Rubraca Discussion Guide

QUESTIONS TO ASK YOUR DOCTOR

Rubraca is a type of targeted treatment called a PARP Inhibitor for men with *BRCA*+ metastatic castration-resistant prostate cancer. If you are interested in Rubraca as a treatment option, it's important to talk to your healthcare team. Before you meet with your doctor, make a list of questions to ask. This guide can help you get started.

- **How would I know if Rubraca is right for me? Would I need to get tested for a *BRCA* gene mutation?**

- **How is Rubraca different from other treatments for my type of prostate cancer?**

- **How could Rubraca help me and how quickly could it work?**

- **How would I take Rubraca and how often would I take it?**

- **Can you explain the potential side effects of Rubraca? How could they impact my day-to-day life?**

- **Who can help me find available financial support programs?**

INDICATION

What is Rubraca used for?

Rubraca® (rucaparib) tablets are a prescription medicine used in adults for the treatment of castration-resistant prostate cancer (prostate cancer that no longer responds to medical or surgical treatment that lowers testosterone):

- that has spread to other parts of the body, and
- has a certain type of inherited (germline) or acquired (somatic) abnormal *BRCA* gene, and you have been treated with certain medicines for your cancer.

Rubraca was approved based on response rate and how long patients' responses lasted. There are ongoing studies to confirm the clinical benefit of Rubraca. 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.

SELECT IMPORTANT SAFETY INFORMATION

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, although MDS or AML was not observed in men with prostate cancer during the clinical study. MDS or AML may lead to death. If you develop MDS or AML, your healthcare provider will stop treatment with Rubraca.

Please see additional Select Important Safety Information on back.

Rubraca®
(rucaparib) 300 mg tablets

Rubraca® (rucaparib) tablets | connections

RubracaConnections has a live, dedicated Access Specialist to support you in getting Rubraca and help you in navigating financial assistance. For more information, please call 1-844-779-7707, Monday through Friday, 8 AM to 8 PM ET, or visit **www.RubracaConnections.com**.

SELECT IMPORTANT SAFETY INFORMATION (continued)

What Warnings should I know about Rubraca? (continued)

If you are a male with a female partner who is pregnant or able to become pregnant, effective birth control should be used during treatment and for 3 months after the last dose of Rubraca. Do not donate sperm during use and for 3 months after the last dose of Rubraca.

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 men in Rubraca clinical studies were weakness/fatigue, nausea, decreased red blood cell count, changes in liver function tests, decreased appetite, constipation, rash, decreased platelet count, vomiting, and diarrhea.

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 on front.

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