FORBES PHARMACY BRIXADI REFERRAL

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In order to initiate Brixadi treatment, please send a valid prescription to Forbes Pharmacy along with a completed referral form.

REFERRAL SOURCE								
Name/Title of Person Completing Form:				Date:				
PATIENT INFORMATION								
Last Name: First Na		ame:		DOB:		Sex: M F	Facility:	
Address:			City:	City:		State:	ZIP:	
Cell Phone:	Home Phone:			Room/Unit N		lumber (Inpatient):		
Emergency Contact:			Pho	Phone Number:				
INJECTION PLAN								
Outpatient Injection CPCDS Clinic Contact			_	Magee Mercy NATP Other				
		inic (Name & Phone Number)						
In accordance with the REMS, Brixadi injection may only be delivered to a healthcare setting and is never dispensed to a patient directly. Brixadi injection will only be used for the patient it is ordered and will NOT be transported to any other location/office/person.								
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CLINICAL INFORMATION								
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CLINICAL INFORMATION	- <i>plea</i> : I with E	se include ap Brixadi in the p mended test (oplicab past? [<i>le clii</i> □ NO	nical chart n	otes e of Last Injection		
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CLINICAL INFORMATION Has the patient been treated Has the patient received the buprenorphine-containing p	- please I with E recom product	se include ap Brixadi in the p mended test (oplicab past? (dose of a	le clii NO	nical chart n YES - Dat g transmucos	otes e of Last Injection al	: NO YES	
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PRESCRIPTION INFORMATION

Please send an electronic prescription to UPMC Forbes Pharmacy (3501 Forbes Ave, Oxford Buillding, Room 756)

The recommended dose of Brixadi is dependent on the current dose of a patient's transmucosal buprenorphine-containing product. Brixadi should be administered in 7-day intervals for those on weekly regimens, and 28-day intervals for those on monthly regimens. Please see the BRIXADI Full Prescribing Information, including Boxed Warning, at brixadihcp.com

- *For abdominal subcutaneous injection only. Do not administer intravenously or intramuscularly.
- Examine the injection site for signs of infection or evidence of tampering or attempts to remove the depot.
- Prescription use of this product is limited by the Drug Addiction Treatment Act (DATA) to prescribers who are authorized to treat opioid dependence and are DATA 2000 waivered.
- Brixadi may only be delivered to a healthcare setting and is NEVER dispensed to a patient directly
- Serious harm or death could result if administered intravenously. Braxadi forms a solid mass upon contact with body fluids and may cause occlusion, local tissue damage and thrombo-embolic events, including life-threatening pulmonary emboli, if administered intravenously
- Because of the risk of serious harm or death that could result from intravenous self-administration, Brixadi is only available through a restricted program called the Brixadi REMS Program. Healthcare settings and pharmacies that order and dispense Brixadi must be certified in this program and comply with the REMS requirements.