THERE ARE LOTS OF REASONS.

For one, scientists can’t figure out a cure without knowing what causes MS. There are lots of clues, and these are like pieces of a jigsaw puzzle that scientists all over the world are trying to put together.

MS attacks a very complicated part of the body, the brain and the spinal cord (called the central nervous system). Scientists need special tools to figure out what’s going on in there. Another complicated part of the body that is involved in MS is the immune system. This system is made to help our bodies fight germs, but something goes wrong in MS and the immune system attacks the central nervous system instead.

Scientists are looking for the puzzle piece that will give them the power to turn off the attack forever.

THE GOOD NEWS is that scientists are making lots of progress filling in pieces of the puzzle. They’re also learning a lot from research being done in other diseases. So, we hope soon that all of the pieces will come together and the answer will be in front of us. Then we won’t have to be patient anymore, because we’ll have a cure.
WHEN A PERSON HAS MS the covering (myelin) that protects the nerves gets damaged. Scars form where the myelin is damaged. As messages travel from the brain, they sometimes get stuck or slowed down by these scars. When this happens, the other parts of the body can’t always do what the brain is telling them to do.

SOMETIMES people with MS have trouble seeing. Sometimes their arms and legs feel weak or their skin feels “tingly” (like pins and needles). Sometimes they lose their balance, feel very tired, or have trouble walking. MS problems like these are called symptoms.

SYMPTOMS of MS can come and go. We don’t know exactly why. Sometimes you don’t even notice the symptoms. At other times they are pretty obvious. It’s hard for a person with MS to know from one day to the next how he or she will feel. That is why we say that MS is unpredictable.
Hi. My name is Hannah and I am 13 years old. My mom has MS and we work together to try and help! Sometimes she needs to use a cane, or when we’re out I will let her use my shoulder. I am the perfect height. People are always staring, but just because sometimes you can’t see it, does not mean you don’t have a disability. But it’s okay because me and my family know and that’s all that matters. She carries a cute notebook to remember things because she forgets stuff sometimes. I love my mom very much. And any kid out there that finds out your mom, dad or family member has MS, just treat them the same. My mom takes me places and shopping even though sometimes she can hardly move. I just love that she tries. She will never let MS rule her life.

That’s my story.

If your mom or dad has an exacerbation, he or she may notice new MS symptoms. Sometimes these symptoms appear and disappear quickly. Other times they last for a long time.

REMEMBER that MS is different for each person, so your mom or dad may never have any of these symptoms.
Michael: “When I had an ear infection last month, I took medicine and it went away in a couple of days. I wish MS were like that.”

Ben: “I bet MS researchers dream the same dream we do.”

Crystal: “Yeah, wouldn’t it be great if all you needed to do was take some medicine and MS would just go away? I dream of that a lot!”
Why did Joe keep his trumpet in the fridge?

Because he liked cool music.

**ANSWER:**

What is the difference between here and there?

**ANSWER:** The letter T.

What part of a ship is made out of cards?

**ANSWER:** The deck.

What happens when you don't dust your mirror?

You get a dirty look.

**ANSWER:**

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**ANSWER:**
Maybe, it will be a vaccine to keep people from ever getting MS. This would be like the shot you get from your doctor so you won’t ever get the measles or mumps. Or, maybe it will be a medicine that makes MS go away once a person has it, maybe it helps cells in the immune system do what they are supposed to do—like fighting germs—rather than damaging the myelin and nerves the way they do in a person who has MS. Or maybe there will be a cure that reverses the damage. Scientists are exploring all these possibilities, and more. And maybe, the cure will turn out to be something that no one has even thought about yet. What do you think it will be? ...???
Here is a "speech" that Sid wrote for his creative writing class at the Doris Eaton School in Walnut Creek, California. In this speech, Sid imagines that he has found a cure for MS. Do you dream about finding the cure for MS?

I am honored to have won this award for finding a cure for multiple sclerosis. It all started when I heard about MS from a doctor at a school assembly. I was very bothered by how awful the disease is and its effect on human lives. So, I decided to find a cure for MS no matter how long it took.

I researched websites, talked to experts, and learned as much as I could about possible cures for this disease. I traveled through every continent, ocean, and sea, but nothing cured my patients. So, I went to space searching. On Mars, I found a glowing herb. I brought the herb back to Earth and tested it on a patient. He recovered fully and became very healthy. I felt so good that the patient was cured and no longer had any problems.

I am now in a program that grows the special herb and produces the medicine called Cure-a-Bunch. I will continue my research and hope that we make medical breakthroughs and find cures for all diseases.

Thank you very much.

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Thank you very much.

TELL US YOUR DREAMS & IDEAS FOR WHAT A CURE WILL LOOK LIKE.

You can send us your stories by e-mail:
keepsmyelin@nmss.org or
KEEP S'MYE LIN NATIONAL MS SOCIETY
900 S Broadway #200 Denver CO 80209.
Families can count on the National MS Society!

To learn more about the resources available to you and your family, please contact an MS Navigator at:

1-800-344-4867

START SEARCHING FOR THE CURE!