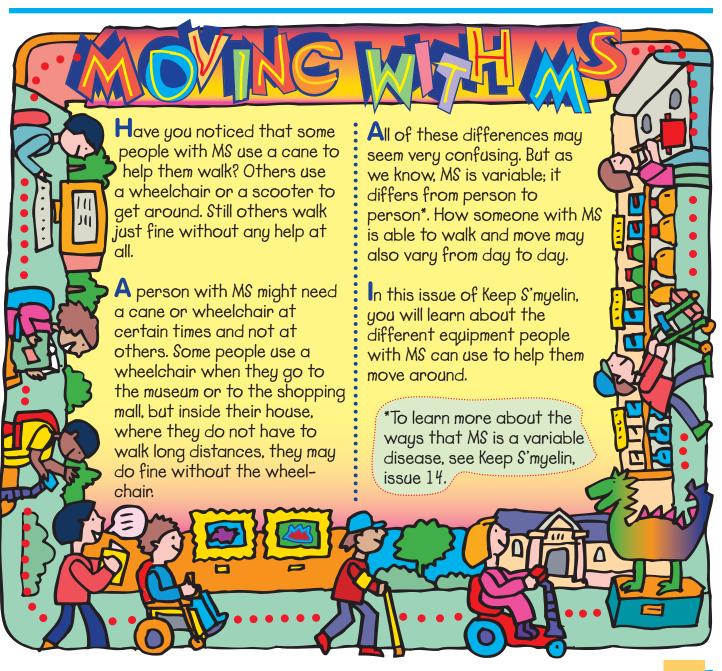
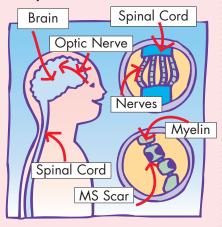


A NEWSLETTER FOR KIDS ABOUT MULTIPLE SCLEROSIS



MS

Multiple sclerosis (MS for short) is a disease that affects the central nervous system (the brain, optic nerve and the spinal cord). The brain is like a computer that tells the body what to do. The spinal cord is like a thick wire attached to the computer. Messages travel from the brain, along the spinal cord, to the other parts of the body.



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, or sometimes it's hard to walk. 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.

Interesting Fact:
Multiple means many.
Sclerosis means scars.
So, multiple sclerosis
means "many scars."

Keep S'myelin Readers: WE WANT YOU!

We love to publish your pictures, stories, and poems about MS. Please send us your work! Tell us how your family learns about MS together, how you feel about having MS in your family, what advice you would give other kids about having a mom or dad with MS, how you help your mom or dad, or whom you talk with about MS.



National MS Society KEEP S'MYELIN BROADWAY STATION

900 South Broadway
Suite 200
Denver, CO 80209
or e-mail to

keepsmyelin@nmss.org



INTERVIEW WITH COURTINE

Thirteen-year-old Courtney lives in Rochester, Minnesota with her mom and dad, two sisters, and a golden retriever named Gypsy. She is home-schooled and loves to jump on her outdoor trampoline with her little sister and play Frisbee golf with her older sister and friends. Courtney's mom has had MS since Courtney was about 7 years old.

KS: What kind of equipment does your mom use to help her get around?
Courtney: At first, she didn't need to use anything. Then after a while, she started using a cane and now she uses a wheelchair to go to church and to go shopping, a walker around the house, and her scooter when she goes into the yard to pick things from the vegetable garden.

KS: Was it hard for you when she started to need these things?

Courtney: At first, she could do lots of stuff with us, and then, when she needed these things, we were sad about it.
But now we're used to it and

we're grateful that she is happy!

KS: How did you learn about MS?
Courtney: When my mom was first diagnosed, we didn't know anything about MS. My mom and dad talked to us about it and helped us understand. Also, now we go to MS Camp* and we see other kids who are in the same situation. We can ask questions there that we might not want to

KS: Have you ever tried your mom's scooter or wheelchair?

because it might hurt their

ask our mom or dad,

feelings.

Gourtney: When we first got the scooter, my mom let us ride it up and down the sidewalk and we tried the wheelchair too. They're

hard to drive!

KS: Have you had to make any changes to your house for the wheelchair and scooter?

Courtney: Our neighbor is building us a ramp from the porch to the yard so my mom can get out

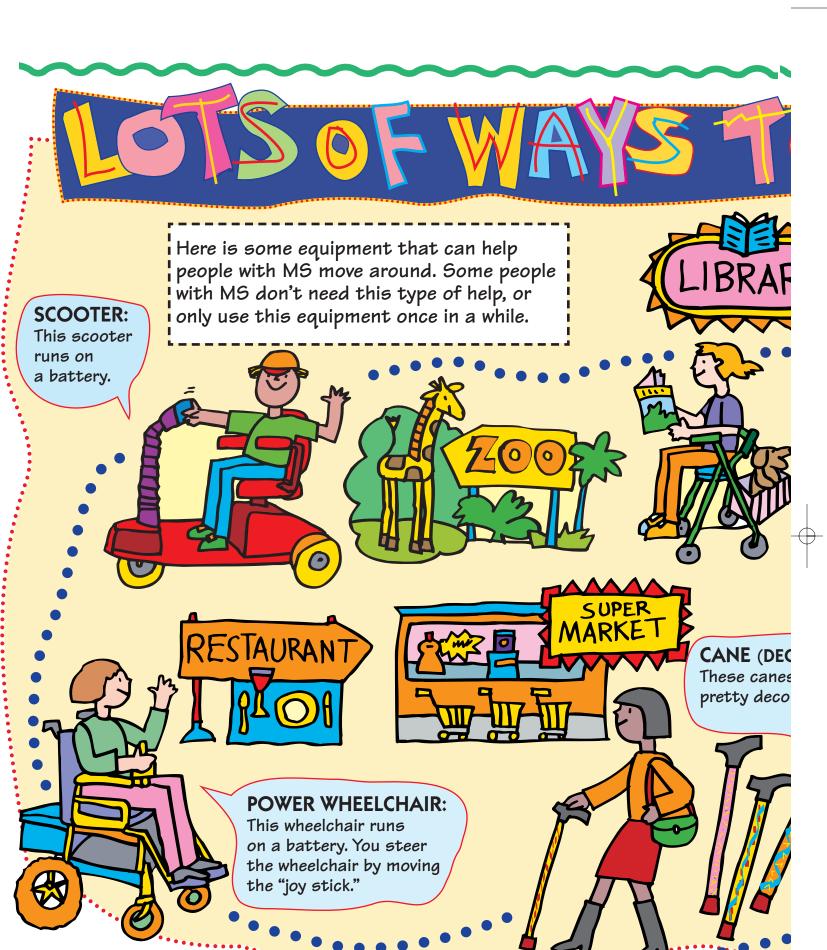
to the garden.
Also, we moved about two years ago to a house that is on one level. It's easier for my mom

to get around.

KS: What advice would you give to a kid who just found out his or her mom or dad has MS?

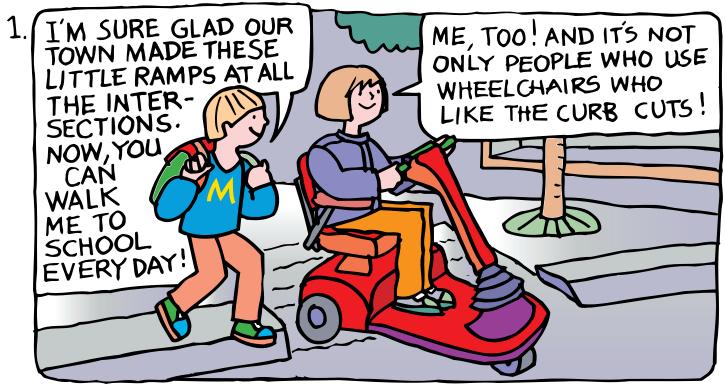
Courtney: It really helps to talk about it. Talk to your family about MS. Remember, things could be a lot worse! Keep your mind set on the good things—MS has really brought our family closer together.

*While not all chapters offer MS Camp many do offer specific programs for children and families. To learn more about these and other types of programs in your area, contact an MS Navigator® at 1-800-344-4867.





MICHAEL BENJAMIN SIMYELIN KIDS IN



ARE ACCESSIBLE TO
EVERYONE. THAT'S A NEW
WORD I LEARNED - IT
MEANS A PLACE WHERE
IT'S EASY FOR EVERYONE
TO MOVE AROUND.









Contact the National MS Society at 1-800-344-4867.



Some days my mom can walk pretty well with just a cane, but other days she can't go very far at all without her scooter. Why isn't her walking the same all the time?

When a person has MS, his or her symptoms can change from one day to the next, and even from morning to afternoon on the same day! These changes seem to be caused by many things, including how hot it is outside or how tired the person is that day. Some days, people just wake up feeling stronger or weaker than they did the day before, and no one really knows why. Your mom is very smart to use whichever piece of equipment helps her feel safe and steady, so she can go everywhere she wants to go.