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**JOURNALSCAN**

**ISCHAEMIC HEART DISEASE**

Clopidogrel for all acute coronary syndromes? JournalScan mentioned the CURE trial when it was presented at the American College of Cardiology meeting earlier this year (Heart 2001;86:15). The full paper is now published. Clopidogrel in addition to aspirin reduces the risk of death/myocardial infarction (MI) and stroke by 20% if given to all patients with non-ST elevation acute coronary syndromes. At present, clopidogrel is only used after stent implantation in the UK. The CURE study showed similar benefits in low, medium, and higher risk patients (such as those with ST depression, positive troponins, or haemodynamic instability). Initiating treatment in all patients with acute coronary syndromes awaiting transfer for in-patient angiography is probably merited. The CURE investigators suggest that this group has a 28% reduction in death/MI at nine months follow up, having had six days pretreatment with the drug.


In-hospital arrest has a survival rate of 23% to discharge: A study of in-hospital resuscitation events suggests that 104 (23%) of 445 patients who received full advanced cardiac life support survived to hospital discharge. Survival was highest for patients with primary cardiac disease (30%), followed by those with infectious diseases (15%), with only 8% of patients with end stage diseases surviving to hospital discharge. Neither sex nor age affected survival. As expected, longer resuscitations were associated with poor survival. Patients who experienced arrests on a nursing unit or intensive care unit had better survival rates than those in other hospital locations. Survival for witnessed arrests (25%) was significantly better than for non-witnessed arrests (7%) (p = 0.005). Non-witnessed arrests in unmonitored beds were uniformly fatal.


TMR—a placebo effect: There are no objective data to suggest sustained benefit from transmyocardial laser revascularisation (TMR). This paper evaluates 98 patients who underwent TMR versus 99 on continued medical treatment for refractory angina. The study was not blinded and there was a 60% crossover from medical to laser therapy! Not surprisingly the TMR group did better with a 44% reduction in angina versus 21% in the medical group. There was no objective evaluation of exercise time, perfusion, etc and the trial was not powered to detect mortality differences. This could all be placebo effect detected by their sensitive health questionnaires.


Total cholesterol: HDL ratio is a better measure of risk than total or LDL cholesterol alone: Populations with lower cholesterol: HDL concentration than the US and UK populations may not reach the threshold targets set by the European Society of Cardiology and American Heart Association/American College of Cardiology guidelines for initiation of statin treatment, but may still be at risk. The same is true of diabetics in the UK, with a low high density lipoprotein (HDL) concentration compound the cholesterol risk. In a case-control study of men with MI compared to controls, the total cholesterol (TC): HDL ratio predicted risk better than TC alone. Algerian patients have a TC of 5.2 mmol/l versus 5.8 mmol/l in France and 6.2 mmol/l in Ireland. The ratio of TC:HDL remained at about 6 for patients with MI from all countries while controls had a ratio of 5.0 or less.


BNP as a risk stratifier in ACS: Brain (B type) natriuretic peptide (BNP) is a neurohormone synthesised predominantly in ventricular myocardium. It provides independent prognostic information in patients with transmural myocardial infarction and can help screen for HF. This trial measured BNP in plasma specimens obtained a mean (SD) of 20 (hours) after the onset of ischaemic symptoms in 2525 patients from the orbofiban in patients with unstable coronary syndromes–thrombolysis in myocardial infarction study 16 study (OPUS-TIMI 16). The baseline level of BNP was correlated with the risk of death, heart failure, and MI at 30 days and 10 months in subgroups of patients who had myocardial infarction with ST segment elevation (p = 0.02), patients who had myocardial infarction without ST segment elevation (p < 0.001), and patients who had unstable angina (p < 0.001). After adjustment for independent predictors of the long term risk of death, the odds ratios for death at 10 months in the second, third, and fourth quartiles of BNP were 3.8 (95% confidence interval (CI) 1.1 to 13.3), 4.0 (95% CI 1.2 to 13.7), and 5.8 (95% CI 1.7 to 19.7). The concentration of BNP was also associated with the risk of new or recurrent myocardial infarction (p = 0.01) and new or worsening heart failure (p < 0.001) at 10 months. Cut off of 80 pg/ml was suggested as a concentration above which risk was substantially increased. Risk prediction was independent of troponins and C reactive protein.


ST analysis of Holter monitors is not a poor man’s exercise test: Asymptomatic ischaemia is known to be associated with worse outcome, and can be diagnosed on ST segment analysis of Holter monitors. This study prospectively assessed 277 patients with chest pain referred for coronary angiography. All patients had a 24 hour ambulatory Holter within 72 hours of the angiogram and were then followed up for the occurrence of death, infarction, hospitalisation for unstable angina, and the need for revascularisation over a five year period. The overall incidence of coronary artery disease was high, at 80%, yet only 20% of patients had significant ST segment deviation on ambulatory monitoring. Comparing patients with and without ST segment changes who were otherwise matched, the incidence of coronary artery disease (CAD) at angiography was similar in the two groups (90% and 79%, respectively) and the occurrence of the combined clinical end point was also similar (48% vs 44%). Overall, the sensitivity of Holter ST segment deviation for diagnosing CAD was only 19% with a specificity of 91%, making its diagnostic accuracy very poor, even among patients with left main stem or triple vessel disease. This study was flawed as the patients were already at high risk for CAD and may have been limiting their activity to reduce
Angiography for patients > 75 with resistant angina: Patients older than 75 years are often denied access to coronary angiography. Now the TIME study addresses this issue by randomising to medical treatment or angiography after two drugs have been tried to relieve symptoms. Of 153 patients in the invasive group, 109 (74%) underwent recanalisation. The remaining 38 (26%) were treated medically because they could not be recanalised (19), because they refused recanalisation (7), or had no significant coronary artery disease (11). Coronary artery bypass graft (CABG) surgery was performed in 25 (20%), and 68 (54%) had percutaneous coronary intervention (PCI) using at least one stent. In the optimum medical group, anti-angiinal medication was increased by a mean (SD) of 0.80 (0.6) drugs per patient, with additional increases in drug dosages in 81 (55%) patients. Death, non-fatal MI, or hospital admission for acute coronary events occurred in 49 (40%) of the medical group with 19% of the invasive group (p < 0.0001). Mortality at six months did not differ between the two groups (8.4% in the invasive group, 4.1% in the medical group), but the study was not powered to detect such a difference. A third of the patients in the medical group needed a recanalisation procedure during follow up for uncontrollable symptoms.

LIMA and RIMA for CABG as the norm? Ten years after CABG, 90–95% of left internal mammary artery (LIMA) grafts are patent and disease free, whereas 75% of vein conduits are patent and disease free, whereas 75% of vein conduits are associated with an increased cataract risk (adjusted odds ratio (OR) 1.14, 95% CI 0.5 to 1.6), nor was use of fibrates or of other lipid lowering drugs (adjusted OR 0.5, 95% CI 0.3 to 1.1; and OR 0.7, 95% CI 0.1 to 5.6, respectively). However, use of simvastatin and an inhibitor of its metabolism, such as erythromycin, did result in increased risk (adjusted OR 2.2, 95% CI 1.2 to 4.1). In a separate study, statins seem to be associated with a reduced risk of macular degeneration (corrected OR 0.9, 95% CI 0.01 to 0.73).

Heart failure nurses reduce admissions and thereby costs of treatment: Admissions with heart failure are common. Excluding those caused by valve disease or acute MI, 165 patients with systolic left ventricular dysfunction were randomised to standard care, or inpatient and subsequent intensive outpatient monitoring by specialist heart failure nurses, with a remit to implement evidence-based treatment. The trial was not designed to take at mortality differences, but the main cost of heart failure treatment is recurrent admission. The combined end point of death or readmission for heart failure was investigated at one year. Only 31 patients (37%) in the intervention group died or were readmitted with heart failure compared with 45 (53%) in the usual care group (hazard ratio 0.61, 95% CI 0.33 to 0.96). Compared with usual care, patients in the intervention group had fewer readmissions for any reason (86 v 114, p = 0.018), fewer admissions for heart failure (19 v 45, p < 0.001), and spent fewer days in hospital for heart failure (mean 3.43 v 7.46 days, p = 0.0051). They also were on larger doses of ACE inhibitors.

Exudates caused by heart failure: Pleural effusions are common in patients with heart failure. Light’s criteria for differentiation between exudates and transudates were used to classify effusions in 81 patients. A total of 54 effusions in 47 patients were found. Of these an obvious cause for the exudates (except heart failure) was found in 22 effusions in 20 patients. Intravenous diuretic treatment in the 24 hours before thoracentesis was significantly more common among patients with exudates without a specific cause.

Statins and the eye: Some studies in dogs suggest that statin use may increase cataract formation. A UK case–control study of 7409 cases and 3727 controls suggests statin use was not associated with an increased cataract risk (adjusted odds ratio (OR) 0.9, 95% CI 0.5 to 1.6), nor was use of fibrates or of other lipid lowering drugs (adjusted OR 0.5, 95% CI 0.3 to 1.1; and OR 0.7, 95% CI 0.1 to 5.6, respectively). However, use of simvastatin and an inhibitor of its metabolism, such as erythromycin, did result in increased risk (adjusted OR 2.2, 95% CI 1.2 to 4.1). In a separate study, statins seem to be associated with a reduced risk of macular degeneration (corrected OR 0.9, 95% CI 0.01 to 0.73).

Heart failure: Failing to deliver heart failure treatment: Previous studies suggest a prevalence of heart failure of 2.9% in those aged < 75 years. In this study of a population aged > 45 years, 1.7% had left ventricular systolic dysfunction. Only 50% of these had symptoms. Of those with definite heart failure, only 30% were on angiotensin converting enzyme (ACE) inhibitors and only 11% on β blockers. Screening for disease is useful when treatment is available, but the treatment has to be prescribed (and taken) to be effective.

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incidence of pulmonary embolism was much higher among passengers travelling more than 5000 km (3100 miles) (1.5 cases per million, as compared with 0.01 case per million among those travelling less than 5000 km). The incidence of pulmonary embolism was 4.8 cases per million for those travelling more than 10 000 km (6200 miles).

5-Fluorouracil can cause coronary vasospasm: The mechanism of 5-fluorouracil (5-FU) cardiotoxicity is not certain. In vitro-studies have shown an endothelial-dependent constriction of vascular smooth muscle cells, involving activation of protein kinase C. Previous case reports suggest very high doses (> 800mg/day) can cause vasospastic angina. Now there is a report of three cases likely to be vasospasm secondary to 5-FU low dose infusion. Symptoms were associated with the initiation of 5-FU treatment, and disappeared after its discontinuation. Although angiographic confirmation of vasospasm was not obtained, ST segment elevation during the early recovery phase of an exercise stress test is considered a hallmark of transmural ischaemia caused by vasospastic origin. Disappearance of these changes with repeat exercise testing after discontinuation of 5-FU treatment confirmed the likely diagnosis.

Syncope—two thirds of patients have an easily identifiable cause: In 611 consecutive patients who presented to the emergency department with syncope as a chief complaint, initial evaluation included history and clinical examination, with carotid sinus massage, ECG, and basic laboratory testing. This yielded a suspected cause of syncope in 69% of the 611 patients, including neurocardiogenic syncope (n = 334, 36%), orthostatic hypotension (n = 156, 24%), arrhythmia (n = 24, 4%), and other diseases (n = 32, 5%). Extensive cardiovascular workups, which were performed in 122 of the 155 patients in whom syncope remained unexplained, provided a suspected cause of syncope in only 30 more patients, including arrhythmias in 18, all of whom had abnormal baseline ECGs. An acute coronary syndrome was the cause in only nine patients.

The importance of revascularization of the transplanted heart: Does a transplanted heart feel pain as it has no nerve supply? This is unexplained, provided a suspected cause of syncope in only 30 more patients, including arrhythmias in 18, all of whom had abnormal baseline ECGs. An acute coronary syndrome was the cause in only nine patients.

HYPERTENSION

Lower the BP after CVA to reduce recurrent CVA risk: Aspirin reduces recurrent cerebrovascular accident (CVA) by about 20%. Lowering systolic blood pressure (BP) by 10 mm Hg lowers recurrent CVA by 28% in patients with ischaemic CVAs or transient ischaemic attacks. Lowering diastolic BP by 5–6 mm Hg lowers risk of first CVA by a third. Now the PROGRESS study suggests that lowering BP by 9/5 mm Hg can reduce recurrent CVAs. Stroke occurred in 307 (10%) individuals assigned active treatment, compared with 420 (14%) assigned placebo (relative risk reduction 28% (95% CI 17% to 38%) over four years follow up). Combination therapy with perindopril plus indapamide reduced BP by 12.5 mm Hg and stroke risk by 43% (95% CI 30% to 54%). The reduction in first MI (38% reduction in non-fatal MI) was in keeping with data from the HOPE trial. Most were on aspirin and half were on antihypertensives. It seems that the lower the blood pressure the better. Interestingly, perindopril alone, which produced a smaller BP drop, was not associated with significant benefit.

1 PROGRESS Collaborative Group. Randomised trial of a perindopril-based blood-pressure-lowering regimen among 6105 individuals with previous stroke or transient ischaemic attack. Lancet 2001;358:1003–11.

BASIC RESEARCH

A case report in a science journal: Nitric oxide (NO) is known to be a vasodilator. Disease states that affect the endothelium display impaired NO dependent vasodilatation. Thus atherosclerosis is associated with abnormal flow mediated brachial artery dilatation. The authors report a rare case of lysinuric protein intolerance (LPI)—an autosomal recessive defect of dibasic amino acid transport caused by mutations in the SLC7A7 gene, resulting in an L-arginine deficiency. They found that the patient’s serum concentration of NO and his flow mediated brachial artery vasodilator response were each approximately 70% lower than in controls. Both of these could be corrected by intravenous infusion of L-arginine. Despite a normal coronary angiogram, exercise testing and positron emission tomography showed evidence of myocardial ischaemia. Again this could be corrected by intravenous administration of L-arginine. This provides more evidence that NO is vital in the control of tissue perfusion.


An animal model for long QT syndrome type III: Abnormal prolongation of repolarization, reflected by a long QT interval, is associated with ventricular arrhythmias in the long QT (LQT) syndrome. An inherited deletion in the cardiac sodium (Na+) channel, encoded by SCN5A, causes type I-a autosomal dominant LQT3 syndrome, associated with fatal ventricular arrhythmias. This report shows that Scn5a–/– mice lacking the same residues as LQT3 patients showed the essential features of the LQT3 syndrome and were highly susceptible to ventricular tachyarrhythmias in response to sudden heart rate accelerations or premature beats. These arrhythmias were attributable to a paradoxical lengthening of the action potential, initiated by an increased peak and late Na+ inward current. Adrenergic agonists normalised the response to rate acceleration in vitro and suppressed arrhythmias upon premature stimulation in vivo. These results show the possible risk of sudden heart rate accelerations. The Scn5a–/– mouse with its predisposition for pacing induced arrhythmias might be useful for the development of new treatments for the LQT3 syndrome.