



# Univers™ Apex Outcomes

## PURPOSE

To report the clinical outcomes of pain, function and quality of life for patients who have undergone total shoulder arthroplasty utilizing Univers Apex prosthesis.

## METHODS

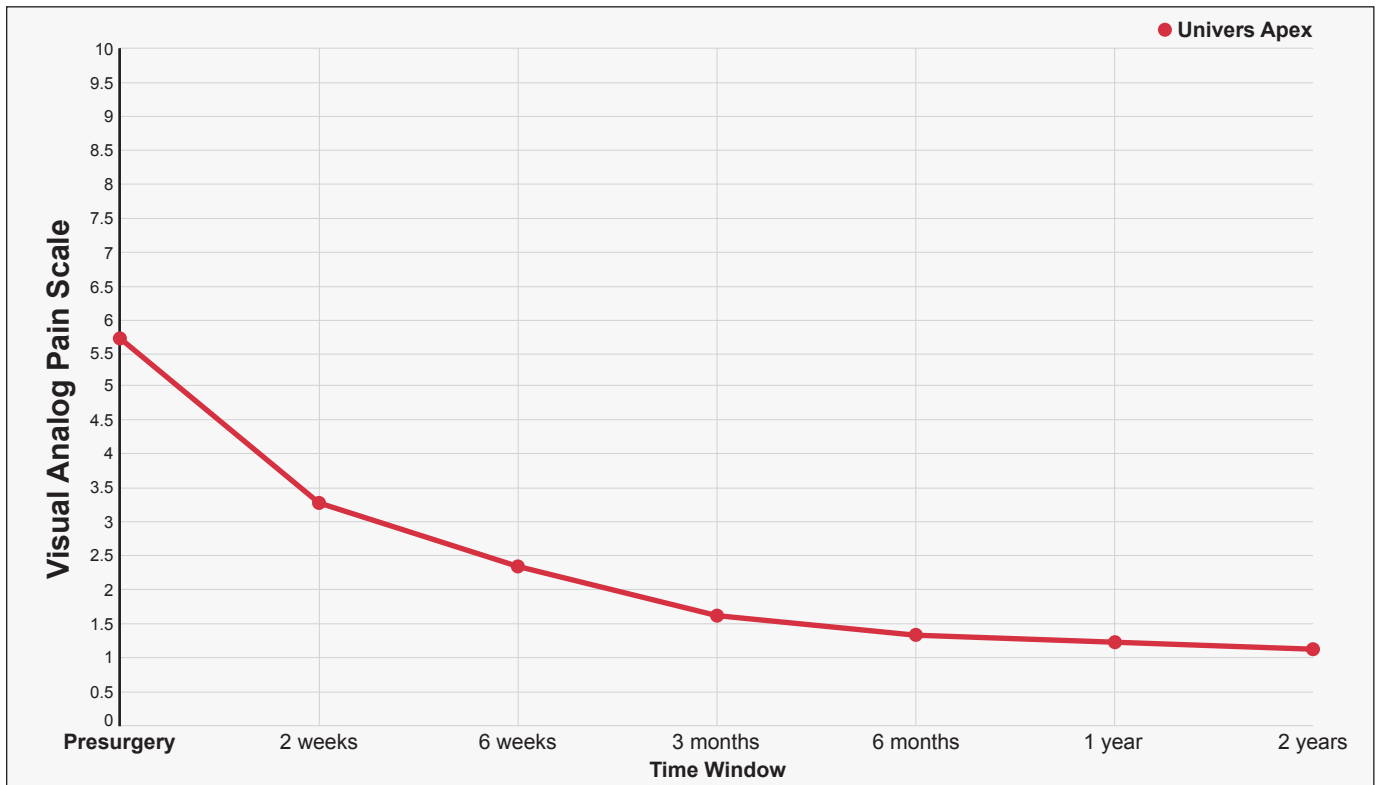
Patients enrolled in the Surgical Outcomes System™ global registry who underwent TSA utilizing Univers Apex prosthesis. Standard patient-reported outcomes questionnaires for VAS, ASES Shoulder Function and Index and SANE were administered at standard time points postoperatively. Results were reported from presurgery out to 2 years postsurgery. The number of patients included per time point is shown to the right.

Time Point	# of Compliant Univers Apex Patients/ Total Patients
Presurgery	354/479
1 year	235/317
2 years	138/198

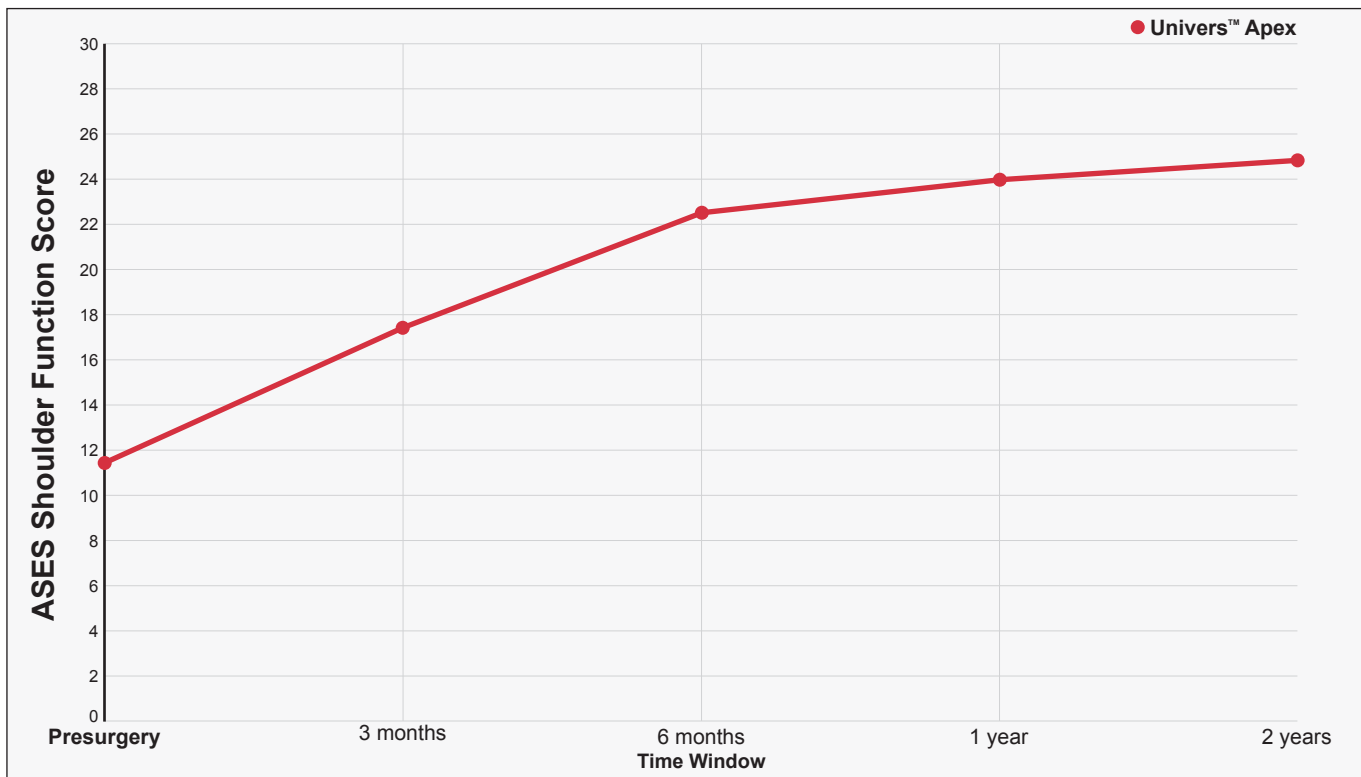
## TREND CONCLUSION

Based on these results, the pain, function and quality-of-life scores for patients undergoing total shoulder arthroplasty with the Univers Apex prosthesis trend towards favorable outcomes. However, no claims can be made on the potential of these results without further statistical analysis to determine statistical significance.

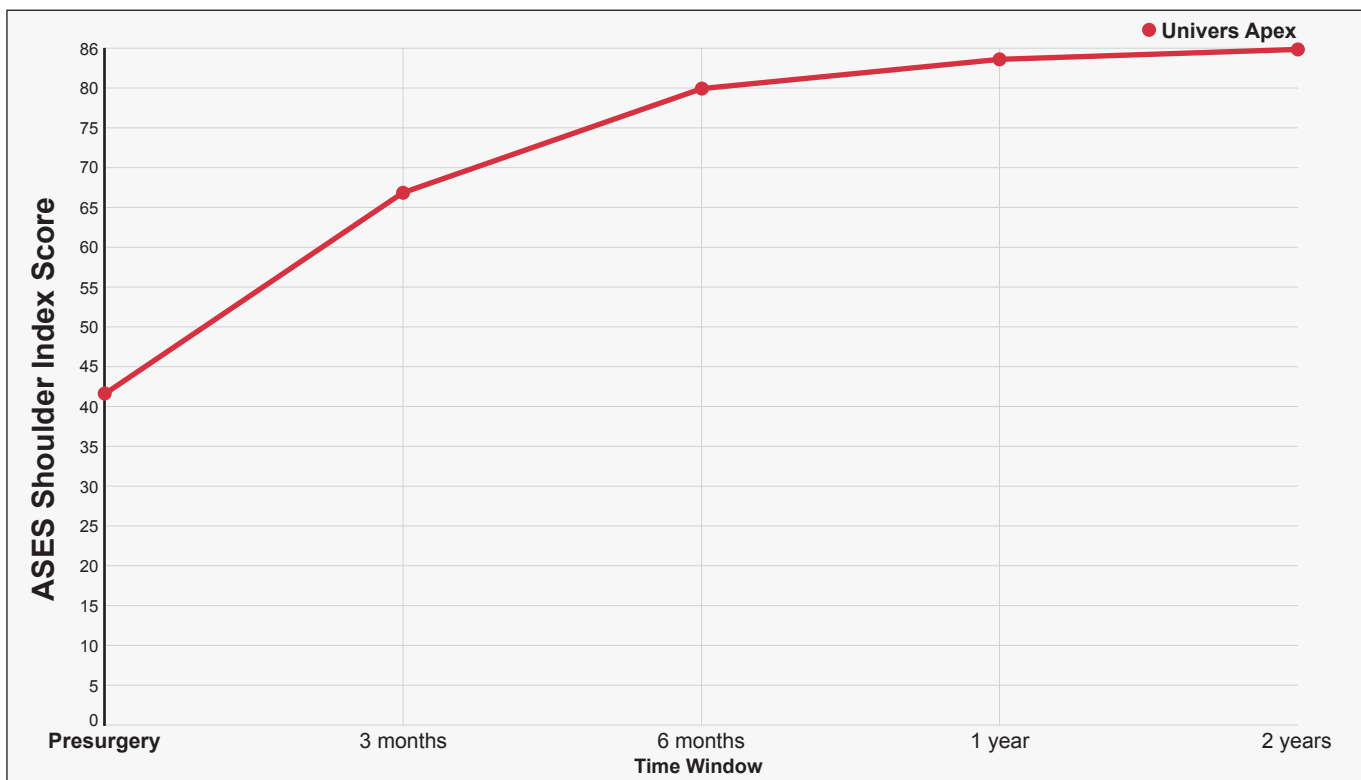
## RESULTS



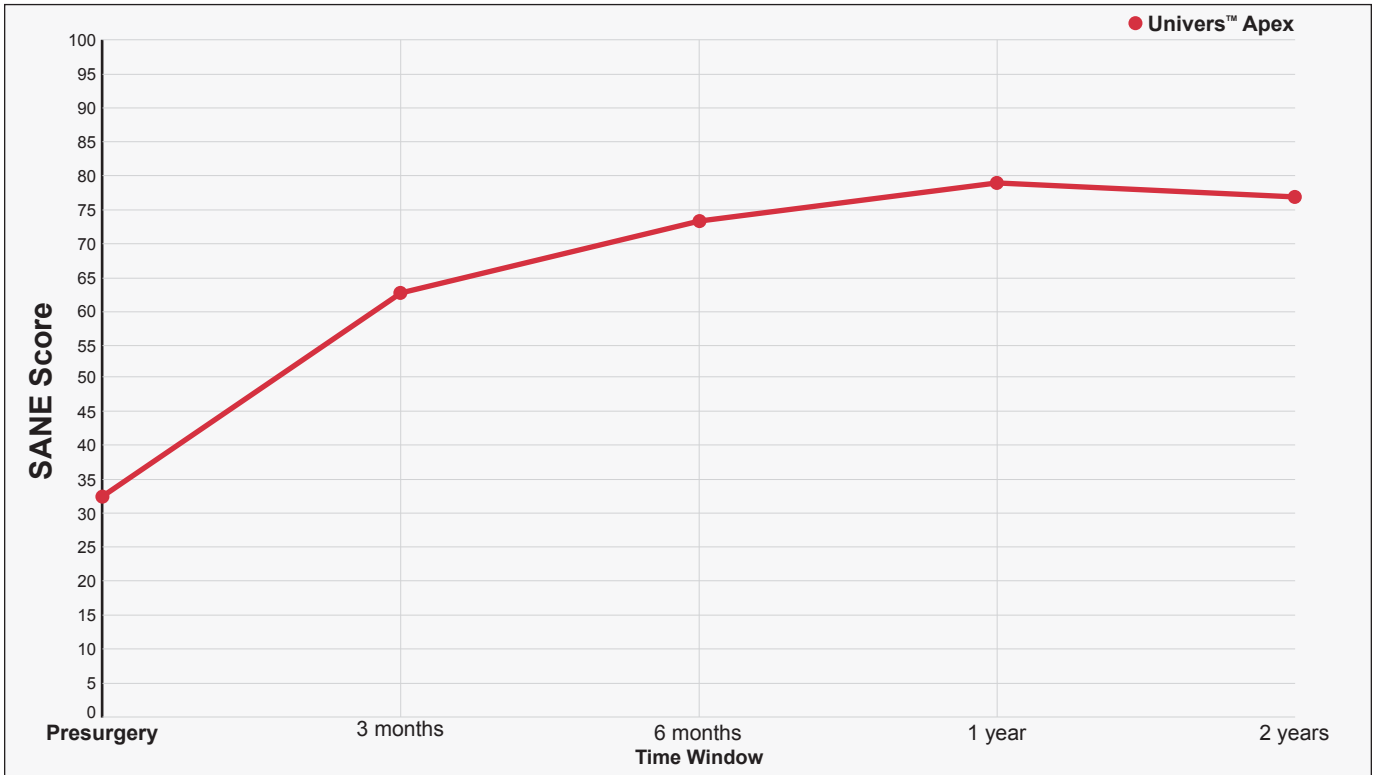
VAS



ASES Shoulder Function



ASES Shoulder Index



SANE

<b>Time Point</b>	<b>Univers™ Apex Avg +/- STD VAS</b>
Presurgery	5.7 +/- 2.3
1 year	1.3 +/- 2.0
2 years	1.1 +/- 1.7

<b>Time Point</b>	<b>Univers Apex Avg +/- STD ASES Function</b>
Presurgery	11.5 +/- 5.2
1 year	24.0 +/- 5.8
2 years	24.6 +/- 5.7

<b>Time Point</b>	<b>Univers Apex Avg +/- STD ASES Index</b>
Presurgery	40.7 +/- 16.6
1 year	83.6 +/- 17.5
2 years	85.6 +/- 16.3

<b>Time Point</b>	<b>Univers Apex Avg +/- STD SANE</b>
Presurgery	32.3 +/- 19.2
1 year	79.3 +/- 21.3
2 years	76.6 +/- 26.0