U.S. IDE 7-YEAR RESULTS

The M6-C[™] disc is the only FDA-approved artificial cervical disc designed to mimic the natural motion of a native cervical disc with six degrees of motion.

<u>Мб-С</u>

Artificial Cervical Disc



Evidence from the U.S. IDE clinical trial continues to demonstrate the safety and efficacy of the M6-C artificial cervical disc

At the 7-year endpoint, the M6-C disc had a

Subsequent Surgical Intervention rate of 6.9%

which is comparable to other FDA-approved artificial cervical discs* year post-op scores for patients were of a mean Shoulder/Arm pain VAS score of

Seven year post-op scores for patients were retained with a mean Shoulder/Arm pain VAS score of 0.5, which was significantly better than the mean of 2.1 observed in the ACDF control group



Decreases in disability as measured by NDI and decreases in Neck and Arm Pain Scores that were observed at prior follow-up periods were retained through seven-years post-op

Patient Focused. Proven Results.

REAL WORLD EVIDENCE

Results of a Kaplan Meier analysis of the M6-C disc based on

16 years

of real world evidence suggest a global cumulative survivorship (percentage of implanted devices that are still intact and functional at a specific time period) of

99% at 10 years



More than **15,000** M6-C discs have been implanted for more than **10 years***

*Data on file

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