

## Abstracts

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#### Long-term predictors of descending aorta aneurysmal change in patients with aorta dissections

Song J-N, Kim S-D, Kim J-H, et al. *J Am Coll Card* 2006;50:799-804.

**Conclusion:** Aneurysmal dilatation as a late manifestation of aortic dissection is most likely to take place in the upper thoracic aorta. Aneurysmal dilatation of the upper thoracic aorta is predicted by a large false lumen diameter on the initial computed tomography (CT) scan.

**Summary:** After acute aortic dissection, late aneurysmal dilatation of the aorta is a significant complication. The authors sought to demonstrate the long-term natural history of descending aorta dilatation after acute aortic dissection. They also sought to identify early predictors of late aneurysmal change in the dissected aorta. Contrast-enhanced CT scans for acute aortic dissection were performed in 100 patients. There were 51 patients with DeBakey type 1 dissections and 49 patients with DeBakey type 3 dissections. The patients with type 1 dissections underwent ascending aortic surgery  $\leq 24$  hours of diagnosis.

Clinical follow-up was for  $53 \pm 26$  months, and CT scans were repeated at a mean of  $31 \pm 27$  months. Aortic dilatation to  $>60$  mm occurred in 14.4% of the upper thoracic descending aortas, 8.2% of the middle descending thoracic aortas, 4.1% in the lower descending thoracic aortas, and in 3.1% of the abdominal aortas. A repeat CT scan was done in 53 patients  $>2$  years after the initial dissection. On the basis of these scans, rates of aortic enlargement per year at the upper, middle, and lower thoracic, and abdominal aorta levels were  $3.43 \pm 3.66$ ,  $3.21 \pm 2.70$ ,  $2.62 \pm 2.19$ , and  $1.93 \pm 3.66$  mm, respectively ( $P < .01$ ). Aneurysms developed in 28%. Predictors of late aneurysmal change included the initial false lumen diameter of the upper thoracic aorta, aortic diameter in the middle thoracic aorta, and Marfan syndrome. A  $\geq 22$ -mm initial false lumen diameter of the upper thoracic aorta predicted late aneurysmal degeneration with a sensitivity of 100% and a specificity of 76%. The 42 patients with initial upper thoracic aorta false lumen diameters  $\geq 22$  mm also had a higher rate of the combination of aneurysm formation and death ( $P < .001$ ).

**Comment:** The study suggests patients with acute aortic dissection with large false lumens of their upper thoracic aorta may be ones where prophylactic stent grafting might be beneficial. A European trial that has either just begun, or will begin soon, is investigating the impact of stent grafting on uncomplicated acute type 3 aortic dissections, the ADSORB (Acute Dissection Stenting or Best Medical Treatment study). Results of this trial will, hopefully, help determine the optimal role of stent grafting for acute aortic dissection.

#### Early results after staged hybrid repair of thoracoabdominal aortic aneurysms

Lee WA, Brown MP, Martin TD, Seeger JM, et al. *J Am Coll Surg* 2007;2005:420-31.

**Conclusion:** The hybrid approach of thoracoabdominal aneurysm (TAA) repair can be performed with relatively few technical complications, but morbidity and mortality rates remain significant.

**Summary:** The authors reviewed the morbidity and mortality associated with the hybrid approach of TAA repair. This is a retrospective review of 17 patients (mean age,  $69 \pm 15$  years, 76% men), who underwent renal and visceral revascularization as a first stage of a hybrid repair of a TAA. The TAAs treated in this study included two Crawford extent type II, eight type III, and seven type IV. Perioperative mortality and complication rates associated with the debranching portion of the procedure were 24% and 25%, respectively. The mean intensive care unit stay and hospital stay were  $7 \pm 12$  days and  $22 \pm 33$  days, respectively. The mean delay between the staged procedures was  $27 \pm 27$  days. The thoracic stent graft (stage 2) was placed in 12 (92%) of the 13 patients who survived the initial portion of the procedure. No additional deaths or postoperative complications were associated with the second stage of the procedure. After the second stage, patients did not require intensive care unit stay, and the overall length of stay after the procedure was  $2 \pm 2$  days. Postoperative follow-up among 11 patients completing both stages was  $8 \pm 12$  months, with no additional deaths during follow-up. Primary patency for renal and visceral grafts was 96% (54 of 56).

**Comment:** The hybrid approach to TAA repair is difficult operation for both the patient and the surgeon. If complete debranching of the aorta is performed, and a conduit is placed for subsequent placing of the thoracic endograft, operative times of 6 to 9 hours are expected. Mortality and morbidity rates are significant. The hybrid approach to TAA repair, although a tribute to surgical technical expertise, is not likely to be the final word in the surgical treatment of patients with TAAs. We await the availability of an

off-the-shelf branched stent graft that can be implanted with a reasonable and not exceptional level of technical expertise.

#### Effects of torcetrapib in patients at high risk for coronary events

Barter PJ, Caulfield M, Eriksson M, and the ILLUMINATE Investigators. *N Engl J Med* 2007;357:2109-22.

**Conclusion:** Torcetrapib therapy, which increases high-density lipoprotein (HDL), results in an increased risk of mortality and morbidity.

**Summary:** Cholesteryl ester transfer protein (CETP) results in transfer of cholesteryl esters from HDL to other lipoproteins. Inhibiting CETP therefore results in increased HDL cholesterol levels and results in decreased low-density lipoprotein (LDL) cholesterol levels. Given these possible antiatherogenic effects, it would seem reasonable that torcetrapib, an inhibitor of CETP, would have a favorable effect on morbidity and mortality. To test this hypothesis, the authors conducted a double-blind randomized study of 15,067 patients considered at high cardiovascular risk. Patients were treated with atorvastatin alone or torcetrapib plus atorvastatin. The primary outcome was time to the first major cardiovascular event, which was defined as death from coronary heart disease, stroke, or hospitalization for unstable angina or nonfatal myocardial infarction.

In patients receiving torcetrapib therapy, after 12 months, there was an increase of 72.1% in HDL and a decrease of 24.9% in LDL cholesterol compared with baseline ( $P < .001$ ), and there was also an increase in systolic blood pressure of 5.4 mm Hg. Serum sodium, bicarbonate, and aldosterone levels all also increased in patients treated with torcetrapib ( $P < .001$ ). Patients treated with torcetrapib had an increased risk of cardiovascular events (hazard ratio, 1.25; 95% confidence interval, 1.09-1.44;  $P = .001$ ) as well as an increase of death from any cause (hazard ratio 1.58; 95% confidence interval, 1.14-2.19;  $P = .006$ ). Post hoc analysis indicated there was an increased risk of death in patients with torcetrapib therapy who had increases in bicarbonate or reductions in potassium levels greater than the median change.

**Comment:** This study documented a number of off-target pharmacologic effects of torcetrapib. The increase of systolic blood pressure, as well as the metabolic effects, may have negated the effects of improvement in HDL and LDL levels induced by torcetrapib. This study, therefore, neither validates nor invalidates the possibility that raising HDL cholesterol levels by inhibition of CETP can be cardio protective. As the authors point out in the discussion of the article, this hypothesis can only be tested by use of a CETP inhibitor that does not share the off-target pharmacologic effects of torcetrapib.

#### Childhood body-mass index and the risk of coronary heart disease in adulthood

Baker JL, Olsen LW, Sorensen TIA. *N Engl J Med* 2007;357:2329-37.

**Conclusion:** Increases in the body mass index (BMI) during childhood are associated with an increased risk of coronary heart disease (CHD) in adulthood.

**Summary:** Children are becoming overweight at progressively younger ages, with 19% of children aged 6 to 11 years in the United States considered overweight (Int J Obes Relat Metab Disord 2002;26[suppl 4]:S2-S4, and *JAMA* 2006;295:1549-55). In this study, the authors sought to determine long-term effects of increased weight in childhood CHD. Authors studied the association between BMI in children aged 7 to 13 years and CHD in adults aged  $>25$  years. Data were analyzed with and without adjustment for birth weight.

Subjects were derived from a cohort of 276,845 school children in Denmark whose height and weight measurements were available. National registries were used to determine CHD events. There were 5,063,622 person-years of follow-up. A total of 10,235 adult men and 4318 adult women had a diagnosis of CHD or died of CHD for whom childhood BMI data were also available. Risk of fatal and nonfatal events and any CHD event in the adults were positively associated with BMI from 7 to 13 years of age for boys and from 10 to 13 years of age for girls. Risk increased across the entire BMI distribution, and associations were linear for each age. Risk also increased as the age of the child increased. Adjustment for birth weight strengthened the results.

**Comment:** The data provide some quantitative information on the adverse effect of childhood obesity on future health. The magnitude of increased risk of CHD is relatively modestly increased in children at 7 years of age and dramatically increases further by age 13. Parents must somehow