Absolute, attributable, and relative risk in the management of coronary heart disease

Absolute risk

Treatment decisions in the prevention and management of coronary heart disease (CHD) often require knowledge about the level of risk. Risk is defined as the probability of encountering a particular event. Information about risk is obtained from cohort studies or other longitudinal studies such as randomised controlled trials. In these studies, risk is simply the incidence of the event in a particular group. For example, the risk (or probability) of death associated with coronary angiography is 0.1% (or 1 in 1000). This is termed the absolute risk.

In order to inform clinical decisions, it is often useful to compare the absolute risk in two or more groups having different exposures or different treatments (including treatment versus no treatment). The two main measures we use when comparing risk are attributable risk (risk difference) and relative risk (risk ratio).

Attributable risk

Attributable risk measures the excess risk accounted for by exposure to a particular factor. This is simply the difference between the absolute risks in the two groups. The term attributable risk is most commonly used in epidemiological studies. In the clinical environment, when comparing a particular treatment with placebo for example, the excess risk associated with treatment (attributable risk) may well be negative, if the treatment is beneficial. This is therefore termed an absolute risk reduction. Absolute risk reduction is increasingly used when reporting clinical trials. The absolute risk reduction enables calculation of the “number needed to treat” (NNT). The NNT is simply the inverse of the absolute risk reduction. The NNT represents the number of patients that need to be treated to prevent one adverse event.

Relative risk

Relative risk is the ratio of two absolute risks. It measures the strength of effect of an exposure (or treatment) on risk. A beneficial treatment will result in a relative risk of less than 1; this can then be subtracted from 1 to give the relative risk reduction. A harmful treatment, or other exposure, than 1; this can then be subtracted from 1 to give the relative risk reduction. If a condition is common the importance of reducing risk is much less than if it is rare. In clinical practice, the relative risk reduction associated with treatment of blood pressure or serum cholesterol is constant at different levels of absolute risk. So, the absolute risk reduction associated with antihypertensive or cholesterol lowering treatment is proportional to the initial absolute risk. This is why guidelines for the primary prevention of CHD require quantitative prediction of absolute risk based on a patient's risk factor profile.

The Joint British Societies' recommend that “as a minimum all individuals with an absolute CHD risk of 30% or more over 10 years should be targeted now for comprehensive risk factor management, which will include, as appropriate, blood pressure and lipid lowering therapy” followed by “a progressive expansion of coronary prevention from 30% down to 15% absolute CHD risk”. If we assume that the relative risk reduction associated with statin treatment is 33%, in patients with an initial risk of 30% the absolute risk will be reduced (by one third) from 30% to 20% (that is, 10%). In those with an initial risk of 15% it will be reduced (by one third again) from 15% to 10% (that is, only 5%). These absolute risk reductions give NNTs of 10 and 20, respectively. So half as many patients with an initial risk of 30% (compared with 15%) need to be treated to prevent one adverse event.

Other guidelines based on the same Framingham equations exist. However, in the recently published comparison study, in primary care, the joint British guidelines appeared to perform at least as well as others. Guidelines differ in their advice on the level of risk at which treatment should be initiated. In the UK the recommended threshold is partly an issue of cost to the National Health Service (NHS). The National service framework for coronary heart disease recommends targeting risk reduction at those with a 10 year CHD risk greater than 30%. This will mean 3% of men aged 30–74 years are targeted. If 15% were chosen as the threshold 28% would be eligible. So, while (initially) focusing on patients at highest risk sensibly targets resources at those who are most likely to benefit, the policy will deny a large proportion of the population effective treatment.

Secondary prevention

Of course, those patients with the greatest risk of a future CHD event are those with CHD already. Secondary prevention is therefore the area with the greatest potential for patient benefit. Absolute risk quantification is probably unnecessary in these patients, as all should be subject to attempts at risk reduction through: lifestyle changes (smoking, diet, and physical activity); the control of blood pressure, lipids, and glucose; and drug treatment (for example, aspirin). The purpose is to reduce the risk of a major cardiovascular event and reduce mortality. In these patients, it is important to realise and communicate the likely benefits of behavioural change in preventing further CHD events, such as the 50% relative risk reduction associated with stopping smoking, and the importance of long term compliance with antihypertensive and cholesterol lowering treatment, and aspirin, each of which are probably associated with relative risk reductions in CHD events of 10–30%.
In order to initiate risk reduction strategies in CHD, a clear understanding of the meanings and appropriate uses of absolute, attributable, and relative risk is required.

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IMAGES IN CARDIOLOGY

Right aortic arch and coarctation: delineation by three dimensional magnetic resonance angiogram

A 14 year old girl returned for follow up. She initially presented with coarctation and right aortic arch at two months of age. Cardiac catheterisation determined the coarctation to be between the right common carotid (RCC) and right subclavian arteries (RSA), with a retro-oesophageal left subclavian artery. The intracardiac anatomy was normal. The patient underwent patch augmentation of the coarctation site at 2 months of age and again at 8 months of age for recurrent coarctation. At 14 years of age cardiac catheterisation demonstrated a widely patent conduit but a residual coarctation between the right carotid and subclavian arteries with a 16 mm Hg gradient. A magnetic resonance angiogram including three dimensional reconstruction provided excellent definition of the isolated coarctation between the RCC and RSA, in addition to identifying additional areas of stenosis in the RSA and upper descending thoracic aorta (below left and right).

Right aortic arch in association with coarctation is extremely rare. This is in keeping with the principle of flow related development of the central great vessels. Coarctation is more likely to occur in situations of right to left shunting through the ductus arteriosus with decreased blood flow across the aortic isthmus. Additionally right aortic arch is strongly associated with right side obstructive lesions, in which there is reversed ductal flow and increased antegrade flow across the isthmus, which makes this entity even rarer.

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(Left) Left lateral view of three dimensional reconstructed gadolinium enhanced magnetic resonance angiogram showing the right aortic arch and the hypoplastic descending aorta. Note the residual coarctation between the right common carotid and subclavian arteries, and the additional areas of stenosis in the right subclavian artery and descending aorta. The mid-conduit stenosis can be clearly seen. There is a retro-oesophageal left subclavian artery. (Right) Posterior view showing the residual coarctation and the mid-conduit stenosis. A, anterior; P, posterior; L, left, R, right, H, head, F, foot.
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