[°] All about

me



This is a booklet about a person living with Alzheimer's disease or other dementia.

Name:

AlzheimerSociety







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Introduction

This booklet is all about you, a person living with Alzheimer's disease or other dementia.

Although you have a form of dementia, you are still the same person you have always been. This booklet is designed to focus on the positive: what you are good at rather than what is no longer possible.

You and your primary caregiver* know what makes you feel comfortable better than anyone. By answering the questions in this booklet, you will have a record of what makes you content and at ease that can be used when your primary caregiver cannot be with you and others need to provide care and support. Anyone can use this booklet to give you the best day possible now and as the disease progresses.

The first section of this booklet is designed to help someone new to supporting you get to know you better. It will also suggest conversation topics that may make you feel more at ease and contribute to more enjoyable times together.

Other sections of this booklet allow you and your caregiver to outline your usual habits: your daily routines, your likes and dislikes and what makes you enjoy each day. This information will help new caregivers maintain the routines that give you a sense of security, comfort and pleasure.

When completing this booklet, always keep in mind the main purpose: to give as clear a picture as possible of you to help others provide care when the person who usually supports you is unavailable.

To help others provide effective care, keep this book in an easy-to-find location. You and your caregivers can review it from time to time to note changes and plan for the future. There are replacement pages at the back that you can use to make any updates or changes.



^{*}The term "caregiver" is used throughout this booklet to mean anyone who supports you.

0	Contact information
	This booklet contains information about:
	Some of the information is provided by:
	Names, phone numbers, email addresses of significant people in my life (family, friends, neighbours):
	Name:
	Relationship:
	Phone number:
	Email address:
	Name:
0	Relationship:
_\/	Phone number:
	Email address:
	Name:
	Relationship:
	Phone number:
	Email address:
	Name:
	Relationship:
	Phone number:

Email address:

•	(name, phone number, address):
Police:	
Poison Control	:
Local Alzheime	er Society:
Home-care serv	vices:
Spiritual or fait	th leader:
Other:	
	ormation ving dementia, are there other medical issues that the caregiver should know
•	any important information on:
Hearing:	
Vision:	
Medications (a	ttach list, if necessary) – include dosage and frequency

0	This section is like a photograph. Try to use as much detail as you can to give readers a real sense of your personality.
	My personal life
	How do you like to be addressed? (e.g. nickname, Mr., Mrs., Miss, first name)
	When were you born?
	Where?
	Single/married/partner/longstanding relationship(s) with
	Name:
	We've been together since(year)
	Where have you lived?
)	Describe this relationship (e.g. loving, difficult, supportive)
	Children (names and where they are now living)
	Are they involved in your life now? If so, how?
	Do you have any pets? If so, what are their names?
)	

Date: _

In the past What kind of jobs did you have? (e.g. homemaker, lawyer, nurse, electrician, teacher)
How do you feel about the job(s) you have done? (e.g. proud, satisfied, indifferent)
If you were asked about the major milestones in your life, what would you likely talk about? (e.g. major life events, favourite places visited)
Likes and dislikes What makes you physically more comfortable? (e.g. always have glasses on, have a hearing aid in, daily lotion to prevent dry skin, toe spacers)
What makes you happy? (e.g. conversation topics, activities, sports, music performances, being around children/animals)
What do you dislike? (e.g. foods, activities, topics of conversation, music, smells)
What comforts you when you're upset?
What frightens you?

A typical day

Routine is important for all of us, but can be especially helpful for a person with dementia.

Writing down your daily routine will help you see how you spend your time and help others who might be providing care.

Try to look at your care through the eyes of someone who has never met you before. Do you like to sleep in, have a bath in the evening, or go for a daily walk?

Use this section to describe regular daytime activities. Include activities you are involved in as well as your caregiver. Include anything that provides pleasure, comfort, or something you particularly don't like.

Morning (usual wake up time). How do you start your day?
Afternoon	
e sata.	
Evening	
Night (usual bedtime)

Date: _____

Enjoying each day

A person living with dementia is just like everyone else, a whole person with likes and dislikes, opinions, values, and experience. Though some skills are lost as the disease progresses, many remain. Here are some activities that may bring you pleasure and will help you continue to live a full life while adding enjoyment to time spent with others.

Do you like to listen to music?	☐ Yes	□ No
If yes, what kind? (e.g. classical, ja	zz, folk, blue	es, or all kinds of music)
What effect does it have on you?		
Do you play an instrument? If yes, what kind of instrument do	☐ Yes o you play? (o	☐ No e.g. guitar, violin, clarinet)
Do you enjoy singing? What effect does it have on you?	☐ Yes	□ No
Reading		
short stories)		☐ No science fiction, romance, adventure, fantasy, ne
Do you like to be read to?	☐ Yes	□ No

Do you like watching TV?	☐ Yes	□ No	
If yes, what are your favourite sh	ows?		
Games			
Do you like to play games?	☐ Yes	□ No	
If yes, what kind of games do you			ı)
Sports			
Sports Are you interested in sports?	☐ Yes	□ No	
If yes, what sports do you like to	_	_	ina)
Hobbies Do you have hobbies that you en	, ,	☐ Yes ☐ No	
	, ,		
Do you have hobbies that you en	. scrapbookin	aration, dusting, sweeping)	Yes
Do you have hobbies that you en If yes, what kind of hobbies? (e.g	g. meal prepare	aration, dusting, sweeping)	Yes 🗆

Date: ____

Help with daily living

How much help, if any, do you need with routine daily activities such as dressing, bathing or getting in and out of bed?

The chart on the next page lists typical routine daily activities. Feel free to change the chart to include activities that apply to you.

Where you are able to be completely independent, write "no help needed." When you need help, note how much help you need.

The "Useful tips" section is a good place for caregivers to note the degree of stress the activity creates and what special approaches might be helpful.

Here is a sample chart to guide you.

Activity	Useful tips	Is help needed?
Tub/shower Usual time: 8:00 a.m. Twice a week	 Prefer shower, don't like bath Enjoy music or conversation during bath time Give lots of time Respect privacy Be patient 	Need help in and out
Dressing	 Can button shirt, put on underwear and socks Need to take dirty clothes away immediately Can dress independently if clothes put on bed in right order Offer help tying shoe laces 	May need help from time to time

Date: _____

Activity	Useful tips	Is help needed?
Tub/shower		
Dressing		
Dental care/dentures		
Eye care/glasses		
Hearing aid		
Treating aid		
Hair care		
Professional style/cut		
Makeup/shave		
In/out of chair		
invoic of chair		
In/out of bed		

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Activity	Useful tips	Is help needed?
On stairs		
Use of toilet		
Use of appliances • e.g. kettle, stove, electric shaver		
 Household tasks e.g. sweeping, dusting, vacuuming, meal preparation, garden work 		
Financial Responsibility with money		
WalkingHabits, usual routes, ability to be independent		
Preparing for bed		



_	n enjoyable breakfast:
Lι	ınch:
_	
Di	nner:
_	
_	
_	
Sr	packs:
Αı	ny particular likes or dislikes?
W	hat assistance, if any, is required?
Cι	utting:
_	
US	se of cutlery:
_	
_	
Н	ot and cold liquids:
_	•

Date: ____

Regular weekly activities calendar

Use this calendar to show regular outings or appointments. You can use pencil so changes can be made every month, or a make a copy for each month.

Month:

Sunday	Monday	Tuesday	Wednesday	Thursday	Friday	Saturday

0

Date: _____

0	Note: The questions in this section are designed to be answered by your primary caregiver. Your input will be valuable to give the best information possible.					
	Alzheimer's disease and other dementias progress over time. As the disease progresses, your abilities will change.					
	The information in this section will help anyone supporting you know what these changes are and how they affect your mood, behaviour and abilities. Your caregiver can suggest ways that help you feel content, engaged, and secure. For example – Is there a special approach that helps? Does your behaviour change only at certain times? Are there warning signs?					
	Below are examples of common situations.					
	The term "family member" is used to mean anyone with dementia whom you support. "He" and "she" are alternated.					
	Unsafe walking (e.g. "wandering")					
	Does your family member walk outside in ways that are unsafe? (e.g. will go out in winter wearing only a dressing gown) \square Yes \square No					
	If yes, what safety precautions do you use? (e.g. camouflaged doors, ID bracelet, regular walks with a neighbour)					
D	Does she become upset when returned home? \square Yes \square No					
	If yes, is there a special approach to use to help her feel calm?					
	Is he registered with the Alzheimer Society's Safely Home 8 program? \Box Yes \Box No					
	If yes, what is the number of your local police station if he becomes lost?					
	What is his Safely Home® ID bracelet #?					
	Night time restlessness					

☐ No

If "yes," what safety precautions do you use? (e.g., nightlight, disconnecting stove or turning off water valves before retiring at night, locking closet door to prevent dressing at odd hours)

☐ Yes

Data

Does this occur?

	elps to re-settle her?
Restless	ness
Does thi	is occur at certain times of the day?
What he	elps to settle him? (e.g. a walk or a distracting activity)
Anger o	r agitation
Does thi	s occur at certain times? (e.g., bathing, meal time)
What us	sually triggers this? (e.g., rushing her, too many instructions given at once)
When a	nger occurs, what responses tend to be helpful?
Does yo	ur family member suspect people of stealing from him? How do you deal with this?
Repetiti	on
When h	e repeats himself over and over, what responses are helpful?

Hiding o	or hoarding articles
Are the	re particular places to check where your family member "stores" specific things?
Does an	ything need to be kept out of reach? (e.g. knives, tools such as electronic drills)
Safety p	precautions
,	additional information that is important for other caregivers. (e.g. doors or cupboards ept locked, such as where toxic cleaning fluids are stored)
Aro any	other safety measures being used? (e.g. alarms, GPS locating devices)
————	other safety measures being used: (e.g. alarms, GF3 locating devices)
Commu	nication
-	amily member has difficulty understanding and following instructions, what do you do anything that helps?
To learn the topi	n more about Communication, please read the Alzheimer Society's <u>Information sheet</u> or c.
	re any other areas of concern and/or tips for care that comfort, reassure, support the
20130111	

Alzheimer's disease follows a number of stages. While these stages can be somewhat predictable, the course of the disease will vary from person to person. Changes in physical condition, such as flu, pneumonia, infection, or constipation can often result in changes in mood and behaviour. By noting changes, the person who supports you may be able to determine a pattern and prevent a situation from getting worse. It is particularly important to keep a record when medications are used. Recording these items in a journal will help caregivers when they are talking with your doctor. Caregivers should take this "All about me" booklet to your appointments.

Anyone providing care can use these pages to record all the events in a particular day.

Here is a sample journal.

Date	Comments		
Mar. 30/12	To bed 8:30 p.m up again 2:00 a.m. Wandered through house, could not settle down. Did not recognize me.		
Mar. 31/12	Another night with no sleep!		
Apr. 1/12	Still won't sleep. Now dozing all day. Doctor's appointment April 4/12.		
Apr. 5/12	On new medication for an infection. Slept till 6:00 a.m. I'm trying to keep her awake during the day. I think things are getting better.		

ate	Comments	

Replacement pages

Date	Changes	

Date: _____



Alzheimer Society

Alzheimer Society of Canada

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