2019 Investor and Analyst Day



Safe Harbor

Forward-Looking Statements

These slides and the accompanying oral presentation contain forward-looking statements and information. The use of words such as "may," "might," "will," "should," "expect," "plan," "anticipate," "believe," "estimate," "project," "intend," "future," "potential," or "continue," and other similar expressions are intended to identify forward looking statements. For example, all statements we make regarding the regulatory approval pathway for bempedoic acid and the bempedoic acid / ezetimibe combination tablet and the therapeutic potential of, clinical development plan for, bempedoic acid and the bempedoic acid / ezetimibe combination tablet, including Esperion's timing, designs, plans and announcement of results regarding its CLEAR Outcomes study and other ongoing clinical studies for bempedoic acid and the bempedoic acid / ezetimibe combination tablet, timing for the review and approval of the NDAs and the MAAs and Esperion's expectations for the market for therapies to lower LDL-C, including the market adoption of bempedoic acid and the bempedoic acid / ezetimibe combination tablet, if approved, Esperion's cash position and financial outlook, and the expected upcoming milestones described in this press release. Any express or implied statements contained in this press release 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 studies, that positive results from a clinical study of bempedoic acid may not be sufficient for FDA or EMA approval or necessarily be predictive of the results of future or ongoing clinical studies, that notwithstanding the completion of Esperion's Phase 3 clinical development program for LDL-C lowering, the FDA or EMA may require additional development in connection with seeking regulatory approval, that DSE is able to successfully commercialize the bempedoic acid/ezetimibe combination tablet and bempedoic acid, if approved, that existing cash resources may be used more quickly than anticipated, and the risks detailed in Esperion's filings with the Securities and Exchange Commission. Esperion disclaims any obligation or undertaking to update or revise any forward-looking statements contained in this press release, other than to the extent required by law.



Investor and Analyst Day Event Program

9:00 a.m. - 9:30 a.m.

Opening Remarks – Tim Mayleben, Mark Glickman

9:30 a.m. - 9:45 a.m.

Positioning for Success – Mark Glickman

9:45 a.m. - 10:05 a.m.

Patients & Unmet Need – Renee Marotta

10:05 a.m. - 10:40 a.m.

Physician Panel, Q&A – Tim Mayleben, Dr. Mintz, Dr. Arulandu

10:40 a.m. - 11:15 a.m.

Managed Care, Q&A – Mark Glickman, Jerry Penn, Roberta Peterson

11:15 a.m. - 11:30 a.m.

U.S. Launch Update – Mark Glickman

11:30 a.m. - 11:40 a.m.

Financial Update, Q&A – Tim Mayleben, Rick Bartram

11:40 a.m. - 11:45 a.m.

Closing Remarks





Tim Mayleben

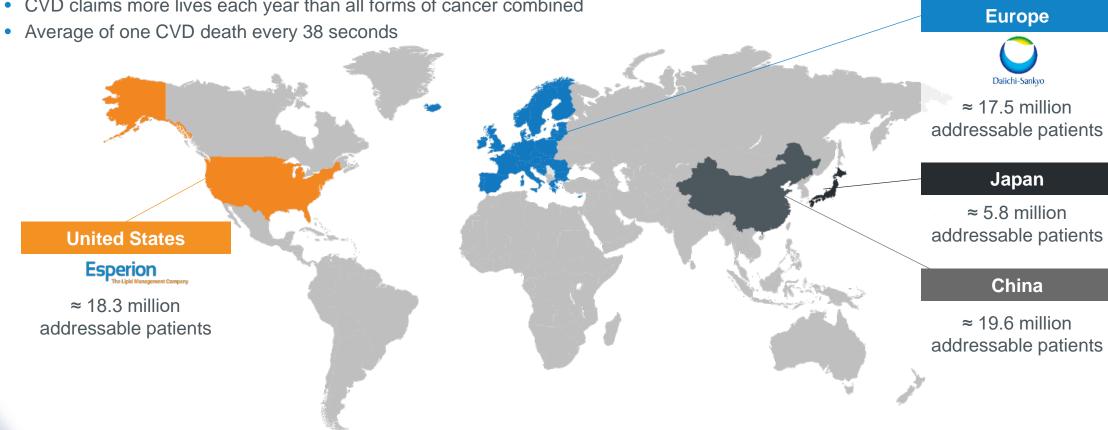
President,
Chief Executive Officer,
Board Member



Addressing a Truly Global Problem

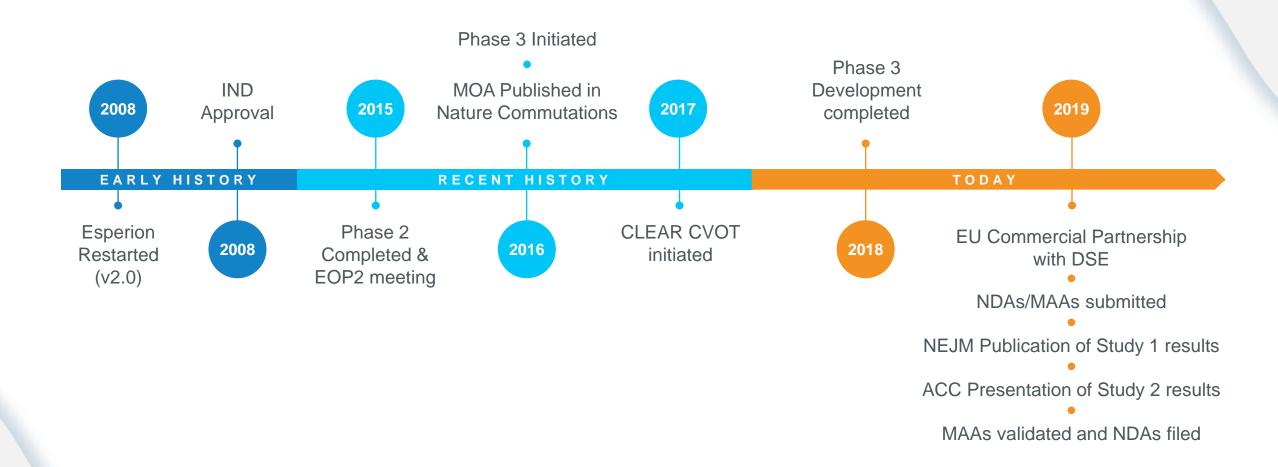
Cardiovascular Disease Remains the #1 Cause of Death Globally

- CVD accounts for ~1 in 3 deaths in the U.S. and Europe
- >800,000 U.S. CVD deaths annually
- CVD claims more lives each year than all forms of cancer combined





Esperion – Delivering History in LDL-C Lowering







Mark Glickman

Chief Commercial Officer



Leveraging Previous Success...the KOS Story



The New York Times

"Abbot Labs Buying Kos Pharmaceuticals in \$3.7 Billion Deal"

MM&M

"Kos Pharmaceuticals and Takeda Pharmaceuticals reach agreement to co-promote Niaspan and Advicor"

MarketWatch

"Abbott to pay \$3.7 billion for Kos Pharma"



"Kos Pharmaceuticals gets \$50 million boost"



"Kos Pharmaceuticals and SkyePharma Inc. Sign Exclusive License Agreement For The Marketing And Distribution of Flutiform™ In The United States; Includes Upfront Licensing Payment of \$25 Million"



Why the Market Needs the Bempedoic Acid Franchise...

of 10 adults on statin therapy reach their LDL-C targets.

Physicians believe that Bempedoic Acid (BA) and Bempedoic Acid/Ezetimibe (EZE) Combination Tablet can provide an efficacious, safe, and convenient therapy to help patients make progress towards their LDL-C goals.

Physicians recognize the strength and value of PCSK9i, but challenges remain.

Marketopportunityfor BempedoicAcid



...Where the Bempedoic Acid Franchise Could Fit...

Bempedoic Acid & Bempedoic Acid / **PCSK9i Inhibitors** Ezetimibe **Statins Combination Tablet** Standard of first-line care in LDL-C reduction Primary prevention Secondary prevention For patients who need LDL-C reductions of 50% or more Recommended by AHA/ACC for patients with high-risk, comorbid ASCVD or HeFH





Nearly 20 Million Patients Could Benefit





Commercial Opportunity& Launch Planning



The Esperion Difference

Driving Toward a Successful Launch

Focus on Physicians

- We listen to our physicians and key opinion leaders and act on all learnings
- Conduct extensive, in depth market research, not "check the box" style
- Challenge ourselves to find the truth

Understand Patients

- Partner with patients and advocacy groups to truly understand patient needs, concerns and treatment preferences / drivers
- Committed to providing effective, affordable lipid management for all patients

Managed Care & Access

- Extensive managed care & access relationships and expertise: ~ 30 Years experience each
- Productive conversations well in advance of launch to ensure the best access for all patients

We listen and learn from ALL of our constituents, because we genuinely care about making the right decisions.



Understanding the Opportunity

Market Research



Market research is critical to our successful launch effort.

- Identifies most critical patient needs
- Outlines physician perceptions and perspectives around new treatment launch that can impact adoption and use
- Drives better understanding of the current treatment paradigm and market access opportunities



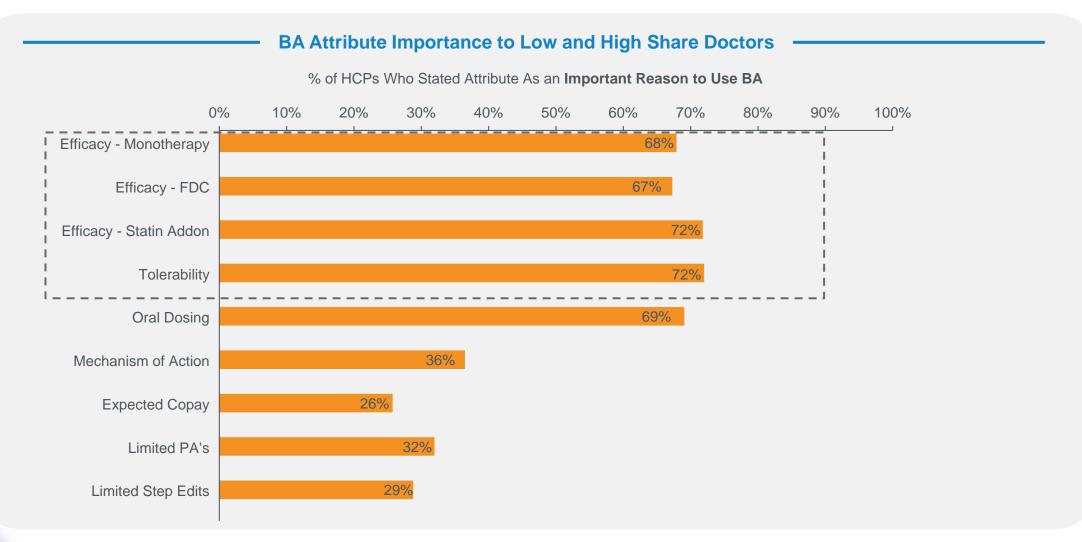
Esperion approach - focused on data-driven decisions.

- Conducted in-depth research and 1x1 interviews with 800 physicians to date
- Have amassed a robust body of research
- Invested in getting to the truth of what patients and physicians need



The Physician Perspective on BA

Efficacy & Tolerability are Key Drivers



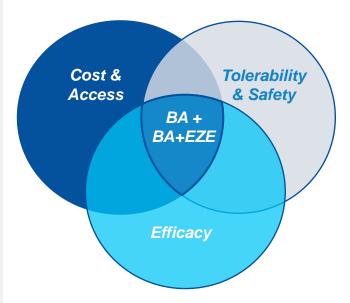


Physicians See an Important Potential Place for BA in the Treatment Paradigm

Feedback on Existing Therapies

	Existing Therapies	Positive	Negative	Neutral
-	Statins	Powerful, effective, amazing, gold standard	Flawed, muscle AEs, limiting, imperfect	What I use, therapy for dyslipidemia, affect LDL in liver
	PCSK9i	Innovative, exciting, effective	Cost prohibitive, only cards can rx with prior authorization, difficult to access	Biologic, just getting a feel for them, another way to affect LDL
	Ezetimibe	Beneficial, some benefits, inexpensive, safe	Weak, not impressive, not powerful, fallback medication	Add-on, 2 nd tier after statin failure

Potential BA and BA+EZE Fit



"I could see this as an alternative med in place of statins to try to get patients to goal. This could be a competitor with PCSK9s, especially because it's safe and non-injectable, though a lot depends on price."

—IM, Chicago

"We need another medication that's as efficacious as statins without the side effects, and I think this totally delivers the efficacy."

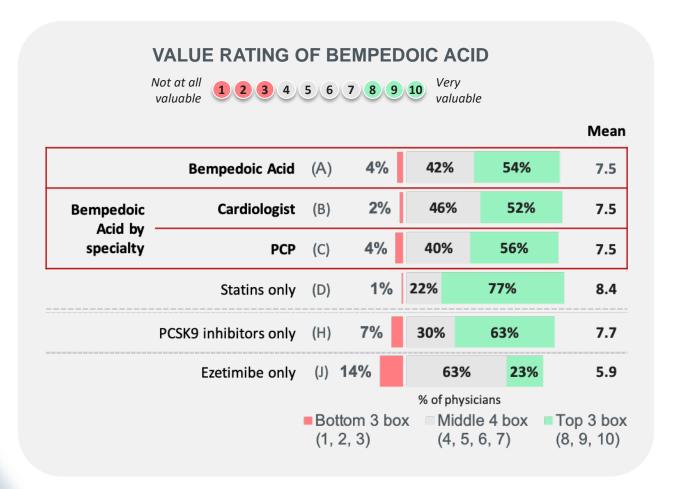
—IM, Philadelphia

"This is stronger than Zetia, but I'm not sure if it's stronger than statins. I could see putting 25% of my patients on Product X."

—Cardiologist, Philadelphia



Physicians: Bempedoic Acid is as Valuable as PCSK9 Inhibitors for Lowering Cholesterol



Nearly **70% of physicians** are likely to prescribe BA because...









Payers: The Unmet Needs, Bempedoic Acid Value and Fit Among Therapeutic Options

Top stated unmet needs in the cholesterol-lowering area are:

- Therapy for those patients who are statin-intolerant
- Efficacy/potency beyond statins
- Oral formulation paired with efficacy, cost effectiveness and outcomes data
- New mechanism of action leading to fewer or no side effects

Bempedoic Acid value to payers falls into "above average category" with 10 being "very valuable"

 Relative to other non-specialty products in large 'primary care' therapy categories, 6 out of 10 is a very good result for payer research

Value of Bempedoic Acid



Payer feedback on Bempedoic Acid fit among therapeutic options

- Most payers see BA fitting clinically as 2nd line agent after generic statins and/or EZE, and in some cases 3rd line to generic statins and/or EZE, however, most definitely before PCSK9i
- Restrictions—if any—could include a 'step to label' through a generic statin



Payers: Bempedoic Acid Has Perceived Advantages Over Existing Cholesterol Medications

Advantages over Statins

- For statin-intolerant
- Liver-enzyme elevations comparable to statins and ezetimibe
- No worsening of blood glucose (improvement with diabetics)
- Does not require titration

Advantages over Ezetimibe

- Better efficacy
- Synergy when dosed with ezetimibe

Advantages over PCSK9is

- Oral, not an injectable
- Once daily, not 1-2 times per month

"And then, ezetimibe, like I said before when I was rating it, has minimal effect. So, I would think Product X, just looking at the two different studies, has a much higher potential for reducing LDL. So, it would be a better value than that."

-Health Plan ~500K lives

"Statins, of course, because of their cost and efficacy are always going to be first line. And then, I think it would be similar to how we have the PCSK9i inhibitors, but perhaps without all of the hoops to jump through with those."

-Health Plan ~500K lives

"I think it's really hard to convince a patient to inject themselves with a medication that is for a more silent condition. It's not like a – rarely apparent to them. It's not a lifestyle medication that they're going to see significant improvement in their day-to-day function, so I think that continues to be a concern. Adherence."

-Health Plan ~800K lives



The Esperion Difference:

Driving Toward a Successful Launch



strong corporate footprint



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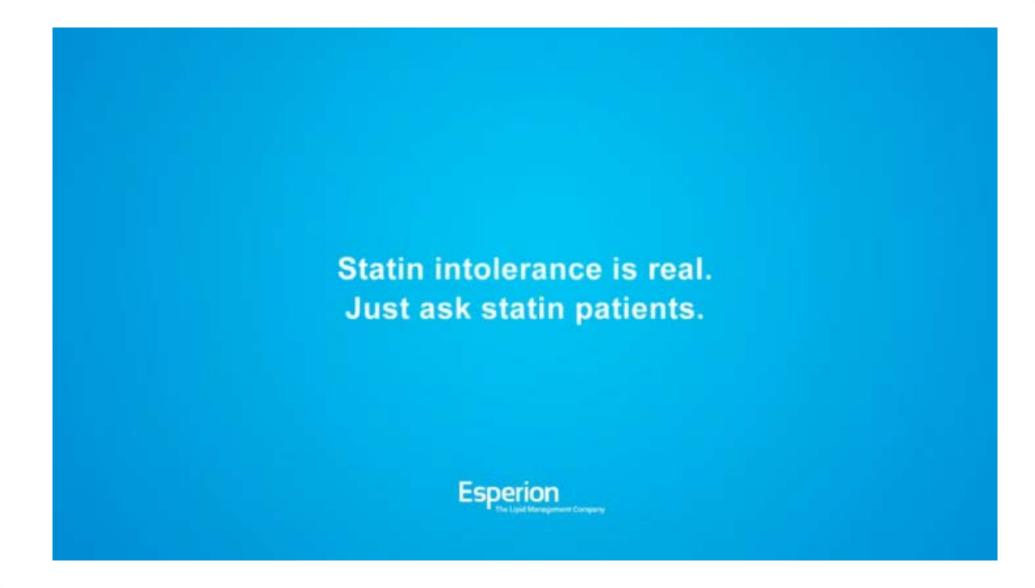


Renée Marotta Head of Marketing



Patients







Despite a wide range of cholesterol-lowering medicines, there remains a large population of patients with high LDL-C ("high cholesterol")

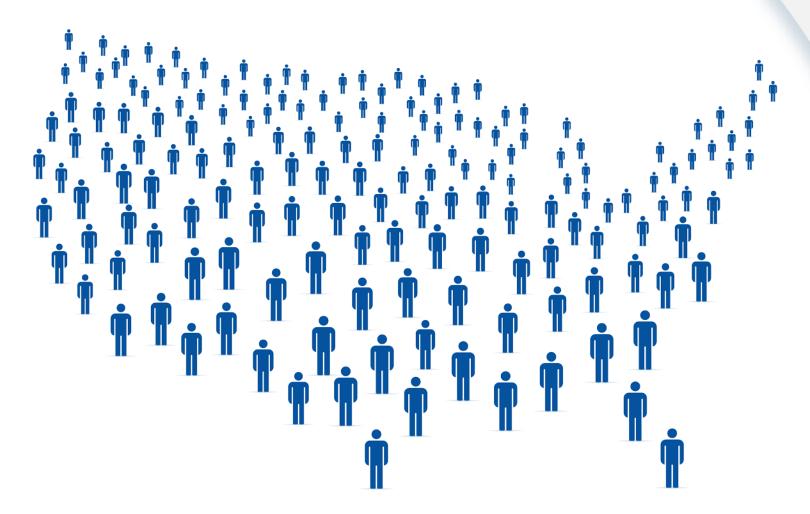


261.8 MILLION ADULTS LIVE IN THE U.S.

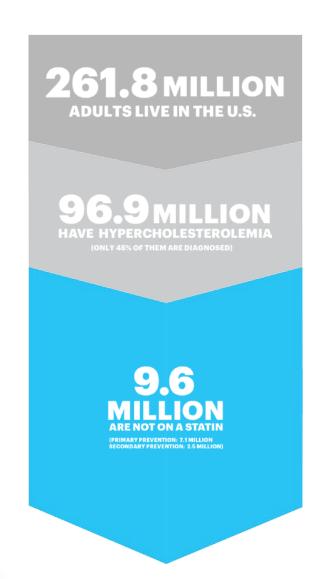


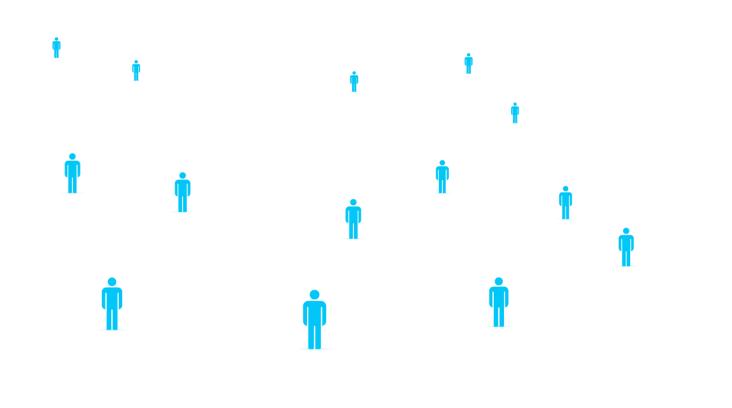




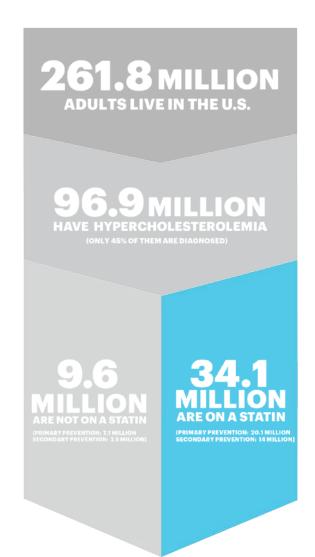


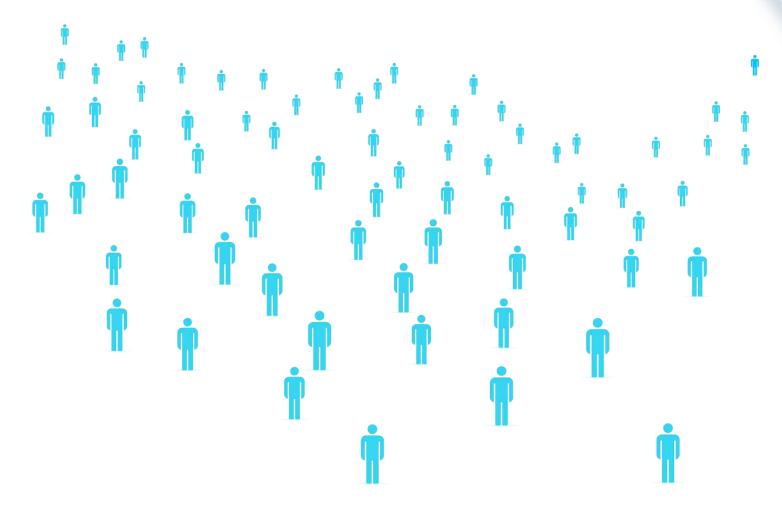




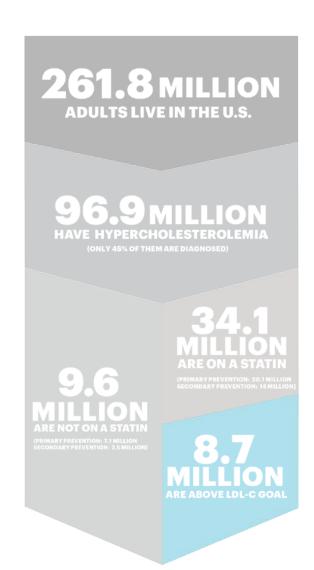


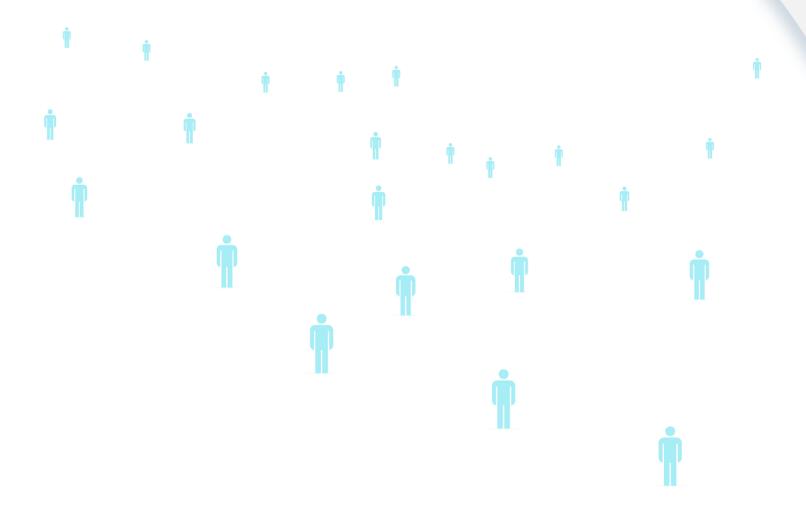




















183 MILLION ARE NOT AT GOAL THESE ARE OUR ELIGIBLE PATIENTS

Esperion is Exclusively Focused on Lipid Management for Everybody

We have been working for more than a decade with 1 mission:

To introduce innovative new therapies to help more patients reach their LDL-C targets

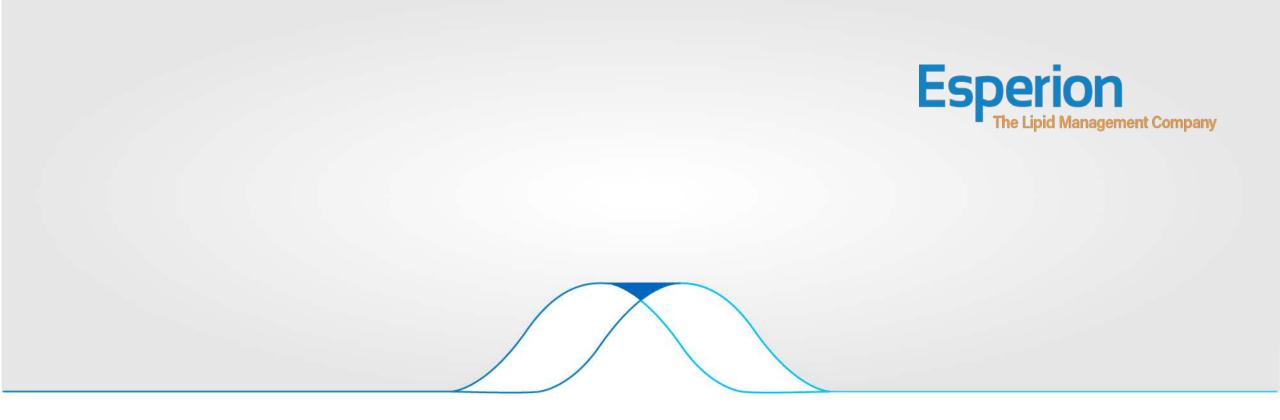
- Committed to helping patients taking statins who still need additional cholesterol lowering
- Dedicated to helping patients access our cholesterol treatments

Our focus and philosophy make Esperion unique among modern pharmaceutical companies

- Our company focus is in 1 area: LDL-C management (the main type of bad cholesterol)
- We strive to be the optimal partner with standard of care
 - We acknowledge that while patients may be getting the most out of first-line statin treatment—
 their best first-line option—it may not be enough to get them to their bad cholesterol targets





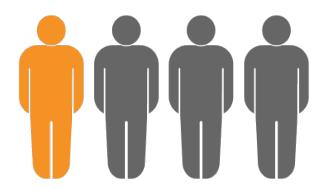


Patients Are at the Center of What We Do

18.3 MILLION PATIENTS ARE NOT AT GOAL, AND...



1 OUT OF 4 PATIENTS ON A STATIN ARE NOT AT GOAL





Patients Face Challenges with Statins

STABBING KNIVES TRIED

MEDICAL CAROUSEL MANY
INABILITY TO CONCENTRATE
SLEEP
DEPRIVATION

DEPRIVATION

LEG CRAMPS

FOGGINESS
DEPRESSION
CAN'T WAIT
FEELS LIKE A
SHARK BITE
MUSCLE
LEG CRAMPS



Why are 18.3 Million Patients Not at Goal?

Many Either Cannot Tolerate a Statin or are Not Controlled on Statin

9.6
MILLION
PATIENTS
ARE NOT ON A STATIN

STATIN AVERSE

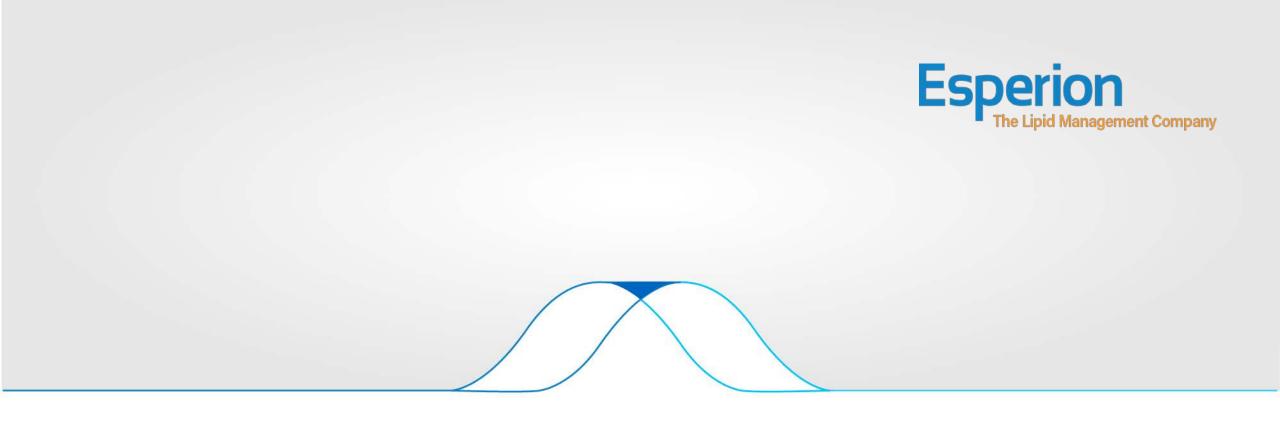
- Difficult to treat with limited options available
 - Ezetimibe is not very potent (lower efficacy)
 - PCSK9 inhibitors are expensive

8.7
MILLION
PATIENTS
ARE ON A STATIN
BUT NOT AT GOAL

ON STATIN, NOT CONTROLLED

- Not getting to goal can cause patient discouragement and lack of compliance
- Side effects can lead to lowering dosages, which limits effectiveness
- A new option is needed between statins and PCSK9 inhibitors
- Getting to goal is critical to avoid negative CV outcomes





Understanding the Healthcare Professional

Partnering with HCPs Has Allowed Us to Uncover the Truths They Face When Treating Patients



How do they view current treatments?

- Statins are the standard of care but have trade-offs
 - Well-established and proven to reduce cardiac events
 - Highly effective but face patient resistance due to lack of tolerance
 - Leave many patients above cholesterol level goals
- Healthcare providers want new options with fewer side effects
 - To prevent the need for increasing the dose of statins or simply switching to another statin and facing the same problems (cycling).
 - Second-line treatment options are well known but unpopular



Despite Statins Being an Established Treatment, There are Still Concerns

Statin concerns

Muscle cramps

25% of statin users reported muscle cramps

Statin cycling

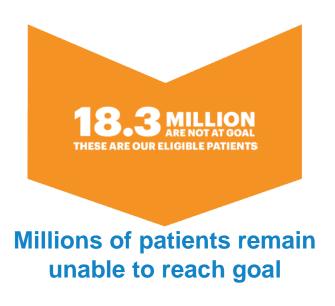
>50% of patients have been on more than 1 statin

Tolerability

28% switched statins due to side effects



The Market is Awaiting a New Treatment for Patients Not at Their Cholesterol Goal

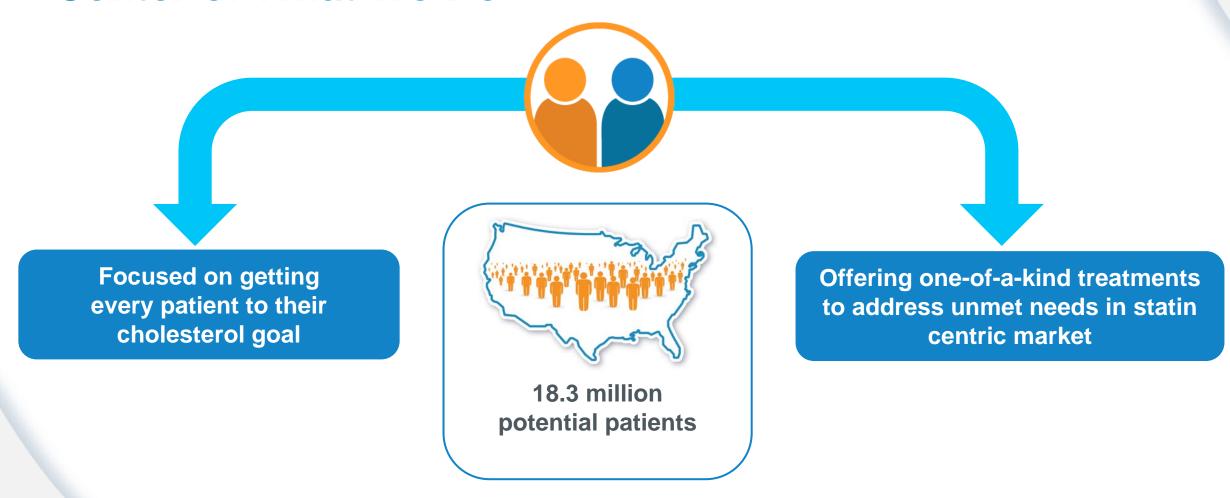




Healthcare providers and patients need additional options to get to goal



At Esperion, Helping Healthcare Providers and Patients Achieve Better Cholesterol Control is at the Center of What We Do





What is the ideal cholesterol medication? Esperion The Lipid Management Company



Physician Panel



Dr. Joseph Arulandu

- Board certifications in Internal Medicine, Clinical Lipidology, and Obesity Medicine
- Recognized as a diabetes expert by the National Committee for Quality Assurance (NCQA) and the American Diabetes Association (ADA)
- Spent the last 18 years as a clinician in private practice dealing with complex cholesterol, diabetes, hypertension, obesity and lifestyle weight management issues, and preventive cardiology
- Worked with the pharmaceutical industry and Madison Ave. on multiple scientific advisory boards, advising new drug "feasibility in real-world clinics", new drug placement and marketing for the past 13 years





Dr. Guy Mintz

- Specialist in Preventive Cardiology and Lipidology
- National leader in the field with more than 25 years of clinical experience
- On the board of directors of the National Lipid Association, and national chairman of the membership and early career development committee
- Fellow of the American College of Cardiology, American College of Physicians, National Lipid Association, and American Society of Preventive Cardiology
- Principal Investigator in The Clear Outcomes Trial with Bempedoic Acid and other lipid lowering trials





Two Non-Statin Oral Tablets That Lower LDL-C and Reduce hsCRP

Complement to Standard of Care LDL-C Lowering Drugs

Bempedoic Acid

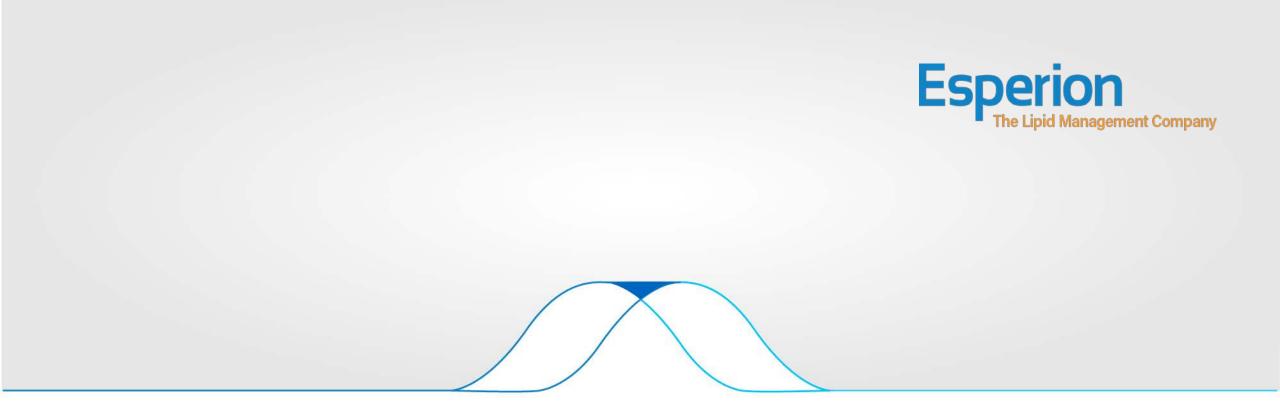
Bempedoic Acid / Ezetimibe Combination Tablet

Shared Benefits:

- Oral, once-daily, convenient, cost-effective therapies
- Did not lead to higher overall adverse events compared with placebo
 - HbA1c lowering and lower rate of new onset/worsening diabetes
- Consistent and complementary LDL-C lowering plus differentiated hsCRP reduction
- 18% LDL-C lowering on maximally tolerated statins, including high-intensity statins
- 28% LDL-C lowering on no background statin
- 19-40% hsCRP reduction; a key marker of inflammation

- Efficacy comparable to injectable PCSK9i monotherapy (~50% LDL-C lowering) – plus differentiated hsCRP reduction
- 29% LDL-C lowering on maximally tolerated statins
- 44% LDL-C lowering on no background statins
- 34% hsCRP reduction; a key marker of inflammation





Question & Answer

Managed Care





Jerry Penn

Head of Managed Care

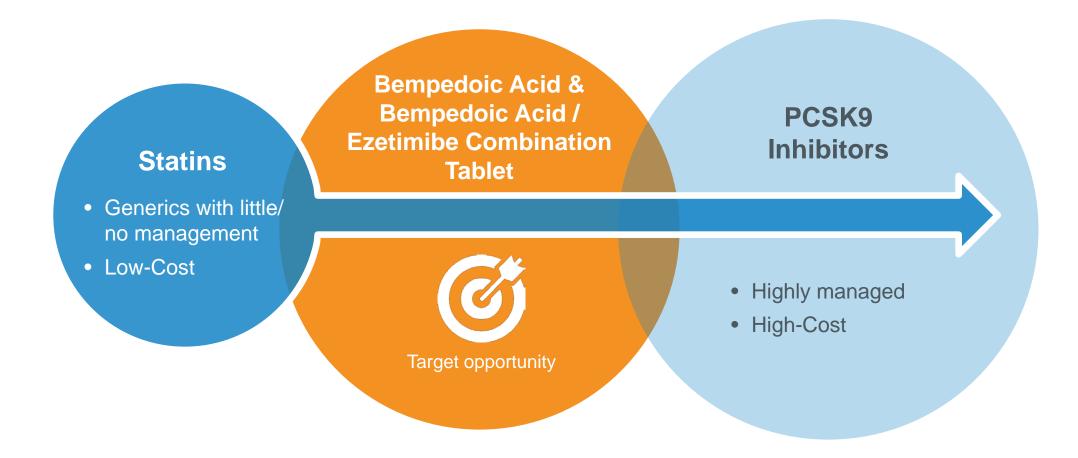


All Roads Lead to Launch – Market Access & Trade Excellence





Bempedoic Acid Payer Opportunity





Foundations for Success

Completed Key Tactics Required for Successful Payer Review and Negotiation



Corporate Presentation

Introductory presentation for payer audiences introducing Esperion, our corporate mission and setting the foundation for pipeline discussions



AMCP Dossier

AMCP formatted clinical and economic evidence in support of formulary consideration.
Used in responding to unsolicited request to support reimbursement and/or formulary placement of a new product



Budget Impact Model

An interactive, user-friendly model for use by/with Payers and PBMs highlighting the budget impact of bempedoic acid in patients with hypercholesterolemia not adequately treated with existing lipid lowering therapies

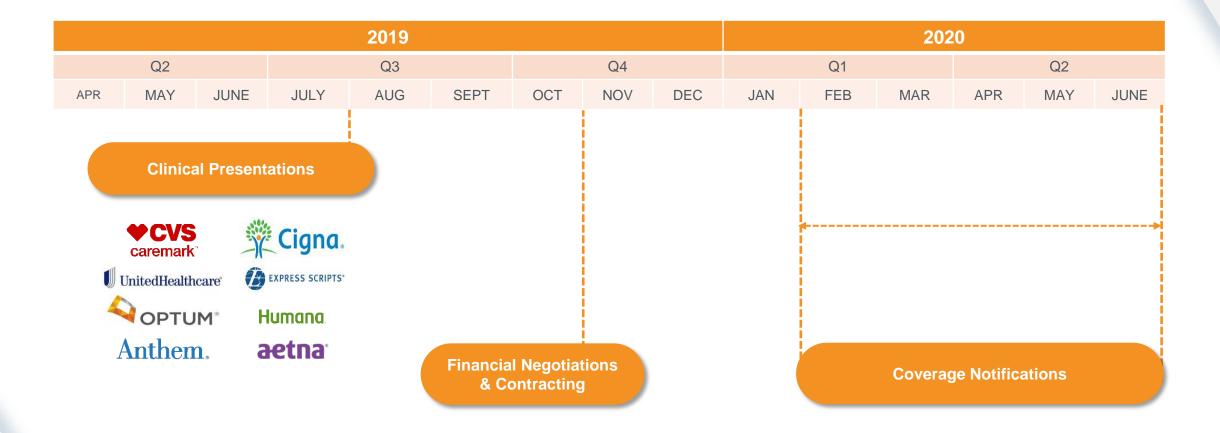


Compendia

NDC compendia submission packet that includes essential information for the update of compendia database to address new therapeutic class information, clinical data, pricing and optimized prescribing access to support immediate product access



Payer Negotiations Overview

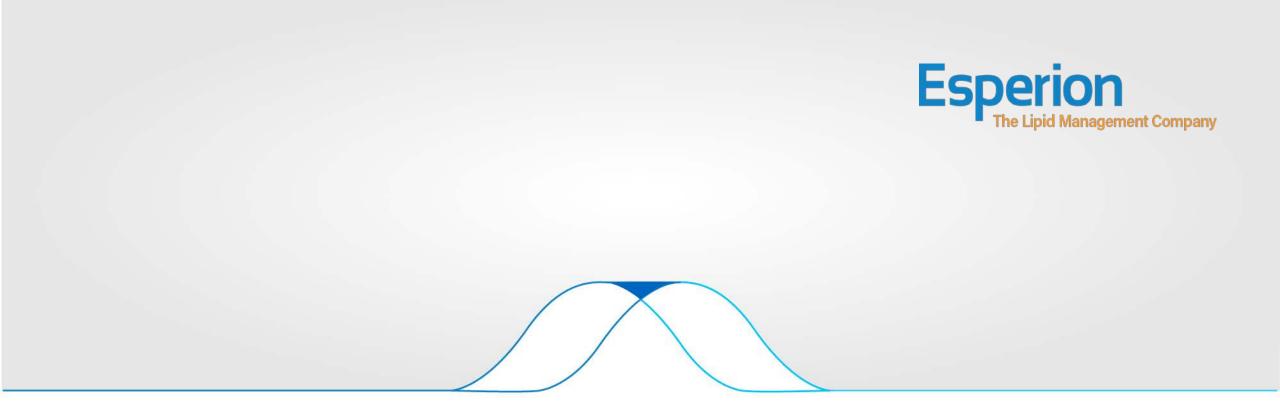




All Roads Lead to Launch – Market Access & Trade Excellence







Question & Answer

BA Sales Force: The Esperion Strategy



Forecasting Process

High-Level Flow of Work Steps

Secondary
Research (data
sourced from
literature reviews)

Population prevalence rates for hyper-cholesterolemia

Primary Research (data sourced from direct interviews)

Percentage of patients
who are eligible to
receive LDL therapy
based on BA and
BA+EZE product
profile, and who will
ultimately receive BA
and BA+EZE (share of
market)

Pricing Assumptions

Gross price,
discounts, net price,
gross-to-net changes
over time, and
therapy compliance
rate all have an
impact on pill volumes
and revenues derived
from forecast

Product Analog Analysis

Launch experience from similar products/ situations can inform the uptake curve for BA and BA+EZE, including time to peak share and the shape of the curve Sales Force
Promotional Effort
Analysis

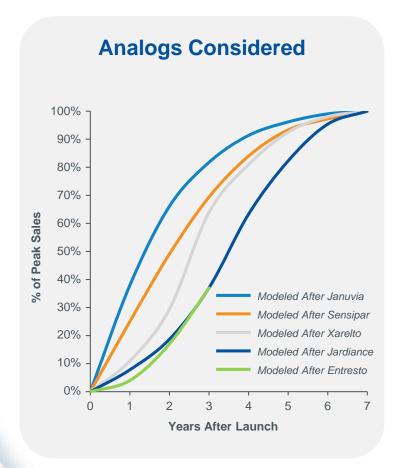
Analysis of market size (prescriptions and writers), profitability per Rx, and analog launch sales force sizes can inform the size of the sales team needed to promote BA and BA+EZE

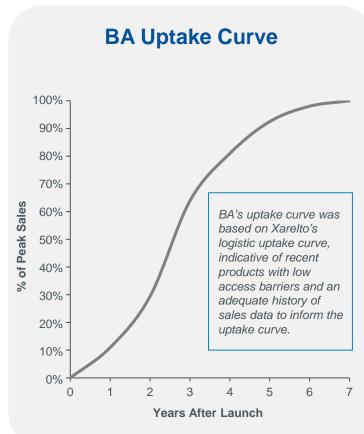
Each of the work steps above combine to create a multi-year patient, prescription, and revenue forecast model which will be reviewed in detail in the following slides



Extensive Launch Analog Research & Modeling

Our Insights into BA Potential & Expected Penetration





Product Profile



First FXa inhibitor approved by FDA to treat embolism. thrombosis, and AFib

- · Molecule: rivaroxaban
- Class: factor Xa inhibitor
- · Treats: embolism, thrombosis (DVT), and atrial fibrillation (AFib)
- · Delivery: once-daily, oral tablet

- Developer: Janssen
- Launched: July 2011
- LOE Date: August 2024
- Notes: Xarelto has a large DTC presence













Chronic

Competitor

MOA

First in Class

Daily Oral

Recently Peaked



Xarelto Case Study

Launched with 300 Reps, Grew to 1,000

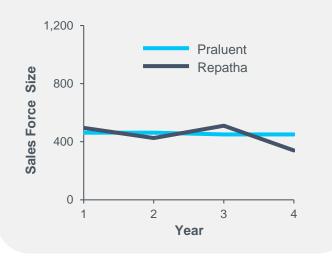
Previous Market Blockbuster SF Sizes

Cholesterol-lowering blockbuster drugs in the early 2000's launched with large salesforce sizes of 1.3K to 2.2K sales reps

	At Launch	At Peak			
Lipitor	~2,200	3,000			
Zetia	~1,500	> 2,000			
Crestor	~1,300	~2,500			

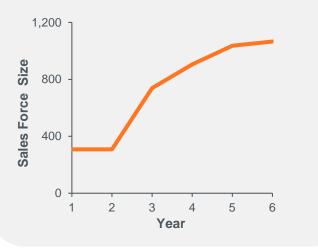
PCSK9 SF Sizes

PCSK9s were launched with ~500 rep salesforces and have remained relatively steady (arguably due in part to aggressive payor controls)



Xarelto SF Size

Xarelto, used as the uptake analog for the BA forecast, launched in 2011 with ~300 reps and increased over time to eventually arrive at optimal size



Based on Xarelto's use as the uptake analog in the forecast, it can also be an effective guide for BA's recommended sales force sizes over time

Salesforce size data is sourced from PharmaForce. SF sizes considered to be 75% accurate, and should only be treated as approximations Lipitor (PharmaBusiness), Zetia (ContractPharma), and Crestor (PharmaLetter) approximate launch sizes acquired from pharma-focused news journals Source: ZS Associates Market Sizing and Forecasting Project, August 2018. N=350 Physicians



Insights & Understanding the U.S. LDL-C Rx Market

How Physician and Rx Counts Impact Field Force Coverage

- ~660,000 health care practitioners write 227 million annual Rx in the US statin/ezetimibe/combination/ PCSK9/bile acid sequestrant market
- Assuming a typical rep can manage 120 writer 'targets,'
 46 reps can cover all decile 10 writers, and it would take
 5,495 reps to cover all targets.
- The recommended sales force size at launch is ~300 reps, which means that ESPR will cover deciles 10, 9, 8, and 7 prescribers representing 40% of market Rx (the 'market opportunity') at launch. These 300 reps will call on ~35,500 prescriber targets.

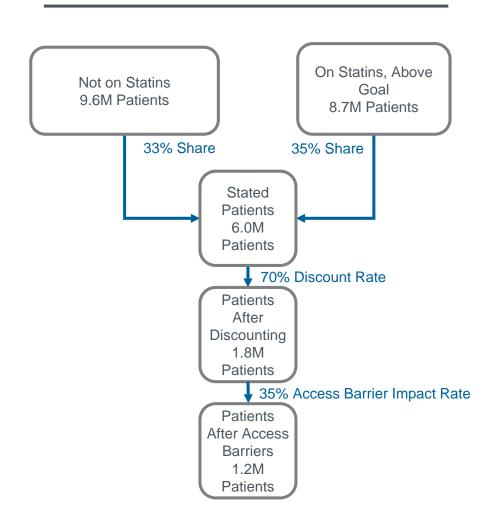
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HCP Writer Count		5,474	8,076	9,976	11,929	14,207	17,094	21,208	28,062	43,962	499,449	
C	Cumulative Writer Count	5,474	13,550	23,526	35,455	49,662	66,756	87,964	116,026	159,988	659,437	
	PCP	4,436	6,655	8,041	9,324	10,538	11,873	13,618	15,924	20,431	136,029	
Specialty Group	Cardiologist	750										
	NP/PA	128	Cumulative Rep Count				4	46		196		295
	Endocrinologist	77	Gairie	ilativo	Top of		46		'		200	
	Other	83	% of Rx Covered					10%				40%
	Annual Market TRx	22,700,000					10			30'	%	
	TRx Per Writer 4,147											
	Sales Rep Count Assum	ing 120 Targe	ts Per Rep									
	Rep Count	46	46 67 83 99 118 142 177 234		234	366 4,16						
	Cumulative Rep Count				295	414	556	733	967	1,333	5,495	
	% of Rx Covered				40.0%	50.0%	60.0%	70.0%	80.0%	90.0%	100.0%	
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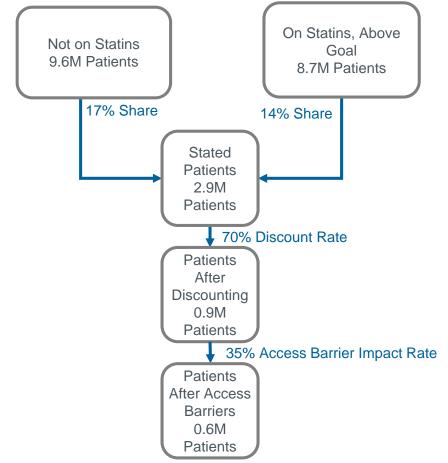


Bempedoic Acid is Forecasted to Have 1.8M Patients on Therapy at Peak

Bempedoic Acid

Bempedoic Acid / Ezetimibe Combination









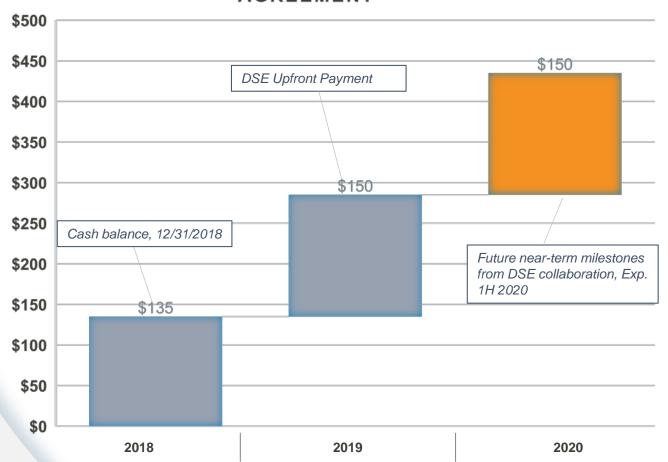
Rick Bartram

Chief Financial
Officer



Esperion is Funded to Deliver Value From the Bempedoic Acid Franchise

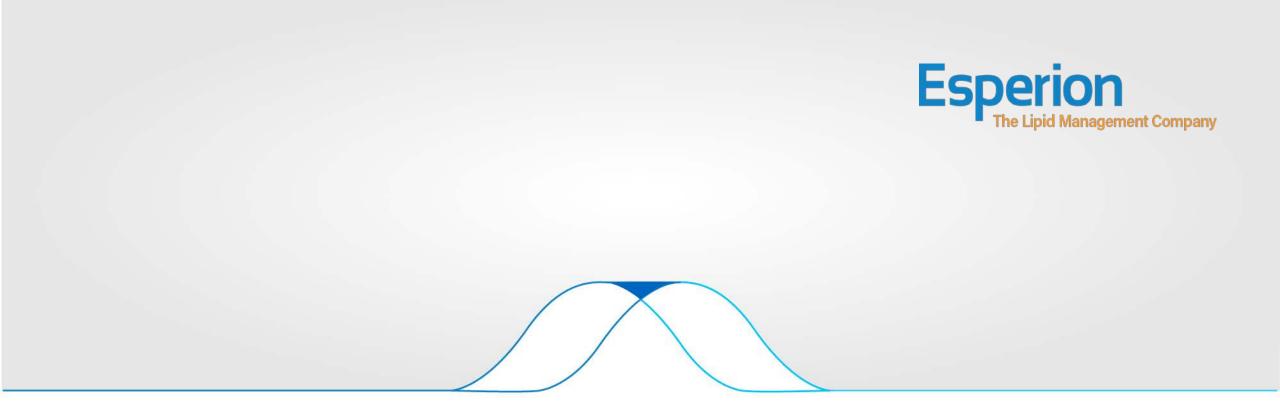
CAPITAL PROCEEDS FROM COLLABORATION AGREEMENT



As of March 31, 2019:

- Cash equivalents \$229.7 million
 - Future near-term payment from DSE collaboration agreement, 1H 2020:
 - \$150 million
- 2019 Net Cash Used in Operations:
 - \$25 \$35 million
- Future capital available upon the completion of both
 - U.S. Net Sales Royalty Deal
 - Ex-US ROW collaboration
- Significant cash balance to fund the US commercial launch in Q1 2020





Question & Answer

Closing Remarks



Thank You for Your Support

