

## ***INTRODUCTION***

In patients with hypertension, a pattern of left ventricular hypertrophy, on the electro cardiogram is associated with a risk of sudden death in excess of the risk attributable to hypertension alone (*James, et al., 1987*).

Left ventricular hypertrophy is an important risk factor for cardiovascular morbidity and mortality (*Kannal and Schatzkin, 1985*).

Persons with electrocardiographic left ventricular hypertrophy are at a high risk for sudden cardiac death, a catastrophic event that is a well recognized consequence of ventricular fibrillation or rapid ventricular tachycardia in most of monitored cases (*Schaffer and Cobba, 1975*).

The QT interval represents the time of total electrical activity of the left ventricle including ventricular depolarization and repolarization (*Schwartz and Wolf, 1978*).

The prolongation of the rate corrected QT interval (Qtc) is considered to be a marker for increased risk for malignant arrhythmias and sudden cardiac death (*Moss, 1993*).

In addition to the inherited forms of prolonged QT interval, QT prolongation had been noted in hypertrophic cardiomyopathy (*Buja et al., 1967*) and in certain drug toxicities (*Devereux and Reichek, 1977*).

There is a significant variation in regional repolarization which is reflected as a variation in the QT interval from lead to lead on body surface electro cardiogram (*Franz et al., 1978*).

QT dispersion which is defined as the difference between the maximum and minimum QT occurring in any of the 12 leads of the electrocardiogram is believed to have a predictive value in the assessment of the risk for ventricular arrhythmias (*Day et al., 1992*).