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Hypertrophic Cardiomyopathy: Foreword by Bernard Gersh and Historical Context by Eugene Braunwald This e book offers a complete scientific evaluate of Hypertrophic Cardiomyopathy (HCM), the commonest genetic dysfunction of the center characterised by dysfunctional contractility on the sarcomere degree, ensuing within the improvement of irregular and infrequently focal hypertrophy on a macroscopic degree.

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Introduction This book provides a comprehensive clinical review of Hypertrophic Cardiomyopathy (HCM), the most common genetic disorder of the heart characterized by dysfunctional contractility at the sarcomere level, resulting in the development of abnormal and occasionally focal hypertrophy on a macroscopic level.

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Abstract Hypertrophic cardiomyopathy (HCM) is the most

common heritable cardiomyopathy, manifesting as left ventricular hypertrophy in the absence of a secondary cause. The genetic underpinnings of HCM arise largely from mutations of sarcomeric proteins; however, the specific underlying mutation often remains undetermined.

Hypertrophic Cardiomyopathy | JACC: Heart Failure

Learn about this heart disorder in which the heart muscle becomes abnormally thick, which makes it harder for the heart to pump blood.

Hypertrophic cardiomyopathy - Doctors and departments

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Folsom California Cardiologist Doctors physician directory - Read about how Hypertrophic cardiomyopathy (HCM) is associated with thickening of the heart muscle, most commonly at the septum between the ventricles, below the aortic valve.

Heart Disease: Hypertrophic Cardiomyopathy

BACKGROUND: Hypertrophic cardiomyopathy (HCM) is the most common inherited cardiac disease, which generally manifests during adolescence. Adolescents may be diagnosed incidentally, following the investigation of symptoms, or during family screening. Early recognition may prevent sudden cardiac death. First-degree relatives of an adolescent with HCM should be screened for the condition.

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