WHAT'S THE PLAN?
MANAGING POTENTIAL SIDE EFFECTS

Rubraca can cause side effects. This guide will help you work with your healthcare provider to manage side effects that may occur during your treatment.

What is Rubraca used for?

Rubraca is a prescription medicine used for:

- The maintenance treatment of adults with ovarian cancer, fallopian tube cancer, or primary peritoneal cancer whose cancer has come back and who are in response (complete or partial response) to a platinum-based chemotherapy
- The treatment of adults with ovarian cancer, fallopian tube cancer, or primary peritoneal cancer who have certain "BRCA" gene mutations, either inherited (germline) or acquired (somatic), and who have been treated with 2 or more chemotherapy medicines for their cancer. Your healthcare provider will perform a test to make sure Rubraca is right for you

It is not known if Rubraca is safe and effective in children.

Please see additional Select Important Safety Information throughout this document.

Rubraca® (rucaparib) 300 mg tablets

What you can expect

Create a dosing schedule to help you remember to take your Rubraca

_____ AM

Two _____ mg tablets
in the morning

About **12** hours
after the first dose

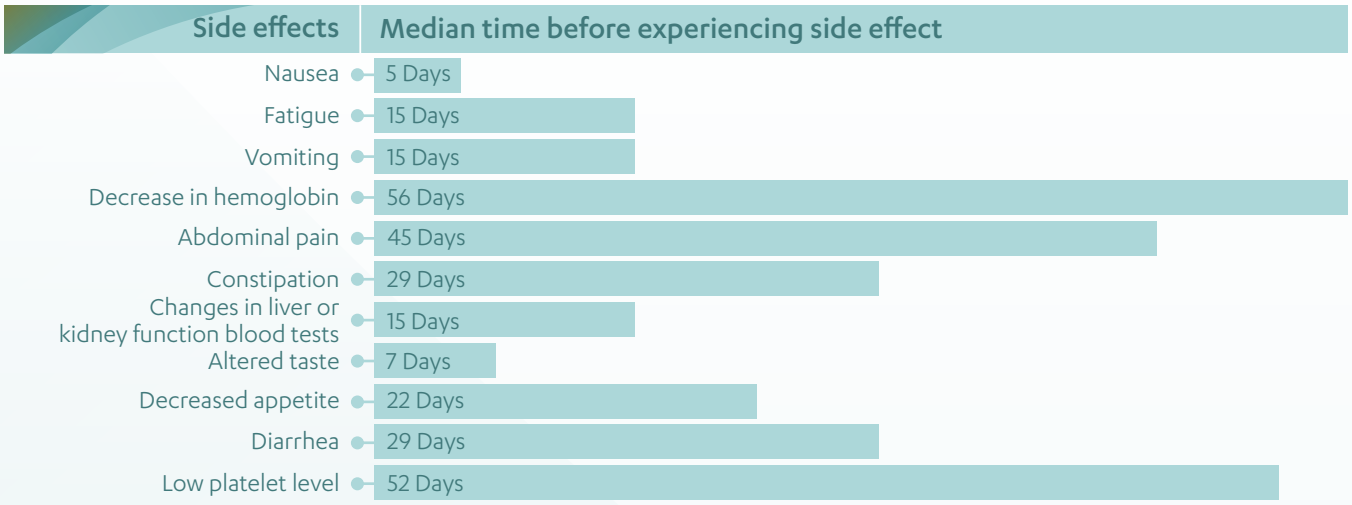
_____ PM

Two _____ mg tablets
in the evening

Rubraca affects each person differently.

The chart below lists the side effects associated with taking Rubraca. Side effects are listed in order starting with the most commonly experienced by patients in clinical trials. Next to each side effect is the amount of time the average patient started experiencing a side effect after starting treatment with Rubraca.

Please note that you may experience other side effects not listed here. It's important to keep your oncology care team updated about any changing health issues.



Median is defined as the middle number in a group of numbers arranged from the lowest to the highest.

Your oncologist may lower your dose or tell you to stop taking Rubraca for a time; this is a way to help manage side effects.

Rubraca tablets come in 3 strengths, each with its own color

300 mg | 250 mg | 200 mg

Images do not reflect actual size of each tablet.

- Take tablets with or without food
- If a dose is missed take the next dose at its scheduled time
- If a dose is vomited, do not retake the dose but wait until the next scheduled time

What warnings should I know about Rubraca?

Rubraca tablets may cause serious side effects including bone marrow problems called Myelodysplastic Syndrome (MDS) or a type of cancer of the blood called Acute Myeloid Leukemia (AML). Some people who have ovarian cancer and who have received previous treatment with chemotherapy or certain other medicines for their cancer have developed MDS or AML during or after treatment with Rubraca. MDS or AML may lead to death. If you develop MDS or AML, your healthcare provider will stop treatment with Rubraca.

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How you can prepare

Ask your healthcare provider for recommendations to help you manage any side effects associated with Rubraca:

Contact your healthcare provider if you experience any of the following: weakness, weight loss, fever, frequent infections, blood in urine or stool, shortness of breath, feeling very tired, or bruising or bleeding more easily.

Healthcare provider: _____

Phone number: _____

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You should not use Rubraca if you:

- Are pregnant or plan to become pregnant. Rubraca can harm your unborn baby and may cause loss of pregnancy (miscarriage). You should not become pregnant during treatment with Rubraca
  - If you are able to become pregnant, your healthcare provider may do a pregnancy test before you start treatment with Rubraca
  - Females who are able to become pregnant should use effective birth control during treatment and for at least 6 months after receiving the last dose of Rubraca
  - Talk to your healthcare provider about birth control methods that may be right for you.
  - Tell your healthcare provider right away if you become pregnant
- Are breastfeeding or plan to breastfeed. It is not known if Rubraca passes into breast milk. Do not breastfeed during treatment and for 2 weeks after the last dose of Rubraca. Talk to your healthcare provider about the best way to feed your baby during this time

What other important information should I know about Rubraca?

Your healthcare provider will do blood tests before, and every month during treatment with Rubraca to monitor your blood cell counts. Weekly blood tests will be performed if you have low blood cell counts for a long time. Your healthcare provider may stop treatment with Rubraca until your blood cell counts improve.

Avoid spending time in sunlight while on Rubraca since your skin may become more sensitive to the sun and may sunburn more easily. You should wear a hat and clothes that cover your skin and use sunscreen to help protect against sunburn if you have to be in the sunlight.

What are the side effects of Rubraca?

The most common side effects for women in the Rubraca clinical studies were nausea, tiredness/weakness, stomach pain, rash, altered taste, decrease in hemoglobin, changes in liver or kidney function blood tests, constipation, vomiting, diarrhea, decrease in platelets, upper respiratory tract infection, mouth sores, decreased appetite, shortness of breath, and decrease in white blood cell count.

What other medications might interact with Rubraca?

Rubraca can increase the amounts of other medications you may be taking which can increase the risk of side effects. Tell your healthcare provider about all of your medical conditions and all medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

You may report side effects to the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch. You may also report side effects to Clovis Oncology, Inc. at 1-415-409-7220 (US toll) or 1-844-CLVS-ONC (1-844-258-7662; US toll-free).

Please see additional Select Important Safety Information throughout this document.

To learn more, please visit Rubraca.com.

CLOVIS ONCOLOGY

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