

Second Quarter 2019 Financial Results

August 5, 2019

Acceleron Forward-Looking Statements

THIS PRESENTATION CONTAINS FORWARD-LOOKING STATEMENTS ABOUT THE COMPANY'S STRATEGY, FUTURE PLANS

and prospects, including statements regarding the development of the Company's compounds, the timeline for clinical development and regulatory approval of the Company's compounds and the expected timing for reporting of data from ongoing clinical trials. The words "anticipate," "believe," "could," "estimate," "expect," "goal," "intend," "may," "plan," "potential," "project," "should," "target," "will," "would," and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words.

ACTUAL RESULTS COULD DIFFER MATERIALLY FROM THOSE INCLUDED IN THE FORWARD-LOOKING STATEMENTS DUE to

various factors, risks and uncertainties, including, but not limited to, that preclinical testing of the Company's compounds and data from clinical trials may not be predictive of the results or success of ongoing or later clinical trials, that the results of any clinical trial may not be predictive of the results or success of other clinical trials of the same product candidate, that the development of the Company's compounds will take longer and/or cost more than planned, that the Company will be unable to successfully complete the clinical development of the Company's compounds, that the Company may be delayed in initiating, enrolling or completing any clinical trials, and that the Company's compounds will not receive regulatory approval or become commercially successful products. These and other risks and uncertainties are identified under the heading "Risk Factors" included in the Company's most recent Annual Report on Form 10-K, and other filings that the Company has made and may make with the SEC in the future.

THE FORWARD-LOOKING STATEMENTS CONTAINED IN THIS PRESENTATION ARE BASED ON MANAGEMENT'S CURRENT

views, plans, estimates, assumptions and projections with respect to future events, and the Company does not undertake and specifically disclaims any obligation to update any forward-looking statements.

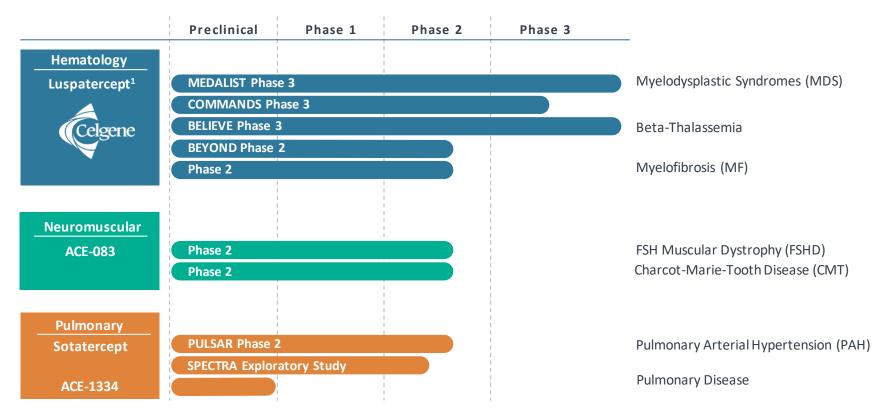




Habib DableChief Executive Officer



Building Therapeutic Area Leadership





US and EU Marketing Applications Accepted



Phase 3

Beta-Thalassemia, **Transfusion-Dependent** **December 4, 2019**

FDA PDUFA Target Action Date



Phase 3

Lower-Risk MDS, RS+

April 4, 2020

FDA PDUFA Target Action Date

EMA decision on MAA expected in 2H 2020



ACE-083 Part 2 of Phase 2 Trials Fully Enrolled

FSHD Part 2

ACE-083

N = 28 (14TA / 14BB)

Randomized 1:1

Placebo

N = 28 (14TA / 14BB)

Topline Outcome Measures:

- Percent change in muscle volume and change in fat fraction
- Percent change in function
 - TA cohort: 4-stair climb, 6-minute walk, 10-meter walk/run
 - BB cohort: performance upper-limb
- Improvement in health-related quality of life
 - FSHD-Health Index
- Safety and tolerability

Double-blind, placebo-controlled, 6-month primary treatment period

CMT Part 2

ACE-083

N = 20

Randomized 1:1

Placebo

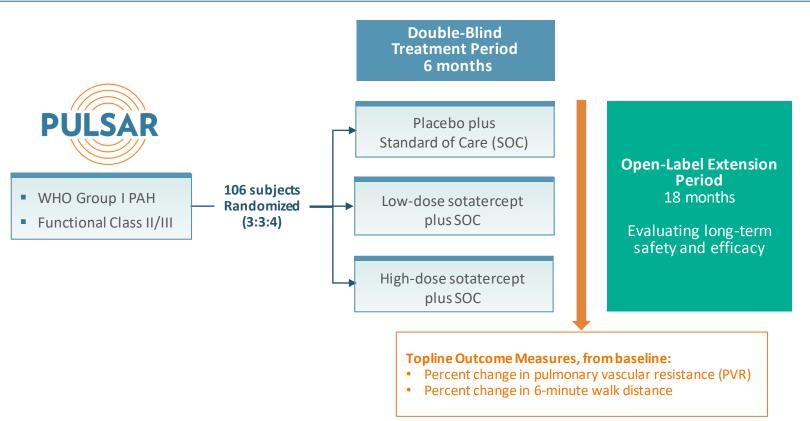
N = 20

Topline Outcome Measures:

- Percent change in muscle volume and change in fat fraction
- Percent change in function
 - 6-minute walk and 10-meter walk/run
- Improvement in health-related quality of life
 - CMT-Health Index
- Safety and tolerability



Sotatercept PULSAR Phase 2 Trial Fully Enrolled







Kevin McLaughlinChief Financial Officer



Q2 2019 Financial Results

Cash	
Cash, cash equivalents and investments	\$500.9M
Revenue	
Collaboration Revenue (includes regulatory milestone)	\$27.7M
Costs and Expenses	
Total Costs and Expenses	\$48.8M
R&D Expenses	\$34.8M
G&A Expenses	\$14.0M
Net Loss	
Net Loss	\$17.9M



Upcoming Corporate Priorities

HEMATOLOGY

- Luspatercept
 - FDA PDUFA target action dates:
 - Beta-thalassemia expected December 2019 and MDS expected April 2020
 - EMA decision on the MAA expected in 2H 2020
 - MEDALIST and BELIEVE Phase 3 trial results planned to be submitted for publication in 2019
 - Myelofibrosis Phase 2 trial results expected by YE 2019
 - BEYOND Phase 2 trial topline results expected by **YE 2020**, COMMANDS Phase 3 trial patient enrollment
 - Potential expansion of clinical program into other indications in 2019

NEUROMUSCULAR

- ACE-083
 - FSHD Part 2 of the Phase 2 trials topline results expected in 2H 2019
 - CMT Part 2 of the Phase 2 trials topline results expected in Q1 2020

PULMONARY

- Sotatercept
 - PULSAR Phase 2 trial topline results now expected in Q1 2020
 - SPECTRA exploratory study preliminary results expected in 2020



Q2 2019: Financial Results Q&A Session

Habib Dable Chief Executive Officer

Kevin McLaughlin Chief Financial Officer

John Quisel, Ph.D., J.D. Chief Business Officer

Sujay Kango Chief Commercial Officer

Todd James, IRC VP, Investor Relations and Corp. Comm.



