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PATIENT INITIALS:

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Board Certified in Pain Management			
NFORMED CONSENT AND PAIN MANAGEMENT AGREEMENT AS REQUIRED BY THE TEXAS MEDICAL BOARD REFERENCE: TEXAS ADMINISTRATIVE CODE, TITLE 22, PART 9, CHAPTER 170 Brd Edition: Developed by the Texas Pain Society (www.texaspain.org)			
NAME OF PATIENT:	DATE:		
TO THE PATIENT: As a patient, you have the right to be informed about your coprocedure or drug therapy to be used, so you may make the informed decision what hazards involved. This disclosure is not meant to scare or alarm you, but rather or withhold your consent/permission to use the drug(s) recommended to you by the use of the word "physician" is defined to include not only my physician but all assistants, nurses, staff, and other health care providers as might be necessary	whether or not to take the drug after knowing the risks and it is an effort to make you better informed so you may give me, as your physician. For the purpose of this agreement lso my physician's authorized associates, technical		
CONSENT TO TREATMENT AND/OR DRUG THERAPY: I voluntarily request recondition which has been explained to me as chronic pain. I hereby authorize are or write prescription(s) for dangerous and/or controlled drugs (medications) as a lt has been explained to me that these medication(s) include opioid/narcotic drug supervision. I further understand these medication(s) may lead to physical depet the practice of medicine, produce adverse side effects or results. The alternative possibilities of complications have been explained to me. I understand that this I common side effects or reactions, and that death is also a possibility as a result	and give my voluntary consent for my physician to administer an element in the treatment of my chronic pain. g(s), which can be harmful if taken without medical endence and/or addiction and may, like other drugs used in the methods of treatment, the possible risks involved, and the listing is not complete, and that it only describes the most		
THE SPECIFIC MEDICATION(S) MY PHYSICIAN PLANS TO PRESCRIBE WI FROM THIS AGREEMENT. THIS INCLUDES THE USE OF MEDICATIONS FO APPROVED BY THE DRUG COMPANY AND THE GOVERNMENT (THIS IS S PRESCRIBING). MY DOCTOR WILL EXPLAIN HIS TREATMENT PLAN(S) FO	OR PURPOSES DIFFERENT THAN WHAT HAVE BEEN COMETIMES REFERRED TO AS "OFF-LABEL"		
I HAVE BEEN INFORMED AND understand that I will undergo medical tests and Those tests include random unannounced checks for drugs and psychological et hereby give permission to perform the tests or my refusal may lead to terminate substances may result in my being discharged from your care.	evaluations if and when it is deemed necessary, and		

For female patients only:

To the best of my knowledge I am NOT pregnant.

If I am not, I will use appropriate contraception/birth control during my course of treatment. I accept that it is **MY responsibility** to inform my physician if I become pregnant.

If I am pregnant or am uncertain, I WILL NOTIFY MY PHYSICIAN IMMEDIATELY.

All of the above possible effects of medication(s) have been fully explained to me and I understand that, at present, there have not been enough studies conducted on the long-term use of many medication(s) i.e. opioids/narcotics to assure complete safety to my unborn (child)ren. With full knowledge of this, I consent to its use and hold my physician harmless for injuries to the embryo / fetus / baby.

I UNDERSTAND THE MOST COMMON SIDE EFFECTS THAT COULD OCCUR IN THE USE OF THE DRUGS USED IN MY TREATMENT INCLUDE BUT ARE NOT LIMITED TO THE FOLLOWING:

- Emotional and physical dependence, which is rare (occurring in approximately 1 in 10,000 patients without a history of substance abuse).
- Constipation, difficulty with urination, nausea, vomiting, itching, slow breathing, reduced sexual function, depression, insomnia and tolerance to medication(s).
- Decreased ability to perform activities such as driving and using machinery.
- Patient is aware taking benzodiazepines (alprazolam, diazepam, lorazepam) with opioids can lead to an overdose.

The alternative methods of treatment, the possible risks involved, and the possibilities of complications have been explained to me, and I still desire to receive medication(s) for the treatment of my chronic pain.

The goal of this treatment is to help me gain control of my chronic pain in order to live a more productive and active life. I realize that I may have a chronic illness and there is a limited chance for complete cure, but the goal of taking medication(s) on a regular basis is to reduce (but probably not eliminate) my pain so that I can enjoy an improved quality of life. I realize that the treatment for some will require prolonged or continuous use of medication(s), but an appropriate treatment goal may also mean the eventual withdrawal for the use of all medication(s). My treatment plan will be tailored specifically for me. I understand that I may withdraw from this treatment plan and discontinue the use of the medication(s) at any time and that I will notify my physician of any discontinued use. I further understand that I will be provided medical supervision if needed when discontinuing medication use.

I understand that no warranty or guarantee has been made to me as to the results of any drug therapy or cure of any condition. The long-term use of medications to treat chronic pain is controversial because of the uncertainty regarding the extent to which they provide long-term benefit. I have been given the opportunity to ask questions about my condition and treatment, risks of non-treatment and the drug therapy, medical treatment or diagnostic procedure(s) to be used to treat my condition, and the risks and hazards of such drug therapy, treatment and procedure(s), and I believe that I have sufficient information to give this informed consent.

SOAPP-R Score:		
ORT Score:	 PATIENT INITIALS:	
Data:		