



DISCLOSURE AND CONSENT MEDICAL AND SURGICAL PROCEDURES

TO THE PATIENT : You have the right as a patient to be informed about your condition and the recommended surgical, medical or diagnostic procedure to be used so that you may make the decision whether or not to undergo the procedure after knowing the risks and hazards involved. This disclosure is not meant to scare or alarm you; it is simply an effort to make you better informed so you may give or withhold your consent to the procedure.
1. I (we) voluntarily request Doctor(s) as my physician(s) and such associates, technical assistants and other health care providers as they may deem necessary, to treat my condition which has been explained to me (us) as (lay terms): Back Pain
2. I (we) understand that the following surgical, medical, and/or diagnostic procedures are planned for me and I (we) voluntarily consent and authorize these procedures (lay terms): Posterior Lumbar Interbody Fusion-the disc space is fused by approaching the spine through the back
3. INTRAOPERATIVE NEUROPHYSIOLOGICAL MONITORING: I (we) understand that intraoperative neurophysiological monitoring (IOM) may be utilized to identify neural structures, aid in performing the surgical procedure, and detect and prevent injury to the nervous system.
Please check appropriate box: □ Right □ Left □ Bilateral □ Not Applicable
4. I (we) understand that my physician may discover other different conditions which require additional or different procedures than those planned. I (we) authorize my physician, and such associates, technical assistants, and other health care providers to perform such other procedures which are advisable in their professional judgment.
 Please initialYesNo I consent to the use of blood and blood products as deemed necessary. I (we) understand that the following risks and hazards may occur in connection with the use of blood and blood products: a. Serious infection including but not limited to Hepatitis and HIV which can lead to organ damage and permanent impairment.
b. Transfusion related injury resulting in impairment of lungs, heart, liver, kidneys and immune system.

- c. Severe allergic reaction, potentially fatal.
- 6. I (we) understand that no warranty or guarantee has been made to me as to the result or cure.
- 7. Just as there may be risks and hazards in continuing my present condition without treatment, there are also risks and hazards related to the performance of the surgical, medical, and/or diagnostic procedures planned for me. I (we) realize that common to surgical, medical and/or diagnostic procedures is the potential for infection, blood clots in veins and lungs, hemorrhage, allergic reactions, and even death. I (we) also realize that the following hazards may occur in connection with this particular procedure: Pain, severe bleeding, infection, weakness, numbness or clumsiness, impaired muscle function or paralysis, incontinence, impotence, or impaired bowel function (loss of bowel/bladder control and/or sexual function), migration of implants (movement of implanted devices), failure of implants (breaking of implanted devices), adjacent level degeneration (breakdown of spine above and/or below the level treated), cerebrospinal fluid leak with potential for severe headaches, meningitis (infection of coverings of brain and spinal cord), recurrence, continuation or worsening of the condition that required this operation (no improvement or symptoms made worse), unstable spine (abnormal movement between bones and/or soft tissues of the spine)





<u>Lumbar Interbody Fusion-Posterior (cont.)</u>

8.	I (we)	understand tha	t Do Not Resuscitat	e (DNR),	Allow N	Natural Dea	th (AND) ar	nd all resu	scitative
restr	ictions	are suspended	during the periopera	ative perio	d and ur	ntil the pos	t anesthesia	recovery	period is
com	plete. A	All resuscitative	measures will be de	etermined 1	by the an	esthesiolog	gist until the	patient is o	officially
discl	narged	from the post a	nesthesia stage of car	e.					

complete. All resuscitative measures will discharged from the post anesthesia stage		logist until the patient is officially
9. I (we) authorize University Medical C use in grafts in living persons, or to other		
10. I (we) consent to the taking of still p during this procedure.	photographs, motion pictures, vide	otapes, or closed circuit television
11. I (we) give permission for a corpor- consultative basis.	ate medical representative to be p	present during my procedure on a
12. I (we) have been given an opportunity and treatment, risks of non-treatment, the benefits, risks, or side effects, including achieving care, treatment, and service goa informed consent.	procedures to be used, and the rise g potential problems related to re	ks and hazards involved, potential ecuperation and the likelihood of
13. I (we) certify this form has been full me, that the blank spaces have been filled	• •	
IF I (WE) DO NOT CONSENT TO ANY OF THI	E ABOVE PROVISIONS, THAT PROV	ISION HAS BEEN CORRECTED.
I have explained the procedure/treatment therapies to the patient or the patient's aut	<u> </u>	significant risks and alternative
A.M. (P.M.)		
Date Time	Printed name of provider/agent	Signature of provider/agent
Date Time A.M. (P.M.)		
*Patient/Other legally responsible person signature	Relationsh	ip (if other than patient)
*Witness Signature	Printed Na	me
 □ UMC 602 Indiana Avenue, Lubbock, TX □ UMC Health & Wellness Hospital 11011 □ OTHER Address: 		Street, Lubbock, TX 79430
Address (Str	reet or P.O. Box)	City, State, Zip Code
Interpretation/ODI (On Demand Interpreting)) □ Yes □ No	

☐ Yes ☐ No_____

Date/Time (if used)

Printed name of interpreter

Alternative forms of communication used

Date procedure is being performed:

Date/Time



	Lubbock,	Texas		
Da	te			

Resident and Nurse Consent/Orders Checklist

Instructions for form completion

Note: Enter "not applicable" or "none" in spaces as appropriate. Consent may not contain blanks.

Section 1:	Enter name of physician(s) responsible for procedure and patient's condition in lay terminology. Specific local of procedure must be indicated (e.g. right hand, left inguinal hernia) & may not be abbreviated.							
Section 2: Section 3:	 Enter name of procedure(s) to be done. Use lay terminology. The scope and complexity of conditions discovered in the operating room requiring additional surgical procedure. 							
B. Procedowith the Section 8:	should be specific to diag Enter risks as discussed we or procedures on List A mu- ures on List B or not address e patient. For these procedures Enter any exceptions to discuss of the control of the co	with patient. It be included. Othe sed by the Texas Me ures, risks may be er isposal of tissue or st	dical Disclosure panel do umerated or the phrase: 'ate "none".	o not require that sp "As discussed with	patient" entered.			
Section 9:	An additional permit with patient's consent for release is required when a patient may be identified in photographs or on video.							
Provider Attestation:	Enter date, time, printed r	name and signature o	f provider/agent.					
Patient Signature:	Enter date and time patient or responsible person signed consent.							
Witness Signature:	Enter signature, printed name and address of competent adult who witnessed the patient or authorized person's signature							
Performed Date:	Enter date procedure is being performed. In the event the procedure is NOT performed on the date indicated, staff must cross out, correct the date and initial.							
	s not consent to a specific prized person) is consenting		sent, the consent should b	oe rewritten to refle	ct the procedure that			
Consent	For additional information	n on informed conse	nt policies, refer to policy	y SPP PC-17.				
☐ Name of th	e procedure (lay term)	☐ Right or left	ndicated when applicable	e				
☐ No blanks	left on consent	☐ No medical a	bbreviations					
Orders								
Procedure 2	Date	Procedure						
☐ Diagnosis		☐ Signed by Pl	nysician & Name stamped	d				
Viirse	Res	zident	Den	nartment				