## Heartbeat: Achieving better medication adherence

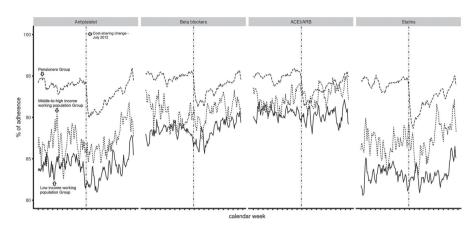
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Catherine M Otto

Even when effective medical therapies are available and prescribed, patient adherence is an important factor in the actual clinical benefit experienced by patients. One postulated reason for poor adherence is patient out-of-pocket medication expenses. To examine the relationship between adherence and patient expenses. González López-Valcárcel and colleagues<sup>1</sup> examined the effects of a change in costsharing in a population based study of over 10 thousand patients with an acute coronary event. A low income working population (no change in medication expenses) was compared with pensioners (a increase from no cost-sharing to 10%) and a middle to high income working population (an increase in cost-sharing from 40% to 50%-60%). For low-cost essential medications, such as antiplatelet therapy and beta-blockers, the change in cost-sharing had no significant effect. However, for higher cost medications, such as statins, adherence temporarily decreased by about 8% in both pensioner and middle-high income populations, who experienced higher out-of-pocket costs (figure 1).

In the accompanying editorial. Mathews<sup>2</sup> comments that these findings are aligned with previous studies showing poor medication adherence in acute coronary syndrome patients. In addition to out-of-pocket expenses, other factors such as health literacy and sociodemographic factors may contribute to non-adherence with prescribed medications. In addition, future studies should consider the effects of nonadherence on clinical outcomes; for example, the rate of coronary stent thrombosis with premature discontinuation of dual anti-platelet therapy. He concludes: 'Medication non-adherence is a widespread and pervasive problem across all disease states and one that spans the breadth of health delivery systems across the globe. It is time for the medication adherence paradigm to shift and be redefined in a place where patients and their providers share a mutual interest in achieving better adherence. Though patients may bear the final responsibility, it is incumbent on us as providers to create

Correspondence to Professor Catherine M Otto, Division of Cardiology, University of Washington, Box 356422, Seattle, WA 98195, USA; cmotto@uw.edu



**Figure 1** Weekly rates of adherence for the drugs considered for the three cohorts. ACEI, ACE inhibitor; ARB, angiotensin II receptor blocker.<sup>1</sup>

systems that place patients in the best position to advocate for their own health. Ultimately, solutions to meet this complex challenge need to be creative, pragmatic and scalable.'

Previous research on out-of-hospital cardiac arrest (OHCA) has focused on predictors of survival to hospital discharge with a paucity of data on factors predicting longer term survival. In this issue of Heart, Andrew and colleagues<sup>3</sup> report a mean survival duration of about 12 years in 3449 patients who survived to hospital discharge after OHCA over a 14 year period in Australia (figure 2). For the first year after OHCA, mortality was 5.6 times higher than expected but was similar to the general population 5 years post-OHCA. The only peri-arrest factor associated with long term survival was transport to a centre capable of percutaneous coronary intervention. Instead, in these patients who survived the initial OHCA event, long term survival was more related to post-discharge return to work and a favourable physical/functional recovery.

Kragholm and Torp-Pedersen<sup>4</sup> suggest that these findings highlight our current knowledge gaps in survivors of OCHA. 'First, further insight into factors that are important for obtaining long survival are necessary to improve the outlook for these patients. Second, the marked result that survival quickly approached the background population should inspire to studies of whether there is early depletion of a high-risk group of patients such that few interventions are necessary after a certain time. Third, and finally, because a larger number of people survive out-of-hospital cardiac arrest and remain alive in long-term outcome assessments, further insights are needed into the longterm quality of life and other indicators of functional status.'

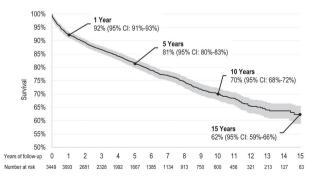


Figure 2 Kaplan-Meier survival estimates of out-of-hospital cardiac arrest survivors to hospital discharge.<sup>3</sup>



## Device oriented composite events

	BVS	5	DES	5		Odds Ratio	Odds Ratio
Study or Subgroup	<b>Events</b>	Total	<b>Events</b>	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI
3.3.1 Randomized							
ABSORB China	10	237	11	237	18.2%	0.91 [0.38, 2.17]	<del></del>
ABSORB II	25	335	7	166	18.9%	1.83 [0.78, 4.33]	<del></del>
ABSORB Japan Subtotal (95% CI)	19	261 833	5	130 533	13.7% 50.9%	1.96 [0.72, 5.38] 1.45 [0.86, 2.45]	•
Total events	54		23				
Heterogeneity: Tau <sup>2</sup> = Test for overall effect:				P = 0.4	2); I² = 0%	6	
3.3.2 Observational							
BVS-EXAMINATION	17	274	14	290	26.4%	1.30 [0.63, 2.70]	
Imori	20	215	10	215	22.7%	2.10 [0.96, 4.61]	-
Subtotal (95% CI)		489		505	49.1%	1.63 [0.95, 2.77]	•
Total events	37		24				
Heterogeneity: Tau <sup>2</sup> = Test for overall effect:				P = 0.3	8); I² = 0%		
Total (95% CI)		1322		1038	100.0%	1.53 [1.06, 2.23]	•
Total events Heterogeneity: Tau <sup>2</sup> = Test for overall effect: Test for subgroup diff	Z = 2.24	(P = 0.0)	12)				0.01 0.1 10 100 Favours (BVS) Favours (DES)

## All-cause death

	BVS		DES			Odds Ratio	Odds Ratio	
Study or Subgroup	<b>Events</b>	Total	<b>Events</b>	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI	
3.1.1 Randomized								
ABSORB China	1	237	6	237	22.9%	0.16 [0.02, 1.37]		
ABSORB II	4	335	1	166	21.8%	1.99 [0.22, 17.98]		
ABSORB Japan Subtotal (95% CI)	4	261 833	0	130 533	13.8% 58.4%	4.56 [0.24, 85.36] 0.98 [0.13, 7.19]		
Total events	9	-	7		001111	0.00 [0.10, 1110]		
Heterogeneity: Tau2 =	1.60; Chi	$i^2 = 4.1$	6, df = 2 (	P = 0.1	2); $I^2 = 52$	%		
Test for overall effect:	Z = 0.02	(P = 0.9)	39)					
3.1.2 Observational								
Imori Subtotal (95% CI)	4	215 <b>215</b>	5	215 215	41.6% 41.6%	0.80 [0.21, 3.01] 0.80 [0.21, 3.01]		
Total events	4	213	5	213	41.0%	0.00 [0.2 1, 5.0 1]		
Heterogeneity: Not ap	plicable							
Test for overall effect:		(P = 0.7)	74)					
Total (95% CI)		1048		748	100.0%	0.86 [0.26, 2.81]		
Total events	13		12					
Heterogeneity: Tau2=	0.42; Chi	$i^2 = 4.1$	7, df = 3 (	P = 0.2	$(4); I^2 = 28$	%	0.01 0.1 1 10 100	
Test for overall effect:	Z = 0.25	(P = 0.8)	30)		1.510)			
Test for subgroup diffe	erences:	Chi <sup>2</sup> =	0.03, df=	1 (P=	$0.86$ ), $I^2 =$	0%	Favours [BVS] Favours [DES]	

**Figure 3** Long-term outcomes in coronary artery disease (CAD) with bioresorbable vascular scaffold (BVS) versus drug-eluting stent (DES).<sup>5</sup>

The best choice for a coronary stent in patients with coronary disease, ranging from silent ischemia to acute coronary syndrome, remains controversial. Nairooz and colleagues<sup>5</sup> performed a meta-analysis outcomes at 2 years in studies comparing bioresorbable vascular scaffolds (BVS) to second generation drug eluting stents (DES). In 2360 patients from five trials,

this meta-analysis showed that that the device-oriented composite endpoint (cardiac mortality and target vessel myocardial infarction or ischemia-driven revascularisation) was lower with DES compared with BVS, although there was no difference in all-cause mortality (figure 3).

In the accompanying editorial, Brugaletta and Sabate<sup>6</sup> raise the concern

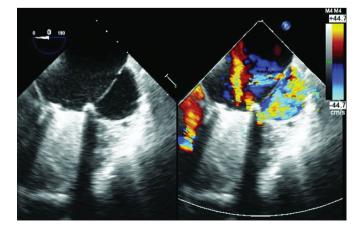


Figure 4

about BVS that 'the present meta-analysis represents a first attempt to understand the real incidence of scaffold thrombosis (ST) problem beyond 1 year after implantation, but many questions remain unanswered. As doctors having the safety of the patient as objective and as scientists with a close critical scrutiny on the clinical performance of these novel devices, we need to understand when ST may occur more frequently, which are the causes, how and if we can solve it.'

The Education in Heart article in this issue provides a summary of the literature, current guidelines and a practical algorithm for selecting the a novel oral anticoagulant in patients with atrial fibrillation. The Image Challenge question asks you to make the diagnosis from an echocardiographic image (figure 4); the online videos are helpful in understanding the anatomy and physiology of this rare complication after mitral valve surgery.

Competing interests None declared.

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