



## Patient Information

### Are you pregnant and have multiple sclerosis (MS)?

Are you interested in contributing to our knowledge about medication use and pregnancy?

Join our observational registry to help us learn more about the health of pregnant women with MS, and the health of their babies.

The registry includes women who have taken Vumerity<sup>®</sup> (diroximel fumarate) and women who have not had MS therapy during pregnancy.

---

This brochure contains information about the BlossMS Pregnancy Registry.



**BlossMS**



## What is a pregnancy registry?

A pregnancy registry is an observational study (a type of medical research study) that helps to answer important questions about an approved medication and its possible effects on pregnancy, delivery, and the health of babies.

Pregnant women are often not allowed to take part in clinical research studies that test potential new medications. Therefore, the effect of a medication on pregnancy is often not known. Pregnancy registries are designed to help healthcare providers learn more about the possible effects of medications when taken during pregnancy.

For women who participate in pregnancy registries, the treatment and medical care they receive are the same as for women who do not participate.

You and/or your doctor(s) will be sharing information with the BlossomMS Pregnancy Registry team based on your routine doctors' appointments and other information you provide. Your personal information, such as your age and date of birth, together with any health information collected about you during the study, will be kept private.



## About the BlossomMS Pregnancy Registry

This registry will help doctors to better understand if Vumerity® (diroximel fumarate) has any effect on pregnancy, delivery, or the health of babies, for women who have recently taken it while pregnant. This information will be compared with information from women who have not had MS therapy during pregnancy. It is not known whether diroximel fumarate will harm unborn babies.



## Why is the registry important?

Many women with multiple sclerosis (MS) are diagnosed at about the age when they may be thinking of becoming pregnant. The information learned from this registry will help to understand more about the use of Vumerity® during pregnancy and may help pregnant women taking Vumerity® in the future.

## What will joining the registry involve?



Between 383 and 454 pregnant women who have taken Vumerity® are expected to contribute to the Blossoms Pregnancy Registry. Women with MS who have not had MS therapy during pregnancy can also take part.

If you take part in the registry, information will be collected until your baby is 12 months old.

You will need to:

- Answer questions about your health and pregnancy:
  - when you enroll
  - during each trimester of your pregnancy.
- Allow your and your baby's doctors to be contacted:
  - when you enroll
  - during your second trimester
  - 4 weeks after your due date
  - when your baby is 1, 3, and 12 months old.



You will only be contacted after your delivery if the registry team cannot contact your or your baby's doctors.



## Who can take part?

You, or someone you know, may be able to take part if you/they:

- are pregnant
- have been diagnosed with MS, and
- have taken Vumerity® (diroximel fumarate) at any time from 2 weeks after the first day of your/their last menstrual period through pregnancy, or
- have not had MS therapy at any time during this pregnancy.

## What else do I need to consider?

- You do not have to take part in the registry if you do not want to.
- If you choose to take part in the registry, you can stop participating at any time.
- There will be **no** additional medical appointments for this registry.
- This registry has been reviewed by an Institutional Review Board (IRB)/Ethics Committee (EC), which protects the rights, safety, and well-being of participants.

---

## How do I get more information?

To find out more, please visit the registry website at [BlossomMSstudy.com](https://BlossomMSstudy.com), or contact the Registry Coordinating Center using the information provided here. Registry participation is voluntary. By contacting us, you are under no obligation to take part in the registry.

