

Patient Enrollment



COMPLETE ALL FIELDS TO AVOID PROCESSING DELAYS. PRESCRIPTION ONLY VALID IF FAXED. FAX COMPLETED FORM TO: 1-877-329-8484.

QUESTIONS? CALL 1-800-VIVITROL (1-800-848-4876), 9AM-8PM (EST).

V2G ID# (Vivitrol2gether Use Only): _____

Admittance Date: _____ Estimated Discharge Date: _____

1. PRESCRIBER OR FACILITY INFORMATION

Prescriber Name* _____
State License # _____ DEA # _____
Prescriber Phone # _____ NPI # _____
Facility Name _____ Fax # _____
Address _____
City _____ State _____ ZIP Code _____
Staff Contact Name _____
Staff Contact Phone # _____
Staff Contact E-mail _____

2. PATIENT INFORMATION

Name (First) _____ (Last) _____
Date of Birth _____ Gender Male Female
Address _____
City _____ State _____ ZIP Code _____
Home Phone # _____ Mobile Phone # _____
Best Number to Call Home Mobile
Best Day to Call M T W TH F
Best Time to Call Morning Afternoon Evening
E-mail Address _____

→ INSTRUCT PATIENT TO LIST ALTERNATE CONTACTS ON PAGE 2.

3. PATIENT DIAGNOSIS—Please complete the diagnosis code(s) you would like to use by filling in the additional digits.

(A list of codes can be found on page 3, section 12)

Alcohol Dependence ICD-10	Opioid Dependence ICD-10	Patient has tried and failed the following medication(s):
F10. _____	F11. _____	_____
F10. _____	F11. _____	_____
F10. _____	F11. _____	Please list any known allergies to medications or other substances:
F10. _____	F11. _____	_____
F10. _____	F11. _____	_____
Other _____	Other _____	Patient's concurrent medications:
<input type="checkbox"/> Check if patient has concurrent medications		_____

6. PRESCRIPTION INFORMATION AND ATTESTATION

*PRESCRIBER SIGNATURE MUST BE THE SAME AS THE PRESCRIBER NAME ABOVE

Patient Name _____
VIVITROL 380 mg x 1 unit Inject 380 mg IM every 4 weeks or every 1 month _____
Refill _____ times (Complete refills to minimize interruption in monthly VIVITROL therapy)
Provider State License # _____

By signing below, I verify that the information provided in this Vivitrol2gether enrollment form is complete and accurate to the best of my knowledge. I understand that Alkermes, Inc., reserves the right at any time and for any reason, without notice, to modify this Vivitrol2gether enrollment form or to modify or discontinue any services or assistance provided through Vivitrol2gether. Finally, I authorize Alkermes, its affiliates, representatives and agents as my designated agents to use and disclose my patient's health information as necessary to verify the accuracy of any information provided, to provide reimbursement services through Vivitrol2gether, to forward the above prescription, by fax or other mode of delivery, to a pharmacy for fulfillment, and (as applicable) to assess my patient's eligibility for co-pay assistance.

Prescriber's Signature _____
(If applicable) _____
Dispense as Written _____
Date of Signature _____
(no stamps allowed) _____
Substitution Permitted _____

PLEASE SEE IMPORTANT SAFETY INFORMATION ON PAGE 4. PLEASE SEE PRESCRIBING INFORMATION AND MEDICATION GUIDE, OR VISIT VIVITROL.COM. PLEASE REVIEW MEDICATION GUIDE WITH PATIENTS.

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PATIENTS SHOULD COMPLETE ALL FIELDS ON THIS PAGE.

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7. ALTERNATE PATIENT CONTACT(S)

By signing below, I authorize my Contact(s), listed below, to receive logistical and administrative information related to my treatment, such as appointment reminders, and to make decisions on my behalf—for which I will remain liable—regarding delivery of VIVITROL® (naltrexone for extended-release injectable suspension). Alkermes is not liable for any decision(s) made by the Contact(s) or actions taken in reliance on such Contact(s) decisions.

Please list any Contacts authorized as set forth above:

Contact Name (1)	Relationship	Phone #	<input type="checkbox"/> Home <input type="checkbox"/> Mobile <input type="checkbox"/> Work
Contact Name (2)	Relationship	Phone #	<input type="checkbox"/> Home <input type="checkbox"/> Mobile <input type="checkbox"/> Work
Patient's Signature X	Date of Signature	Phone #	<input type="checkbox"/> Home <input type="checkbox"/> Mobile <input type="checkbox"/> Work

8. PATIENT AUTHORIZATION FOR USE/DISCLOSURE OF HEALTH INFORMATION

By signing and printing my name below, I authorize: **1.** my prescribing healthcare provider, **2.** the healthcare provider who will administer VIVITROL to me, **3.** the pharmacy(ies) to which my VIVITROL prescription is sent for fulfillment (the "Pharmacy"), and **4.** my health plans and insurers (collectively, my "Healthcare Entities") to use and disclose to: **1.** Alkermes, Inc. and the companies working with Alkermes, Inc. to provide the VIVITROL patient support services I request, which are United BioSource Corporation, OPUS Health, LASH Group, Human Care Systems (collectively, "Alkermes") and **2.** my Contact(s) listed above (together with Alkermes, the "Recipients") health information related to my medical condition, including information about my drug or alcohol addiction, my mental health condition(s), my treatment with VIVITROL, my insurance coverage, as well as the information requested in this form (taken together, "Information") **for the specific purposes** of allowing Alkermes to facilitate: **1.** ordering, delivering and administering VIVITROL, **2.** conducting reimbursement verification and obtaining payment from my health plan(s) and insurer(s), **3.** providing me with educational and therapy support services by mail, text-messaging, e-mail, and/or telephone, which may include sending me product information materials, treatment reminders, and motivational messages, **4.** referring me to, or determining my eligibility for, other programs, foundations or alternative sources of funding or coverage to help me with the costs of VIVITROL and **5.** reviewing and analyzing fulfillment of VIVITROL prescriptions. **Information May Be Further Disclosed:** I understand that Information disclosed pursuant to this authorization could be re-disclosed by a Recipient and may no longer be protected by federal privacy law (HIPAA).

I understand that signing this authorization is voluntary and if I do not sign this authorization it will not affect my ability to obtain treatment, insurance or insurance benefits from my Healthcare Entities. I understand, however, that if I do not sign this authorization, I will not be eligible to receive the educational, patient support or other services described above, which are being provided by, or on behalf of, Alkermes. I will consult with my healthcare provider before making any treatment decisions. I understand I have the right to receive a copy of this authorization after I sign. I understand that the Pharmacy may receive payment from Alkermes, Inc. in exchange for Information.

I may withdraw this authorization at any time by mailing or faxing a written request to Vivitrol2gether, 852 Winter Street, Waltham, MA 02451. Withdrawal of this authorization will end my consent to further disclosures of Information authorized herein by my Healthcare Entities when they receive notice of my withdrawal, but will not affect previous disclosures and uses pursuant to this authorization or as permitted by applicable law. This authorization expires on the earlier of **(1)** five years from the date of signature below or **(2)** the maximum period permitted by applicable state law, unless I withdraw it earlier as set forth above.

Patient's Signature X	Print Name	Date of Signature
Guardian/Legal Representative Signature+ X	Authority/Relationship to Patient	

+If patient does not have capacity to act alone under state law, signature of guardian or authorized legal representative is required.

9. CO-PAY INFORMATION FOR ELIGIBLE PATIENTS†

- (Check if "yes") I would like to receive co-payment assistance from Alkermes.
- I certify that I am at least 18 years old, and I am being treated for opioid dependence after detox or alcohol dependence.

Please confirm that you understand the eligibility requirements for the VIVITROL® Co-pay Savings Program:

Co-payment assistance for VIVITROL is not valid for prescriptions that are purchased (in full or in part) by any federal, state or government funded program. Such programs include, but are not limited to:

- Medicare, including Medicare Part D and Medicare Advantage plans
- Medicaid, including Medicaid Managed Care and Alternative Benefit Plans ("ABPs") under the Affordable Care Act
- Medigap
- Veterans Administration ("VA")
- Department of Defense ("DoD")
- TRICARE®
- State-funded programs such as medical or pharmaceutical assistance programs

I agree that I understand the eligibility requirements described above. I certify that I am not using, and I will not use, any federal or state funded program to help pay for my VIVITROL prescription. I understand that if I do use any benefits from state, federal or other government funded program to help pay for my VIVITROL prescription, I will no longer be eligible for co-pay support for VIVITROL. I agree to comply with any requirements of my insurer(s) regarding co-pay support, including disclosure of the amount of co-pay support I receive to my insurer(s).

By signing below I agree that I understand, and will comply with, the terms of this VIVITROL® Co-pay Savings Program.

<input type="checkbox"/> YES <input type="checkbox"/> NO	Patient's Signature X	Date of Signature
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†Eligibility for Sponsored Co-pay Assistance: Please see page 3, section 11.

PLEASE SEE IMPORTANT SAFETY INFORMATION ON PAGE 4. PLEASE SEE PRESCRIBING INFORMATION AND MEDICATION GUIDE, OR VISIT VIVITROL.COM. PLEASE REVIEW MEDICATION GUIDE WITH PATIENTS.

10. INJECTION PROVIDER/SPECIALTY PHARMACY SELECTION INFORMATION (AS APPLICABLE)

If you have requested injection services for your patient, Vivitrol2gether will identify several injectors based on geographic proximity to your patient's address listed on the enrollment form (from closest to farthest from such address) and obtain information from Alkermes on pharmacy capabilities and time/rate of fulfillment for VIVITROL prescriptions.

These injection providers are listed on the VIVITROL Provider Locator[®] at VIVITROL.com.

These options will be provided to you for your patient. We will also contact the selected injection services provider to help coordinate injection services.

Upon request, prescriptions of patients enrolled in Vivitrol2gether are routed to qualified pharmacies based on insurance plan requirements, provider selection, patient preference and information obtained by Alkermes on pharmacy capability and performance in dispensing VIVITROL prescriptions. Participation is open to all qualified pharmacies free of charge. Interested pharmacies may contact 1-800-VIVITROL (1-800-848-4876).

11. ELIGIBILITY FOR SPONSORED CO-PAY ASSISTANCE

Offer valid only for prescriptions for FDA-approved indications. Patients must be at least 18 years old. If patients are purchasing their VIVITROL prescriptions with benefits from Medicare, including Medicare Part D or Medicare Advantage plans; Medicaid, including Medicaid Managed Care or Alternative Benefit Plans ("ABPs") under the Affordable Care Act; Medigap; Veterans Administration ("VA"); Department of Defense ("DoD"); TRICARE[®]; or any similar state funded programs such as medical or pharmaceutical assistance programs, they are not eligible for this offer. Void where prohibited by law, taxed or restricted. Alkermes, Inc. reserves the right to rescind, revoke or amend these offers without notice.

12. PATIENT DIAGNOSIS CODES

Alcohol Dependence:

(ICD-10)		
F10.2	Alcohol dependence	F10.27 With alcohol-induced persisting dementia
- F10.20	Uncomplicated	
- F10.21	In remission	
- F10.22	Alcohol dependence with intoxication	F10.28 Alcohol dependence with other alcohol-induced disorders
- F10.220	Uncomplicated	
- F10.221	Delirium	
- F10.229	Unspecified	- F10.280 Alcohol dependence with alcohol-induced anxiety disorder
F10.23	Alcohol dependence with withdrawal	- F10.281 Alcohol dependence with alcohol-induced sexual dysfunction
- F10.230	Uncomplicated	
- F10.231	Delirium	- F10.282 Alcohol dependence with alcohol-induced sleep disorder
- F10.232	With perceptual disturbance	
- F10.239	Unspecified	- F10.288 Alcohol dependence with other alcohol-induced disorder
F10.24	With alcohol-induced mood disorder	
F10.25	Alcohol dependence with alcohol-induced psychotic disorder	F10.29 With unspecified alcohol-induced disorder
- F10.250	With delusions	
- F10.251	With hallucinations	
- F10.259	Unspecified	
F10.26	With alcohol-induced persisting amnesic disorder	

Opioid Dependence:

(ICD-10)		
F11.2	Opioid dependence	F11.28 Opioid dependence with other opioid-induced disorder
- F11.20	Uncomplicated	
- F11.21	In remission	- F11.281 Opioid dependence with other opioid-induced sexual dysfunction
- F11.22	Opioid dependence with intoxication	
- F11.220	Uncomplicated	- F11.282 Opioid dependence with other opioid-induced sleep disorder
- F11.221	Delirium	
- F11.222	With perceptual disturbance	- F11.288 Opioid dependence with other opioid-induced disorder
- F11.229	Unspecified	
F11.23	With withdrawal	F11.29 With unspecified opioid-induced disorder
F11.24	With opioid-induced mood disorder	
F11.25	Opioid dependence with opioid-induced psychotic disorder	
- F11.250	With delusions	
- F11.251	With hallucinations	
- F11.259	Unspecified	

Inclusion in the Locator is voluntary and free of charge to qualified healthcare providers and, along with the provider-specific information in the Provider Locator, is based on healthcare provider responses. Inclusion in the Locator does not imply a referral, recommendation, or endorsement by Alkermes. Alkermes has not independently verified the qualifications of any healthcare provider included in the Locator. We recommend that you research the credentials, qualifications, and experience of each provider before confirming an appointment. Alkermes shall not be liable to you or to anyone for any decision made or action taken in reliance on this information.

IMPORTANT SAFETY INFORMATION FOR VIVITROL® (NALTREXONE FOR EXTENDED-RELEASE INJECTABLE SUSPENSION)

INDICATIONS

VIVITROL is indicated for:

- Treatment of alcohol dependence in patients who are able to abstain from alcohol in an outpatient setting prior to initiation of treatment with VIVITROL. Patients should not be actively drinking at the time of initial VIVITROL administration.
- Prevention of relapse to opioid dependence, following opioid detoxification.
- VIVITROL should be part of a comprehensive management program that includes psychosocial support.

CONTRAINDICATIONS

VIVITROL is contraindicated in patients:

- Receiving opioid analgesics
- With current physiologic opioid dependence
- In acute opioid withdrawal
- Who have failed the naloxone challenge test or have a positive urine screen for opioids
- Who have exhibited hypersensitivity to naltrexone, polylactide-co-glycolide (PLG), carboxymethylcellulose, or any other components of the diluent

WARNINGS AND PRECAUTIONS

Vulnerability to Opioid Overdose:

- After opioid detoxification, patients are likely to have a reduced tolerance to opioids. VIVITROL blocks the effects of exogenous opioids for approximately 28 days after administration. As the blockade wanes and eventually dissipates completely, use of previously tolerated doses of opioids could result in potentially life-threatening opioid intoxication (respiratory compromise or arrest, circulatory collapse, etc.).
- Cases of opioid overdose with fatal outcomes have been reported in patients who used opioids at the end of a dosing interval, after missing a scheduled dose, or after discontinuing treatment. Patients and caregivers should be told of this increased sensitivity to opioids and the risk of overdose.
- Although VIVITROL is a potent antagonist with a prolonged pharmacological effect, the blockade produced by VIVITROL is surmountable. The plasma concentration of exogenous opioids attained immediately following their acute administration may be sufficient to overcome the competitive receptor blockade. This poses a potential risk to individuals who attempt, on their own, to overcome the blockade by administering large amounts of exogenous opioids.
- Any attempt by a patient to overcome the VIVITROL blockade by taking opioids may lead to fatal overdose. Patients should be told of the serious consequences of trying to overcome the opioid blockade.

Injection Site Reactions:

- VIVITROL injections may be followed by pain, tenderness, induration, swelling, erythema, bruising, or pruritus; however, in some cases injection site reactions may be very severe.
- Injection site reactions not improving may require prompt medical attention, including, in some cases, surgical intervention.
- Inadvertent subcutaneous/adipose layer injection of VIVITROL may increase the likelihood of severe injection site reactions.
- Select proper needle size for patient body habitus, and use only the needles provided in the carton.
- Patients should be informed that any concerning injection site reactions should be brought to the attention of their healthcare provider.

Precipitation of Opioid Withdrawal:

- When withdrawal is precipitated abruptly by administration of an opioid antagonist to an opioid-dependent patient, the resulting withdrawal syndrome can be severe. Some cases of withdrawal symptoms have been severe enough to require hospitalization, and in some cases, management in the ICU.

- To prevent occurrence of precipitated withdrawal, opioid-dependent patients, including those being treated for alcohol dependence, should be opioid-free (including tramadol) before starting VIVITROL treatment:
 - An opioid-free interval of a minimum of 7-10 days is recommended for patients previously dependent on short-acting opioids.
 - Patients transitioning from buprenorphine or methadone may be vulnerable to precipitated withdrawal for as long as two weeks.
- If a more rapid transition from agonist to antagonist therapy is deemed necessary and appropriate by the healthcare provider, monitor the patient closely in an appropriate medical setting where precipitated withdrawal can be managed.
- Patients should be made aware of the risk associated with precipitated withdrawal and be encouraged to give an accurate account of last opioid use.

Hepatotoxicity:

- Cases of hepatitis and clinically significant liver dysfunction have been observed in association with VIVITROL. Warn patients of the risk of hepatic injury; advise them to seek help if experiencing symptoms of acute hepatitis. Discontinue use of VIVITROL in patients who exhibit acute hepatitis symptoms.

Depression and Suicidality:

- Alcohol- and opioid-dependent patients taking VIVITROL should be monitored for depression or suicidal thoughts. Alert families and caregivers to monitor and report the emergence of symptoms of depression or suicidality.

When Reversal of VIVITROL Blockade Is Required for Pain Management:

- For VIVITROL patients in emergency situations, suggestions for pain management include regional analgesia or use of non-opioid analgesics. If opioid therapy is required to reverse the VIVITROL blockade, patients should be closely monitored by trained personnel in a setting staffed and equipped for CPR.

Eosinophilic Pneumonia:

- Cases of eosinophilic pneumonia requiring hospitalization have been reported. Warn patients of the risk of eosinophilic pneumonia and to seek medical attention if they develop symptoms of pneumonia.

Hypersensitivity Reactions:

- Patients should be warned of the risk of hypersensitivity reactions, including anaphylaxis.

Intramuscular Injections:

- As with any IM injection, VIVITROL should be administered with caution to patients with thrombocytopenia or any coagulation disorder.

Alcohol Withdrawal:

- Use of VIVITROL does not eliminate nor diminish alcohol withdrawal symptoms.

ADVERSE REACTIONS

- Serious adverse reactions that may be associated with VIVITROL therapy in clinical use include severe injection site reactions, eosinophilic pneumonia, serious allergic reactions, unintended precipitation of opioid withdrawal, accidental opioid overdose, and depression and suicidality.
- The adverse events seen most frequently in association with VIVITROL therapy for alcohol dependence (ie, those occurring in ≥5% and at least twice as frequently with VIVITROL than placebo) include nausea, vomiting, injection site reactions (including induration, pruritus, nodules, and swelling), muscle cramps, dizziness or syncope, somnolence or sedation, anorexia, decreased appetite or other appetite disorders.
- The adverse events seen most frequently in association with VIVITROL in opioid-dependent patients (ie, those occurring in ≥2% and at least twice as frequently with VIVITROL than placebo) were hepatic enzyme abnormalities, injection site pain, nasopharyngitis, insomnia, and toothache.

You are encouraged to report side effects to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

PLEASE SEE [PRESCRIBING INFORMATION](#) AND [MEDICATION GUIDE](#), OR VISIT VIVITROL.COM. PLEASE REVIEW MEDICATION GUIDE WITH PATIENTS.



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