Patient Name:	Date of Birth:
	nt or legal guardian if the patient is a minor must initial st sign and date the informed consent form.
I,, und for specific medical conditions and/or syn Health.	derstand that Medical Cannabis is offered as treatment opportunity as designated by the Oklahoma Department of
<u>PLEASE II</u>	NITIAL EACH SECTION
	MD is a qualified physician who is registered with the thority and may order medical cannabis for my medical build benefit from this medical decision.
I understand that Dr. Amer Nouh , should be a substitute for any other treatm	MD is not implying or suggesting that medical cannabis ent prescribed by another physician.
I understand that Medical marijua contain unknown quantities of active ingre	na is not regulated by the USFDA and therefore may dients, impurities and/or containments.
	liance has not been issued under the Food and Drug he safety and effectiveness of marijuana as a drug. I
I am aware that medical marijuana understand that medical marijuana has no	has not been approved under federal regulations, and I t been deemed legal under federal law.
	ks associated with the use of marijuana are not fully may involve risks that have not been identified. I accept
I agree that if I am a female patient think about becoming pregnant. I acknowle	that I will contact my attending physician if I become or edge that the use of medical
marijuana creates passthrough problems breastfeeding.	s to a fetus during pregnancy and to a baby during
I should not drive a vehicle while driving under the influence.	using medical marijuana and that I can get a DUI for
substance. The federal government has substance. Schedule I substances are de (2) no currently accepted medical use accepted safety for use under medical	sification of marijuana as a Schedule I controlled as classified marijuana as a Schedule I controlled ifined, in part, as having (1) a high potential for abuse; in treatment in the United States; and (3) a lack of supervision. Federal law prohibits the manufacture, even in states, such as Oklahoma, which have modified dicine.
	the influence of medical marijuana, the patient or the edical marijuana use registry identification card in his or

Patient Name:	Date of Birth:
drug. Therefore, the "manufact standards, quality control, or oth ingredients, which may vary in	approved by the Food and Drug Administration for marketing as a ture" of marijuana for medical use is not subject to any federal ner oversight. Marijuana may contain unknown quantities of active potency, impurities, contaminants, and substances in addition to noactive chemical component of marijuana.
dependence on, or addiction to	nat the use of marijuana by individuals may lead to a tolerance to, marijuana. I understand that if I require increasingly higher doses if I think that I may be developing a dependency on marijuana, I or my primary care.
think, judge and reason. Driving which escalates if alcohol is als not drive, operate heavy machi respond quickly and I should	in affect coordination, motor skills and cognition, i.e., the ability to g under the influence of cannabis can double the risk of crashing, so influencing the driver. While using medical marijuana, I should nery or engage in any activities that require me to be alert and/or not participate in activities that may be dangerous to myself or rive while under the influence of marijuana, I can be arrested for
following: dizziness, anxiety, comemory, euphoria, difficulty in system, may affect the product concentrate, impaired motor sland/or restlessness. Marijuana disorder. In addition, the use of my perception of time and spause of medical marijuana, es	from the use of marijuana include, but are not limited to, the onfusion, sedation, low blood pressure, impairment of short term completing complex tasks, suppression of the body's immune ction of sex hormones that lead to adverse effects, inability to kills, paranoia, psychotic symptoms, general apathy, depression may exacerbate schizophrenia in persons predisposed to that f medical marijuana may cause me to talk or eat in excess, alter ce and impair my judgment. Many medical authorities claim that pecially by persons younger than 25, can result in long-term y, learning, drug abuse, and schizophrenia.
	ng marijuana while consuming alcohol is not recommended. Ome present when using both alcohol and marijuana.
any of the side effects listed thoughts, or experience crying	mer Nouh or my primary care or go to nearest ER if I experience above, or if I become depressed or psychotic, have suicidal spells, experience respiratory problems, changes in my normal gue, increased irritability, or begin to withdraw from my family
restlessness, agitation, loss of tiredness. Symptoms of mariju hacking cough, disturbances in	n include: feelings of depression, sadness, irritability, insomnia, appetite, trouble concentrating, sleep disturbances and unusual ana overdose include, but are not limited to, nausea, vomiting, heart rhythms, numbness in the hands, feet, arms or legs, anxiety experience these symptoms, I agree to contact Dr. Amer Nouh or e nearest emergency room.
known. Some mixtures of med	nown to interact with marijuana and not all drug interactions are ications can lead to serious and even fatal consequences. I will physician(s) of my use of medical marijuana.

Therapeutic Health Clinic

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Consent Form

Patient Name:	Date of Birth:
liver enzymes, and other bodily sys	risk of bleeding, low blood pressure, elevated blood sugar, stems when taken with herbs and supplements. I agree to y or go to the nearest emergency room if these symptoms
	marijuana may have serious risks and may cause low babies. I will advise Dr. Amer Nouh or my PCP if I become be breastfeeding.
The current state of research set forth in this section:	on the efficacy of marijuana to treat the qualifying conditions
effective treatment for cancers, include (and the endocannabinoid system of processes. Due to a lack of recent, effectiveness of cannabis or cannabis	oral cannabinoids are effective antiemetics in the treatment of vomiting. There is insufficient evidence to support or refute re an effective treatment for cancer-associated anorexia-
effective treatment for epilepsy. Frandomized controlled trials evaluating Currently available clinical data there	support or refute the conclusion that cannabinoids are an Recent systematic reviews were unable to identify any ng the efficacy of cannabinoids for the treatment of epilepsy. If ore consist solely of uncontrolled case series, which do not icacy. Randomized trials of the efficacy of cannabidiol for completed and await publication.
intraocular pressure associated with glaucoma treatments. Non- random have shown short-term reductions	cannabinoids are an ineffective treatment for improving glaucoma. Lower intraocular pressure is a key target for ized studies in healthy volunteers and glaucoma patients in intraocular pressure with oral, topical eye drops, and ing the potential for therapeutic benefit. A good-quality

Crohn's Disease:

limited potential for cannabinoids in the treatment of glaucoma.

• There is insufficient evidence to support or refute the conclusion that dronabinol is an effective treatment for the symptoms of irritable bowel syndrome. Some studies suggest that marijuana in the form of cannabidiol may be beneficial in the treatment of inflammatory bowel diseases, including Crohn's disease.

systemic review identified a single small trial that found no effect of two cannabinoids, given as an oro-mucosal spray, on intraocular pressure. The quality of evidence for the finding of no effect is limited. However, to be effective, treatments targeting lower intraocular pressure must provide continual rather than transient reductions in intraocular pressure. To date, those studies showing positive effects have shown only short-term benefit on intraocular pressure (hours), suggesting a

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Human Immunodeficiency Virus & Acquired Immunodeficiency Syndrome (Positive Status):

• There is limited evidence that cannabis and oral cannabinoids are effective in increasing appetite and decreasing weight loss associated with HIV/AIDS. There does not appear to be good-quality primary literature that reported on cannabis or cannabinoids as effective treatments for AIDS wasting syndrome.

Post-Traumatic Stress Disorder (PTSD):

• There is limited evidence (a single, small fair-quality trial) that cannabinoids are effective for improving symptoms of posttraumatic stress disorder. A single, small crossover trial suggests potential benefit from the pharmaceutical cannabinoid nabilone. This limited evidence is most applicable to male veterans and contrasts with non-randomized studies showing limited evidence of a statistical association between cannabis use (plant derived forms) and increased severity of posttraumatic stress disorder symptoms among individuals with posttraumatic stress disorder. There are other trials that are in the process of being conducted and if successfully completed, they will add substantially to the knowledge base.

Amyotrophic Lateral Sclerosis (ALS):

• There is insufficient evidence that cannabinoids are an effective treatment for symptoms associated with amyotrophic lateral sclerosis. Two small studies investigated the effect of dronabinol on symptoms associated with ALS. Although there were no differences from placebo in either trial, the sample sizes were small, the duration of the studies was short, and the dose of dronabinol may have been too small to ascertain any activity. The effect of cannabis was not investigated.

Parkinson's Disease:

• There is insufficient evidence that cannabinoids are an effective treatment for the motor system symptoms associated with Parkinson's disease or the levodopa- induced dyskinesia. Evidence suggests that the endocannabinoid system plays a meaningful role in certain neurodegenerative processes; thus, it may be useful to determine the efficacy of cannabinoids in treating the symptoms of neurodegenerative diseases. Small trials of oral cannabinoid preparations have demonstrated no benefit compared to a placebo in ameliorating the side effects of Parkinson's disease. A seven-patient trial of nabilone suggested that it improved the dyskinesia associated with levodopa therapy, but the sample size limits the interpretation of the data. An observational study demonstrated improved outcomes, but the lack of a control group and the small sample size are limitations.

Multiple Sclerosis (MS):

• There is substantial evidence that oral cannabinoids are an effective treatment for improving patient-reported multiple sclerosis spasticity symptoms, but limited evidence for an effect on clinician-measured spasticity. Based on evidence from randomized controlled trials included in systematic reviews, an oral cannabis extract, nabiximols, and orally administered THC are probably effective for reducing patient-reported spasticity scores in patients with MS. The effect appears to be modest. These agents have not consistently demonstrated a benefit on clinician-measured spasticity indices.

Patient Name:	Date of Birth:
The qualifying physician has provided	the patient or the patient's caregiver a summary of marijuana to treat the patient's medical
 Terminal conditions diagnosed by a physician certification The qualifying physician has provided of the current research on the efficace 	nysician other than the qualified physician issuing the patient or the patient's caregiver a summary by of marijuana to treat the patient's terminal
condition.	
The majority of studies on pain evaluated nabixing studies have evaluated the use of cannabis in cannabis in flower form provided by the National cannabis products that are sold in state-regulate that are available for research at the federal letopical forms. While the use of cannabis for the	an effective treatment for chronic pain in adults. mols outside the United States. Only a handful of a the United States, and all of them evaluated Institute on Drug Abuse. In contrast, many of the d markets bear little resemblance to the products vel in the United States. Pain patients also use treatment of pain is supported by well- controlled cy, dose, routes of administration, or side effects annabis products in the United States.
I have access to up to date information marijuana	regarding Oklahoma laws surrounding medical
	today is for my physician is to determine if it is mendation for medical marijuana in the State of
recommendation for a medical marijuana card i	oility to determine the appropriateness for a is based on my medical history, current medical sical exam, and medical records. I CERTIFY ALL ATE.
expect from their primary care provider. I under factors/conditions relating to a recommendation f	comprehensive medical evaluation as one would stand this evaluation is intended to focus on the for a medical marijuana recommendation. It is not reatment or recommendation of my primary care
I understand that it is my responsibility to approves of my intentions regarding the use of m	ensure my primary care provider is aware of and nedical marijuana.
I understand that my recommendation physician has the right to reverse a recommendation	is valid for two years after the issue date. My tion decision at his discretion.
I understand that the physician providing care for any and all matters related to my use of	this medical evaluation is available for follow up medical marijuana.
I have, or will, discuss my use of marijua	ana with my primary medical provider(s) before I

Patient Name:	Date	of Birth:
I do not have medication	abuse or drug abuse problems.	
I have not engaged in tra	afficking drugs or in drug diversion	on and will not do so
questions regarding anything I	may not understand or that I Nouh MD has informed me o	s with the physician and to ask believe needed to be clarified. I f the nature of a recommended arding medical marijuana.
benefits and risks have been d	iscussed. I understand no fees d to any insurance plan. Myself	questions have been addressed; associated with care or obtaining or my legal representative prior to
Signature:		_
Print Name:		_
Legal representative:		_
<u>Date</u> :		
Witness		
Witness: Print name:		
		_
<u>Date</u> :		
Provider:	· · · · · · · · · · · · · · · · · · ·	
Amer Nouh, MD		
Date:		