

Specialty Pharmacy Request Form

Complete the form and fax to chosen Specialty Pharmacy.
Please give page 2 to the patient.

SPECIALTY PHARMACY (Choose one)			
Specialty Pharmacy	Fax	Phone	Hours of Operation
<input type="checkbox"/> Biologics by McKesson	1-855-215-5315	1-888-275-8596	Mon-Fri 9:00 AM - 6:00 PM ET
<input type="checkbox"/> CenterWell Pharmacy (formerly Humana Pharmacy)	1-877-405-7940	1-800-486-2668	Mon-Fri 8:00 AM - 11:00 PM ET, Sat 8:00 AM - 6:30 PM ET
<input type="checkbox"/> Giannotto's Specialty Pharmacy	1-844-587-9626	1-973-482-8220 ext. 319	Mon-Fri 9:00 AM - 7:00 PM ET, Sat 9:00 AM - 4:00 PM ET

PATIENT INFORMATION	PRESCRIBER INFORMATION
Patient Name: _____ Address: _____ City: _____ State: _____ Zip: _____ Home Phone: _____ Cell Phone: _____ Date of Birth: _____ <input type="checkbox"/> See Attached Demographic Sheet	Prescriber Name: _____ State Lic #: _____ NPI #: _____ Specialty: _____ Facility Name: _____ Address: _____ City: _____ State: _____ Zip: _____ Ship To Address (Required): City: _____ State: _____ Zip: _____ Prescriber's Phone: _____ Prescriber's Fax: _____ PREFERRED COMMUNICATION Office Contact Name: _____ Direct Phone Number: _____ Direct Email Address: _____ Direct Fax: _____

INSURANCE INFORMATION (Please attach copies of front & back of cards)

N/A (Patient Self-Pay)

Primary Insurance	Secondary Insurance	Rx Card (PRM)
City: _____ State: _____	City: _____ State: _____	PBM BIN: _____
Plan #: _____	Plan #: _____	City: _____ State: _____
Group #: _____	Group #: _____	Group #: _____
Phone #: _____	Phone #: _____	Phone #: _____
Subscriber Name (First/Last): _____	Subscriber Name (First/Last): _____	Subscriber Name (First/Last): _____
ID #: _____	ID #: _____	ID #: _____
Employer: _____	Employer: _____	Employer: _____

PRESCRIPTION INFORMATION	DIAGNOSTIC INFORMATION (ICD-10 Code)
<input type="checkbox"/> PAR T380A with Single Hand Inserter – QTY 1 Paragard (intrauterine copper contraceptive) to be inserted one time by prescriber.	<input type="checkbox"/> Z30.430: Encounter for insertion of intrauterine contraceptive device. <input type="checkbox"/> Other: Please Specify _____ <input type="checkbox"/> Z30.014: Encounter for initial prescription of intrauterine contraceptive device. This code includes the initial prescription of the IUD, counseling, and advice, but excludes the IUD insertion. _____

If patient is a minor and is signing the authorization on the following page on her own behalf, please affirm that:

This patient has the capacity to consent to treatment with Paragard under the law of the state in which I practice (and the consent of a parent or guardian is not required), or

This patient's parent or guardian has consented to the patient's treatment with Paragard, as required by applicable state law.

I understand that my signature will be used as an approval allowing the Specialty Pharmacy to dispense Paragard. If I have a financial responsibility for obtaining Paragard, I understand that the selected specialty pharmacy will contact me prior to the dispense.

Patient Signature: _____ **Date:** ____/____/____

Prescriber Signature: _____ **Date:** ____/____/____

If Applicable for NP and PA with collaborative physician agreement: _____ **Date:** ____/____/____

Dear Patient,

Your healthcare provider has ordered Paragard through the following specialty pharmacy. This specialty pharmacy may contact you regarding Paragard, or you may contact them directly if you have any questions.

Specialty Pharmacy	Phone Number
<input type="checkbox"/> Biologics by McKesson	1-888-275-8596
<input type="checkbox"/> CenterWell Pharmacy (formerly Humana Pharmacy)	1-800-486-2668
<input type="checkbox"/> Giannotto's Specialty Pharmacy	1-973-482-8220 ext. 319

To learn more visit [Paragard.com](https://www.paragard.com)

INDICATIONS AND USAGE

Paragard is a copper-containing intrauterine system (IUS) indicated for prevention of pregnancy in females of reproductive potential for up to 10 years.

CONTRAINDICATIONS

The use of Paragard is contraindicated when one or more of the following conditions exist:

- Pregnancy or suspicion of pregnancy, abnormalities of the uterus resulting in distortion of the uterine cavity, acute pelvic inflammatory disease (PID), postpartum or postabortal endometritis in the past 3 months, known or suspected uterine or cervical malignancy, uterine bleeding of unknown etiology, untreated acute cervicitis or vaginitis or other lower genital tract infection, conditions associated with increased susceptibility to pelvic infections, Wilson's disease, a previously placed IUS that has not been removed, hypersensitivity to any component of Paragard including to copper or any of the trace elements present in the copper component of Paragard.

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS

Ectopic Pregnancy: Evaluate for possible ectopic pregnancy in any female who becomes pregnant while using Paragard.

Intrauterine Pregnancy: Failure to remove Paragard increases the risk of miscarriage, sepsis, premature labor, and premature delivery.

Sepsis: Severe infection or sepsis, including Group A Streptococcal Sepsis (GAS), have been reported following insertion of IUSs, including Paragard.

Pelvic Inflammatory Disease and Endometritis: Remove Paragard in cases of recurrent PID or endometritis, or if an acute pelvic infection is severe or does not respond to treatment.

Embedment: Partial penetration or embedment of Paragard in the myometrium can make removal difficult; surgical removal may be necessary. Breakage of an embedded Paragard during non-surgical removal has been reported.

Perforation: Partial or total perforation of the uterine wall or cervix may reduce contraceptive efficacy and result in pregnancy.

Delayed detection or removal of Paragard may result in migration outside the uterine cavity, adhesions, peritonitis, intestinal penetration, intestinal obstruction, abscesses and/or damage to adjacent organs. Increased risk when the uterus is fixed, retroverted or not completely involuted during the postpartum period. If perforation does occur, locate and remove Paragard promptly.

Expulsion: Partial or complete expulsion of Paragard has been reported, resulting in the loss of contraceptive protection. The risk of expulsion may be increased when the uterus is not completely involuted at the time of insertion. Remove a partially expelled Paragard.

Wilson's Disease: Paragard may exacerbate Wilson's disease.

Bleeding Pattern Alterations: Paragard can alter the bleeding pattern and result in heavier and longer menstrual cycles with intermenstrual spotting.

Magnetic Resonance Imaging (MRI) Safety Information: Non-clinical testing has demonstrated that Paragard is MR Conditional.

Medical Diathermy: Avoid using high medical RF transmitter devices in females with Paragard.

ADVERSE REACTIONS

Adverse reactions reported in clinical trials include anemia, backache, dysmenorrhea, dyspareunia, expulsion (complete or partial), prolonged menstrual flow, menstrual spotting, pain and cramping, and vaginitis.