Medicure Inc. Investor Presentation

August 25, 2022





Forward Looking Statement

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Medicure Inc. undertakes no obligation to update publicly or otherwise revise any forward-looking information as a result of new information, future results or other such factors which affect this information, except as required by law.



Our Vision

We want to be a leading provider of pharmaceutical and healthcare products to U.S. customers

We have a vertically integrated team, with a pipeline and acquisitions in place to build growth





What Sets Us Apart

Our approach to engagement and service with healthcare professionals and patients

Focusing on value for hospitals, prescribers, and patients



Why Invest in Medicure Inc.

- Profitable and growing profit
- Steady sales of Aggrastat
- Zypitamag sales growing
- Marley Drug E-Commerce pharmacy
- MC-1 pediatric voucher upon completion of phase 3 trial of 10 PNPO patients
- No Debt







Medicure's Acquisition of Marley Drug Pharmacy:

The Industry Game-Changer

Slides 25-28







Our Products and Pipeline

	PRODUCT	CATEGORY	DEVELOPMENT REGULATORY APPROVED / CLINICAL APPROVAL / MARKETED
AGGR STAT® (tirofiban hydrochloride) Injection	AGGRASTAT®	Hospital IV Injectable	Branded antiplatelet, grew from 2% to 65% market share
Zypitamag (pitavastatin) tablets	ZYPITAMAG®	Prescription Consumer	Branded statin with large market potential
	Sodium Nitroprusside	Hospital IV Injectable	Generic mainstay of hospital formularies
	ANDA	Hospital IV Injectable	Cardiovascular Generic with few competitors
	NDA	Prescription Consumer	MC-1 for Rare orphan disease
marleydrug	Marley Drug®	Retail Pharm	acy for marketing of generic and branded medications





Strategic Approach

1 Maintain positive earnings and cash flow through existing products and innovation in direct-to-consumer sales

3 Acquisitions of synergistic companies or cardio products for US market

PROFITABILIY

DEVELOPMENT

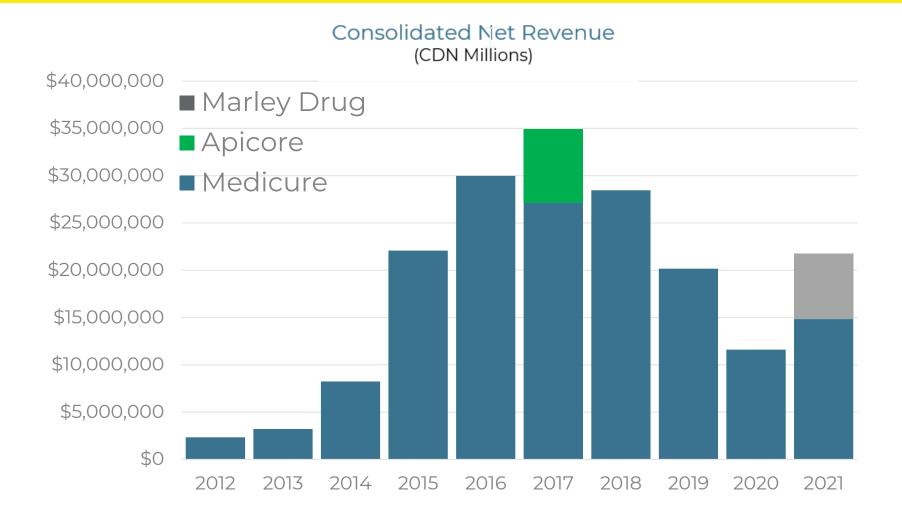
ACQUISITION

2 Expand product offering through development of high margin IV and solid dose products





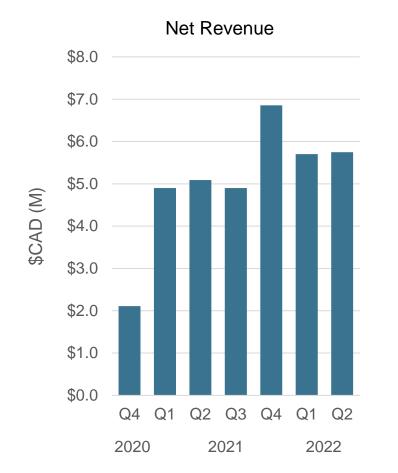
Net Revenue Growth

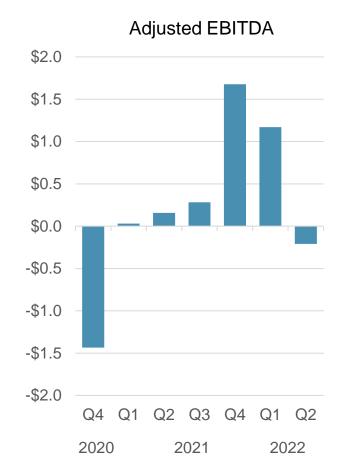


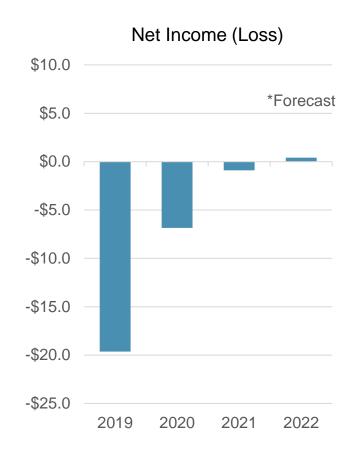




Key Financial Highlights









Key Financial Highlights

Capital Structure as of August 24, 2022

•	Issued Shares	10,251,313
•	Fully Diluted Total	10,977,313
•	Share Price	C\$1.10

Financial Highlights

- 2022 Q2 Net Revenue
- 2022 Q2 R&D Expenses
- 2022 Q2 Adj. EBITDA
- 2021 Net Revenue
- 2021 Adj. EBITDA
- Cash

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\$5.7M vs \$5.7M in Q1 \$1.3M vs \$0.3M in Q1 (\$0.2M) \$21.7M vs \$11.6M in 2020 \$2.6M vs (\$3.9M) in 2020 \$4.8M – No Debt

• Completed substantial issuer bid in 2019 returning \$26M in cash to investors









Medicure Leadership Team



Albert D. Friesen Chief Executive Officer & Chairman of the Board

- Founded Medicure in 1997
- Created and developed multiple companies, including ABI Biotechnology (Apotex Fermentation), The Winnipeg Rh Institute, DiaMedica and Genesys Venture Inc.



Neil Owens President & Chief Operating Officer

- Joined Medicure in 2014 in Medical Affairs, named as President & COO in 2019
- Responsible for the execution of strategic plans and oversight of operations



Reuben Saba Vice President, Scientific and Medical Affairs

- Joined Medicure in 2014
 in Medical Affairs
- Responsible for the development of new business opportunities for Medicure and management of Medical Affairs

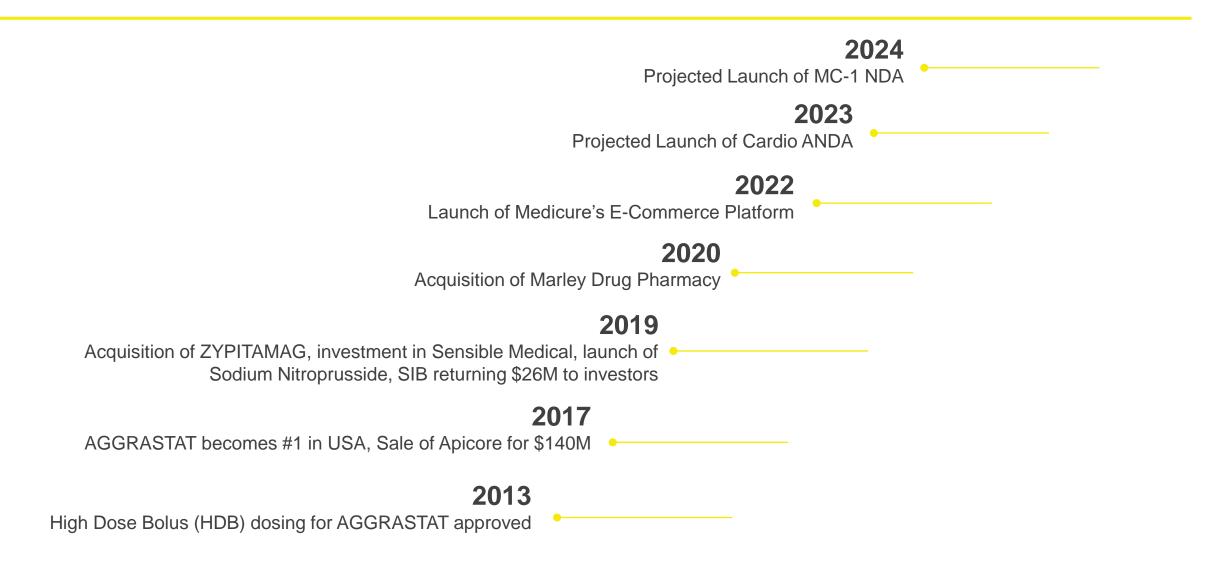


Haaris Uddin Chief Financial Officer

- Joined Medicure in 2022
 as CFO
- Responsible for financial reporting activities and oversight of finance systems and strategy for Medicure



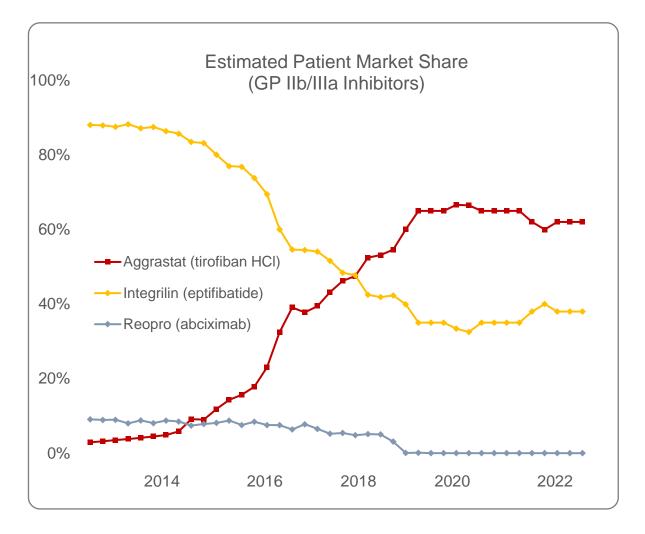
Medicure's History and Trajectory







AGGRASTAT[®] (tirofiban hydrochloride) Profile



AGGR/STAT® (tirofiban hydrochloride) Injection



Background:

- Hospital product (GP IIb/IIIa inhibitor) for acute cardiovascular care
- US rights acquired in 2006
- High Dose Bolus (HDB) dosing approved in 2013
- New 15 mL bolus vial format approved in 2016
- Consistently used in more than 1,200 US hospitals

The Story:

- Increase in market share came as a result of overcoming significant skepticism of the efficacy of the product
- Explained correct HDB dosing and value proposition to interventional cardiologist and pharmacy managers
- Developed partnerships with Key Opinion Leaders who helped drive change





AGGRASTAT Sales Update



Aggrastat Net Revenue By Quarter Aggrastat Units Per Quarter Analysis (\$ CAD Millions) Unit Demand in Q2 2022: ٠ 35,000 \$3.5 1% higher compared to Q1 2022 • 30,000 4% lower compared to Q2 2021 • \$3.0 25,000 \$2.5 Net Revenue in Q2 2022: • 20,000 3% higher compared to Q1 2022 \$2.0 . 3% higher compared to Q2 2021 . 15,000 \$1.5 10,000 \$1.0 Overall there has been consistent demand ٠ 5,000 \$0.5 \$0.0 0 Q2 Q3 Q4 Q1 Q1 Q2 Q1 Q2 Q3 Q4 Q1 Q2 2021 2022 2021 2022



SAVI-PCI Study Met Its Primary Endpoint





Randomized clinical study evaluating AGGRASTAT (short and long infusions) compared to INTEGRILIN[®] (eptifibatide)

Variable, n (%)	AGGRASTAT® Short infusion n=209	AGGRASTAT® Long infusion n=124	Integrilin [®] Long infusion n=202
Death	0 (0.0%)	0 (0.0%)	1 (0.5%)
uTVR	1 (0.5%)	1 (0.8%)	0 (0.0%)
REPLACE-2 major bleeding	0 (0.0%)	4 (3.2%)	1 (0.5%)
Periprocedural myonecrosis*	69 (34.2%)	45 (36.6%)	60 (30.9%)
Primary outcome*	69 (34.2%)	48 (39.0%)	60 (30.9%)

These results fill a gap in clinical evidence and provide an exciting development to bolster the contemporary use of AGGRASTAT

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Please refer to IMPORTANT SAFETY INFORMATION on slide 34



Strategic Approach for AGGRASTAT







ZYPITAMAG[®] (pitavastatin) Profile





Key Information

A New Type of Statin:

- Lowers cholesterol, with some benefits over other statins
- Metabolized differently from most other statins
- Well tolerated with low rates of muscle pain



Please refer to IMPORTANT SAFETY INFORMATION on slide 35



Clinical Differentiation

Zypitamag is different from most other statins (minimally processed by CYP enzymes)

Reduced risk of certain drug-drug interactions and drug-food interactions

Well tolerated: low overall rates of muscle pain side effect

Similar efficacy to statins with majority (69%) of market share: atorvastatin and simvastatin

Superior efficacy to pravastatin (10% of statin market)

Target Patient Groups

Of 17 million Americans taking a statin, 5-10% have side effects

Direct competitor Livalo currently has 0.37% of the statin market

Three main target patient groups are:

- Difficulty tolerating their current statin
- Currently taking Livalo
- Taking multiple medications that can interact with their statin

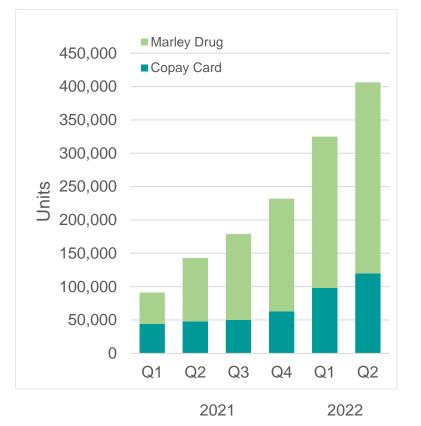


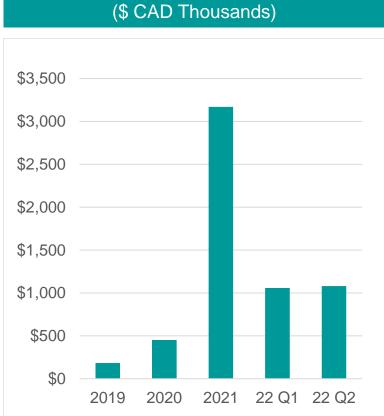




ZYPITAMAG Sales Update

Zypitamag Units Dispensed per Quarter*





Zypitamag Net Revenue per Year

Analysis

- Units dispensed via Marley Drug and Copay Cards in Q2 2022:
 - 25% higher compared to Q1 2022
 - 2.8x higher compared to Q2 2021
- Net Revenue in Q2 2022:
 - Consistent with Q1 2022
- Number of new patients and new prescribers continues to grow
- Net Revenue in Q2 was impacted by returns through traditional wholesaler sales channel



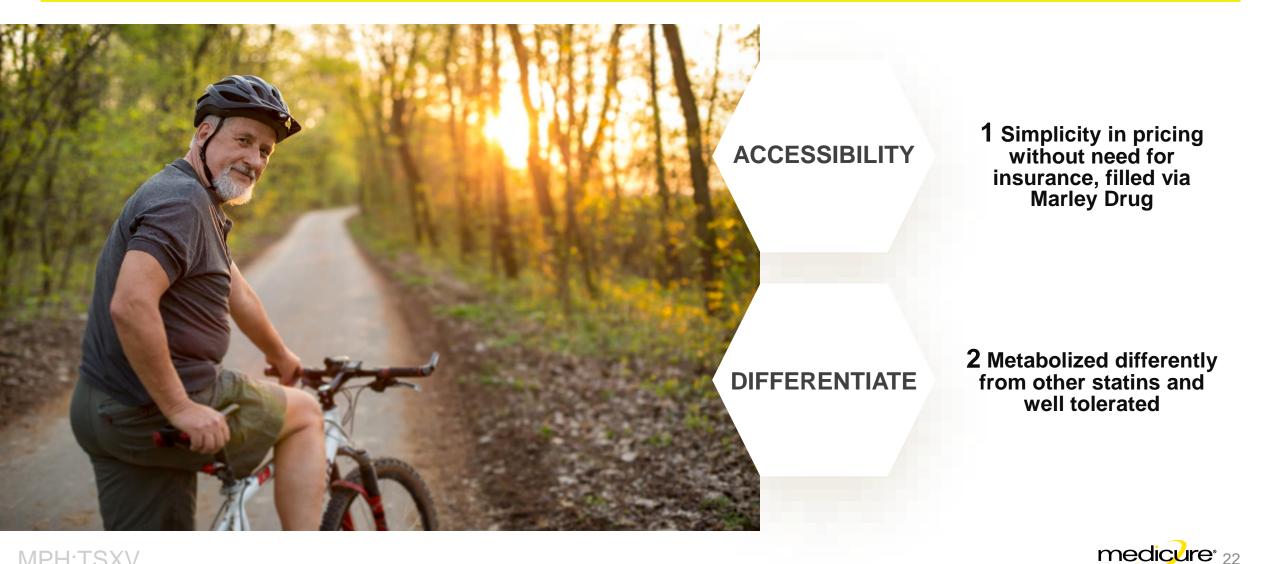
*Not captured here are sales to government covered recipients (e.g. VA and Medicare Part D)

*Captures all lines of business.



Marketing Focus





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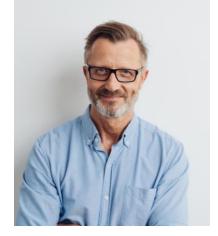
Please refer to IMPORTANT SAFETY INFORMATION on slide 35

Customer Quotes





"ZYPITAMAG just works for me. And, it's affordable"



"The best lipid levels I have had in treating cholesterol... I have not had any side effects"



"I personally tried multiple statins and had intolerable side effects. Because ZYPITAMAG is metabolized differently, it's a statin I can take"



"With ZYPITAMAG my cholesterol levels are down and I haven't experienced any leg pain"



These are stock photos representing actual testimonials. Personal experiences with ZYPITAMAG may differ. ZYPITAMAG like all statins, can cause side effects, see **IMPORTANT SAFETY INFORMATION** on Slide 35



Strategic Approach









Medicure Subsidiary - Marley Drug Pharmacy





- A US pharmacy licensed to ship medications to all 50 states, Washington D.C. and Puerto Rico
- Net Revenue in Q2 2022 of \$1.78M, compared to \$1.86M in Q1 2022
- Thousands of existing customers and proven success in marketing based on pricing of generic drugs & focus on cash price without use of insurance
- Opportunity for direct-to-consumer marketing, distribution and improved profit margin for ZYPITAMAG



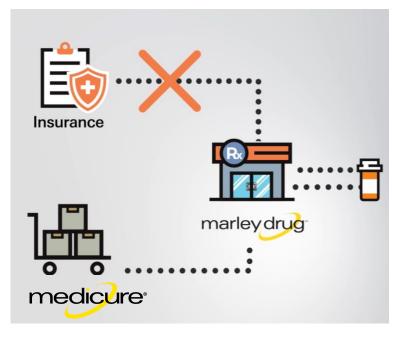
marley drug[®]

What this provides to Medicure

- 1. A more efficient distribution and marketing vehicle for ZYPITAMAG
 - a Branded drug
- 2. A marketing vehicle for other Branded drugs challenged to get insurance coverage
- 3. Saves on high fees to wholesalers and pharmacy benefit managers

What this provides Consumers

- Low cost generic medications and direct access to Medicure's branded consumer products
- Home delivery, with no added pharmacy fees targeting cash paying customers, which is a growing market







Factors Driving Americans to Pay Cash for Pharmaceuticals





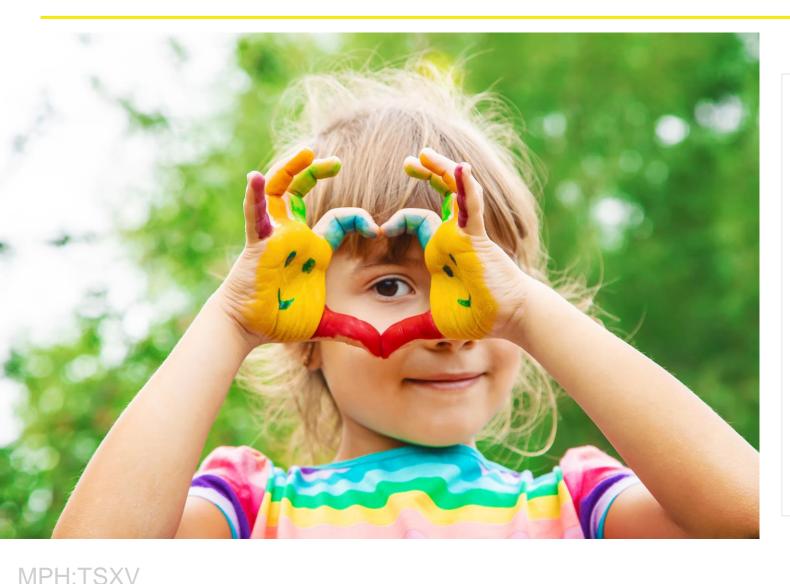
E-Commerce Platform and Go Forward Plans







Development of MC-1 for Rare Pediatric Disease



- PNPO deficiency is a rare pediatric disease, which leads to seizures and is fatal if untreated
- The disease leads to the inability to produce a critical cofactor required for normal development
- Currently there is orphan drug status, with no approved treatment
- Medicure is seeking NDA approval for treatment of PNPO using its pharmaceutical product MC-1
- Approval requires a Phase 3 study, which is planned to start in late 2022



New Products - Our Target Product Profile

In-Hospital	Build on the experience, relationships and success of AGGRASTAT	Products that can be licensed or acquired with low or no initial investment, and if licensed can be acquired with set milestones	Growth Opportunity
Primary Care	Build on the experience,	Products that can be marketed	Branded Products
	relationships and success of	and accessed directly to	for Direct-to-
	ZYPITAMAG	consumers through Marley Drug	Consumer Access



Previous Investments and Returns to Shareholders

APICORE (2014-2017)	SENSIBLE MEDICAL (2019)	SIB (2019)
 July 2014 – Acquired 5% interest in Apicore with a 3-year option to purchase the remaining shares 	USD \$10M investment in Sensible Medical, and still hold the investment	 Completed substantial issuer bid (SIB) in 2019 for \$26M in cash
 December 2016 – Increased ownership to 60% with CDN \$60 million loan 	 Markets ReDS device, used to measure lung fluid level in patients with Heart Failure 	 Purchased and cancelled 4 million shares at a set price of \$6.50
 July 2017 – Increased ownership to 92% with Apicore funds 	Medicure held U.S. marketing rights from March 2019 to July 2020	 Returned cash to investors, to reflect Medicure's intent to return value to shareholders
 November 2017 – Sold Apicore business for in excess of CDN \$140 million 	 Rights returned to Sensible because of low margin and long sales cycle, investment remains 	

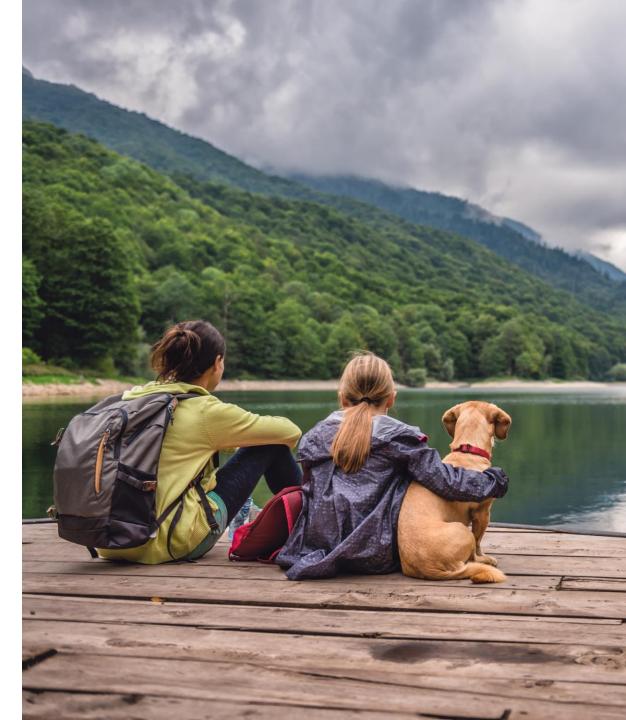


5 Key Takeaways About Medicure

- 1. Medicure's focus is on pharmaceuticals and healthcare products for the US market
- 2. Proven success with AGGRASTAT, building sales of ZYPITAMAG
- 3. Focus on direct consumer sales for growth and profitability through Marley Drug subsidiary
- 4. Pipeline of products including MC-1 for Rare Disease

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5. We believe in innovation and returning profits to shareholders



Further Information

Visit our Websites

Medicure.com AggrastatHDB.com Zypitamag.com Marleydrug.com

Investor Relations

ir@medicure.com 1.888.435.2220 (Ext. 228)

Follow us

Linkedin.com/company/medicure Twitter.com/MedicureInc Instagram.com/medicureinc

Contact a Product Specialist

1.800.509.0544





Indication: Aggrastat is indicated to reduce the rate of thrombotic cardiovascular events (combined endpoint of death, myocardial infarction, or refractory ischemia/repeat cardiac procedure) in patients with non-ST elevation acute coronary syndrome (NSTE-ACS).

Contraindications: Known hypersensitivity to any component of Aggrastat; history of thrombocytopenia with prior exposure to Aggrastat; active internal bleeding, or history of bleeding diathesis, major surgical procedure or severe physical trauma within the previous month.

Warnings and Precautions: Aggrastat can cause serious bleeding. If bleeding cannot be controlled discontinue Aggrastat; thrombocytopenia: discontinue Aggrastat and heparin.

Adverse Reactions: Bleeding is the most commonly reported adverse reaction.

For additional information, refer to the <u>full Prescribing Information</u>. You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.FDA.gov/medwatch or call 1-800-FDA-1088.



Important Safety Information for ZYPITAMAG (pitavastatin) tablets

INDICATIONS & USAGE

ZYPITAMAG is indicated as an adjunctive therapy to diet in adult patients with primary hyperlipidemia or mixed dyslipidemia to reduce elevated total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), apolipoprotein B (Apo B), triglycerides (TG), and to increase high-density lipoprotein cholesterol (HDL-C). Limitations of Use: The effect of ZYPITAMAG on cardiovascular morbidity and mortality has not been determined.

CONTRAINDICATIONS: ZYPITAMAG is contraindicated in patients with a known hypersensitivity to product components, in patients with active liver disease (which may include unexplained persistent elevations in hepatic transaminase levels), in women who are pregnant or may become pregnant, in nursing mothers, or in co-administration with cyclosporine.

WARNINGS & PRECAUTIONS

- Myopathy and Rhabdomyolysis: Risk factors include age 65 and greater, renal impairment, inadequately treated hypothyroidism, concomitant use of certain drugs, and higher doses of ZYPITAMAG. ZYPITAMAG is contraindicated in patients taking cyclosporine and not recommended in patients taking gemfibrozil. The following drugs when used concomitantly with ZYPITAMAG may also increase the risk of myopathy and rhabdomyolysis: lipid-modifying dosages of niacin (>1 g/day), fibrates, and colchicine. Discontinue ZYPITAMAG if markedly elevated CK levels occur or myopathy is diagnosed or suspected. Temporarily discontinue ZYPITAMAG in patients experiencing an acute or serious condition at high risk of developing renal failure secondary to rhabdomyolysis; e.g., sepsis; shock; severe hypovolemia; major surgery; trauma; severe metabolic, endocrine, or electrolyte disorders; or uncontrolled epilepsy. Inform patients of the risk of myopathy and rhabdomyolysis when starting or increasing the ZYPITAMAG dosage. Instruct patients to promptly report any unexplained muscle pain, tenderness or weakness particularly if accompanied by malaise or fever.
- Immune-Mediated Necrotizing Myopathy (IMNM): There have been rare reports of IMNM, an autoimmune myopathy, associated with statin use. IMNM is
 characterized by: proximal muscle weakness and elevated serum creatine kinase, which persist despite discontinuation of statin treatment; positive anti-HMG CoA
 reductase antibody; muscle biopsy showing necrotizing myopathy; and improvement with immunosuppressive agents.
- Hepatic Dysfunction: Increases in serum transaminases can occur. Rare postmarketing reports of fatal and non-fatal hepatic failure have occurred. Consider liver enzyme testing before initiating therapy and as clinically indicated thereafter. If serious hepatic injury with clinical symptoms and/or hyperbilirubinemia or jaundice occurs, promptly discontinue ZYPITAMAG.
- Increases in HbA1c and Fasting Serum Glucose Levels: Increases of each have been reported with statins, including ZYPITAMAG. Optimize lifestyle measures, including regular exercise, maintaining a healthy body weight, and making healthy food choices.

ADVERSE REACTIONS: The most frequent adverse reactions (rate ≥ 2%) are myalgia, back pain, diarrhea, constipation and pain in extremity. This is not a complete list of all reported adverse events.

For additional information, refer to full <u>Prescribing Information</u>. You are encouraged to report negative side effects of prescription drugs to the FDA. Visit <u>www.FDA.gov/medwatch</u> or call 1-800-FDA-1088

