APPOINTED PHARMACY CONSENT

(buprenorphine HCl/naloxone HCl dihydrate) sublingual tablet or film (buprenorphine HCl) sublingual tablet, naltrexone (oral or extended-release injectable)

		-
<u></u>	do hereby: (check all that apply)
	Patient Name (Print)	
	Authorize at the above add treatment for opioid use disorder to employees of the pharmacy spectreatment disclosure most often includes, but may not be limited to medications with the pharmacist, and faxing/calling in my buprenor prescriptions directly to the pharmacy.	, discussing my
	Agree to purchase all buprenorphine/naloxone, and any other media treatment from the pharmacy specified below.	cations related to my
	Agree not to use any pharmacy other than the one specified below the treatment with the physician specified above, unless specific arrangemade with the physician.	
	Agree to make payment arrangements with the pharmacy specified treatment, so that my buprenorphine/naloxone prescriptions can be delivered to the physician's office address given above or picked-up same.	e filled and either
	I understand that I may withdraw this consent at any time, either very except to the extent that action has been taken in reliance on it. This I am being treated for opioid use disorder by the physician specified withdraw my consent during treatment. This consent will expire 365 my treatment, unless the provider specified above is otherwise notice.	s consent will last while above, unless I days after I complete
tre cor und (42	nderstand that the records to be released may contain information per eatment and/or substance use disorder treatment. These records may infidential information about communicable diseases including HIV/AI derstand that these records are protected by the Code of Federal Reg 2 CFR Part 2), which prohibits the recipient of these records from mak acclosures to third parties, without the express written consent of the p	also contain DS or related illness. I gulations Title 42 Part 2 ing any further
tre	cknowledge that I have been notified of my rights pertaining to the coeatment information/records under 42 CFR Part 2, and I further acknoderstand those rights.	•
Sig	gnature of patient	Date

Signature of parent/guardian/authorized signature	Date	
Signature of witness		Date
APPOINTED PHARMACY:		
NAME:	PHONE:	
ADDRESS:		

CONFIDENTIALITY OF SUBSTANCE USE DISORDER PATIENT RECORDS

THE CONFIDENTIALITY OF SUBSTANCE USE DISORDER PATIENT RECORDS MAINTAINED BY THIS PRACTICE/PROGRAM IS PROTECTED BY FEDERAL LAW AND REGULATIONS. GENERALLY, THE PRACTICE/PROGRAM MAY NOT SAY TO A PERSON OUTSIDE THE PRACTICE/PROGRAM THAT A PATIENT ATTENDS THE PRACTICE/PROGRAM, OR DISCLOSE ANY INFORMATION IDENTIFYING A PATIENT AS HAVING OR HAVING HAD A SUBSTANCE USE DISORDER UNLESS:

- 1. THE PATIENT CONSENTS IN WRITING;
- 2. THE DISCLOSURE IS ALLOWED BY A COURT ORDER, OR
- 3. THE DISCLOSURE IS MADE TO MEDICAL PERSONNEL IN A MEDICAL EMERGENCY OR TO QUALIFIED PERSONNEL FOR RESEARCH, AUDIT, OR PRACTICE/PROGRAM EVALUATION.

VIOLATION OF THE FEDERAL LAW AND REGULATIONS BY A PRACTICE/PROGRAM IS A CRIME. SUSPECTED VIOLATIONS MAY BE REPORTED TO APPROPRIATE AUTHORITIES IN ACCORDANCE WITH FEDERAL REGULATIONS. THE REPORT OF ANY VIOLATION OF THESE REGULATIONS MAY BE DIRECTED TO THE ATTORNEY GENERAL FOR YOUR STATE.

FEDERAL LAW AND REGULATIONS DO NOT PROTECT ANY INFORMATION ABOUT A CRIME COMMITTED BY A PATIENT, EITHER AT THE PRACTICE/PROGRAM OR AGAINST ANY PERSON WHO WORKS FOR THE PRACTICE/PROGRAM OR ABOUT ANY THREAT TO COMMIT SUCH A CRIME.

FEDERAL LAWS AND REGULATIONS DO NOT PROTECT ANY INFORMATION ABOUT SUSPECTED CHILD ABUSE OR NEGLECT FROM BEING REPORTED UNDER STATE LAW TO THE APPROPRIATE STATE OR LOCAL AUTHORITIES.

After completion, scan form into patient record and provide a copy to the patient.

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