



satellos

REGENERATING MUSCLE FROM WITHIN™

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Satellos: A New Chapter in Treating Muscle Diseases with SAT-3247



- Disruptive science
- Novel approach with potential to be disease modifying
- Well tolerated, easy-to-administer medication
- Areas of enormous unmet need

Corporate Highlights

1.

Clinical stage company

2.

Strong balance sheet

3.

Upcoming flow of milestones

4.

Accomplished leadership team

5.

Elite institutional investors

Grounded in science. Inspired by need.

Our mission is to improve the treatment of Duchenne muscular dystrophy and other devastating muscle diseases by restoring the body's ability to regenerate muscle.



CHARLIE, age 7,
living with Duchenne

SAT-3247 is Totally Different: Not Just Stabilizing — Regenerating

Genetic approaches

- Administered by infusion or injection
- Gene therapy (DNA) or exon skipping (RNA)
- Aims to slow disease progression by stabilizing muscle
- Limited by exon status or potential immunogenicity concerns



Satellos's approach

- Oral, once-daily pill
- Designed to be safe and tolerable
- Aims to reverse disease progression by regenerating muscle
- Intended for all patients regardless of exon or immune status
- Novel mechanism with potential as stand-alone or add-on therapy

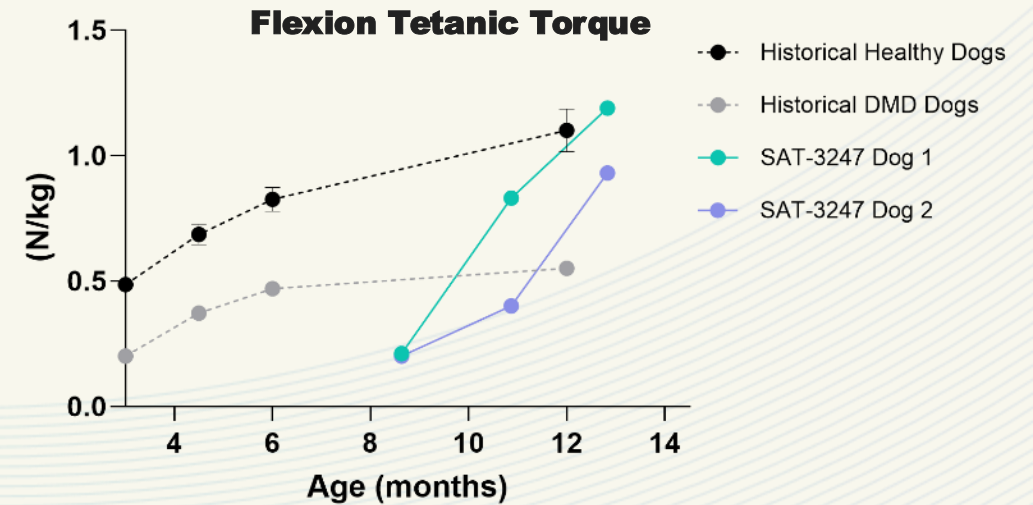
SAT-3247 Restores Muscle Strength in Canine Model of DMD to Normative Levels

Satellos Canine Study



- 4-month study
- 2 dogs ~9 months of age to ~13 months
- Retriever cross with natural DMD mutation
- Oral daily dosing with SAT-3247 (4x/week)
- Strength at 2 and 4 months

SAT-3247 treated vs. historical controls



Contraction force generated by tarsal joint flexion and extension in dogs with golden retriever muscular dystrophy

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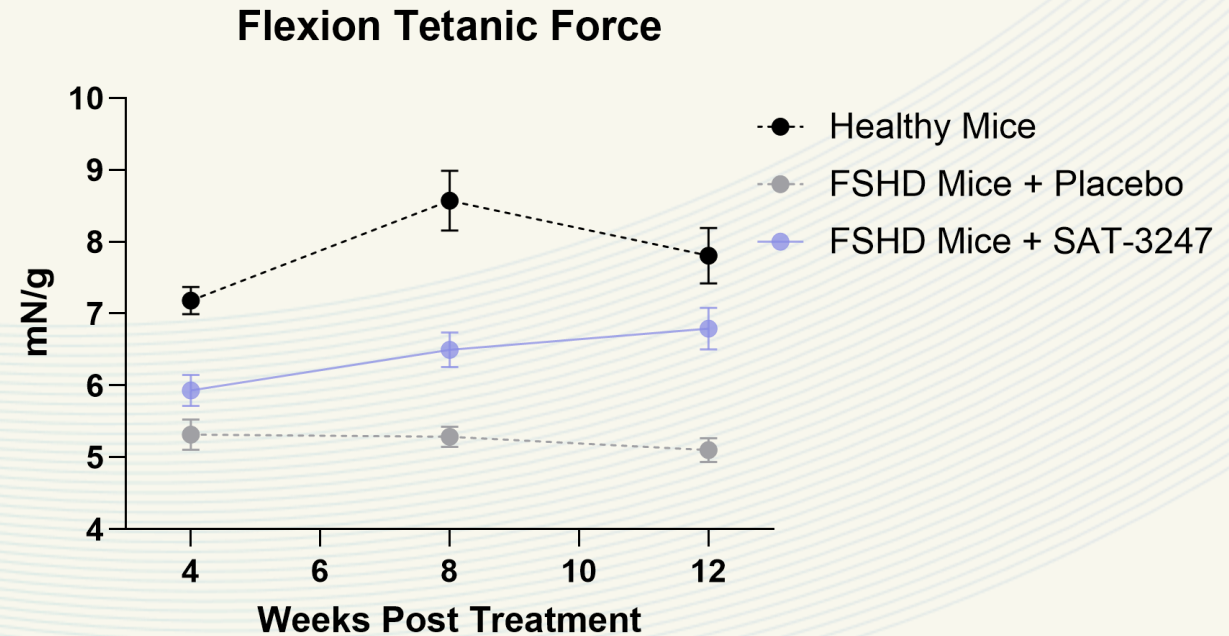
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SAT-3247 Restores Muscle Strength in Murine Model of FSHD to Near Normative Levels

Satellos FSHD Canada Joint Study

- 3-month study
- FLExDux4 mouse model of FSHD
- Oral daily dosing with SAT-3247 (4x/week)
- Strength at 4, 8, 12 weeks post administration

SAT-3247 vs placebo vs healthy animals



Advancing SAT-3247 into Clinical Development



SAT-3247 Phase 1 First-in-Human Clinical Study

Healthy Volunteers (n=72)			Adults with DMD
SAD	MAD	Crossover Food Effect	Multiple Dosing
<ul style="list-style-type: none">10 - 400 mg5 cohorts of 8Single dose / 1 day	<ul style="list-style-type: none">60 - 240 mg5 cohorts of 8Single dose / 7 days	<ul style="list-style-type: none">150 mg1 cohort of 8Single dose / 1 day	<ul style="list-style-type: none">60 mg1 cohort of up to 10Single dose for 5 consecutive weekdays 4 wks
Complete			Ongoing

Endpoints

Primary: Safety & tolerability

Secondary: Pharmacokinetics (NB., DMD adults also receiving their prescribed steroid treatment)

Exploratory*: Grip strength, 99th maximum effort, forced vital capacity, serum markers

*SAD: single ascending dose; MAD: multiple ascending dose; *adults with DMD*

SAT-3247 Appears Safe and Tolerable in Healthy Volunteer Study¹

Single Ascending Dose

No clinically significant findings observed in 10, 50, 150, 300 and 400 mg doses.

- ✓ Labs
- ✓ Vitals
- ✓ Physical Exam
- ✓ ECG

Drug-related adverse events, reversible:

- **400 mg:** mild nausea (1); mild abdominal pain (1)

Multiple Ascending Dose

No clinically significant findings observed in **60**, **120**, 180, 240 mg doses:

- ✓ Labs (mild transaminase elevations at 180 and 240 mg)
- ✓ Vitals
- ✓ Physical Exam
- ✓ ECG

Drug-related adverse events, reversible:

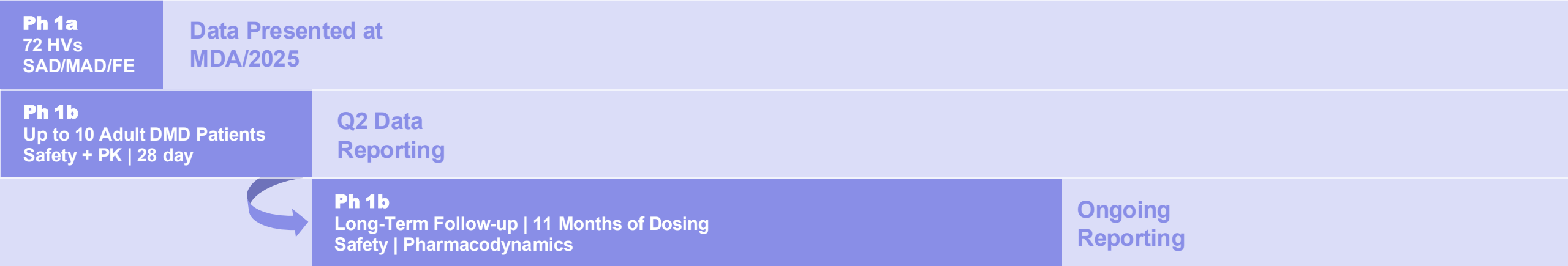
- **120 mg:** mild lightheadedness (1)
- **180 mg:** intermittent mild epigastric pain (1); mild sleepiness (1); mild lethargy (1)

¹ Based on 20 Feb 2025 data cut-off; study is ongoing

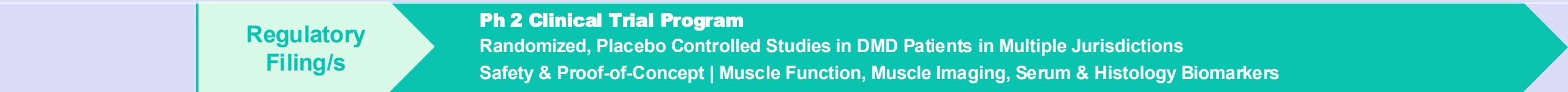
Clinical Program Overview

2025				2026			
Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4

Phase 1 Program (Underway: Australia)



DMD Phase 2 Program (Planned: USA, Canada, AUS, EU)



Ph 2 Clinical Program Designed to Produce Data Readouts at Pre-Defined Intervals

Inside Satellos

Our company and the team leading it forward



Satellos Has Built a World-Caliber Team



Frank Gleeson, MBA
Co-Founder & Chief
Executive Officer



Michael Rudnicki, OC, PhD, FRS
Co-Founder & Chief
Discovery Officer



Liz Williams, CPA, CA
Chief Financial Officer



Courtney Wells
SVP – Clinical
Development Operations



Philip Lambert, PhD
Chief Scientific Officer



Ryan Mitchell, PhD
SVP – Medical &
Scientific Affairs

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Institute of Child Health

Perry Shieh, MD, PhD, FAAN
Professor of Neurology, UCLA David
Geffen School of Medicine

Ronald Cohn, MD, PhD
President and CEO, Hospital for Sick
Children, Toronto

Reimagine

how muscle degeneration is treated.

Regenerate

with small molecule medicines.

Realize

the next horizon to improve lives.