FAX this page

PATIENT AUTHORIZATION FOR VA PATIENTS





Complete these forms if you are electing Circle of Care™ Services.	injection, for subcutaneous use * 4.9 mg/mL			
Patient Name:	DOB:			
Please read the following carefully, then sign and date below.				
I. Patient Authorization for Release of Health Information Form				
By signing this Authorization, I authorize my healthcare provider, my health ("Healthcare Entities") and each of their respective representatives, emploinformation relating to my medical condition, treatment, and prescription Pharmaceuticals and its agents, contractors, and other partners, including as Circle of Care TM (collectively, "Supernus") for Supernus to (i) provide me of Supernus' products, including, but not limited to, educational support padherence support ("Programs"), (ii) conduct data analytics, market resea limited to, evaluating the Programs provided, and (iii) provide me with info and other topics of interest for marketing, educational or other purposes.	oyees, and agents (collectively "Providers") to disclose details ("Protected Health Information" or "PHI") to Supernus companies working with Supernus, which may be branded with support and related information and materials on any provided in-person, online or by telephone, and medication rich and other internal business activities including, but not			
For purposes of clarification, Supernus includes but is not limited to brand specific support through a hub service provider, specialty pharmacy service providers, Circle of Care Clinical Nurse Navigators, as well as other entities under contract with Supernus to support these or similar aspects of the Programs. For purposes of providing support through the Programs, I thereby authorize Supernus to contact me via text messaging, phone, fax, and/or mail – including texts and calls made using an automatic telephone dialing system or prerecorded or artificial voice messages – and to leave a detailed message that includes reference to ONAPGO treatment, as needed.				
Once my PHI has been disclosed to Supernus, I understand that state and federal privacy laws no longer protect the information. However, Supernus agrees to protect my PHI by using and disclosing it only for purposes authorized in this Authorization or as required by law or regulations. I understand that my pharmacy provider may receive remuneration from Supernus in exchange for the PHI and/or for any support services provided to me.				
I understand that I am not required to sign this Authorization and that my lenrollment, or eligibility for benefits on whether I sign this Authorization. Trequired by state law, unless I revoke it sooner by writing Circle of Care/Si I understand that revoking my Authorization will not affect any use of my in am entitled to a copy of this signed authorization. I certify the information of my knowledge and I understand that Supernus can revise, change or te	his Authorization will expire in 10 years or a shorter period if upernus, c/o PharmaCord, PO Box 5490, Louisville, KY 40255. Information that occurred before my request was processed. I provided on this form is complete and accurate, to the best			
I have read and understand the Patient Authorization for Release of Hea is required in order to receive Supernus services which are not a part of Veterans Affairs.				
SN HERE				
Signature of Patient	Date			
In addition, I authorize the disclosure of my Protected Health Inform	ation to the following designated individual(s) (ontional):			
in addition, rabilionize the disclosure of my Protected Health informs	ation to the following designated marriadal(s) (optional).			
Designated Individual (print name)	Relationship			
II. Marketing/Other Communications Opt-In (optional)				
I further authorize Supernus, and companies working with Supernus, any o "Supernus"), to contact me by mail, email, fax, telephone call, and text me with information about Supernus' products, services, and programs or oth ask me about my experience with or thoughts about such topics. I undersused by Supernus to help develop new products, services, and programs. data to any unrelated third party for marketing purposes without my expressions. I have read and understand the Marketing/Other Communications Opt-	essage for marketing purposes or otherwise provide me her topics of interest, conduct market research or otherwise tand and agree that any information that I provide may be Note that Supernus will not sell or transfer my personal less permission.			
Signature of Patient:	_ Date:			
Designated Individual (print name):	_ Relationship:			

NP.ONA.2025-0001

CIRCLE OF CARE™ PROGRAM SUPPORT



Complete these forms if you are electing Circle		(apomorphine injection, for subcutaneous	
PATIENT INFORMATION			
Patient Name:	Date of Birth:	Sex: Male	Female
address:	City:	State: ZIP:	
Daytime Phone:	Evening Phone:		
mail:	Preferred Patient Language: Engl	ish Spanish Other:_	
are Partner/Alternate Contact:			
elationship to Patient:	Best time to	o contact: Morning	Afternoo
Prescribing Doctor Information			
irst Name:	Last Name:		
/A Facility:			
City: State: ZIF	P: Phone:		
The Circle of Care Program is optional and not reselect A or B below.	equired for ONAPGO. You are not required to e	nroll. If you choose to enr	oll,
will occur in your doctor's office. Prior to expectations, work to establish proficience daily administration and setup, and train a prescribers' orders. Ongoing Support and time you are on ONAPGO. The Clinical Nurse Naministration, and setup. Clinical Nurse Naministration.	ect A, your ONAPGO Start (Initiation) and subs starting ONAPGO, your Circle of Care Clinical N by with the ONAPGO device, review the Patient and educate on skin health. You will be taught had Education will be provided through Dose Titratures Navigator will continue to educate on device Navigators do not give medical advice about a patient of that specific medical advice. Education	Jurse Navigator will review Instructions for Use includ ow to use ONAPGO per yo tion and ongoing for the e e proficiency, skin health, patient's personal treatmen	ling our ntire daily nt plan
Education and Support as outlined above Visits will occur in your home. Your Circle Start appointment. During your Start appo with your prescribers' orders. The Clinical Nurse Navigator will educate on skin health Titrations, the Clinical Nurse Navigator will prescribers' orders. Dose Titration will occur	in box A, and your ONAPGO Start (Initiation) at of Care Clinical Nurse Navigator will schedule are interest, you will be instructed and trained on a Nurse Navigator will record your response to ONA, future dose titration and optimization, and dail schedule and coordinate your ONAPGO Titration or in your home and the ONAPGO continuous do ions B and C on the prescription form. Titration of the prescription form.	nd subsequent Dose Titrated coordinate your ONAPG dministering ONAPGO con NAPGO. In addition, your Cily administration. For your on Visits consistent with yoursages and extra dose will	tion GO nsistent Clinical r Dose our be

If in-home Start and Dose Titration support and education by the Supernus Circle of Care Clinical Nurse Navigator is requested by the patient, a copy of the ONAPGO Prescription Form for VA Patients must be faxed to 1-888-525-2431.

dose has been reached. Device setting will be set by your prescriber and/or a Supernus/MDD US Operations Clinical Nurse

SIGN	HERE		
	Signature	Date	

Navigator based on selections your prescriber made in sections B and C on the prescription form.

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PATIENT AUTHORIZATION FOR VA PATIENTS



Patient Copy

I. Patient Authorization for Release of Health Information Form

By signing this Authorization, I authorize my healthcare provider, my health insurance company, and my pharmacy providers ("Healthcare Entities") and each of their respective representatives, employees, and agents (collectively "Providers") to disclose information relating to my medical condition, treatment, and prescription details ("Protected Health Information" or "PHI") to Supernus Pharmaceuticals and its agents, contractors, and other partners, including companies working with Supernus, which may be branded as Circle of CareTM (collectively, "Supernus") for Supernus to (i) provide me with support and related information and materials on any of Supernus' products, including, but not limited to, educational support provided in-person, online or by telephone, and medication adherence support ("Programs"), (ii) conduct data analytics, market research and other internal business activities including, but not limited to, evaluating the Programs provided, and (iii) provide me with information about Supernus' products, services, and programs and other topics of interest for marketing, educational or other purposes.

For purposes of clarification, Supernus includes but is not limited to brand specific support through a hub service provider, specialty pharmacy service providers, Circle of Care Clinical Nurse Navigators, as well as other entities under contract with Supernus to support these or similar aspects of the Programs. For purposes of providing support through the Programs, I thereby authorize Supernus to contact me via text messaging, phone, fax, and/or mail – including texts and calls made using an automatic telephone dialing system or prerecorded or artificial voice messages – and to leave a detailed message that includes reference to ONAPGO treatment, as needed.

Once my PHI has been disclosed to Supernus, I understand that state and federal privacy laws no longer protect the information. However, Supernus agrees to protect my PHI by using and disclosing it only for purposes authorized in this Authorization or as required by law or regulations. I understand that my pharmacy provider may receive remuneration from Supernus in exchange for the PHI and/or for any support services provided to me.

I understand that I am not required to sign this Authorization and that my Providers will not condition my treatment, payment, enrollment, or eligibility for benefits on whether I sign this Authorization. This Authorization will expire in 10 years or a shorter period if required by state law, unless I revoke it sooner by writing Circle of Care/Supernus, c/o PharmaCord, PO Box 5490, Louisville, KY 40255. I understand that revoking my Authorization will not affect any use of my information that occurred before my request was processed. I am entitled to a copy of this signed authorization. I certify the information provided on this form is complete and accurate, to the best of my knowledge and I understand that Supernus can revise, change or terminate the Programs at any time.

I have read and understand the Patient Authorization for Release of Health Information Form and agree to the terms. A signature is required in order to receive Supernus services which are not a part of, endorsed by, or administered by the U.S. Department of Veterans Affairs.

II. Marketing/Other Communications Opt-In (optional)

I further authorize Supernus, and companies working with Supernus, any of which may be branded as Circle of Care (collectively "Supernus"), to contact me by mail, email, fax, telephone call, and text message for marketing purposes or otherwise provide me with information about Supernus' products, services, and programs or other topics of interest, conduct market research or otherwise ask me about my experience with or thoughts about such topics. I understand and agree that any information that I provide may be used by Supernus to help develop new products, services, and programs. Note that Supernus will not sell or transfer my personal data to any unrelated third party for marketing purposes without my express permission.

I have read and understand the Marketing/Other Communications Opt-In and agree to the terms.



USE

ONAPGO is a prescription medicine used to treat motor fluctuations (OFF episodes) in adults with advanced Parkinson's disease (PD). It is not known if ONAPGO is safe and effective in children.

IMPORTANT SAFETY INFORMATION

Do not take ONAPGO if you are:

- taking certain medicines to treat nausea (ondansetron, granisetron, dolasetron, palonosetron) and alosetron. People taking ondansetron with apomorphine had very low blood pressure and lost consciousness (blacked out).
- allergic to apomorphine or to any ingredients in ONAPGO including sulfite. Sulfites can cause severe, life-threatening allergic reactions, especially in people with asthma.

Call your healthcare provider or get emergency help right away if you have any of the following symptoms of severe life-threatening allergic reaction:

• hives • itching • rash • swelling (eyes, tongue, lips, or mouth) • chest pain • throat tightness • trouble breathing or swallowing.

Before you start using ONAPGO, tell your healthcare provider about all of your medical conditions, including:

• difficulty staying awake during the daytime • dizziness, fainting spells, or low blood pressure • asthma • allergies to any medicines containing sulfites • heart problems • a history of stroke or other brain problems • kidney problems • liver problems • a mental problem called a major psychotic disorder • drinking alcohol • if you are pregnant or plan to become pregnant, or breastfeeding or plan to breastfeed. It is not known if ONAPGO will harm your unborn baby or pass into your breast milk.

Tell your healthcare provider about all the medicines you take, including prescription and non-prescription (over-the-counter) medicines, vitamins, and herbal supplements. ONAPGO and certain other medicines may affect each other and cause serious side effects.

• If you take nitroglycerin under your tongue (sublingual) while using ONAPGO, your blood pressure may decrease and cause dizziness. If possible, lie down before taking it and then try to continue lying down for at least 45 minutes after.

What should I avoid while using ONAPGO?

- Do not drink alcohol. It can increase your chance of developing serious side effects.
- Do not take medicines that make you sleepy.
- Do not drive, operate machinery, or do other dangerous activities until you know how ONAPGO affects you.
- Do not change your position too fast, get up slowly from sitting or lying. ONAPGO can lower blood pressure and cause dizziness or fainting.

What are the possible side effects of ONAPGO?

ONAPGO may cause serious side effects, including:

- blood clots. Infusing ONAPGO into a vein (intravenous) can cause blood clots. Do not infuse ONAPGO in your vein.
- nausea and vomiting are common. May be serious or severe. Your healthcare provider may prescribe medicine (trimethobenzamide) to help decrease nausea/vomiting. Follow your healthcare provider's instructions on how to take/when to stop this medicine.
- sleepiness or falling asleep during the day is common and may be serious. Some people may get sleepy during the day or fall asleep without warning while doing everyday activities such as talking, eating, or driving.
- dizziness is common and may be serious. ONAPGO can lower your blood pressure and cause dizziness. Dizziness can happen when treatment is started or when the dose is increased. **Do not** get up too fast from sitting or lying down, especially if you have been sitting or lying down for a long time.
- falls. Changes that can happen with PD, and effects of some PD medicines, including ONAPGO, as well as trimethobenzamide, can increase your risk of falling.
- infusion site reaction is common and may be serious. Reactions and infections including infusion site nodules, redness, bruising, swelling, rash, and itching may happen.
- hallucinations or psychotic-like behavior. ONAPGO can cause/worsen psychotic-like behavior including hallucinations (seeing or hearing things that are not real), confusion, excessive suspicion, aggressive behavior, agitation, delusional beliefs (believing things that are not real), and disorganized thinking.
- sudden uncontrolled movements (dyskinesia) are common and may be serious. Some people with PD may get sudden, uncontrolled movements after treatment with some PD medicines. ONAPGO can cause/make dyskinesia worse.
- low red blood cells (hemolytic anemia). Tell your healthcare provider if you have: become pale, fast heartbeat, feel more tired or weaker than usual, skin or eyes look yellow, chest pain, shortness of breath or trouble breathing, dark-colored urine, fever, dizziness, or confusion.
- strong (intense) urges. New or increased gambling urges, sexual urges, and other intense urges have been reported.
- heart problems. If you have shortness of breath, fast heartbeat, or chest pain, call your healthcare provider or get emergency help right away.
- serious heart rhythm changes (QT prolongation). Tell your healthcare provider right away if you have a change in your heartbeat (a fast or irregular heartbeat), or faint.
- allergic reaction. Tell your healthcare provider or get medical help right away if you get hives, itching, rash, swelling of the eyes and tongue, or trouble breathing.
- tissue changes (fibrotic complications). Some people have had changes in the tissues of their pelvis, lungs, and heart valves when taking medicines called non-ergot derived dopamine agonists like ONAPGO.
- prolonged painful erections (priapism). May occur. If you have an erection that lasts more than 4 hours, call your healthcare provider or go to the nearest hospital emergency room right away.

Other common side effects of ONAPGO include headache and trouble falling asleep or staying asleep (insomnia).

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

Patients and care partners must receive complete instructions on the proper use of ONAPGO. Please see Patient Information and talk to your healthcare provider.



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