

Real-life clinical outcomes with the use of an ultrathin sirolimus-eluting stent in Sweden: A report from the Swedish coronary angiography and angioplasty registry

Sergio Buccheri, MD, Giovanna Sarno, MD, PhD, David Erlinge, MD, PhD, Bo Lagerqvist, MD, PhD, Per Grimfjärd, MD, Nils Witt, MD, PhD, Ole Fröbert, MD, PhD, Jonas Persson, MD, PhD, Felix Böhm, MD, PhD, Stefan James, MD, PhD



SCAAR registry

We sought to assess the performance of a **sirolimus-eluting ultrathin DES** (Orsiro, Biotronik AG, Bülach) in a large cohort of consecutive patients undergoing PCI in Sweden.

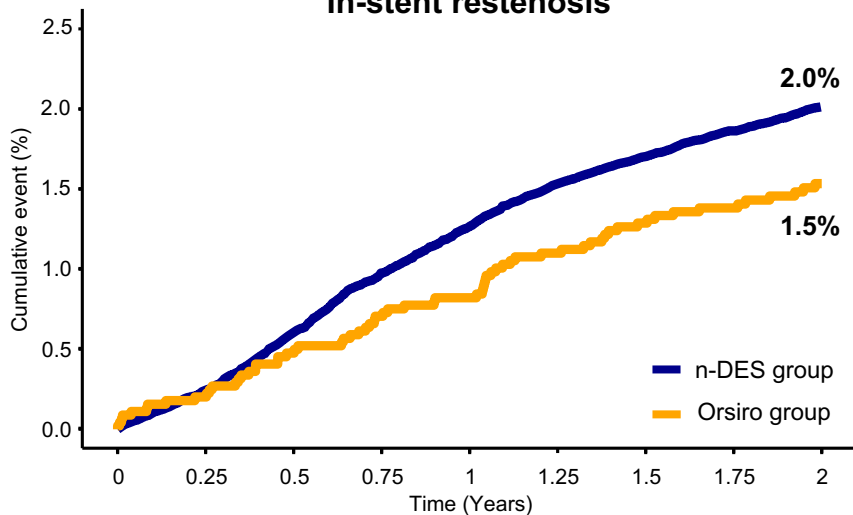
All patients in this study were registered in the **Swedish Coronary Angiography and Angioplasty Registry (SCAAR)**.

SCAAR is a nationwide, prospective, multicenter registry which collects the clinical and procedural characteristics of *all* consecutive patients undergoing PCI in Sweden.

Outcomes of interest: all-cause death, myocardial infarction, definite stent thrombosis, clinically relevant restenosis and target lesion revascularization by PCI up to two years.

What are the essential results?

In-stent restenosis



Numbers at risk

N-DES group 69,570 66,992 66,245 65,539 64,926 64,302 63,754 61,424 57,914

Orsiro group 4,561 4,378 4,342 4,305 4,278 4,227 4,189 4,036 3,741

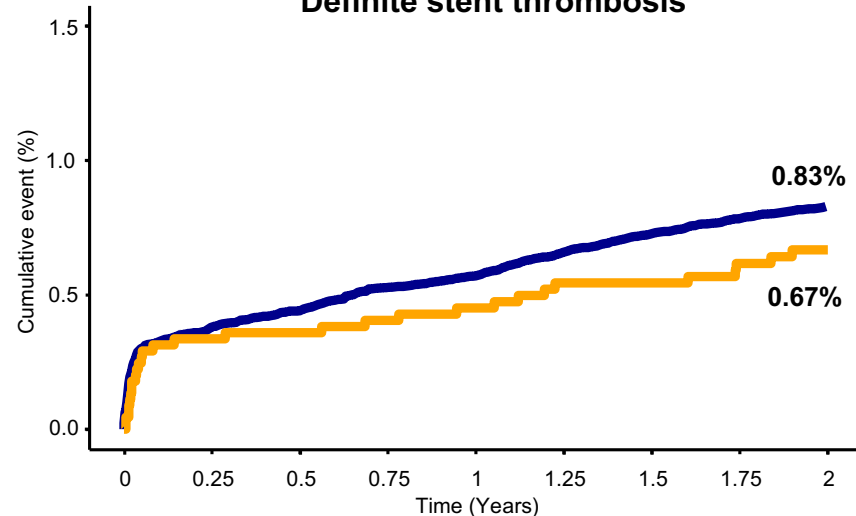
Cumulative number of events

N-DES group 3 164 408 653 846 1,021 1,131 1,234 1,325

Orsiro group 1 9 21 31 36 48 56 60 66

Adjusted HR 0.81, 95% CI 0.63-1.03; p=0.09

Definite stent thrombosis



Numbers at risk

N-DES group 69,570 67,101 66,510 65,933 65,418 64,883 64,379 62,070 58,578

Orsiro group 4,561 4,385 4,357 4,326 4,300 4,253 4,219 4,066 3,768

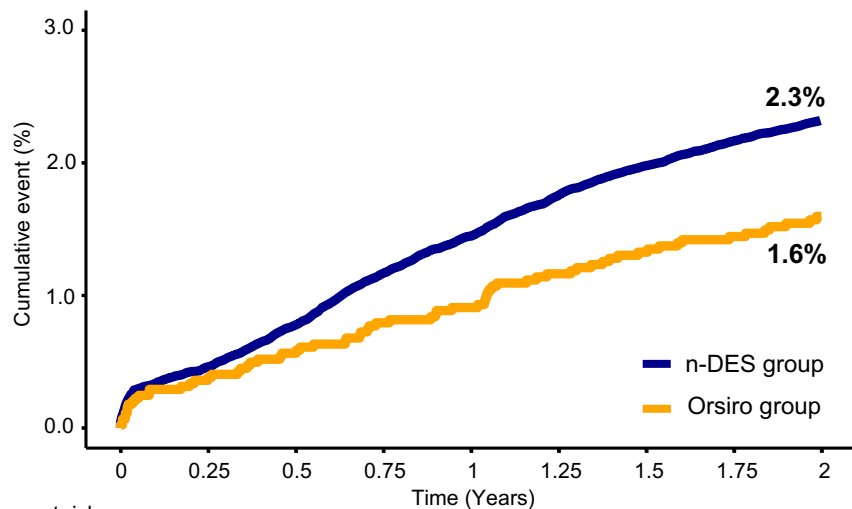
Cumulative number of events

N-DES group 15 260 301 359 387 445 490 527 553

Orsiro group 0 15 16 18 20 24 24 27 29

Adjusted HR 0.90, 95% CI 0.55-1.46; p=0.66

Target lesion revascularization by PCI



Adjusted HR 0.75, 95% CI 0.60-0.94; p=0.013

Numbers at risk

N-DES group 69,570 67,082 66,403 65,729 65,150 64,548 64,007 61,684 58,183

Orsiro group 4,561 4,384 4,354 4,319 4,294 4,249 4,212 4,059 3,759

Cumulative number of events

N-DES group 25 316 528 788 974 1,179 1,323 1,444 1,538

Orsiro group 1 17 25 35 40 51 58 63 69

To date, there is a paucity of data regarding the performance of **ultrathin DES in the real world**. Moreover, clinical outcomes beyond one year have not been systematically collected.

In this large nationwide analysis with a follow-up extended up two years, **Orsiro, a sirolimus-eluting ultrathin DES** was associated with low rates of definite stent thrombosis, numerically lower rates of in-stent restenosis and a significantly lower risk of target lesion revascularization as compared to other n-DES.

These findings **corroborate the results of randomized clinical trials** showing a potential incremental clinical benefit with ultra-thin metallic DES use during PCI.

Why?

There is a paucity of data regarding the performance of ultrathin DES platforms in the real world.

What?

This was a nationwide, registry-based analysis looking at the comparative performance of modern DES in the real world.

How?

We included a large cohort of consecutive patients undergoing PCI with modern generation DES in Sweden.

What are the results?

After addressing confounding, an ultrathin sirolimus-eluting DES use during PCI was associated with low rates of definite ST, restenosis and a significantly lower risk of target lesion revascularization by PCI.

Why is this important?

Our results corroborate the emerging evidence showing incremental benefits with ultrathin DES use during PCI.