Participant ID:	Date:
C-DEMQOL A measure of carer quality of life (formatted for self-report)	
DIIDDOCE	

This questionnaire is designed to find out about your dementia.	life and experiences of being a carer of a person with
course, there might have been good and bad days, b	have felt in your caring role in the past 4 weeks . Of but we want to capture a snapshot of how you have felt estion, choose one option that best describes how you wwn x
Here is an example question:	
My own physical health in the past 4 weeks has been	
 □ very good □ mostly good □ satisfactory □ quite poor □ very poor 	
 Please remember: There are no right or wrong answers, so please be as the best of your judgement. If you feel that a question does not apply to you or to Applicable) next to the question. 	open as you can, and try to answer all the questions to the person you care for, write "N/A" (stands for Not
CARER RESPONSIBILITIES AND PERSO	NAL NEEDS
First, we would like to know how you feel about your caring responsibilities , which may include managing medications, providing help with washing and dressing, medical visits or social outings, and also anything you need to do if the person with dementia is in a care home. We would also like to know how your caring responsibilities affect your ability to have time for yourself , and doing the things that you enjoy or want to do, in particular activities that are important to your quality of life .	
1. Carrying out my caring tasks and responsibilities takes up	2. Being a carer for the person with dementia takes up
 very little of my energy some of my energy a considerable amount of my energy most of my energy all of my energy 	 very little of my time some of my time a considerable amount of my time most of my time all of my time

3. Meeting my own needs (for doing things I enjoy) whilst also caring has been	4. Thinking of my ability to do things I enjoy, I have felt
 not a problem for me quite easy a problem at times quite difficult very difficult 	 ☐ free to do them when I want ☐ restricted in a little way by my caring duties ☐ restricted to some extent by my caring duties ☐ restricted a lot by my caring duties ☐ unable to do them due to my caring duties
5. In terms of freedom to do things I enjoy, my caring responsibilities have had	6. Thinking of my ability to do activities I enjoy whilst also caring, I feel that
 no impact on my freedom little impact on my freedom some impact on my freedom a significant impact on my freedom restricted my freedom completely 	 □ all of my needs are met □ most of my needs are met □ some of my needs are met □ only a few of my needs are met □ none of my needs are met
CARER WELLBEING	
Next, we would like to find out about your wellbeing Think of how you have felt in the past 4 weeks.	and how this has been affected by your caring role.
7. I consider the emotional demands that caring places on me	8. In terms of my wellbeing, meeting the demands of caring has been
very light quite light moderate quite heavy very heavy	 not at all stressful a little stressful quite stressful stressful very stressful
9. I have been having emotional problems (such as feeling sad, stressed or anxious)	10. Whenever I need to get away from the person I care for, I feel
 none of the time a little of the time some of the time most of the time all of the time 	 not at all guilty a little guilty quite guilty guilty very guilty
11. Thinking of changes in the person I care for due to dementia (such as changes in ability, memory, or behaviour), I find them	12. Thinking of the physical and emotional demands of caring, I feel that they have
 not at all upsetting a little upsetting quite upsetting upsetting very upsetting 	 no impact on my health little impact on my health some impact on my health a significant impact on my health a very significant impact on my health

CARER ROLE

Now we would like to know how you feel about **being a carer** for a person living with dementia. This includes how you feel about your **relationship with the person you care for**, and your feelings about him or her.

13. Considering all the demands that caring places on me, I feel that overall I have	14. When thinking of my present role as a carer for a person with dementia, I feel
coped very well coped quite well coped OK coped quite poorly coped very poorly	 not at all resentful a little resentful quite resentful resentful very resentful
15. Since the person I care for developed dementia, our relationship has	16. My relationship with the person I care for in the past 4 weeks has been
improved a lot improved a little not changed deteriorated a little deteriorated a lot	very good quite good fair quite poor very poor
17. I feel appreciated by the person I care for	18. I feel frustration toward the person I care for
always or almost always often sometimes once in a while never	 none of the time a little of the time some of the time most of the time all of the time
FEELINGS ABOUT FUTURE	
Now we would like to know how you feel about your cyou care for. Again, think of how you have felt in the	own future as a carer, and the future for the person e past 4 weeks.
19. When thinking about meeting the future care needs of the person with dementia, I	20. I worry about how I will be able to cope emotionally as the dementia gets worse in the future
 ☐ feel relaxed ☐ have only a few worries ☐ have some worries ☐ have significant worries ☐ worry a lot 	 none of the time a little of the time some of the time most of the time all of the time
21. I worry about what may happen to the person I care for in the future	22 I anticipate that trying to meet the needs of the person with dementia in the future will be
 none of the time a little of the time some of the time most of the time all of the time 	 not at all difficult a little difficult quite difficult difficult extremely difficult

23. When thinking of making important caring decisions in the future, I	24. When thinking about the future impact of caring on my finances, I	
 ☐ feel relaxed ☐ have only a few worries ☐ have some worries ☐ have significant worries ☐ worry a lot 	 ☐ feel relaxed ☐ have only a few worries ☐ have some worries ☐ have significant worries ☐ worry a lot 	
CARER SUPPORT		
Lastly, we would like to know how you feel about the help and support that you may need as a carer. This includes the help and support given by family and friends , as well as any professional support provided by health and social services, or other groups or organisations. Of course, some members of your family may help a lot and some not at all. The same may be true for your friendship group or other groups. However, we want to capture how you feel about the help provided by these groups on balance , perhaps considering only the help from members who you expect to help .		
25. Overall, my needs for support have been met	26. The level of professional support offered or provided to me has met my expectations	
completely considerably to some extent a little not at all	completely considerably to some extent a little not at all	
27. I feel supported by professionals (for example, doctors, social workers, care workers)	28. I feel supported by family members	
completely considerably to some extent a little not at all	completely considerably to some extent a little not at all	
29. I feel supported by my friends	30. I feel supported by community groups (for example, charities, faith groups)	
completely considerably to some extent a little	completely considerably to some extent a little	

Thank you for taking the time to complete the questionnaire. Your responses are invaluable and greatly appreciated.