ESPERION 1Q20 CONFERENCE CALL

May 6, 2020



SAFE HARBOR

FORWARD-LOOKING STATEMENTS

This presentation contains forward-looking statements that are made pursuant to the safe harbor provisions of the federal securities laws, including statements regarding the clinical development and commercialization plans for bempedoic acid tablet and the bempedoic acid / ezetimibe fixed dose combination tablet, including Esperion's timing, designs, plans for announcement of results regarding its CLEAR Outcomes study and other ongoing clinical studies for bempedoic acid tablet and the bempedoic acid / ezetimibe combination fixed dose tablet, timing for the review and approval of expanded indications for their effect on cardiovascular events, and Esperion's expectations for the market for medicines to lower LDL-C, including the commercial launch and market adoption of bempedoic acid tablet and the bempedoic acid / ezetimibe fixed dose combination tablet in the United States and European Union. Any express or implied statements contained in this presentation that are not statements of historical fact may be deemed to be forward-looking statements. Forward-looking statements involve risks and uncertainties that could cause Esperion's actual results to differ significantly from those projected, including, without limitation, delays or failures in Esperion's brand commercialization plans, or approval of expanded indications, that existing cash resources may be used more quickly than anticipated, the impact of COVID-19 on our business, clinical and commercial supply activities and commercial development plans, and the risks detailed in Esperion's filings with the Securities and Exchange Commission. Any forward-looking statements contained in this presentation speak only as of the date hereof, and Esperion disclaims any obligation or undertaking to update or revise any forward-looking statements contained in this presentation, other than to the extent required by law.







We salute all healthcare workers for their incredible commitment



Our overall health and long-term conditions are more in focus than ever before







Our Medicines are Now Approved in Europe

NILEMDO™

(bempedoic acid) Tablet
Lowered LDL-C Up to 28%*

NUSTENDI™

(bempedoic acid and ezetimibe) Tablet
Lowered LDL-C by a Mean of 38%

*Compared to Placebo, in 4 Pivotal Trials Including Over 3600 Patients



Refer to our Last Public Call for U.S. NEXLETOLTM & NEXLIZETTM Information







\$900M+

Milestones Plus Royalties

Largest EU Agreement in Last Decade



\$600M+

Milestones and Development Costs Plus Royalties

Largest Japan Agreement

COMMERCIAL OVERVIEW

Mark Glickman CCO





Experienced Team Delivering Access & Affordability

Payor Coverage

50+% Commercial Coverage 20%+ Medicare Part D Coverage + Additional in 2020

Preferred brand medicines
Tier 2 and Tier 3

Healthcare providers

Simplified Prior Authorizations Minimized Process

Patients

~\$10/Rx Commercial Co-Pay \$45/Rx Medicare Part D Co-Pay

Conscientious, Virtual, Customized Launch

Accelerated Digital Asset Development

Territory Managers Completed Virtual Training Week of April 13th

Full Deploy of Digital Assets and Virtual Sales Team Week of April 20th

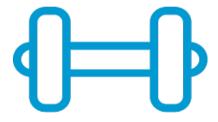
Adapted 7-day Blister Pack Sample Program to Allow for One-Touch On-Line Ordering and Direct Shipment to HCPs



Commercial Team Considerations







New "Normal"

Measured approach based on geography and health care provider specialty and circumstances

It's Early

Just beginning to scratch the surface in terms of reaching target healthcare providers

Strong Position

Prepared to assist healthcare providers and their patients in every appropriate way possible

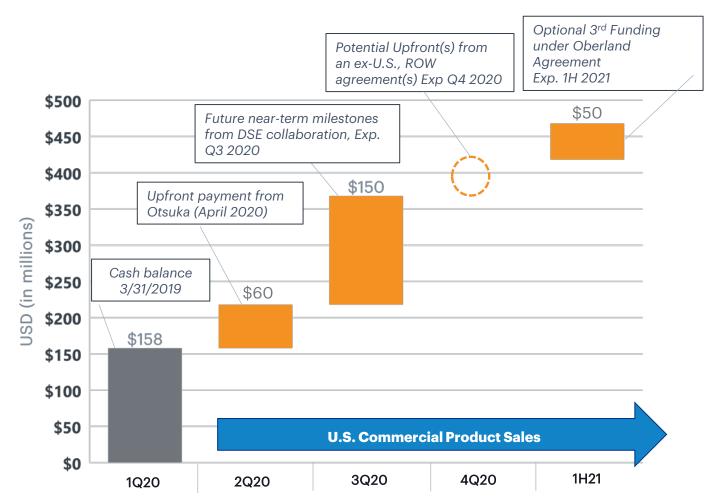
FINANCIAL OVERVIEW

Rick Bartram CFO



FINANCIAL INFORMATION: FULLY FUNDED THROUGH PROFITABILITY

Near-Term Capital Proceeds from Collaboration Agreements



Key Financial Data	
FY 2019 Revenue	\$148.4M
FY 2020 Revenue	No Guidance Provided
FY 2020 Milestone Cash Proceeds	> \$235M • Future capital available upon the completion of an ex-US ROW collaboration (4Q 2020)
FY Op Ex Guidance*	\$335 - \$355M
Anticipated Cash Balance at Dec. 31, 2020	> \$100M
Common Shares Outstanding	Basic 27.5M; Fully Diluted 32.4M



^{*}Excludes \$30M of non-cash stock based compensation expense



Thank You Questions?



Tim Mayleben
CEO & President



Mark Glickman CCO



Rick Bartram CFO