

Professional Intracept Claim Form



HEALTH INSURANCE CLAIN APPROVED BY NATIONAL UNIFORM CLAIM COMMI		Black = Re Blue = Site		I/Required,	if applic	cable		
PICA							PICA	
1. MEDICARE MEDICAID TRICARE (Medicare#) (Medicaid#) (ID#/DoD#)	(Member ID#) (IDI	OUP ALTH PLAN BLK LUNK (ID4)	(ID#)	1a. INSURED'S I.D. N 12345678	9		(For Program in Item	1)
2. PATIENT'S NAME (Last Name, First Name, Middle Jones, Macie 5. PATIENT'S ADDRESS (No., Street)	02 6, PATIEN	O1 1961	F X	INSURED'S NAME INSURED'S ADDRE			Middle Initial)	
27 Wyoming Drive		Spouse Child						
New Richmond	WI	VED FOR NUCC USE		ZIP CODE	TTE	LEGHONE	(Include Area Code)	
54071 (651)555	5-5555					()	
9. OTHER INSURED'S NAME (Last Name, First Name	, Middle Initial) 10. IS PAT	IENT'S CONDITION RELAT	TED TO:	None None	Y GROUP OR	FECA NU	MBER	
a. OTHER INSURED'S POLICY OR GROUP NUMBER		YMENT? (Current or Previo	us)	a. INSURED'S DATE O	YY	м	SEX F	
b. RESERVED FOR NUCC USE	b. AUTO A	YES NO	LACE (State)	b. OTHER CLAIM ID (
c. RESERVED FOR NUCC USE	c. OTHER	ACCIDENT? YES NO		c. INSURANCE PLAN	NAME OR PRO	OGRAM N	AME	
d. INSURANCE PLAN NAME OR PROGRAM NAME Medicare	10d. CLAIN	I CODES (Designated by N	UCC)	d. IS THERE ANOTHE			NY? a items 9, 9a, and 9d.	
	FORE COMPLETING & SIGNING TURE (authorize the release of an	y medical or other informatio	n necessary	13. INSURED'S OR AL	JTHORIZED PE I benefits to the	ERSON'S S	SIGNATURE I authorize ed physician or supplie	
signed Signature on file		ATE			nature	on fi	le	
14. DATE OF CURRENT ILLNESS, INJURY, or PREG	NANCY (LMP) 15. OTHER DAT	E MM DD	YY	16. DATES PATIENT I			JRRENT OCCUPATION	N Y
17. NAME OF REFERRING PROVIDER OR OTHER S Dr. Timothy Jones	-301	123456789			DATES RELA	TED TO C	URRENT SERVICES	′
19. ADDITIONAL CLAIM INFORMATION (Designated		120 1007 00	$\overline{}$	20. OUTSIDE LAB?	NO I	\$ CH	IARGES	
Enter place of service code: 24 – Ambulatory Surgery Center			Ì	22. RESUBMISSION ORIGINAL REF. NO.				
E F.	22 – On Campus	Outpatient Hos	23. PRIOR AUTHORIZATION NUMBER A123456789					
24. A. DATE(S) OF SERVICE B Floor To PAC O MM DD YY MM DD YY SER SE	C. D. PROCEDURES, SEI (Explain Unusual C EMG CPT/HCPCS	RVICES, OR SUPPLIES Circumstances) MODIFIER	E. DIAGNOSIS POINTER	F. S CHARGES	G. H. DAYS EPSI OR Fami UNITS Plin	L ID. GUAL	J. RENDERING PROVIDER ID.	
01 07 22 01 07 22	64628		Α	XXXX	1	NPI	12345678	390 ⁻
01 07 22 01 07 22	64629		Α	XXXX	1-2	NPI	12345678	90
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	destruction of intra-			erve, inclusive	e of all im	naging		
	destruction of intraction distribution distr				of all im	aging		
guidanos, <u>sacira</u>	- TOTONIA	, idilibe	01 0001	ω		Name -		
25. FEDERAL TAX LD. NUMBER SSN EIN 123456789	26. PATIENT'S ACCOUNT N	O. 27. ACCEPT ASS	IGNMENT? see back	28. TOTAL CHARGE \$ XXXX	29. AM	OUNT PAI	D 30. Revd for NU	JCC Use
31. SIGNATURE OF PHYSICIAN OR SUPPLIER INCLUDING DEGREES OR CREDENTIALS	32. SERVICE FACILITY LOC Main Stree			33. BILLING PROVIDE		١.)	
 () certify that the statements on the reverse apply to this bill and are made a part thereof.) 	12345 Ma	in Street		12:	John Si 34 Main	Street		
SIGNED DATE	a. New Richi	mond WI 54071		a. NP	W Richm	nond V	VI 54071	
NUCC Instruction Manual available at: ww	NI DUICE OFG PI	EASE PRINT OR T	VPF	APPR/	OVED OMB	-0938-1	197 FORM 1500	(02-12

Indications for Use: The IntraceptTM Intraosseous Nerve Ablation System is intended to be used in conjunction with radiofrequency (RF) generators for the ablation of basivertebral nerves of the L3 through S1 vertebrae for the relief of chronic low back pain of at least six months duration t hat has not responded to at least six months of conservative care, and is also accompanied by features consistent with Type 1 or Type 2 Modic changes on an MRI such as infla mmation, edema, vertebral endplate changes, disruption and fissuring of the endplate, vascularized fibrous tissues within the adjacent marrow, hypointensive signals (Type 1 Modic change), and changes to the vertebral body marrow including replacement of normal bone marrow by fat, and hyperintensive signals (Type 2 Modic change). Contraindications - Use of the Intracept Intraosseous Nerve Ablation System is contraindicated in: Patients with severe cardiac or pulmonary compromise, pat ients with active implantable pulse generators (e.g. pacemakers, defibrillators), patients where the targeted ablation zone is < 10 mm away from a sensitive structure not intended to be ablated, including the vertebral foramen (spinal canal), patients with active systemic infection or local infection in the area to be treated, p atients who are pregnant, and/or skeletally immature patients (generally ≤ 18 years of age). Refer to the Instructions for Use provided with the Intracept Procedure or www.relievant.com/intracept/ for potential adverse effects, warnings, and precautions prior to using this product.

Caution: U.S. Federal law restricts this device to sale by or on the order of a physician.

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