

Arthrex Distal Radius Plate Early Outcomes

Purpose

To report the clinical outcomes of pain, function, and quality of life for patients who underwent a distal radius / ulna fracture procedure with an Arthrex volar plate.

Methods

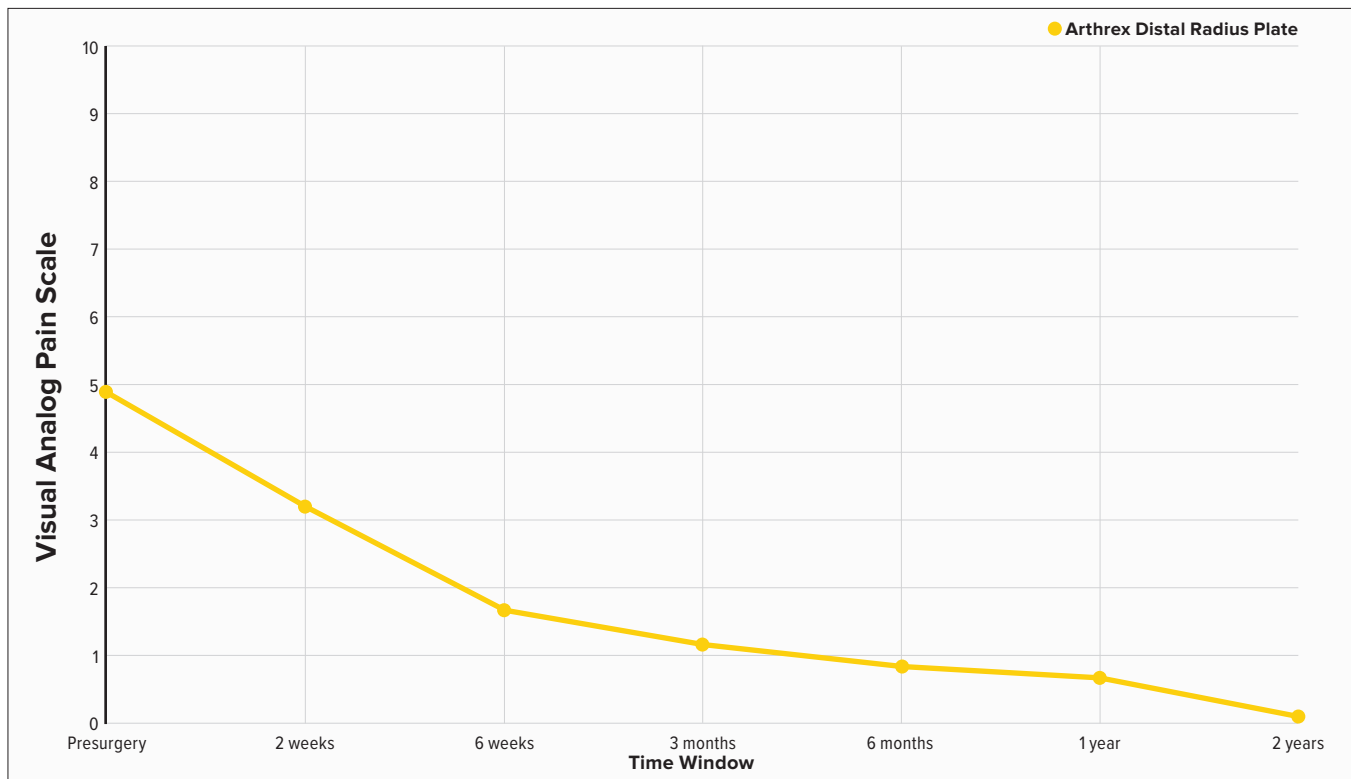
The inclusion criteria for this analysis were patients enrolled in the Surgical Outcomes System™ registry who underwent a distal radius /ulna fracture procedure with an Arthrex volar plate. Standard patient-reported outcomes questionnaires for VAS, VR-12, and QuickDash were administered at standard time points postoperatively. Results were reported from presurgery to 2 years postsurgery. The number of patients included per time point is shown to the right.

Time Point	# of Compliant Patients/ Total # of Patients
Presurgery	27/48
6 months	28/47
1 year	20/42
2 years	10/19

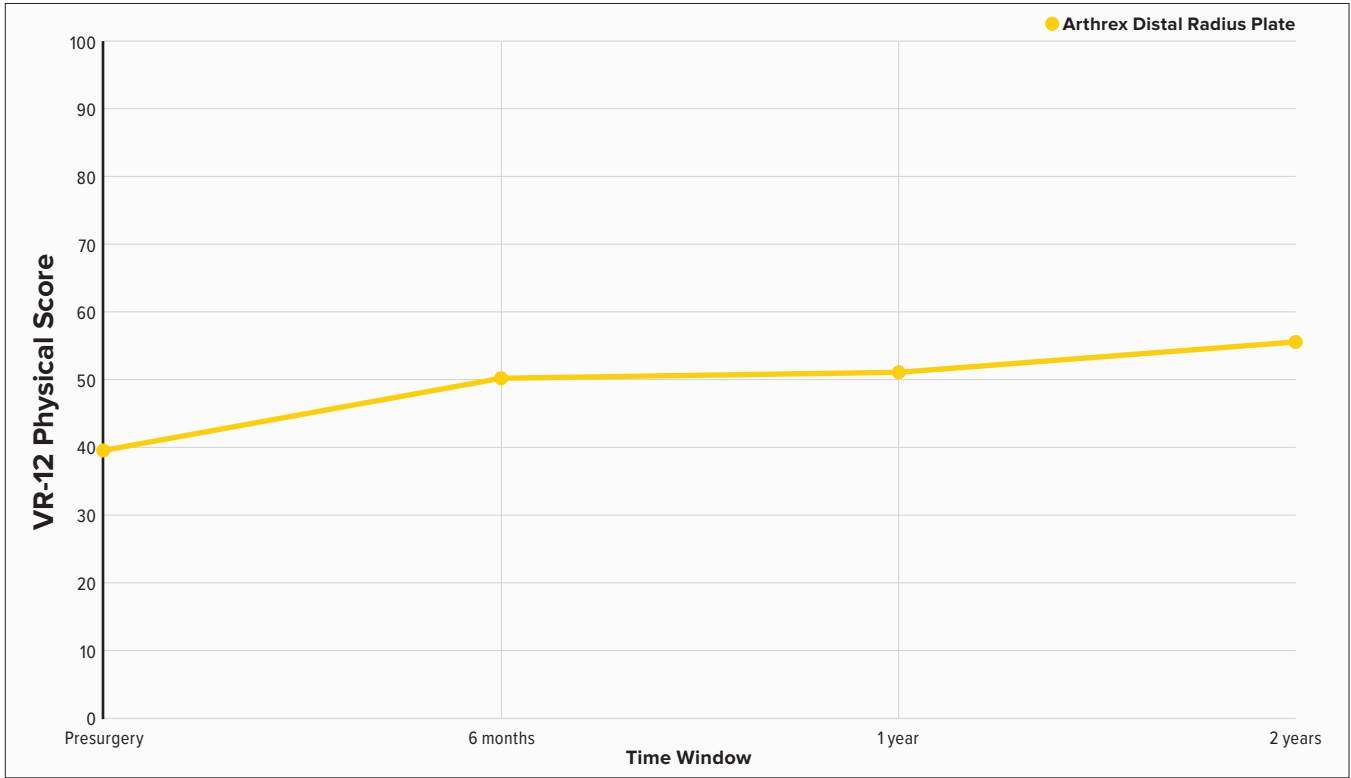
Trend Conclusion

No claims can be made on the potential of these results without further analysis to determine statistical significance.

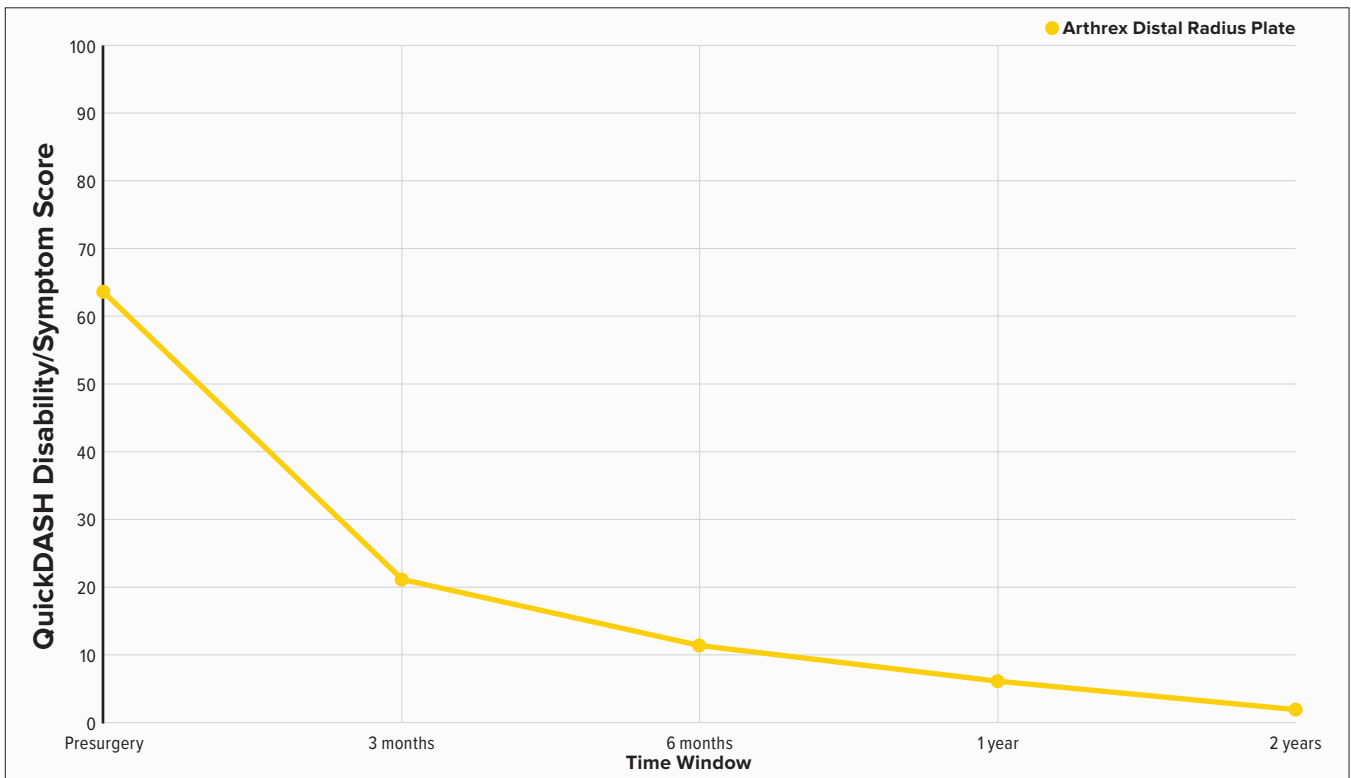
Results



VAS



VR-12



QuickDash

Time Point	Distal Radius Plate Avg ± STD VAS
Presurgery	4.9 ± 2.5
6 months	0.8 ± 1.2
1 year	0.7 ± 1.0
2 years	0.07 ± 0.13

Time Point	Distal Radius Plate Avg ± STD VR-12
Presurgery	39.3 ± 8.5
6 months	50.3 ± 6.9
1 year	51.0 ± 8.1
2 years	55.8 ± 4.0

Time Point	Distal Radius Plate Avg ± STD QuickDash
Presurgery	63.7 ± 20.4
6 months	11.8 ± 12.8
1 year	5.6 ± 8.4
2 years	2.3 ± 4.0