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Valvular Heart Disease

PHENOTYPES OF MITRAL VALVE PROLAPSE AS A CAUSE OF SUDDEN DEATH: A CASE-CONTROL STUDY

Poster Contributions

Poster Hall B1

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Background: Description of Sudden Death (SD) complicating Mitral Valve Prolapse (MVP) come mainly from case reports or autopsy series. Our aim was to identify MVP phenotypes associated with SD.

Methods: This international multicenter study included 42 patients who received an implantable cardioverter-defibrillator after surviving SD for which MVP was the only detectable potential structural cause. Clinical, echocardiographic and electrocardiographic features were compared to 84 matched MVP controls.

Results: Twenty-eight cases (67%) were female of age 45 ± 15 years. At MVP diagnosis, Left Ventricle Ejection Fraction was $62\pm 5\%$, 29 patients (69%) had mild to moderate mitral regurgitation. Compared to controls, more SD cases had 1) familial SD history (4 vs 19%, $P=0.006$); 2) syncope (4 vs 59%, $P<0.0001$) and palpitations symptoms (33 vs 93%, $P<0.0001$); 3) bileaflet MVP (56% vs 88%, $P=0.0003$), myxomatous diffuse disease (67% vs 100%, $P<0.0001$), mitral annulus disjunction (12% vs 100%, $P<0.0001$) and deep MVP (0.6 ± 0.3 vs 1.3 ± 0.3 cm, $P<0.0001$); 4) more frequent ventricular ectopic beats ($2\pm 4\%$ vs $11\pm 8\%$, $P=0.002$), couplets (43 ± 120 vs 681 ± 128 , $P=0.003$) and non-sustained ventricular tachycardia (2 ± 4 vs 57 ± 55 , $P=0.002$) on 24 hours Holter interrogation; 5) more premature ventricular contractions from posterior papillary muscle area ($10\pm 22\%$ vs $64\pm 30\%$, $P<0.0001$) and less from right/left ventricular outflow tracts ($47\pm 46\%$ vs $8\pm 21\%$, $P=0.0003$); 6) greater corrected QT dispersion (27 ± 9 ms vs 62 ± 17 ms, $P<0.0001$) and 7) left diastolic (51 ± 5 mm vs 55 ± 5 , $P<0.0001$) and systolic (33 ± 4 mm vs 37 ± 5 mm, $P<0.0001$) ventricular enlargement.

Conclusion: We identified MVP phenotype associated with SD, including symptomatic women with familial history of SD, bileaflet and deep myxomatous MVP, associated with annular disjunction and ventricular enlargement in a context of frequent and complex ventricular arrhythmias arising from papillary muscle or fascicular origin.