

Artek Total Conventional Hip Investigation

Note: This analysis compares the Artek acetabular prosthesis with all other total conventional hip prostheses.

This prosthesis has been identified as having a significantly higher rate of revision. For a detailed explanation of the process used by the Registry that results in identification of prostheses that have a higher than anticipated rate of revision please refer to the Prostheses with Higher than Anticipated Rates of Revision chapter of the most recent AOANJRR Annual Report, <https://aoanjrr.sahmri.com/annual-reports-2026>.

Note: Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator. Procedures using prostheses with no recorded use in 2025 are excluded from the comparator.

TABLE 1

Revision Rate of Primary Total Conventional Hip Replacement

The revision rate of the Artek total conventional hip prosthesis is compared to all other total conventional hip prostheses.

Table 1: Revision Rates of Primary Total Conventional Hip Replacement

Component	N Revised	N Total	Obs. Years	Revisions/100 Obs. Yrs (95% CI)
Artek	77	179	2569	3.00 (2.37, 3.75)
Other Total Conventional Hip	19701	568612	3623071	0.54 (0.54, 0.55)
TOTAL	19778	568791	3625640	0.55 (0.54, 0.55)

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

TABLE 2

Yearly Cumulative Percent Revision of Primary Total Conventional Hip Replacement

The yearly cumulative percent revision of the Artek total conventional hip prosthesis is compared to all other total conventional hip prostheses.

Table 2: Yearly Cumulative Percent Revision (95% CI) of Primary Total Conventional Hip Replacement

CPR	1 Yr	2 Yrs	3 Yrs	4 Yrs	5 Yrs	6 Yrs
Artek	2.8 (1.2, 6.7)	5.7 (3.1, 10.3)	8.0 (4.8, 13.1)	12.6 (8.5, 18.5)	16.1 (11.4, 22.5)	20.3 (15.0, 27.1)
Other Total Conventional Hip	1.7 (1.7, 1.8)	2.1 (2.1, 2.2)	2.5 (2.4, 2.5)	2.8 (2.7, 2.8)	3.0 (3.0, 3.1)	3.3 (3.2, 3.3)

CPR	7 Yrs	8 Yrs	9 Yrs	10 Yrs	11 Yrs	12 Yrs
Artek	21.5 (16.1, 28.4)	22.1 (16.6, 29.1)	24.6 (18.8, 31.8)	27.1 (21.0, 34.4)	29.5 (23.3, 37.1)	33.5 (26.8, 41.2)
Other Total Conventional Hip	3.5 (3.5, 3.6)	3.8 (3.7, 3.9)	4.1 (4.0, 4.2)	4.4 (4.3, 4.5)	4.7 (4.6, 4.8)	5.1 (5.0, 5.2)

CPR	13 Yrs	14 Yrs	15 Yrs	16 Yrs	17 Yrs	18 Yrs
Artek	38.1 (31.2, 46.1)	38.8 (31.8, 46.8)	41.0 (33.8, 49.1)	41.0 (33.8, 49.1)	41.7 (34.5, 49.8)	43.3 (36.0, 51.5)
Other Total Conventional Hip	5.4 (5.3, 5.5)	5.8 (5.7, 5.9)	6.2 (6.1, 6.3)	6.6 (6.5, 6.7)	7.0 (6.8, 7.1)	7.3 (7.1, 7.5)

CPR	19 Yrs	20 Yrs	21 Yrs	22 Yrs	23 Yrs	24 Yrs	25 Yrs
Artek	43.3 (36.0, 51.5)	44.2 (36.8, 52.4)	45.1 (37.6, 53.4)	46.0 (38.5, 54.4)	47.9 (40.2, 56.4)	47.9 (40.2, 56.4)	
Other Total Conventional Hip	7.7 (7.5, 7.9)	8.0 (7.8, 8.2)	8.6 (8.3, 8.9)	9.0 (8.7, 9.4)	9.6 (9.1, 10.0)	9.6 (9.2, 10.1)	10.3 (9.4, 11.2)

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

FIGURE 1

Yearly Cumulative Percent Revision of Primary Total Conventional Hip Replacement

The yearly cumulative percent revision of the Artek total conventional hip prosthesis is compared to all other total conventional hip prostheses. In addition, hazard ratios are reported.

Hazard ratios are reported for specific time periods during which the hazard ratio is constant. This is done to enable more specific and valid comparisons of the risk of revision over time. The pattern of variation in risk has important implications with respect to the underlying reasons for any difference.

Figure 1: Cumulative Percent Revision of Primary Total Conventional Hip Replacement

***** Proportionality Not Achieved :-(*****

var1 VS var2 (Artek VS Other Total Conventional Hip) is non-proportional in time period 0_1Yr

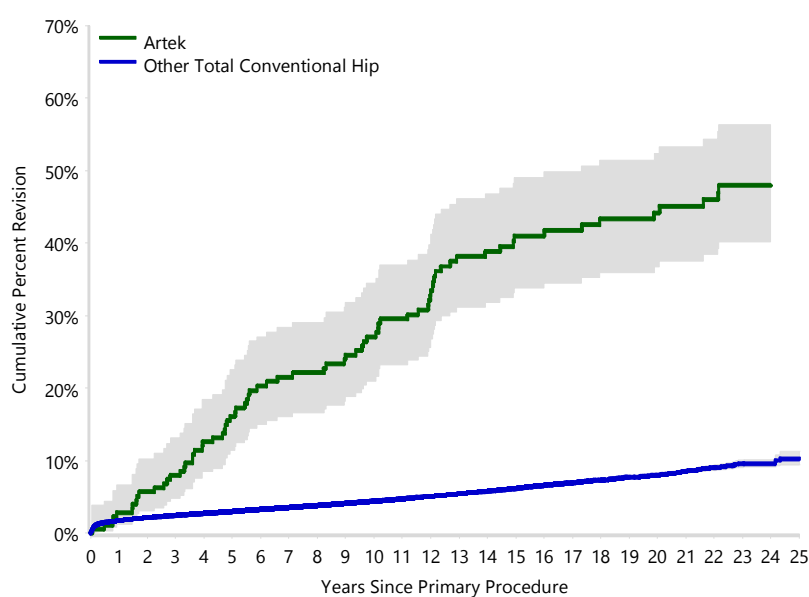
The algorithm is not able to select a time point. Please specify time periods manually

The model will be fit over this entire period instead

var	timepoint	ProbChiSq
var1	0_1Yr	0.0402
var1	1Yr_1p5Yr	0.1774
var1	1p5Yr_2p5Yr	0.2043
var1	2p5Yr_4Yr	0.4007
var1	4Yr_4p5Yr	0.6531
var1	4p5Yr_6Yr	0.6902
var1	6Yr_max	0.1307

The PHREG Procedure

Dummy Text



HR - adjusted for age and sex

Artek vs Other Total Conventional Hip

0 - 1Yr: HR=1.54 (95% CI 0.64, 3.71), p=0.331

1Yr - 1.5Yr: HR=4.96 (95% CI 1.24, 19.85), p=0.023

1.5Yr - 2.5Yr: HR=6.18 (95% CI 2.32, 16.47), p<0.001

2.5Yr - 4Yr: HR=15.25 (95% CI 8.43, 27.59), p<0.001

4Yr - 4.5Yr: HR=4.99 (95% CI 0.70, 35.48), p=0.108

4.5Yr - 6Yr: HR=19.34 (95% CI 10.95, 34.16), p<0.001

6Yr+: HR=6.80 (95% CI 4.98, 9.27), p<0.001

Number at Risk	0 Yr	1 Yr	2 Yrs	3 Yrs	4 Yrs	5 Yrs	6 Yrs	7 Yrs
Artek	179	171	165	160	151	141	133	130
Other Total Conventional Hip	568612	501603	445433	391564	344389	299354	259516	220948

Number at Risk	8 Yrs	9 Yrs	10 Yrs	11 Yrs	12 Yrs	13 Yrs	14 Yrs	15 Yrs	16 Yrs
Artek	126	122	118	111	100	92	87	81	80

Number at Risk	8 Yrs	9 Yrs	10 Yrs	11 Yrs	12 Yrs	13 Yrs	14 Yrs	15 Yrs	16 Yrs
Other Total Conventional Hip	185798	153739	124848	99575	79802	63343	49730	38033	28355

Number at Risk	17 Yrs	18 Yrs	19 Yrs	20 Yrs	21 Yrs	22 Yrs	23 Yrs	24 Yrs	25 Yrs
Artek	75	71	66	63	59	57	54	48	14
Other Total Conventional Hip	20437	14638	10400	7396	4859	2863	1455	560	95

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

TABLE 3**Primary Diagnosis for Revised Primary Total Conventional Hip Replacement**

This table identifies the diagnosis of the primary procedure which was subsequently revised. This information is provided as there is a variation on outcome depending on the primary diagnosis. It is therefore important when considering the reasons for a higher than anticipated rate of revision that there is identification of the primary diagnosis. This information should be compared to the primary diagnosis for the revisions of all other total conventional hip prostheses.

Table 3: Primary Diagnosis for Revised Primary Total Conventional Hip Replacement

Primary Diagnosis	Artek		Other Total Conventional Hip	
	Number	Percent	Number	Percent
Osteoarthritis	73	94.8	16296	82.7
Fractured Neck Of Femur			1521	7.7
Osteonecrosis	3	3.9	844	4.3
Developmental Dysplasia			321	1.6
Rheumatoid Arthritis	1	1.3	201	1.0
Failed Internal Fixation			165	0.8
Tumour			157	0.8
Other Inflammatory Arthritis			102	0.5
Fracture/Dislocation			60	0.3
Other			21	0.1
Arthrodesis Takedown			13	0.1
TOTAL	77	100.0	19701	100.0

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

TABLE 4**Reasons for Revision**

This is reported in two ways: a percentage of primary procedures revised and as a percentage of all revision procedures.

% Primaries Revised: This shows the proportional contribution of each revision diagnosis as a percentage of the total number of primary procedures. This percentage can be used to approximate the risk of being revised for that diagnosis. Differing percentages between groups, with the same distribution of follow up time, may identify problems of concern.

% Revisions: The number of revisions for each diagnosis is expressed as a percentage of the total number of revisions. This shows the distribution of reasons for revision within a group but cannot be used as a comparison between groups.

Table 4: Primary Total Conventional Hip Replacement - Reason for Revision

Revision Diagnosis	Number	Artek		Other Total Conventional Hip		
		% Primaries Revised	% Revisions	Number	% Primaries Revised	% Revisions
Infection	9	5.0	11.7	5017	0.9	25.5
Prosthesis Dislocation/Instability	7	3.9	9.1	4382	0.8	22.2
Fracture	1	0.6	1.3	4349	0.8	22.1
Loosening	33	18.4	42.9	3629	0.6	18.4
Pain	2	1.1	2.6	340	0.1	1.7
Leg Length Discrepancy	2	1.1	2.6	307	0.1	1.6
Malposition				282	0.0	1.4
Implant Breakage Stem				211	0.0	1.1
Lysis	7	3.9	9.1	200	0.0	1.0
Implant Breakage Acetabular Insert				128	0.0	0.6
Wear Acetabular Insert				112	0.0	0.6
Incorrect Sizing				96	0.0	0.5
Metal Related Pathology	16	8.9	20.8	93	0.0	0.5
Implant Breakage Acetabular				73	0.0	0.4
Wear Head				49	0.0	0.2
Tumour				46	0.0	0.2
Heterotopic Bone				26	0.0	0.1
Implant Breakage Head				26	0.0	0.1
Wear Acetabulum				6	0.0	0.0
Osteonecrosis				4	0.0	0.0
Synovitis				2	0.0	0.0
Progression Of Disease				1	0.0	0.0
Other				322	0.1	1.6
N Revision	77	43.0	100.0	19701	3.5	100.0
N Primary	179			568612		

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

FIGURE 2**Cumulative Incidence Revision Diagnosis of Primary Total Conventional Hip Replacement**

This figure details the cumulative incidence of the most common reasons for revision. The five most common reasons for revision are included as long as each of these reasons account for more than 10 procedures or at least 5% of all revisions for the Artek total conventional hip prosthesis. A comparative graph is provided of the cumulative incidence for the same reasons for revisions for all other total conventional hip prostheses.

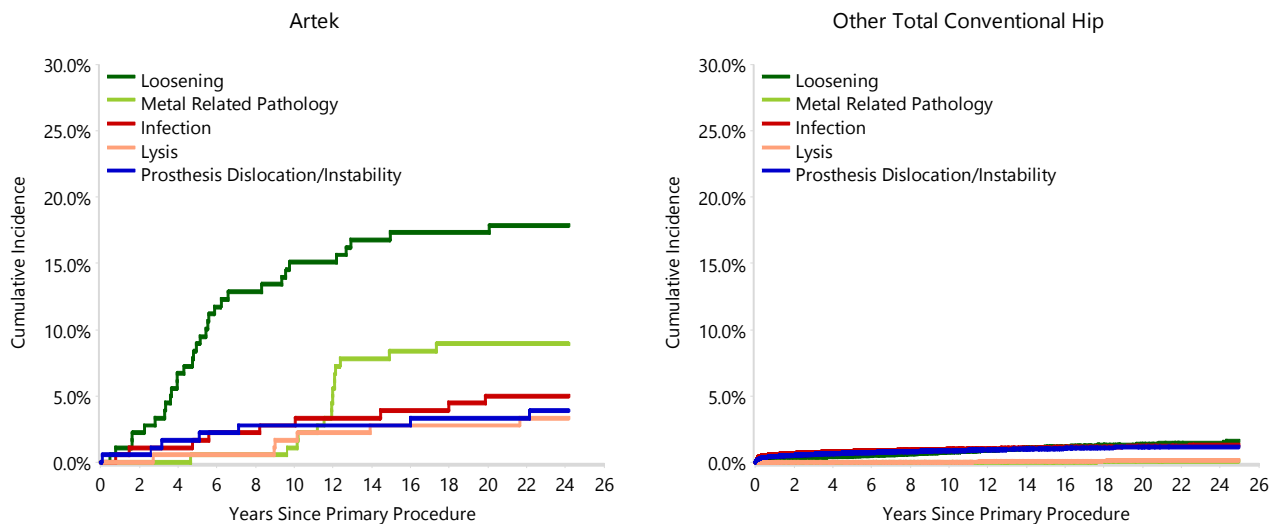
Figure 2: Cumulative Incidence Revision Diagnosis for Primary Total Conventional Hip Replacement

TABLE 5

Type of Revision Performed for Primary Total Conventional Hip Replacement

This analysis identifies the components used in the revision of the Artek total conventional hip prosthesis and compares it to the components used in the revision of all other total conventional hip prostheses.

The reason this analysis is undertaken is to identify whether there is one or more components which are being replaced that differ from the components replaced for revisions of all other total conventional hip prostheses i.e. is there a difference in the type of revision undertaken for the Artek total conventional hip prosthesis compared to all other total conventional hip prostheses.

Table 5: Primary Total Conventional Hip Replacement - Type of Revision

Type of Revision	Artek		Other Total Conventional Hip	
	Number	Percent	Number	Percent
Femoral Component	1	1.3	6645	33.7
Acetabular Component	62	80.5	3311	16.8
THR (Femoral/Acetabular)	8	10.4	2305	11.7
Cement Spacer	3	3.9	556	2.8
Removal of Prostheses			93	0.5
Reinsertion of Components			27	0.1
Total Femoral			12	0.1
Bipolar Head and Femoral			10	0.1
N Major	74	96.1	12959	65.8
Head/Insert	1	1.3	5372	27.3
Head Only	2	2.6	914	4.6
Minor Components			285	1.4
Insert Only			168	0.9
Bipolar Only			1	0.0
Cement Only			1	0.0
Head/Neck/Insert			1	0.0
N Minor	3	3.9	6742	34.2
TOTAL	77	100.0	19701	100.0

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

TABLE 6**Revision Rates of Artek Primary Total Conventional Hip Replacement by Fixation**

This analysis is provided as some prostheses have more than one fixation option. Additionally there are prostheses where an alternative to the recommended approach to fixation was used e.g. a cementless prosthesis that has been cemented or vice-versa.

Table 6: Revised Number of Artek Primary Total Conventional Hip Replacement by Fixation

Fixation	N Revised	N Total
Cementless	70	160
Hybrid (Femur Cemented)	6	17
Reverse Hybrid (Femur Cementless)	1	2
TOTAL	77	179

TABLE 7**Revision Rates of Artek Primary Total Conventional Hip Replacement by Bearing Surface**

This analysis is provided as some prostheses are combined with a variety of bearing surfaces. All bearing surfaces used with this prosthesis are listed.

Table 7: Revised Number of Artek Primary Total Conventional Hip Replacement by Bearing Surface

Bearing Surface	N Revised	N Total
Metal/Metal	77	179
TOTAL	77	179

TABLE 8

Revision Rates of Primary Total Conventional Hip Replacement by State

This enables a state by state variation to be identified for the Artek total conventional hip prosthesis and provides the comparative data for each of the states for all other total conventional hip prostheses.

The purpose of this analysis is to determine if the higher than anticipated rate of revision has widespread distribution between states. If there is widespread distribution then the reason for the higher than anticipated rate of revision is unlikely to be surgeon specific. If the prosthesis has been used in only a small number of states it is not possible to distinguish if the higher than anticipated rate of revision is related to the prosthesis, surgeon, technique or patient.

Table 8: Revised Number of Primary Total Conventional Hip Replacement by State

Component	State	N Revised	N Total
Artek	NSW	9	26
	VIC	35	79
	QLD	3	7
	SA	30	66
	ACT/NT	0	1
Other Total Conventional Hip	NSW	5507	167002
	VIC	4853	146558
	QLD	3968	104359
	WA	2510	64946
	SA	1887	52084
	TAS	439	18269
	ACT/NT	537	15394
TOTAL		19778	568791

Note: Prostheses no longer used in 2025 are excluded from the comparator. Procedures using metal/metal prostheses with head size larger than 32mm are excluded from the comparator.

TABLE 9**Number of Revisions of Artek Primary Total Conventional Hip Replacement by Year of Implant**

This analysis details the number of prostheses reported each year to the Registry for the Artek total conventional hip prosthesis. It also provides the subsequent number of revisions of the primaries reported in that year.

Primary procedures performed in later years have had less follow up time therefore the number revised is expected to be less than the number revised in earlier years. For example, a primary procedure performed in 2025 has a maximum of one year to be revised, whereas a primary procedure performed in 2023 has a maximum of three years to be revised.

Table 9: Number of Revisions of Artek Primary Total Conventional Hip Replacement by Year of Implant

Year of Implant	Number Revised	Total Number
1999	7	12
2000	19	34
2001	43	111
2002	8	22
TOTAL	77	179

TABLE 10

Revision Rates of Artek Primary Total Conventional Hip Replacement by Catalogue Number Range

Many prostheses have a number of catalogue ranges. The catalogue range is specific to particular design features; more than one catalogue range usually indicates a minor difference in design in a particular Artek prosthesis.

This analysis has been undertaken to determine if the revision rate varies according to the catalogue number range.

Model	Catalogue Range	Catalogue Description	Cement	Material
Acetabular				
Artek	91362854-91363858	ARTEK CUP UNCEMENTED METASUL (METAL)	NO	METAL

Table 10: Revised Number of Artek Primary Total Conventional Hip Replacement by Catalogue Number Range

Acetabular Range	N Revised	N Total
91362854-91363858	77	179
TOTAL	77	179

TABLE 11**Revision Rates of Artek Primary Total Conventional Hip Replacement by Component**

A prosthesis may be combined with multiple components. This analysis has been undertaken to determine if the revision rate varies according to the component with which it is combined.

Table 11: Revised Number of Artek Primary Total Conventional Hip Replacement by Femoral Stem Component

Femoral Stem Component	N Revised	N Total
APR	22	39
Alloclassic	29	68
CLS	5	16
MS 30	2	3
Natural Hip	19	53
TOTAL	77	179