**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 Dr.\_\_\_\_\_as my physician, and such associates as he/she may deem necessary (for example anesthesia providers, educational assistants, and other health care providers who are identified and their professional role explained to me) to treat my condition. My <u>condition</u> has been explained to me as:

I (we) understand that the following surgical, medical, and/or diagnostic procedures are planned for me and I (we) voluntarily consent and authorize these <u>procedure(s)</u>:

I (we) understand that my physician may discover other or different conditions which require additional procedures than those planned. I (we) authorize my physician, and any associates, technical assistants and other health care 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 to the surgery such as opening or closing incisions, harvesting or dissecting tissue, altering tissue, implanting devices, tissue removal or photography during procedures.

Initial

I (we) **Do Do Not C** 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.

PATIENT IDENTIFICATION

Initial

Texas Medical Disclosure HEMATIC AND LYMPHATIC SYSTEM

1. Transfusion of blood and blood components.

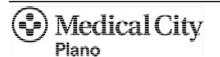
1. Fever.

- 2. 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



Initial

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, such as the potential for infection, blood clots in veins and lungs, hemorrhage, allergic reactions and even death. I (we) also realize that the following specific risks and hazards may occur in connection with this particular procedure(s):

I (we) **Do Do Not C** consent to have students watch my procedure with my doctor for medical education, with the exception of:

I (we) **Do Do Not C** consent to have one or more manufacturer's technical representatives, as requested by my physician, in the room during the procedure. I understand that one or more representatives from the equipment and/or supply company for the products that the physician will use during my procedure, may be present for the procedure but will not perform any portion of the procedure. I further understand that all manufacturer's technical representatives present have confidentiality agreements and that none of my personal health information will be disclosed to anyone other than my care givers within this hospital.

I (we) **Do Do Not C** consent to my physician taking photographs during my procedure as long as my name or identity is not shown to anyone.

I (we) consent to the disposal by hospital authorities of any tissue or parts which may be removed.

I (we) have been given the opportunity to ask questions about my current condition(s), the proposed procedure(s), the benefits, the likelihood of success, the possible problems related to recovery, the possible risks of nontreatment of my condition, and other alternative forms of treatment, and the risks and benefits of alternatives involved. I (we) understand that no warranty or guarantee has been made to me as to result or cure. Any professional/business relationship between my health care providers, the hospital and educational institutions has been explained to me.

I (we) certify this form has been fully explained to me, that I (we) have read it or have had it read to me (us), that the blank spaces have been filled in, and that I (we) understand its contents. I (we) believe that I (we) have sufficient information to give this informed consent and I (we) request the procedure(s) to be done.

	nitials					
Patient's Signature	Date	Time Oth	ner Legally Responsible Person's Signature	Relationship	Date	Time
		_	Medical City Plano, 3901 West 15th Street, Pla Other:	ano, Texas 75075		
Witness Signature/Title/Position	Date	Time	Witness Work Address			
		Re	ason:			
Interpreter						
Date: Time:	Physician Sig	gnature: <u>X</u>			Physician Ide	entifier
Medical City     Plano	3901 West 15 Plano, Texas (972) 596-0	75075	PATIENT IDENTIFICATION			
DISCLOSURE AND CONSENT: UI BLOOD/ BLOOD PRODUC		• • •				

## **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. Initial 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. Initial **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. Additional 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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PATIENT IDENTIFICATION

DISCLOSURE AND CONSENT: NESTHESIA and/or PERIOPERATIVE PAIN MANAGEMENT



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					
Patient's Signature	Date	Time	Other Legally Responsible Person's Signature Rel	lationship	Date	Time
			<ul> <li>☐ Medical City Plano, 3901 West 15th Street, Plano, TX 75075</li> <li>☐ Medical City Frisco, 5500 Frisco Square Blvd., Frisco, TX 75034</li> <li>☐ Other:</li> </ul>			
Witness Signature/Title/Position	Date	Time	Witness Work Address			
Interpreter			Reason:			

The risks, benefits, and alternatives have been explained and the patient/family understand(s) and agree(s) to the procedure

Physician / Proceduralist Responsible for Anesthesia: X

Physician Identifier Time

Date



## LIST A TEXAS MEDICAL DISCLOSURE

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

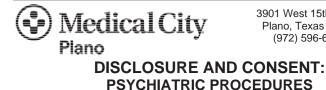
Procedures requiring full disclosure (List A). The following treatments and procedures require full disclosure by the physician or health care provider to the patient or person authorized to consent for the patient.

## Patient to initial appropriate square.

## **PSYCHIATRIC PROCEDURES**



- 1. Electroconvulsive therapy with modification by intravenous muscle relaxants and sedatives.
- A. Memory changes of events prior to, during, and immediately following the treatment.
- B. Fractures or dislocations of bones.
- C. Significant temporary confusion requiring special care.



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PATIENT IDENTIFICATION