

OCREVUS INFORMED CONSENT FORM

Your physician has recommended an OCREVUS infusion for your treatment. Before agreeing to this treatment, it is important that you read and understand the following explanation of the Ocrevus infusion process. This statement will describe the procedure, benefits, discomforts, risks and precautions.

OCREVUS is a medication that keeps the immune system from attacking healthy tissues in the body. The immune system also defends the body against infections caused by bacteria and viruses. Ocrevus interferes with an important step in this attack. By decreasing the immune system's attack on normal tissues.

Indications

Ocrevus is indicated and FDA approved for relapsing and primary progressive multiple sclerosis.

Dosage and Administration

Ocrevus is administered by intravenous infusion. Each infusion could take 3 to 5 hours. Proper medical supervision is essential during your infusion. We cannot be responsible for your care if you leave the suite unsupervised during the course of your infusion. You will not be allowed to leave the suite unmonitored.

A full clinical response may not be evident for several weeks following the infusion. You should continue to take all of your other medications, unless otherwise indicated by your physician.

Potential Adverse Reactions

Ocrevus is generally well tolerated and most reported reactions are mild to moderate, transient and manageable. The following adverse reactions are the most commonly reported:

- Upper respiratory infection
- Viral infection
- Diarrhea
- Back pain
- Flushing
- Rash

Some more severe side effects have been reported including:

- Serious infection
- Anemia and low white count
- Allergic reactions. These reactions are usually mild or moderate, but can be severe.



- There have been rare cases of certain kinds of cancer in patients receiving OCREVUS. The role of OCREVUS in the development of cancer is not known.
- A rare and serious brain infection called progressive multifocal leukoencephalopathy (PML)
 caused by the JC virus has occurred in patients having received immunosuppressants including
 OCREVUS.

Prior to using OCREVUS, please tell y	our physician or nurse if you (please initial):
Have has Hepatitis B	or C Infection
Active cancer	
Pregnant or breast for	eeding
Have new symptoms	or medical problems
	ccinations recently (These include measles, mumps, rubella, oral polio vaccine is not live), oral typhoid (typhoid injection is not live),
Any Fever or Current	Signs/Symptoms of an Infection
Any Acute Cut, Wou	nd or Rash
I understand all possible side effects as new information becomes availab	from this medication may not have been completely identified and le I will be notified.
,, , , , , , , , , , , , , , , , , , , ,	ad, understand and discuss this consent form with my physician nt, I am agreeing to treatment with OCREVUS.
Patient's Name	Witness' Name
Patient's Signature	Witness' Signature
Date	

