

**AIN SHAMS UNIVERSITY
INSTITUTE OF POST GRADUATE
CHILDHOOD STUDIES**

***PSYCHOLOGICAL DISTURBANCES IN PREADOLESCENTS
AND ADOLESCENTS WITH MITRAL VALVE PROLAPSE***

By

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M.Sc. Pediatrics Faculty of Medicine
Ain Shams University

Supervised By

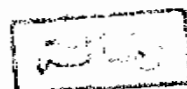
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DISCUSSION AND JUDGMENT COMMITTEE

The vice-president for higher studies and research of
Ain-Shams University has approved to form the following
committee for the discussion of Mr.....

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in Ain Shams University Hospital Member ... Mohamed Ghanem

Dedicated

*To those who cared and still care
To those who gave with love and still do
To my Mother and to the memory of my Father*

*To the one who carried the burden with no
Complaint and shared the hardships with
More encouragement To My Husband*

*To the joy of my life and the stardust blown from
The Lord's hands to the humble me
To Ahmed, Aya, Alaa*

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Contents

CONTENTS

	Page
- Introduction and Aim of The Work.....	i
- Review of Literture	1
• Embryology and anatomy of mitral valve.....	1
• Pathogenesis and pathology of mitral valve prolapse.....	11
• Definition and etiology of mitral valve prolapse.....	15
• Clinical recognition of mitral valve prolapse.....	24
- Incidence and prevalence.....	24
- Symptoms and signs.....	26
- Prognosis.....	41
• Investigations.....	42
- Roentgenography.....	42
- Electrocardiogram.....	42
- Echocardiography: M-mode	45
Two-dimensional.....	56
Doppler study.....	60
- Correlation between clinical and echocardiographic findings of mitral valve prolapse.....	66
- New guidelines for clinical diagnosis of mitral valve prolapse.....	69
• Psychological disturbances in mitral valve prolapse.....	71
- Introduction.....	71
- Anxiety disorders and mitral valve prolapse.....	73

	Page
- Depressive disorders and mitral valve prolapse.....	102
- Subjects and Methods.....	108
- Results.....	129
- Discussion and Recommendation.....	199
- Summary and Conclusion.....	229
- References.....	237
- Appendix.....	271
- Arabic Summary.....	

INTRODUCTION AND AIM OF THE WORK

Mitral valve prolapse is defined as the abnormal motion of one or both leaflets of the mitral valve into the left atrium during left ventricular systole.

Early observations in children and adolescents, an increasing awareness of the clinical features of this condition, and widespread availability of echocardiography have suggested a frequency in the pediatric population similar to that reported for adults (6.3%) (Bisset et al., 1980).

Boudoulas et al. (1990) proposed a clinically useful classification of patients with mitral valve prolapse. Mitral valve prolapse-anatomic which includes patients with a wide spectrum of floppy or myxomatous mitral valvular abnormalities from mild to severe due to expansion of the mitral valve leaflet area with elongated chordae and dilated mitral annuli.

The symptoms, physical findings and laboratory abnormalities in these patients are directly related to mitral valve dysfunction and complications associated with progressive mitral regurgitation. The term mitral valve prolapse syndrome refers to the occurrence of symptoms that result from various forms of neuroendocrine

Introduction and Aim of The Work

or autonomic dysfunction in patients with mitral valve prolapse in whom the symptoms can not be explained on the basis of valvular abnormality alone (Boudoulas et al., 1990).

The diagnostic criteria for mitral valve prolapse include:-

- ♦ Clinical symptoms as: palpitation, chest pain, dyspnea, fatigue, light headedness, dizziness and syncope.
- ♦ Extra cardiac signs as: pectus or spinal deformity, high arched palate and hyper-extensibility of joints. Auscultatory findings such as: apical mid-to late systolic click and apical mid or late systolic murmur.
- ♦ Electrocardiogram shows ST-T wave abnormalities (flattening or inversion in the inferoapical leads).
- ♦ Echocardiogram (two-dimensional) reveals prolapse of one leaflet in two views, prolapse of two leaflets in one view, and prolapse of one leaflet unequivocally in one view (Liberthson et al., 1986).
- ♦ A connection between neuropsychiatric disorders, notably panic disorders, and the heart-specifically, mitral valve prolapse has been recognized for more than a century and well before the clinical description of mitral valve prolapse.

Anxiety, panic attacks and neurotic behavior are often considered to be components of mitral valve prolapse syndrome whether the two syndromes are identical, are

Introduction and Aim of The Work

separate and distinct, or to a variable degree overlap still remains a matter of debate (Liberthson et al., 1986).

At present there is still widespread disagreement as to the degree of association between mitral valve prolapse and psychological symptoms.

AIM OF THE WORK:

The aim of this work is to study the prevalence of psychological disturbances in preadolescent and adolescent patients with mitral valve prolapse.

Also to evaluate the relationship between the severity of mitral valve prolapse and the presence of psychological disturbances in those patients.

Introduction and Aim of The Work

Review of Literature

EMBRYOLOGY OF THE MITRAL COMPLEX

A) Formation of endocardial cushions

In the earliest stages of the development of the embryonic heart the cardiac tube is lined by endothelium which is widely separated from an outer myoepicardial cover by an acellular fluid, the cardiac jelly. By about 34 days, localized masses of mesenchymal cell have accumulated within the cardiac jelly and encircle the common atrioventricular canal. These masses protrude at two locations, superiorly and inferiorly, into the lumen of the embryonic heart as the superior and inferior endocardial cushions. When these endocardial cushions fuse in the midline, the freely communicating chambers are then divided into a left and a right atrioventricular canal (Fig 1). The superior and inferior endocardial cushions also contribute to the formation of atrioventricular valves, the lower atrial septum, and the membranous ventricular septum (Van Mierop et al., 1962).

By the seventh week, the superior and inferior endocardial cushions have fused together uniting with interventricular and interatrial septum (Alley et al., 1962).

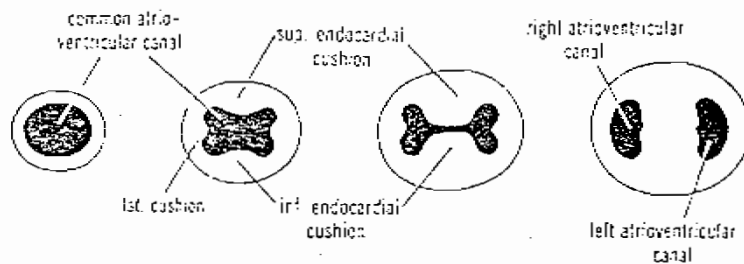


Fig (1): Formation of the septum in the atrioventricular canal. The initial circular opening becomes gradually widened in transverse direction
(Langman, 1980).

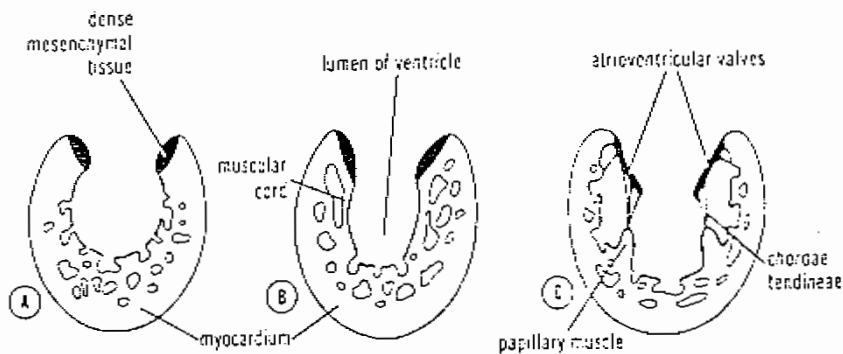


Fig (2): Formation of the atrioventricular valves and chordae tendineae. The valves are hollowed out from the ventricular side, but remain attached to the ventricular wall by the chordae tendineae (Langman, 1980).