

Evaluation of Mavacamten in Symptomatic Patients With Cardiomyopathy

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Modelling genetic diseases for drug development: Hypertrophic cardiomyopathy. <i>Pharmacological Research</i> , 2020, 160, 105176.	3.1	12
2	Hypertrophic Cardiomyopathy 2020. <i>Current Cardiology Reports</i> , 2020, 22, 154.	1.3	9
3	2020 AHA/ACC Guideline for the Diagnosis and Treatment of Patients With Hypertrophic Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2020, 76, e159-e240.	1.2	364
4	2020 AHA/ACC Guideline for the Diagnosis and Treatment of Patients With Hypertrophic Cardiomyopathy. <i>Circulation</i> , 2020, 142, e558-e631.	1.6	263
5	Left ventricular dysfunction in heart failure with preserved ejection fraction—molecular mechanisms and impact on right ventricular function. <i>Cardiovascular Diagnosis and Therapy</i> , 2020, 10, 1541-1560.	0.7	14
6	Unraveling the Genotype–Phenotype Relationship in Hypertrophic Cardiomyopathy: Obesity–Related Cardiac Defects as a Major Disease Modifier. <i>Journal of the American Heart Association</i> , 2020, 9, e018641.	1.6	16
7	Commentary: In a heart of stone beats a muscle of pure flint. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2020, , .	0.4	0
8	Effects of mavacamten on Ca ²⁺ sensitivity of contraction as sarcomere length varied in human myocardium. <i>British Journal of Pharmacology</i> , 2020, 177, 5609-5621.	2.7	36
9	Characterization and validation of a preventative therapy for hypertrophic cardiomyopathy in a murine model of the disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 23113-23124.	3.3	7
10	Mavacamten for treatment of symptomatic obstructive hypertrophic cardiomyopathy (EXPLORER-HCM): a randomised, double-blind, placebo-controlled, phase 3 trial. <i>Lancet, The</i> , 2020, 396, 759-769.	6.3	481
11	Mavacamten: treatment aspirations in hypertrophic cardiomyopathy. <i>Lancet, The</i> , 2020, 396, 736-737.	6.3	8
12	Mavacamten: a novel small molecule modulator of β^2 -cardiac myosin for treatment of hypertrophic cardiomyopathy. <i>Expert Opinion on Investigational Drugs</i> , 2020, 29, 1171-1178.	1.9	13
13	Preventative therapeutic approaches for hypertrophic cardiomyopathy. <i>Journal of Physiology</i> , 2020, 599, 3495-3512.	1.3	6
15	Small Molecules Acting on Myofilaments as Treatments for Heart and Skeletal Muscle Diseases. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9599.	1.8	36
16	Avoiding Burnout From Hypertrophic Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2020, 75, 3044-3047.	1.2	5
17	Mavacamten decreases maximal force and Ca ²⁺ sensitivity in the N47K-myosin regulatory light chain mouse model of hypertrophic cardiomyopathy. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2021, 320, H881-H890.	1.5	25
18	Editor-in-Chief’s Top Picks From 2020. <i>Journal of the American College of Cardiology</i> , 2021, 77, 937-997.	1.2	0
19	Emerging Medical Treatment for Hypertrophic Cardiomyopathy. <i>Journal of Clinical Medicine</i> , 2021, 10, 951.	1.0	18

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41	Hypertrophic Cardiomyopathy – A Heterogeneous and Lifelong Disease in the Real World. <i>Circulation Journal</i> , 2020, 84, 1218-1226.	0.7	15
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56	Effect of Mavacamten on Echocardiographic Features in Symptomatic Patients With Obstructive Hypertrophic Cardiomyopathy. <i>Journal of the American College of Cardiology</i> , 2021, 78, 2518-2532.	1.2	59
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