

+ Evidence in focus

POLAR3[◇] Total Hip Solution

POLARSTEM[◇]/R3[◇]/OXINIUM[◇] - Primary THA

Summarised in this PRISM asset and evidence grids:

25
publications*

>129,000
total patients studied†



- 1 randomised controlled trial (RCT)
- 1 prospective comparative study
- 5 registry reports
- 4 registry analyses
- 4 retrospective comparative studies
- 8 case series/single-arm cohort
- 2 cost/efficiency analyses

See how **RI.HIP NAVIGATION** reduces revision risk compared to conventional THA by clicking or scanning the QR code:

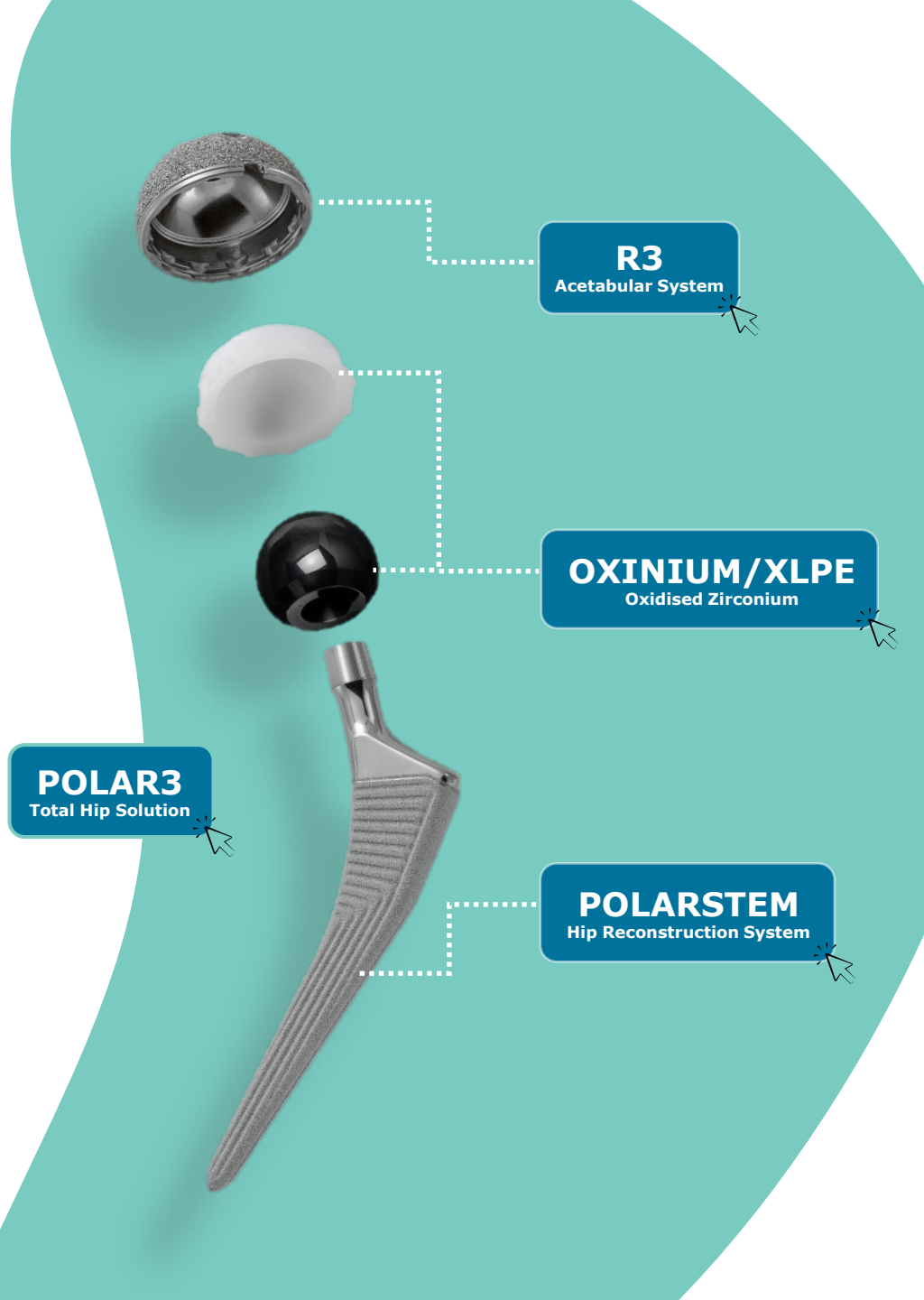


Smith+Nephew

*Including registry reports.

†Patients from included studies who received POLAR3 or any POLAR3 component.

Abbreviations: RCT = randomised controlled trial; THA = total hip arthroplasty.





POLAR3 ♦
Total Hip Solution
Product summary

Access the POLAR3 clinical evidence summary by clicking or scanning the QR code:

6 studies reporting on POLAR3

10 years

Excellent survivorship shown by real-world evidence in the UK NJR*¹

98.1% survivorship

53.4% lower revision risk than class average

POLARSTEM ♦/R3 ♦ is the **second most implanted** cementless THA prosthesis in the UK NJR

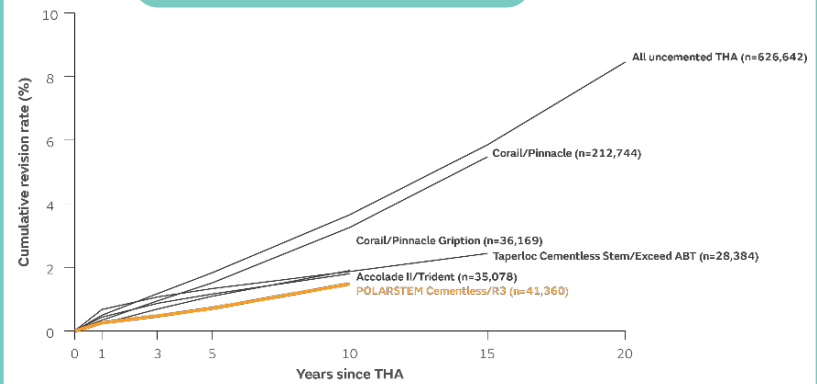


Figure. Cumulative revision of top five most implanted cementless THA prostheses

Improved patient outcomes versus all other stems in the UK NJR²

Significantly improved 6-month PROMs ($p < 0.001$)

Measure	Other THAs	POLAR3
EQ-VAS	66.6%	73.7%
EQ-5D	89.7%	91.6%
OHS	97.3%	98.3%

% improvement (pre-op vs 6 months)

Significantly higher patient satisfaction ($p < 0.001$)

Lower risk of revision due to aseptic loosening[†]

Reduced operating costs and length of stay versus other THAs³

In a propensity score matched study of patients from the Premier PINC AI Healthcare database (n=6,448):

\$1,121
Lower cost of stay
(\$16,760 vs \$17,881; $p < 0.001$)

44%
Higher odds of being discharged home
(92.2% vs 89.1%, OR 1.44; $p < 0.001$)

34%
Lower risk of 30-day readmission
(1.9% vs 2.9%, OR 0.66; $p < 0.001$)

15%
Lower length of stay
(1.52 vs 1.78 days; $p < 0.001$)

*POLARSTEM Cementless/R3 only. †Compared to expected revisions, probability of lower revision risk: > 0.999 .
Abbreviations: EQ-5D = EuroQol 5D; EQ-VAS = EuroQol Visual Analogue Scale; NJR = National Joint Registry; OHS = Oxford Hip Score; OR = odds ratio; PROMs = patient reported outcome measures; THA = total hip arthroplasty.



POLARSTEM[◇]

Hip Reconstruction System

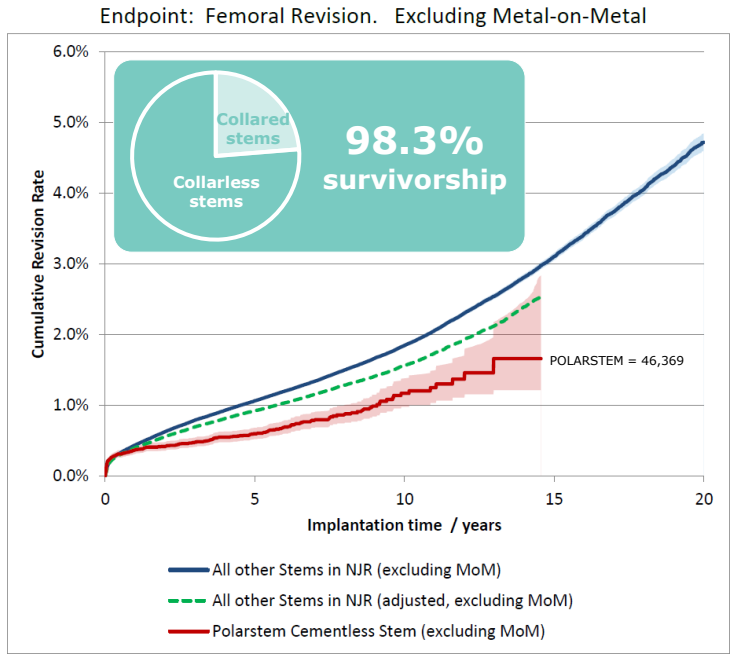
Product summary

Access the POLARSTEM clinical evidence summary by clicking or scanning the QR code:

9 studies reporting on POLARSTEM

Excellent survivorship shown by real-world evidence in the UK NJR⁴

30% lower revision risk than all other stems (HR 0.70; p<0.001)[†]



Improved patient outcomes versus all other stems in the UK NJR (p<0.001)⁴

Significantly improved 6-month PROMS

Measure	All other stems	POLARSTEM Cementless
EQ-VAS	66.6%	72.7%
EQ-5D	89.7%	91.7%
OHS	97.3%	98.1%

% improvement (pre-op vs 6 months)

Significantly higher patient satisfaction

Excellent clinical performance with triple-taper design

Lower risk of revision due to aseptic loosening^{‡4}

Excellent survivorship with both collared and collarless stems^{§5}

ODEP 10A*⁶
POLARSTEM Cementless

ODEP 7A⁶
POLARSTEM Cemented

[†]HR varies with time. Adjusted for age, sex, ASA, BMI, Op.date and indications. [‡]Compared to expected revisions, probability of lower revision risk: >0.999.
[§]No significant difference in 3-year revision rate with POLARSTEM Collared vs Collarless in the AOANJRR (98% vs 97.4%; p=0.409).
Abbreviations: EQ-5D = EuroQol 5D; EQ-VAS = EuroQol Visual Analogue Scale; HR = hazard ratio; MoM = metal-on-metal; NJR = National Joint Registry; OHS = Oxford Hip Score; PROMS = patient reported outcome measures; THA = total hip arthroplasty.



R3
Acetabular System
Product summary

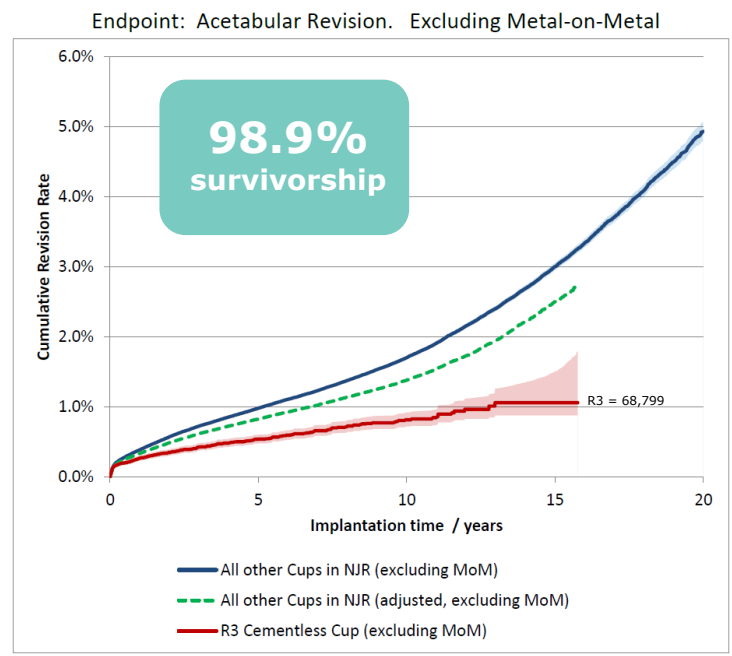
Access the R3 clinical evidence summary by clicking or scanning the QR code:

8 studies reporting on R3

Excellent survivorship shown by real-world evidence in the UK NJR⁷



39% lower revision risk than all other cups
(HR 0.61; p<0.001)[†]



Improved patient outcomes versus all other cups in the UK NJR (p<0.001)⁷



Significantly improved 6-month PROMS

Measure	Other THAs	R3
EQ-VAS	66.6%	70.6%
EQ-5D	89.7%	90.7%
OHS	97.3%	97.7%

% improvement (pre-op vs 6 months)

Significantly higher patient satisfaction

Excellent clinical performance

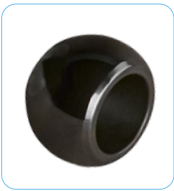
Lower risk of revision due to aseptic loosening or liner dissociation^{‡7}

Significantly lower liner dissociation rate compared with Pinnacle
(0% vs 1.12%; p=0.038)⁸

ODEP 15A^{*6}
R3 HA-coated

ODEP 10A⁶
R3 non-HA coated

[†]HR varies with time. Adjusted for age, sex, ASA, BMI, Op.date and indications. [‡]Compared to expected revisions, probability of lower revision risk: >0.999.
Abbreviations: EQ-5D = EuroQol 5D; EQ-VAS = EuroQol Visual Analogue Scale; HR = hazard ratio; MoM = metal-on-metal; NJR = National Joint Registry; OHS = Oxford Hip Score; PROMs = patient reported outcome measures; THA = total hip arthroplasty.



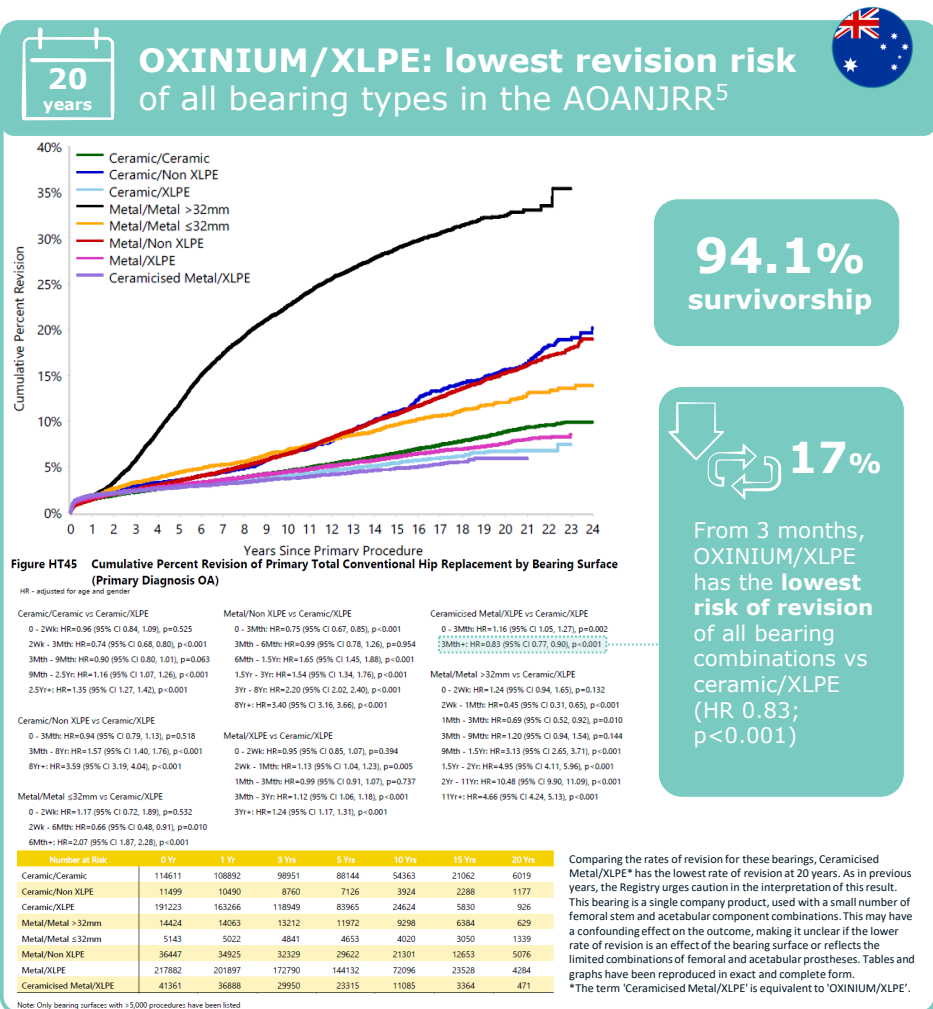
OXINIUM[®]/XLPE

Oxidised Zirconium

Product summary

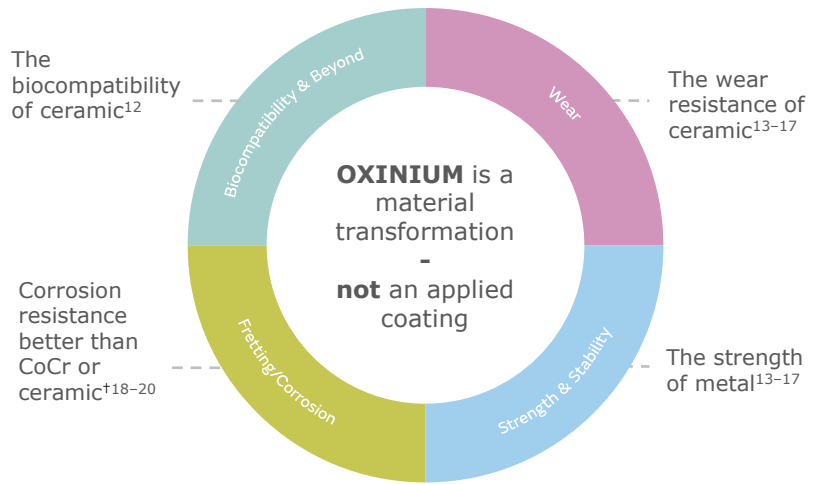
Access the OXINIUM clinical evidence summary by clicking or scanning the QR code:

6 studies reporting on OXINIUM



Data from four joint registries has shown OXINIUM heads have the **highest survivorship***5,9-11

During manufacturing, the zirconium alloy core undergoes a transformation upon heating to form a ~5µm thick ceramicised surface, offering a strong, resilient implant material.¹²



*vs all other reported bearing combinations; NJR data from 2004-2016; RIPO data from 2000-2015; LROI data from 2007-2016; AOANJRR 2025 annual report.
†Aldinger et al. (2017) shows OXINIUM demonstrates a markedly reduced susceptibility to taper fretting compared to CoCr and ceramic during in vitro testing.²⁰
Abbreviations: AOANJRR = Australian Orthopaedic Association National Joint Replacement Registry; CoCr = cobalt chrome; HR = hazard ratio; OA = osteoarthritis; XLPE = highly cross-linked polyethylene.

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*The data used for this analysis was obtained from the National Joint Registry ("NJR"), part of the HQIP, the NJR and/or its contractor, NEC Software Solutions (UK) Limited ("NEC") take no responsibility (except as prohibited by law) for the accuracy, currency, reliability and correctness of any data used or referred to in this report, nor for the accuracy, currency, reliability and correctness of links or references to other information sources and disclaims all warranties in relation to such data, links and references to the maximum extent permitted by legislation including any duty of care to third party readers of the data analysis.

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