



Pregnant women with multiple sclerosis (MS) may be able to join a pregnancy registry.

Help us learn more about the health of pregnant women with MS and the health of their babies.

This booklet contains information about the OCREVUS® Pregnancy Registry. This registry is for women who are pregnant and have not received OCREVUS® (ocrelizumab). This information should help you decide whether you or someone you know may want to take part in the registry.

How can I take part in the registry?

You can enroll in the registry in any of the following ways:

- Visit the registry website at www.ocrevuspregnancyregistry.com for registration information and how to contact the Registry Coordinating Center (RCC).
- Call the RCC toll-free number 1 (833) 872-4370 and speak to a registry staff member.
- Ask your healthcare provider about the registry and how to contact the RCC.



What is the OCREVUS® Pregnancy Registry?

The OCREVUS® Pregnancy Registry is a global registry of women with multiple sclerosis (MS) who are pregnant and either have received ocrelizumab or have not received ocrelizumab. The goal of the registry is to learn more about the health of pregnant women with MS who have received ocrelizumab and the health of their babies, in comparison with those who have not received ocrelizumab. The registry will collect information about the health of participants until they give birth and will continue to collect information about the health of participants' babies until they are 12 months old.

What we learn through the registry will help us learn more about the use of ocrelizumab during pregnancy and may help pregnant women who may receive ocrelizumab in the future.

Who can participate in the registry?

You may be eligible to take part in the comparator group of the registry if:

- you are pregnant
- have MS
- **did not receive** ocrelizumab or any other disease-modifying therapy (except glatiramer acetate) since 6 months before your last menstrual period.

What will I have to do?

If you are willing to take part in the registry, you will be asked to give informed consent. Informed consent means that you receive information about what the registry involves, have a chance to ask any questions, and then agree that you want to take part.

Any questions you may have will be addressed by a registry representative in a personal telephone interview. If you give your consent, you will be contacted about 8 times from the time when you join the registry until your baby is about 12 months old. The registry representative will ask for information about your medical condition, your general health during pregnancy, and your baby's health.

A registry representative will also contact the doctor overseeing your pregnancy care, as well as the doctor who is treating your MS and your baby's doctor. This will be done to collect information about your MS, your general health during pregnancy, and your baby's health.

If you participate in the registry, you will **not** have to:

- attend any extra doctor visits
- take any extra medical tests
- take additional medications.

