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.

I (we) voluntarily request Dras my physician, and such associates as	he/she
may deem necessary (for example anesthesia providers, educational assistants, and other health care providers are identified and their professional role explained to me) to treat my condition. My condition has explained to me as:	oviders
I (we) understand that the following surgical, medical, and/or diagnostic procedures are planned for me and I (voluntarily consent and authorize these <u>procedure(s)</u> :	we)
I (we) understand that my physician may discover other or different conditions which require additional process than those planned. I (we) authorize my physician, and any associates, technical assistants and other health of providers to perform such other procedures which are advisable in their professional judgment. I (we) understand that these qualified medical practitioners may be performing significant tasks related surgery such as opening or closing incisions, harvesting or dissecting tissue, altering tissue, implanting detissue removal or photography during procedures.	are to the
Initial I (we) Do Do Not consent to the use of blood and blood products as considered necessary. Benefits, risks, alternatives and the risks and benefits of alternatives have been discussed and I (we) have been given the opportunity to ask questions.	ve
Initial	
Texas Medical Disclosure HEMATIC AND LYMPHATIC SYSTEM	
1. Transfusion of blood and blood components.	
 Fever. Transfusion reaction which may include kidney failure or anemia. 	
3. Heart failure.	
4. Hepatitis.	
5. AIDS (Acquired Immune Deficiency Syndrome).	
6. Other infections.	



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DISCLOSURE AND CONSENT: UNIVERSAL PROCEDURE(S)
BLOOD/ BLOOD PRODUCT ADMINISTRATION



PATIENT IDENTIFICATION

C-300A (Rev. 06/17)

Page 1 of 2

Initial			
risks and hazards related to the such as the potential for infection	e performance of the on, blood clots in v	ne surgi eins and	uing my present condition without treatment, there are also cal, medical, and/or diagnostic procedures planned for me, d lungs, hemorrhage, allergic reactions and even death. I ards may occur in connection with this particular
I (we) Do Do Not Constitute constitute exception of:	ent to have studer	nts wat	ch my procedure with my doctor for medical education, with
by my physician, in the room equipment and/or supply comp for the procedure but will not	n during the processary for the produce perform any portent have confident	cedure. cts that tion of t tiality ag	nanufacturer's technical representatives, as requested I understand that one or more representatives from the the physician will use during my procedure, may be present the procedure. I further understand that all manufacturer's greements and that none of my personal health information hin this hospital.
I (we) Do Do Not Considentity is not shown to anyone		an takinç	g photographs during my procedure as long as my name or
I (we) consent to the disposal b	y hospital authorit	ties of a	ny tissue or parts which may be removed.
the benefits, the likelihood of s of my condition, and other alte understand that no warranty of	uccess, the possilernative forms of toor guarantee has	ble prob reatmen been m	s about my current condition(s), the proposed procedure(s), elems related to recovery, the possible risks of nontreatment at, and the risks and benefits of alternatives involved. I (we) hade to me as to result or cure. Any professional/business tal and educational institutions has been explained to me.
blank spaces have been filled	in, and that I (we d consent and I (w	e) under	at I (we) have read it or have had it read to me (us), that the stand its contents. I (we) believe that I (we) have sufficient est the procedure(s) to be done.
Patient's Signature	Date	Time	Other Legally Responsible Person's Signature Relationship Date Time
			☐ Medical City Plano, 3901 West 15th Street, Plano, Texas 75075 ☐ Other:
Witness Signature/Title/Position	Date	Time	Witness Work Address
Interpreter			Reason:
I have provided the patient/parent/grmy area of expertise.	uardian with informatic	on on risks	s, benefits, and alternatives to treatment as outlined in the above within
Date: Time:	Physician Sign	ature: <u>X</u>	Physician Identifier



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DISCLOSURE AND CONSENT: UNIVERSAL PROCEDURE(S) BLOOD/ BLOOD PRODUCT ADMINISTRATION



PATIENT IDENTIFICATION

C-300B (Rev. 06/17)

ANESTHESIA CONSENT

TO THE PATIENT: You have the right, as a patient, to be informed about your condition and the recommended anesthesia/analgesia to be used so that you may make the decision whether or not to receive the anesthesia/analgesia 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 that you may give or withhold your consent to the anesthesia/analgesia.

I voluntarily request that anesthesia and/or perioperative pain management care (analgesia) as indicated below be administered to me (the patient). I understand it will be administered by an anesthesia provider and/or the operating practitioner, and such other health care providers as necessary. Perioperative means the period shortly before, during and shortly after the procedure.

I (we) understand that anesthesia involves additional risks and hazards, but I (we) request the use of anesthetics/analgesia for the relief and protection from pain or anxiety during the planned and additional procedures. I (we) realize the type of anesthesia/analgesia may have to be changed possibly without explanation to me (us).

I understand that serious, but rare, complications can occur with all anesthetic/analgesic methods. Some of these risks are breathing and heart problems, drug reactions, nerve damage, cardiac arrest, brain damage, paralysis, or death.

I also understand that other complications may occur. Those complications include but are not limited to:

Have	the patient/other legally responsible person initial the planned anesthesia/analgesia method(s).
	Initial General Anesthesia - Injury to Vocal Cords, Teeth, Lips, Eyes; Awareness during the procedure; Memory Dysfunction/Memory Loss; Permanent Organ Damage; Brain Damage.
	Regional Block Anesthesia/Analgesia - Nerve Damage; Persistent Pain; Bleeding/Hematoma; Infection; Medical necessity to convert to general anesthesia; Brain Damage.
	Initial Spinal Anesthesia/Analgesia - Nerve Damage; Persistent Back Pain; Headache; Infection; Bleeding/Epidural Hematoma; Chronic Pain; Medical necessity to convert to general anesthesia; Brain Damage.
	Initial Epidural Anesthesia/Analgesia - Nerve Damage; Persistent Back Pain; Headache; Infection; Bleeding/Epidural Hematoma; Chronic Pain; Medical necessity to convert to general anesthesia; Brain Damage.
	Deep Sedation - Memory Dysfunction/Memory Loss; Medical necessity to convert to general anesthesia; Permanent Organ Damage; Brain Damage.
	Initial Moderate Sedation - Memory Dysfunction/Memory Loss; Medical necessity to convert to general anesthesia; Permanent Organ Damage; Brain Damage.
Addit	tional comments/risks:
] Initial
	Prenatal/Early Childhood Anesthesia- Potential long-term negative effects on memory, behavior, and learning with prolonged or repeated exposure to general anesthesia/moderate sedation/deep sedation during pregnancy and in early childhood.



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DISCLOSURE AND CONSENT:
ANESTHESIA and/or PERIOPERATIVE PAIN
MANAGEMENT

PATIENT IDENTIFICATION

I (we) have been given an opportunity to ask questions about my anesthesia/analgesia methods, the procedures to be used, the risks and hazards involved, and alternative forms of anesthesia/analgesia. I (we) believe that I (we) have sufficient information to give this informed consent.

I (we) certify this form has been fully explained to me, that I (we) have read it or have had it read to me, that the blank spaces have been filled in, and that I (we) understand the contents.

I (we) understand that no promises have been made to me as to the result of anesthesia/analgesia methods.

Initials					
Date	Time	Other Legally Responsible Person's Signature	Relationship	Date	Time
		· · · · · · · · · · · · · · · · · · ·	,		
Date	Time	Witness Work Address			
		Reason:			
,		tient/family understand(s) and agree(s) to	' <u></u>		····
	Date Date	Date Time Date Time	Date Time Other Legally Responsible Person's Signature Medical City Plano, 3901 West 15th Street, Pland Medical City Frisco, 5500 Frisco Square Blvd. Other: Witness Work Address Reason: We been explained and the patient/family understand(s) and agree(s) to the patient of the patient	Date Time Other Legally Responsible Person's Signature Relationship Medical City Plano, 3901 West 15th Street, Plano, TX 75075 Medical City Frisco, 5500 Frisco Square Blvd., Frisco, TX 75034 Other: Witness Work Address Reason:	Date Time Other Legally Responsible Person's Signature Relationship Date Medical City Plano, 3901 West 15th Street, Plano, TX 75075 Medical City Frisco, 5500 Frisco Square Blvd., Frisco, TX 75034 Other: Witness Work Address



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DISCLOSURE AND CONSENT:
ANESTHESIA and/or PERIOPERATIVE PAIN
MANAGEMENT

MANAGEMENT MANAGEMENT

PATIENT IDENTIFICATION

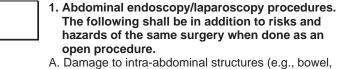
LIST A TEXAS MEDICAL DISCLOSURE

(EFFECTIVE: JANUARY 1, 2012, AMENDED: APRIL 1, 2012)

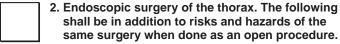
Procedures requiring full disclosure (List A). The following treatments and procedures require full disclosure by the physician or health care provider to the patient or person authorized to consent for the patient.

Patient to initial appropriate square.

ENDOSCOPIC SURGERY



- A. Damage to intra-abdominal structures (e.g., bowel, bladder, blood vessels, or nerves).
- B. Intra-abdominal abscess and infectious complications.
- C. Trocar site complications (e.g.,hematoma/bleeding, leakage of fluid, or hernia formation).
- D. Conversion of the procedure to an open procedure.
- E. Cardiac dysfunction.



- A. Postoperative pneumothorax.
- B. Subcutaneous emphysema.
- C. Conversion of the procedure to an open procedure.



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DISCLOSURE AND CONSENT: ENDOSCOPIC SURGERY



PATIENT IDENTIFICATION

C-305 (Rev. 06/17) Page 1 of 1