## Sample Informed Consent Form

## Introduction

This sample informed consent form can be used with patients who are beginning long-term therapy with opioid analgesics to help ensure they understand the side effects, risks, conditions, and purpose of their treatment. This document can help facilitate clear communication between patients and healthcare providers about long-term opioid therapy, clearly define treatment expectations, and resolve any questions or concerns patients may have before treatment initiation.

## Opioid Therapy for Chronic Pain: Sample Informed Consent\*

Please review the information listed here and put your initials next to each item when you have reviewed it with your provider and feel you understand and accept what each statement says. My provider is prescribing opioid pain medications for the following conditions(s): When I take these medications, I may experience certain reactions or side effects that could be dangerous, including sleepiness or sedation, constipation, nausea, itching, allergic reactions, problems with thinking clearly, slowing of my reactions, or slowing of my breathing. When I take these medications it may not be safe for me to drive a car, operate machinery, or take care of other people. If I feel sedated, confused, or otherwise impaired by these medications, I should not do things that would put other people at risk for being injured. When I take these medications regularly, I will become physically dependent on them, meaning that my body will become accustomed to taking the medications every day, and I would experience withdrawal sickness if I stop them or cut back on them too quickly. Withdrawal symptoms feel like having the flu, and may include abdominal pain, nausea, vomiting, diarrhea, sweating, body aches, muscle cramps, runny nose, yawning, anxiety, and sleep problems. I may become addicted to these medications and require addiction treatment if I cannot control how I am using them, or if I continue to use them even though I am having bad or dangerous things happen because of the medications. Anyone can develop an addiction to opioid pain medications, but people who have had problems with mental illness or with controlling drug or alcohol use in the past are at higher risk. I have told my provider if I or anyone in my family has had any of these types of problems. —Taking too much of my pain medication, or mixing my pain medications with drugs, psychiatric medicine, or other medications that cause sleepiness, such as benzodiazepines, barbiturates, and other sleep aids, could cause me to be dangerously sedated or to overdose and stop breathing. I understand that taking certain medications such as buprenorphine (Suboxone®, Subutex®, naltrexone (ReVia®), nalbuphine (Nubain®), pentazocine (Talwin®), or butorphanol (Stadol®) will reverse the effects of my pain medicines and cause me to go into withdrawal. It is my responsibility to tell any provider that is treating me or prescribing me medications that I am taking opioid pain medications so that they can treat me safely and do not give me any medicines that may interact dangerously with my pain medicines. \*Adapted from the American Academy of Pain Medicine http://www.painmed.org/Workarea/DownloadAsset.aspx?id=321

	ossible risks and benefits of taking op nd have discussed the possibility of or :	
These medications are controlled my pain well enoug	being prescribed to me because othe gh.	r treatments have not
These medications are pain completely.	to be used to decrease my pain but the	ney will not take away my
and my family, and meet othe	to be used to help improve my ability or goals that I have discussed with my neet those goals, they will be stopped.	provider, but if these
For Men: Taking opio and affect sexual function.	oid pain medications chronically may	cause low testosterone levels
pregnant or if I am thinking al medications and continue to ta	responsibility to tell my provider imr bout getting pregnant. If I become pr ake the medicines during the pregnan me of birth and may require withdraw	egnant while taking these cy, the baby will be physically
	ny provider and have had the chance to as tre and by signing give my consent for tre	• •
Patient signature	Patient name printed	Date
Provider signature	Provider name printed	Date
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