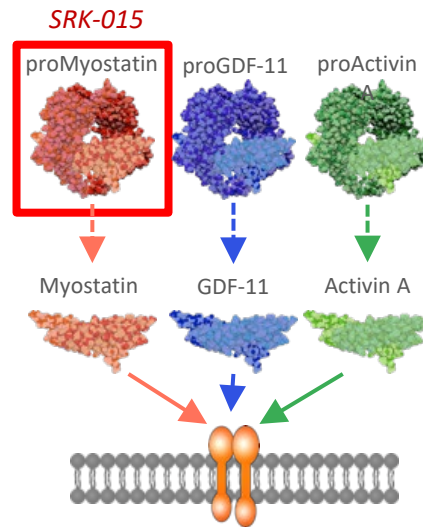
## Traditional Approaches Lack Selectivity

- Most inhibitors of active myostatin also inhibit GDF11 and may inhibit other growth factors as well
- Antibodies to ActRIIb and ligand trap approaches inhibit signaling of multiple ligands



## Scholar Rock Approach

### Exquisite Selectivity By Targeting Precursor Form of Myostatin



# Strategic collaboration focused on fibrosis

*Gilead has exclusive options to license worldwide rights to product candidates from 3 TGF $\beta$  programs:*

- *Inhibitors that target activation of latent TGF $\beta$ 1*
- *Inhibitors that selectively target activation of latent TGF $\beta$ 1 localized to extracellular matrix*
- *Undisclosed TGF $\beta$  discovery program*



SCHOLAR ROCK

## Potent and Selective Inhibitors of Latent TGF $\beta$ Activation Can Offer Novel Approach to Fibrotic Diseases

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- TGF $\beta$ -driven signaling has been broadly implicated as a central regulator of fibrosis<sup>†</sup>
  - Scholar Rock's potent and highly selective TGF $\beta$  inhibitors act locally in the disease microenvironment
  - Demonstrated preclinically that potent and selective inhibitors of TGF $\beta$ 1 signaling prevent the activation of the growth factor in the fibrotic matrix
  - Achieved first milestone and earned \$25 million payment with demonstration of efficacy in in vivo proof-of-concept studies
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***Advance collaboration towards product candidate selection***

<sup>†</sup>Kim KK, Sheppard D, Chapman HA (2018). TGF- $\beta$ 1 Signaling And Tissue Fibrosis. Cold Spring Harb Perspect Biol 10: a022293