

# LET'S TALK:

## CHOOSING CARE PARTNERS

**Everyone living with epilepsy should have a care partner—not just at home, but also at work and at school.**

For your own safety, it's important to have someone close by who knows what to do in case of a seizure. Here are some ideas on how to start the conversation.

### STEP 1

## Choose a good care partner

### Think about where you spend the most time

- Consider choosing a care partner wherever think you might need one, whether at work, at school, or away from home
- Look for an adult who is responsible, reliable, and capable—someone you can trust

### Familiarize yourself with seizure first aid to help decide who would be the best person to ask

- Find seizure first aid information [here](http://www.cdc.gov/epilepsy/about/first-aid.htm) ([www.cdc.gov/epilepsy/about/first-aid.htm](http://www.cdc.gov/epilepsy/about/first-aid.htm))

### STEP 2

## Start the conversation

### Ask to meet somewhere you can talk in private

- If you're uncomfortable giving a reason, you can say, *"I'd like your input on something personal"*

### You may feel awkward beginning the conversation, so prepare an opening ahead of time. You might say something like:

- *"Maybe you noticed I've been absent a few days this month"*
- Or *"Have you ever known anyone with epilepsy?"*

### Stay positive, try to sound relaxed

**After you've broken the ice, you'll find it's easy to continue**

SEE STEPS 3-6 ON NEXT PAGE

STEP  
3

## Explain why you need a care partner

**You can say,** “I have epilepsy, and it’s possible that I might have a seizure sometime when you’re around”

**Then ask,** “Are you willing to learn seizure first aid so you can help me if that happens?”

STEP  
4

## Get down to specifics

**The first question will most likely be,** “What will I have to do?”

**Don’t overwhelm them with details; offer a few facts about basic seizure first aid**

- Find seizure first aid information [here \(www.cdc.gov/epilepsy/about/first-aid.htm\)](http://www.cdc.gov/epilepsy/about/first-aid.htm)

**Don’t pressure for an answer right away. Arrange to meet again in a few days**

STEP  
5

## Follow-up

**If they say no, thank them and ask someone else**

**If they say yes, offer to answer any questions they may have**

**Tell them what signals to watch out for, what a seizure looks like, and what they should be prepared to do**

**Tell them how to learn more online. You might suggest** [www.epilepsy.com](http://www.epilepsy.com)

STEP  
6

## Expand your care circle

**It’s a good idea to have more than one care partner**

- Care partners will be more willing to accept the role if they know the responsibility is shared
- If more people know about seizure first aid, more people will be prepared to help you
- You may want care partners for different aspects of your life

**Consider talking to Human Resources, the school nurse, or Campus Health Services about an epilepsy awareness program**

- They may be able to recruit and train care partners for you
- You might also ask them to display a seizure first aid poster in a common space



**Once you start sharing outside the home, you may be surprised to see how many people are ready to join your circle of support.**

