Safety and Efficacy of Bioabsorbable Cervical Spacers and Low-Dose rhBMP-2 for Multilevel ACDF



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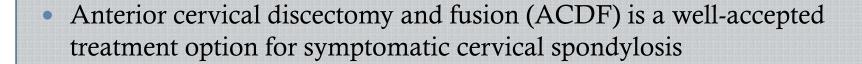
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Disclosures

- FDA off-label usage
 - o rhBMP-2 (INFUSE, Medtronic Sofamor Danek)
- NuVasive, Inc.
 - o Consultant
- All products discussed, spacer, infuse, and plate, all sold by Medtronic
- No relationship with Medtronic



Introduction



- Single-level ACDF:
 - o high fusion rates, regardless of graft choice or instrumentation
- Multi-level ACDF:
 - o variable fusion rates (56-100%)
 - o highly dependent upon graft choice, interbody spacer, instrumentation, # of levels, host factors, & definition



Options

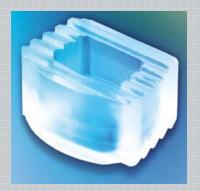
- Autograft is gold standard, but used less today
- Allograft may result in lower fusion rates in multilevel cases
- Synthetic spacers filled with graft material
 - Metal options
 - o PEEK: most commonly used
 - **Radiolucent**
 - Modulus of elasticity similar to bone

Bioabsorbable cervical spacers (BCS)

- Radiolucent; Modulus of elasticity similar to bone
- Rigid during implantation to provide immediate biomechanical stability, steadily degrades over time allowing gradual transfer of stresses to the graft material, degrades completely when fusion matured
- No issues with particulate debris or retained foreign body like PEEK
- The safety of polymers and their degradation products has been adequately demonstrated in the plastic surgery and orthopedic literature (suture, suture anchors, fracture fixation screws, etc.)



- Cornerstone HSR used in this study
 - o Medtronic Sofamor Danek, Memphis, TN
 - Noncrystalline polylactide copolymer of polylactide, 70:30 poly(L-lactide) to poly(D,L-lactide)
 - Degraded slowly to lactic acid → pyruvic acid → (via Krebs cycle) is excreted as CO2 and water

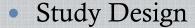




Graft options

- Many graft options exist:
- rhBMP-2 (Infuse, Medtronic Sofamor Danek, Memphis, TN)
 - Achieves fusion at least as well as autograft
 - NOT FDA approved in C-spine
 - Significant complications reported
 - o Great deal of scrutiny lately
- The combination of BCS with low-dose rhBMP-2 may be beneficial in aiding high fusion rates with acceptable morbidity in multi-level ACDF

Study Overview



- Retrospective review of a prospective observational cohort
 - **▼** IRB approved, prospective registry (ProSTOS, PhDx)

Inclusion Criteria

- \circ Consecutive patients treated between 2007-2012 (n=72)
- Multi-level (2+) ACDF with BCS and low-dose rhBMP-2 (& local bone if available)
- o Failure of conservative treatments and available for follow-up

• 72 patients, 187 levels

- o 37 (51%) 2-level cases
- o 27 (38%) 3-level cases
- 0 8 (11%) 4-level cases

Primary Diagnosis:

- Spondylosis: 40
- o Deformity/subluxation: 13
- o HNP: 8
- o Non-Union: 7
- o ASD: 4

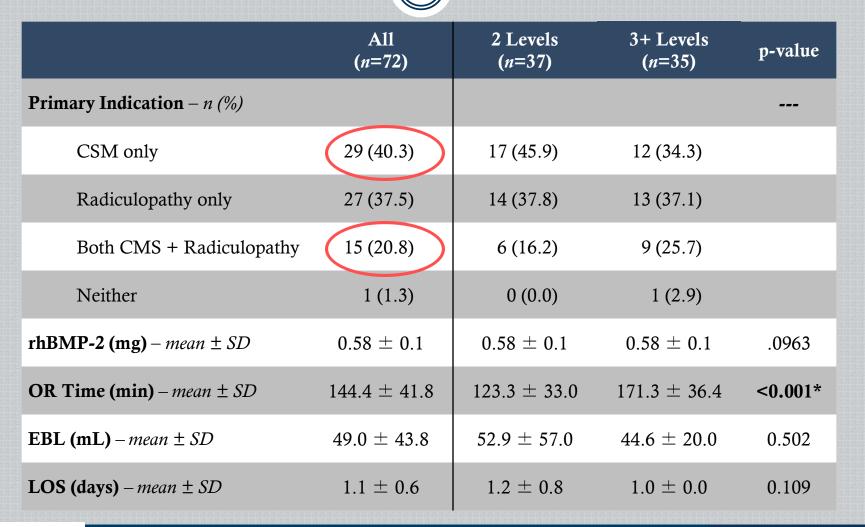
Most (65%) patients had > 1 primary diagnosis



Patient Samples

	A11 (n=72)	2 Levels (n=37)	3+ Levels (n=35)	p-value
Follow-Up (months) – mean ± SD	13.8 ± 6.8	14.2 ± 7.0	13.3 ± 6.6	0.574
Age (years) – mean ± SD	55.3 ± 10.4	51.8 ± 9.5	59.0 ± 10.2	0.003*
Female – <i>n</i> (%)	51 (70.8)	26 (70.3)	25 (71.4)	0.977
BMI (kg/ m^2) – mean \pm SD	28.4 ± 5.6	27.9 ± 5.6	28.8 ± 5.7	0.497
Tobacco Use – n (%)	24 (33.3)	16 (43.2)	8 (22.9)	0.067
Previous Cervical Surgery – n (%)	12 (16.7)	8 (21.6)	4 (11.4)	0.246

Surgical Summary





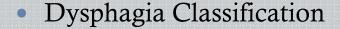
Clinical Outcomes

- o NDI
- o NRS (neck & arm, 0-10)
- SF-36 (PCS & MCS)
- Patient satisfaction
- o Did NOT use myelopathy outcome measures (MDI, JOA, Nurick, etc)

Imaging Studies

- Initial x-rays within 24 hours of surgery
- AP and lateral x-rays at 4-6 weeks
- O Serial x-rays with flexion/extension at 3, 6, 12, 24 months
- o CT obtained only if uncertainty regarding fusion status



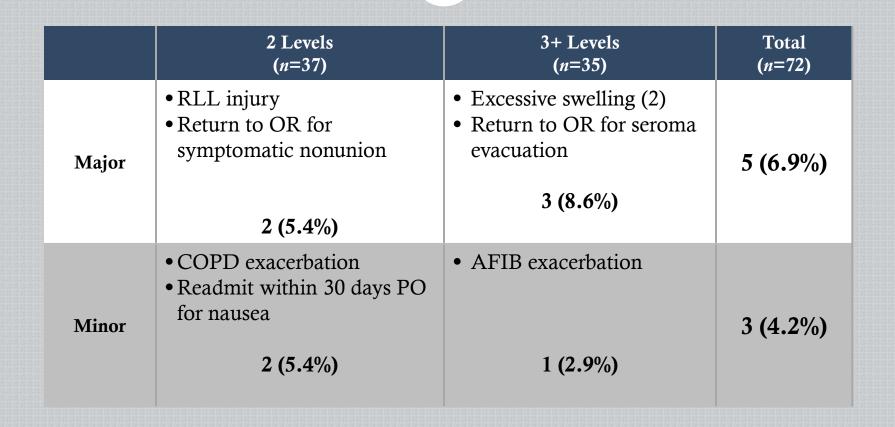


- O Quantified if symptoms were unresolved by 2 weeks PO or hospital readmission and additional treatment were required
- \circ "Prolonged" dysphagia: present ≥ 1 month po visit
- \circ "Persistent" dysphagia: present ≥ 3 month po visit

Analysis

- Chi-squared/Fishers' Exact tests and one-way ANOVA
- Significance accepted for p \leq 0.05

Adverse Events



1 pt with symptomatic pseudoarthrosis (1.4%)



Adverse Events: Dysphagia

- 38 patients (53%) had dysphagia that was not resolved by 2 week PO
 - o 21 patients had their dysphagia symptoms resolved by 1 month
 - o 13 patients had their dysphagia symptoms resolved by 3 months
 - o 2 patient had their dysphagia symptoms resolved by 6 months
 - o 2 patients had dysphagia at 1 & 3 months then lost to f/u
- 2 of these patients required readmission and observation/IV steroids

Prolonged dysphagia 17/72 = 24%

Persistent dysphagia 4/72 = 6%

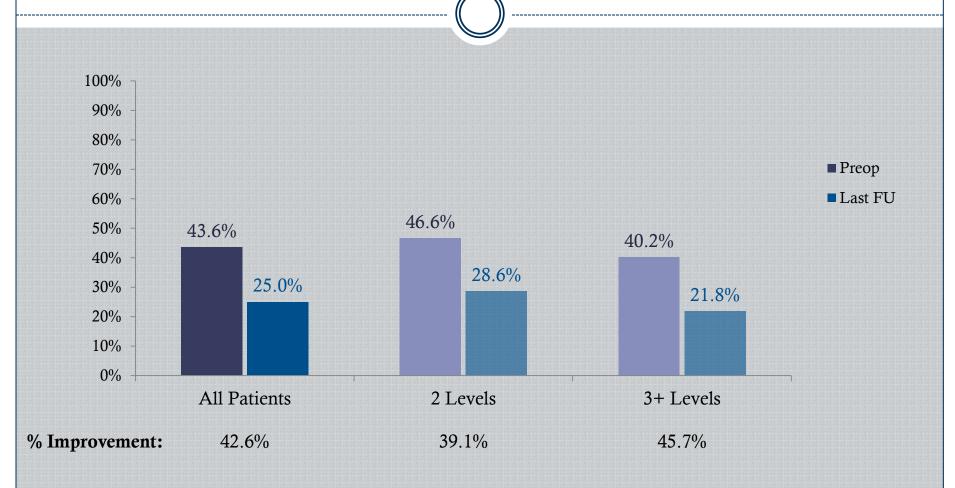


	Dysphagia	No dysphagia	Total
2 levels	18	19	37
3 levels	13	14	27
4 levels	7	1	8

P-value = 0.064

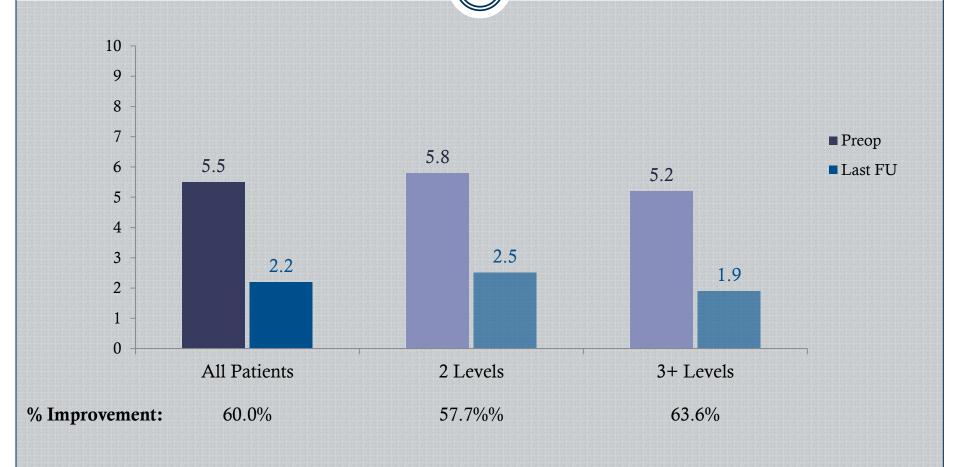


Clinical Outcomes: NDI



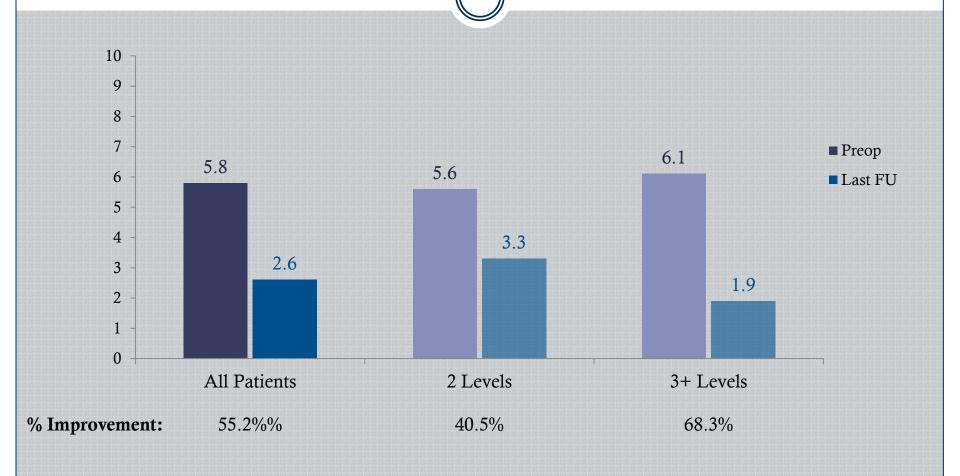








Clinical Outcomes: NRS Arm



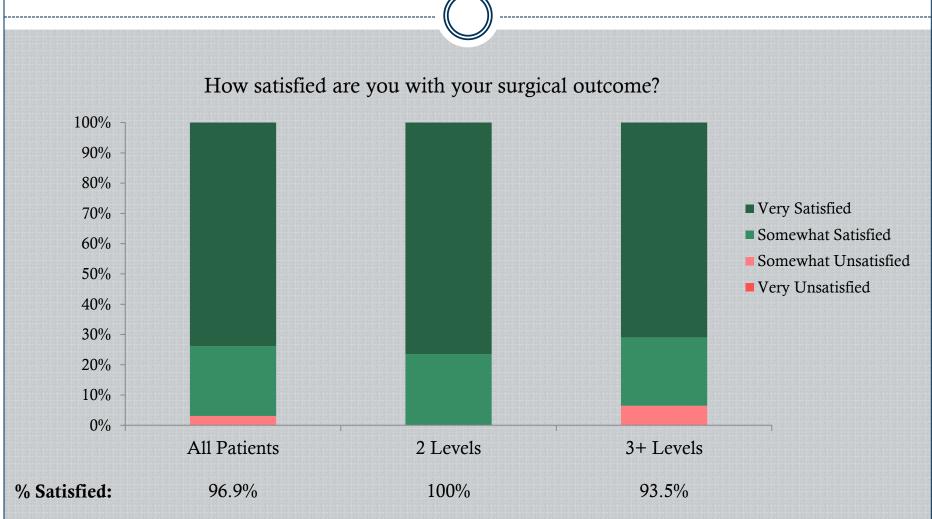


Clinical Outcomes: SF-36 PCS



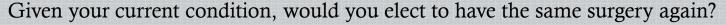


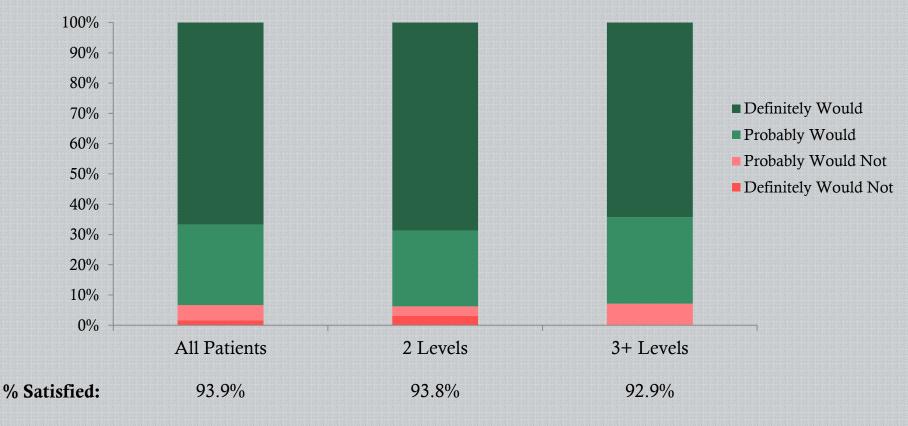
Patient Satisfaction





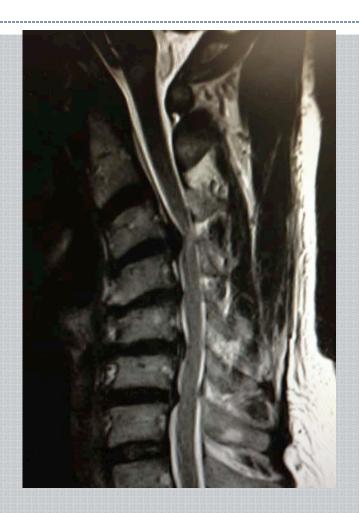
Patient Satisfaction







ACDF x 2 for myelopathy

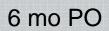




54 yo M CSM, Rt hemibody N/T 10 mo; inc DTRs, Hoffmans, clonus





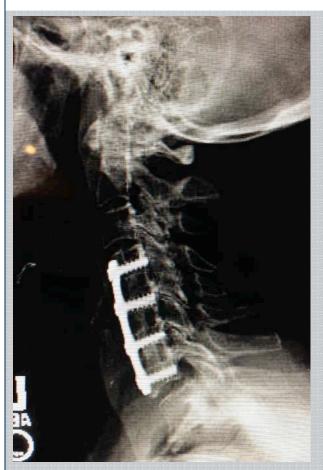




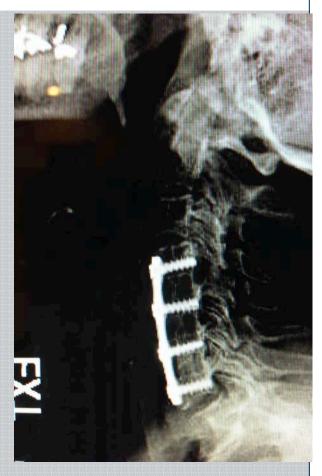




ACDF x 3 for myelopathy & radiculopathy







79 yo F with CSM & radic







ACDF x 4 for CSM





77 yo F CSM, N/C hands 5 yrs, lermittes, intrinsic hand weakness, hoffmans signs



Immediate PO





1 year PO

5 years PO







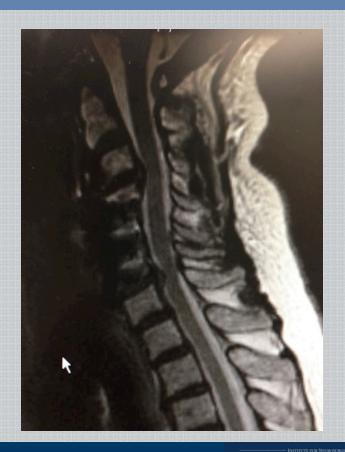
ACDF x 2 for ASD

- 63 y/o female
- PMHx
 - 2 years post C4-6 ACDF with PEEK cages 2009
- Presents 2010 with increasing NP,
 L shoulder pain
- In 2011 develops Left C7 radic











- Felt she had ASD C3-4 and C6-7
- C3-4/C6-7 ACDF July 2011
 - o BCS
 - 0.5mg rhBMP-2 per level + crushed local autograft



- Mild, transient prolonged dysphagia resolved by 1MO PO
- 6 MO PO
 - Solid C3-4 fusion
 - C6-7 symptomatic pseudoarthrosis
 - o Tried external electrical stim unit







- Return to OR 11MO PO for repair of C6-7 nonunion
 - o C6-7 posterior arthrodesis
 - C6-7 instrumented lateral mass screw rod fixation
- Solid fusion by 6 months PO







Discussion: Comparative Studies Fusion rates vs other graft/spacer options

- Current study: BCS + low-dose rhBMP-3, 2-4 levels
 - o 98.6% fusion rate; 1 pt with symptomatic pseudoarthrosis (1.4%)
- Fountas KN, Kapsalaki EZ, Nikolakakos LG, et al. Anterior cervical discectomy and fusion associated complications. *Spine (Phila Pa 1976)* 2007;32:2310-7.
 - Retrospective, autograft or allograft, +/- plate, 1-3 levels, 1015 patients
 - o 94% fusion rate for 2-level cases, 91% for 3-level cases
- Miller LE, Block JE. Safety and effectiveness of bone allografts in anterior cervical discectomy and fusion surgery. *Spine (Phila Pa 1976)* 2011;36:2045-50.
 - Systematic literature review
 - Allograft or autograft: fusion rate = 91% levels treated
 - Cage + morselized autologous bone: fusion rate = 97% levels treated



Discussion

Comparative Studies: Fusion rates vs other BCS papers

- Current study: BCS + low-dose rhBMP-3, 2-4 levels
 - o 98.6% fusion rate; 1 pt with symptomatic pseudoarthrosis (1.4%)
- Lanman TH, Hopkins TJ. Early findings in a pilot study of anterior cervical interbody fusion in which recombinant human bone morphogenetic protein-2 was used with poly(L-lactide-co-D,L-lactide) bioabsorbable implants. *Neurosurg Focus* 2004;16:6.
 - o BCS + rhBMP-2, 100% fusion rate (x-ray & CT), 3 mo f/u

Discussion

Comparative Studies: Complications / Dysphagia Rates

- Current study: BCS + low-dose rhBMP-3, 2-4 levels
 - o 11% complication rate (4% major, 7% minor)
 - o 24% prolonged & 4% persistent dysphagia rate (all resolved by 6 months)
- Fountas KN, Kapsalaki EZ, Nikolakakos LG, et al. Anterior cervical discectomy and fusion associated complication. *Spine (Phila Pa 1976)* 2007;32:2310-7.
 - 1-3 levels, 9% complication rate, 11% prolonged dysphagia rate (resolved by 4 weeks)
- Riley LH, Skolasky RL, Albert TJ. Dysphagia after anterior cervical decompression and fusion. Spine 2005;30:2564-9
 - Retrospective review, 454 pts/23 sites, telephone interviews
 - O Dysphagia at 3 mo: 20% 1 level, 33% 2 levels, 39% for 3+ levels



Study Strengths / Limitations

Strengths

- All consecutive patients included
- Outcomes all prospectively collected

Limitations

- o 72 pts/187 levels still small, short f/u
- Fusion definition based on x-rays
- O Dysphagia self reported. Telephone interviews may yield a higher % of patients with persistent dysphagia
- o Did NOT use myelopathy outcome measures (MDI, JOA, Nurick, etc)

Conclusions

- Multilevel ACDF can be a very effective treatment for symptomatic cervical spondylosis, although complications rates, particularly dysphagia, can be high, as can pseudoarthrosis rates
- Although the use of rhBMP-2 is both off-label and controversial, the results of the current study suggest that the combination of low-dose rhBMP-2 with BCS seems to be an excellent treatment option with acceptable complication rates, high fusion rates, and good clinical improvements in patients undergoing multi-level ACDF.

Thank you!



Minimally Invasive

Maximum Results

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ACDF x 2 for cerv radiculopathy

- 71 y/o female
- CC:
 - 3+ years neck pain + left C6 radiculopathy
 - Left 4+/5 deltoid weakness
 - Left 4/5 bicep weakness
 - o Left 4-/5 BR and WE weakness
- Cervical radiculopathy
 - o C4-5 DDD and subluxation
 - o C5-6 severe DDD



• C4-6 ACDF

- o BCS
- 0.5mg rhBMP-2 per level + crushed local autograft
- o C4-6 anterior cervical plating



- Expected dysphagia immediately PO, resolved by 2 weeks PO
- Clinical outcomes (24MO)

o NDI $52 \rightarrow 16$

o NRS Neck $8 \rightarrow 0$

o NRS Arm $6 \rightarrow 0$

o SF-36 PCS 41.4 → 59.1

o SF-36 MCS 25.3 → 45.3

- Patient satisfaction
 - Very satisfied with surgical outcome
 - Would definitely do again





ACDF x 2 for pseudo and ASD



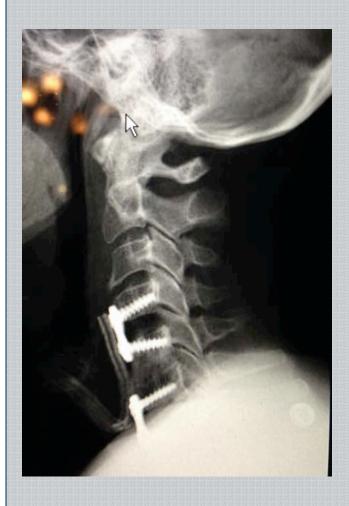




45 yo F s/p C5-6 ACDF 2004, with chronic NP & Left C7 radic



ACDF C45, C6-7







4 years PO







Results

Clinical Outcomes: SF-36 MCS



