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The Alzheimer Society of Canada
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Charitable registration number: 11878 4925 RR0001
First steps for families

A diagnosis of Alzheimer’s disease may be upsetting for both the individual and those who care about him. Most likely you have been worried about the changes you have been seeing and you may also be anxious about the future.

If you are the primary caregiver or an involved family member, you can take steps right now that might make life a little easier. Note: This brochure is a companion piece to “First Steps” for people recently diagnosed with Alzheimer’s disease.

1. **Recognize that you are going through a variety of emotions**

   The news of the diagnosis and the changes it will bring may cause you to feel anger, denial, embarrassment, frustration, fear, sadness and guilt. These emotions are normal and common among caregivers. Sometimes people get depressed if your feelings are overwhelming and won’t go away, talk to your doctor.

   Be aware that the person with the disease and other family members may also be experiencing the same types of emotions.

2. **Learn about Alzheimer’s disease**

   Learn as much as you can about the disease and providing care. Find out how the disease can affect a person, what changes you can expect, and how you can help to maintain her independence and quality of life. Share this information with those closest to her, such as family members, friends and co-workers to help them understand.

   Contact your local Alzheimer Society for useful information and resources or visit our website (www.alzheimer.ca).

3. **Recognize that the disease affects a person’s abilities**

   Alzheimer’s disease progresses over time. It will affect how the person functions daily. Learn about the changes the disease will cause so that you have realistic expectations of his abilities. Ask him how you can help him stay independent and maintain a sense of control. One tip often heard from caregivers is that you must learn to be patient, though it isn’t always easy.

4. **Don’t lose sight of the person**

   No matter how the disease affects the individual, it is important to treat her with dignity and respect. Although certain abilities will be lost, her emotions and feelings will remain, as will the need for companionship and belonging. Provide activities and interactions that bring a sense of joy and celebration.

5. **Explore treatment options**

   Currently there is no cure for Alzheimer’s disease. But medications are available that can help some people with some of the symptoms. Discuss their risks and benefits with the person’s doctor. As well, your local Alzheimer Society will have up-to-date information about new treatments that might be available through drug trials. If the individual chooses to participate in such research, you may be asked to help.

6. **Recognize that caregiving can take its toll**

   While caring for a person with Alzheimer’s disease can be rewarding, caregivers are often at risk of physical and emotional problems. If you are the caregiver, make sure you maintain your physical health, stay active and make healthy food choices. Find time for activities you enjoy. See the brochure “Reducing caregiver stress” available from the Alzheimer Society.

7. **Seek help**

   Call your local Alzheimer Society to find out what help is available in your area. Community agencies may offer practical services like help with household or caregiving tasks. You may also have a network of family and friends who are willing to lend support.

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Step 9

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