

Health-related quality of life in people with advanced dementia: a comparison of EQ-5D-5L and QUALID instruments

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Abstract

Background Assessing health-related quality of life (HRQOL) in people with advanced dementia is challenging but important for informed decision-making. Proxy measurement of this construct is difficult and is often rated lower than self-report. Accurate proxy rating of quality of life in dementia is related to identification of concepts important to the person themselves, as well as the sensitivity of the measures used. The main aim of this study was to compare the performance of two instruments—QUALID and EQ-5D-5L—on measuring HRQOL in people with advanced dementia.

Methods In a sub-study nested within a cluster-RCT we collected proxy(nurse)-completed EQ-5D-5L and QUALID measures at baseline, 3, 6, 9 and 12 months' follow-up for people with advanced dementia, residing in 20 nursing homes across Australia. Spearman's rank correlations, partial correlations and linear regressions were used to assess the relationship between the HRQOL instrument scores and their changes over time.

Results The mean weight from 284 people for the EQ-5D-5L and QUALID at baseline were 0.004 (95% CI -0.026, 0.033) and 24.98 (95% CI 24.13, 25.82), respectively. At 12 months' follow-up, 115 participants remained alive. EQ-5D-5L weights and QUALID scores at baseline and at follow-up were moderately correlated (r=-0.437; p<0.001 at 12 months). Changes within QUALID and EQ-5D-5L across the same follow-up periods were also correlated (r=-0.266; p=0.005). The regression analyses support these findings.

Conclusion Whilst these quality of life instruments demonstrated moderate correlation, the EQ-5D-5L does not appear to capture all aspects of quality of life that are relevant to people with advanced dementia and we cannot recommend the use of this instrument for use within this population. The QUALID appears to be a more suitable instrument for measuring HRQOL in people with severe dementia, but is not preference-based, which limits its application in economic evaluations of dementia care.

Keywords Dementia · Health-related Quality of Life · QALY · QUALID · EQ-5D-5L · Proxy · Advanced dementia

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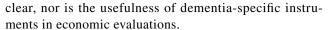


Introduction

Health-related quality of life (HRQOL) measures are used both clinically and in the decision-making process of healthcare resources allocation [1]. The challenging nature of measuring HRQOL is further emphasised by a degree of cognitive impairment, such as with dementia. It is widely recognised that the best source of data for HRQOL assessment is the person themselves. It is possible for people with mild to moderate dementia to give clear reports of their quality of life [2, 3] using dementia-specific measures [4] and through one-on-one interviews [5]. It is more difficult for people with advanced dementia to report on HRQOL with non-dementia-specific instruments such as the EQ-5D [6]. The nature of advanced dementia eliminates the possibility of getting first-hand experience from the patient, as self-reporting HRQOL requires a level of cognition and self-awareness which is unattainable by people with dementia in advanced stages of the disease [7, 8]. To address these issues, HRQOL data can be obtained by proxy, such as a relative or professional caregiver [9].

The choice of instrument for assessing HRQOL in people with dementia is important. Generic measures, such as the EQ-5D, are widely used and are often favoured by decision makers as they are quick to complete, adaptable to a large number of disease conditions and easily translated into Quality-Adjusted Life-Years (QALYs) [10]. However, a European consortium on outcome measures in dementia care has concluded that the 3-level version of the EQ-5D is unsuitable for use in people with dementia and has made the recommendation that dementia-specific measures should be used, in particular when assessing the effect of psychosocial interventions for people with dementia [11]. Since this recommendation, the 5-level EQ-5D version has been introduced for use in dementia [12], with the aim of improving the instrument's sensitivity and reducing the ceiling and floor effects [13]. However, a study on inclusion of a cognitive dimension in the EQ-5D for evaluating HRQOL in people with cognitive impairment found no added benefit of changing the instrument [14].

Dementia-specific instruments potentially capture broader and more relevant aspects of HRQOL in people with dementia, compared to generic measures, such as the EQ-5D [10]. At least 16 dementia-specific instruments exist for measuring HRQOL [15–17]. These differ in breadth and focus, although some common features have been reported: social relations or interaction, self-esteem and mood [10]. Dementia-specific instruments also target different stages of dementia, and some have been specifically designed for self- or proxy-completion [4, 18]. However, the relationship between dementia-specific measures and generic preference-based QOL measures is not always



We investigated the correlations between the two instruments and assessed the sensitivity of the Quality of Life in Late-stage Dementia (QUALID) to changes in EQ-5D-5L weights in the sub-study. In this paper we compare the performance of proxy-completed QUALID and EQ-5D-5L instruments for measuring HRQOL in people with advanced dementia. To the best of our knowledge, there has not been a study comparing the performance of QUALID and EQ-5D-5L instruments for people with advanced dementia thus far.

Methods

Data

This is a secondary analysis of data obtained from a substudy conducted within the IDEAL trial, conducted as a pragmatic parallel cluster randomised controlled trial investigating the effectiveness of facilitated family case conferencing compared to usual care in 20 Australian nursing homes [12, 19]. Participants included in the trial had a Functional Assessment Staging Tool (FAST) [7] in dementia score of 6a or higher and an Australia-modified Karnofsky Performance Status (AKPS) [20] score of \leq 50. These criteria were chosen because a FAST stage 7c combined with functional dependency (measured here by the AKPS) is predictive of an average survival of < 6 months, and the study's primary end-point focused on end-of-life care [21]. The FAST tool was initially designed for assessment of disease severity in people with Alzheimer's disease, it has been used to assess severity in people with broader dementia diagnoses in other studies (e.g. [22, 23]). The detailed inclusion criteria are outlined in the protocol paper [19].

In the intervention arm, registered nurses were trained as Palliative Care Planning Coordinators (PCPCs). The PCPCs' role was to identify optimal time-points for case conferencing, as well as to organise case conferences between the person with dementia, their family, multi-disciplinary nursing home staff and community health professionals involved in their care. The primary end-point was symptom management, comfort and satisfaction with care at the end of life, rated with the End of Life in Dementia (EOLD) Scales [24]. Secondary outcomes included the person's HRQOL (via the EQ-5D-5L and QUALID), resource use and staff attitudes and knowledge of dementia care. Eligible participants were followed up every 3 months for 12 months. Recruitment took place between February 2013 and August 2015.

We found no difference in the EOLD scale or in QUALID in the IDEAL trial [12]; therefore, for the purposes of this analysis, participants from both arms were combined



(n=284). Proxy-HRQOL was measured by specially-trained nurses familiar with participants, using the EQ-5D-5L and QUALID instruments at baseline, 3 months, 6 months, 9 months and 12 months [12]. Measures for each participant were completed by the same nurse at each time point, but may have been completed by different nurses across the study timeline.

Ethics approval was granted by the Human Research Ethics Committee (HREC) of the University of New South Wales, Australia and ratified by HRECs at the University of Technology Sydney and Queensland University of Technology, Australia.

EQ-5D-5L and QUALID instruments

The EQ-5D-5L is a five level, five-item version of the world's most widely used multi-attribute utility instrument EQ-5D [6], which has been revised to include a larger number of severity levels among its response options (therefore giving greater sensitivity). The EQ-5D-5L assesses HRQOL across five dimensions: mobility, self-care, usual activities, pain/discomfort and anxiety [13]. The respondent (or proxy) rates each domain on a 5-point Likert scale from 'no problems with a task' to 'unable to perform a task' [13]. Raw EQ-5D-5L utilities were weighted with Australian-specific weights [25], where one represents full health, zero represents death and negative weights represent states worse than death.

The 11-item QUALID scale is an advanced dementia-specific HRQOL measure that has been specifically designed for proxy-rating by caregivers. QUALID focuses on the HRQOL of the person over the span of the last 7 days by asking the proxy to rank 11 statements on a five-point scale. Possible scores range from 11 to 55, with lower scores representing the highest quality of life [26]. Total QUALID scores were calculated by summing the proxy responses to each of the 11 questions. The Quality of Life in Late-stage Dementia (QUALID) scale was developed for assessing QOL in people with advanced dementia who may be unable to communicate coherently on discrete aspects of QOL. It specifically focuses on measuring aspects that are more relevant to the person with dementia, rather than other more widely recognised contributors to QOL in the general population [26]. QUALID was designed to be administered by a caregiver or family member of a person with advanced dementia residing in long-term care facilities [26]. It has been reported to have strong internal consistency, with high test-retest reliability [15, 26]. However, it is has not been found to be correlated with measures of cognition, or activities of daily living [15, 26]. It is unclear how QUALID compares to generic QOL measures, and whether it is suitable for economic evaluation.

The QUALID was selected due to being designed purposefully for proxy-rating and its high internal validity, as

specified in the protocol paper [19]. The EQ-5D-5L was chosen for comparison with the QUALID due to the potential to undertake economic evaluation, as the QUALID cannot be used to derive Quality-Adjusted Life-Years (QALYs), and also in order to validate the EQ-5D-5L instrument in this population using proxy-rating.

Statistical analyses

Since assessment focused on changes in the participant's HRQOL over time with exposure to the case conference intervention, we used Spearman's rank correlation to assess participant HRQOL and compared QUALID and EQ-5D-5L weights at each follow-up interval. We compared 3-month changes (from baseline to 6 to 9 to 12 months) in QUALID and EQ-5D-5L using partial correlation, controlling for age, gender and baseline EQ-5D-5L and QUALID measurements. We considered correlations to be strong with scores of 0.5 and higher, and moderate with scores of 0.25–0.49 [27]. Correlations less than and equal to 0.24 were considered weak [27].

We used linear regressions to investigate the relationship between changes in EQ-5D-5L and QUALID over 3-month intervals, and controlled for age, gender, as well as the EQ-5D-5L and QUALID scores at the start of each interval being assessed. All statistical analyses were carried out using STATA version 14 [28].

Results

Of the 284 participants with dementia 179 (63%) were female and 105 (37%) male (Table 1). Forty-five percent were widowed, and 43% were married, with the rest divorced. A third (33%) had primary school certificate as their highest level of education attainment; a further 35% and 15% had completed high school and higher school certificates, respectively. Thirteen percent had some form of diploma or tertiary qualification, and 3% had no formal education.

The participants' FAST (level of function) scores ranged between levels 6a and 7f, with 50% of the study population being in groups 7c and d, corresponding to 'Ambulatory ability lost (cannot walk without personal assistance)' and 'Ability to sit up without assistance lost (e.g. the individual will fall over if there are no lateral rests [arms] on the chair)', respectively. Less than half the study population (115 of 284) remained alive at the 12-month follow-up (Table 2). No statistically significant differences in demographics, survival or HRQOL outcomes between control and intervention groups were observed (data not presented) [12].

The mean QUALID score at baseline was 24.98 (SD=7.23), whilst the mean EQ-5D-5L weight was 0.004

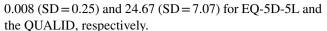


Table 1 Demographic characteristics and FAST scores of study population at baseline and 12 months follow-up

	At baseline		Surviving to 12 months	
Mean age at enrolment [mean, n (SD)]	85 (8)	284	85 (9)	115
Gender, n (%)				
Female	179	63%	79	69%
Male	105	37%	36	31%
Marital status, n (%)				
Married	121	43%	48	42%
Separated	3	1%	2	2%
Divorced	15	5%	7	6%
Widowed	129	45%	51	44%
De-facto	2	1%	1	1%
Never married	8	3%	3	3%
Unknown	6	2%	3	3%
FAST score at enrolment, n (%)				
6a	1	0%	0	0%
6b	2	1%	2	2%
6c	6	2%	3	3%
6d	9	3%	4	3%
6e	60	21%	20	17%
7a	28	10%	11	10%
7b	15	5%	8	7%
7c	75	26%	27	23%
7d	67	24%	29	25%
7e	9	3%	6	5%
7f	12	4%	5	4%
Highest education attained, n (%)				
No formal education	8	3%	3	3%
Primary school certificate	78	33%	28	27%
School certificate	84	35%	39	38%
Higher school certificate	37	15%	11	11%
Diploma	20	8%	13	13%
Bachelor degree	8	3%	6	6%
Postgraduate degree	4	2%	3	3%

(SD=0.25) (Table 2). The mean QUALID score declined slightly to 24.23 (SD=6.14) at 12 months, but the difference was not statistically significant. The EQ-5D-5L weights also experienced a consistent and statistically insignificant reduction with a mean of -0.045 (SD=0.22) at 12-month follow-up.

For participants remaining at the final follow-up (n=115), the mean QUALID score increased from 24.23 (SD=6.14) at baseline to 25.43 (SD=7.45) at 12 months, although the increase was not statistically significant (p=0.104). Similarly, the change in the EQ weights from -0.003 (SD=0.24) at baseline to -0.045 (0.22) at 12 months was not statistically significant either (p=0.083). For participants who died before 12 months' follow-up, mean baseline scores were



The correlations between QUALID and EQ-5D-5L were moderate for each time point (Table 3). Correlations were all highly statistically significant (p < 0.001) and ranged between -0.3 at 9 months and -0.44 at 12 months. Intraclass correlation coefficients for residential facilities, resulting from two-way mixed effects model regression, were also very low, ranging between 0.00 and 0.05.

We assessed the correlation in changes (i.e. compared EQ-5D-5L change between 3 months and baseline to the change in QUALID over the same period) using partial correlations, whilst controlling for age, gender and the actual EQ-5D-5L weights, as well as using a mixed-model regression, also accounting for variations within each residential facility All changes at the 3-month intervals were moderately correlated (ranging between -0.33 and -0.38) (Fig. 1 in Appendix) and highly statistically significant (Table 4). The correlation between 12 months and baseline was not as strong (-0.27) but was statistically significant. The intraclass correlations were very weak (0.00-0.07) indicating low or no levels of correlations between individuals.

We carried out linear regressions to assess the correlation between changes in EQ-5D-5L and changes in QUALID across 3-month intervals, as well as over the entire follow-up period (Table 5). We controlled for age, baseline EQ-5D-5L and QUALID scores. For each one point increase in EQ-5D-5L weight from baseline to 3 months, the QUALID score experienced an 11-point reduction (p < 0.0001). Similar relationships were found in the other follow-up periods.

In order to investigate the nature of the relationship between QUALID and EQ-5D-5L, we correlated the scores/utilities for each of the five domains of EQ-5D-5L with QUALID at baseline (Table 6). The anxiety/depression and pain/discomfort domain scores were moderately correlated with QUALID scores at baseline (r = 0.46, p < 0.001; r = 0.30, p < 0.001, respectively). Baseline QUALID scores were weakly correlated with the 'selfcare' (r = 0.12, p = 0.043) and 'usual activities' (r = 0.13, p = 0.027) domains of the EQ-5D-5L, while 'mobility' did not appear to have any correlation with QUALID scores. Bland–Altman plots were also constructed to explore the convergence between the two instruments, but did not reveal any additional information.

Discussion

Both the QUALID scores and EQ-5D-5L weights increased from baseline to 12-month follow-up in study participants with exposure to the case conference intervention, although the increases were not statistically significant. The main interest of this sub-study, however, was how these



Table 2 Mean QUALID scores and EQ-5D-5L weights at baseline and follow-up for all study participants

	QUALID			EQ-5D-5L		
	\overline{n}	Mean	SD	\overline{n}	Mean	SD
Baseline	284	24.98	7.23	278	0.004	0.25
3 months	249	25.27	7.53	244	-0.004	0.27
6 months	210	25.81	7.35	208	-0.006	0.21
9 months	174	24.83	7.06	174	-0.005	0.24
12 months	115	24.23	6.14	115	-0.045	0.22
Mean QUALID 12 months	scores and EQ	-5D-5L weights	at baseline and	follow-up for pa	articipants survivin	g to
Baseline	115	25.43	6.14	114	-0.003	0.24
3 months	115	24.32	7.14	114	-0.009	0.25
6 months	115	25.43	7.81	114	0.013	0.22
9 months	114	24.6	7.00	114	-0.015	0.21
12 months	115	24.23	7.45	115	-0.045	0.22

Table 3 Correlations between QUALID and EQ-5D-5L for each time point

	Spearman's rank correlation coefficient (p value)	Mixed-model regression coefficient for effect of EQ-5D-5L on QUALID* (p value)	Intraclass correlation coef- ficient for facility	n
Baseline	-0.385 (<0.0001)	-12.03 (<0.0001)	0.05	278
3 months	-0.432 (< 0.0001)	-10.88 (<0.0001)	0.01	244
6 months	-0.324 (< 0.0001)	-12.32 (<0.0001)	0.05	208
9 months	-0.300 (< 0.0001)	-8.12 (< 0.0001)	0.03	174
12 months	-0.437 (<0.0001)	-11.48 (<0.0001)	0.00	115

Table 4 Partial correlations and mixed-model regression between differences within QUALID and EQ-5D-5L at each time point

	Partial correlation QUALID—EQ-5D-5L* (p value)	Mixed-model regression coefficient for effect of EQ-5D-5L on QUALID* (p value)	Intraclass correlation coefficient for facility
3 months - baseline	-0.351 (<0.001)	-11.10 (<0.001)	0.00
6 months - 3 months	-0.378 (<0.001)	-11.65 (<0.001)	0.06
9 months - 6 months	-0.369 (<0.001)	-13.57 (< 0.001)	0.07
12 months −6 months	-0.331 (= 0.004)	-11.25 (<0.001)	0.00
12months - baseline	-0.266 (=0.0048)	-11.35 (0.004)	0.00

^{*}Controlled for each measure (EQ and QUALID) and age

Table 5 Linear regressions to assess the effect of changes in EQ-5D-5L on changes in QUALID

	Effect of EQ-	95% CI	p	R^2	n
Baseline to 3 months	-10.91	(-14.70 to -7.12)	< 0.001	0.430	234
6 months to 3 months	- 10.91 - 10.72	(-14.46 to -6.99)	< 0.001	0.430	197
9 months to 6 months	-12.71	(-17.64 to -7.77)	< 0.001	0.379	169
12 months to 9 months	-11.07	(-17.06 to -5.07)	< 0.001	0.343	114
12 months to baseline	-11.13	(-18.79 to -3.47)	0.01	0.095	114

^{*}Controlled for age, gender, baseline EQ-5D-5L and QUALID scores



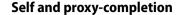
Table 6 Correlations between QUALID and EQ-5D-5L health state dimensions at baseline

EQ-5D-5L domains	Spearman's rank cor- relation coefficient	p	n
Mobility	0.047	0.433	283
Self-care	0.120	0.043	283
Usual activities	0.131	0.027	283
Pain/discomfort	0.299	< 0.001	280
Anxiety/depression	0.455	< 0.001	281

dementia-specific quality of life scales performed in relation to each other. The results indicate a consistent moderate correlation between the proxy-completed EQ-5D-5L and the QUALID across five different time-points over a 12-month follow-up period (Table 3, Appendix See Fig. 1 in Appendix). The strength of the correlations between the two instruments appears to be on par with other instances of comparisons of preference-based measures with non-preference-based ones [29].

Changes within each measure were also moderately correlated across these time-points (Tables 4, 5). This indicates further consistency in the correlation between the two instruments. This relationship, unsurprisingly, occurs between the QUALID scores and the 'pain/discomfort' and 'anxiety/ depression' domains of EQ-5D-5L, and, to a lesser extent, the association with the 'self-care' and 'usual activities' domains (Table 6). The QUALID does not measure mobility, thus explaining the lack of correlation with that domain of the EQ-5D-5L. Overall, there is insufficient evidence to support using the EQ-5D-5L as a proxy instrument for this population. Previous research found the EQ-5D-5L to be suitable for use in people with mild dementia [14, 30, 31]. In this sub-study, however, the EQ-5D-5L weights were very close to zero, indicating that it was not sensitive to changes in HRQOL in study participants in the advanced stage of dementia (Table 2). The majority of participants had significant cognitive impairment, and many were in palliative stages of care (as suggested by a 60% mortality rate over the year-long follow-up), which possibly made it difficult for proxies to rate HRQOL based on EQ-5D-5L constructs.

QUALID provided a broader scope for proxy assessment of HRQOL, using constructs more relevant to advanced and end-stage dementia such as comfort and happiness [26]. Nevertheless, QUALID is subject to the usual drawbacks of a disease-specific measure of HRQOL, in that it is does not reflect preferences for health states. It is, therefore, not possible to obtain utility values with QUALID, which restricts cross-study comparisons and decision-making on investing in dementia-relevant services. As well, QUALID has not proven superior, or any more reliable, than other dementia-specific quality of life scales [2, 32].



Proxy measurements of intangible concepts like QOL and HRQOL are difficult when the person with dementia has deteriorated to such the level that they have lost the capacity to give expression to most indicators being assessed. The reason that proxy-rating of QOL is sought from others in regular contact with the person, is that the person may have lost the capacity to provide a subjective response to how they feel about their life, and may not be able to evaluate their current life quality by weighing up both positive and negative experiences [33–36]. For people with dementia who are able to provide information on their QOL, they often provide higher ratings than the proxy responders, unless proxies have a very good appreciation of what the person's current aspirations and experiences are [33–36].

To obtain information on aspects of HRQOL that are meaningful to the person with dementia, it is important that the proxy informant is able to make unbiased judgements on the concepts being assessed [37]. Proxy-HRQOL rating bias is related to personal HRQOL, financial situation and age [38]. Differences have also been found in the proxy-HRQOL ratings of informal and formal dementia caregivers, since it is necessary for the proxy to reflect on the meanings of QOL concepts in the dementia context, which some caregivers find difficult to do [37, 39].

Proxy ratings require detailed attention to the process of dementia and its effects on peoples' identities, self-perceptions, capacities and value preferences. Another issue is the time-factor, i.e. assessment of HRQOL at a given time or over longer periods, whereby proxies are asked to make judgments on goodness of life as a whole, rather than on goodness in relation to a particular period of time [40]. This suggests that proxy-rating of HRQOL in a person with dementia is a factor of proxy insight of what HRQOL means for the person with dementia, the strength and quality of their relationship with the person and the constancy of their presence with the person in the period that measurement occurs.

In this sub-study, proxy measurement of HRQOL was provided by formal caregivers who had close associations with individual participants with dementia, which may have helped to avoid some of the potential bias with reliance on informal caregivers who spent less time with the participant during the assessment period. The analyses investigated the intra-class correlations of different residential facilities, demonstrating very low or no correlations between individuals, suggesting poor reliability between raters, [41] or a potential lack of variability among raters, as evidenced by little variation in the data.



QUALID and preference-based measures for dementia

To progress OOL assessment in dementia it would be valuable to have access to a preference-based measure based on QUALID. Rowen et al. [42] have made such an attempt with the development of DEMQOL-U and DEMQOL-proxy-U on the basis of DEMQOL and DEMQOL-proxy scales. The DEMQOL-proxy-U is limited to four domains: positive emotion, memory, appearance and negative emotion, each with four levels [42]. These are relatively narrow in scope, and omit any function or physical wellbeing subscales. Furthermore, there is some evidence suggesting that DEMQOL-proxy and DEMQOL-proxy-U are more responsive to changes in depression and delirium symptoms rather than physical symptoms [34, 43]. While vastly different from EQ-5D-5L, it could be argued that DEMQOL-proxy-U may yield similar scores to EQ-5D-5L in populations with dementia as advanced as in this study - i.e. close to zero. Development of a preference-based measure using HRQOL concepts contained in QUALID could potentially address this issue.

Strengths and limitations

One of the strengths of this sub-study was the use of two validated QOL scales in measuring HRQOL in people with advanced dementia until their death, as rated by proxies. The QOL measurements were taken at five different timepoints, further adding to the strengths and robustness of the dataset. The quality and reliability of the data across multiple time-points helped to track and compare changes in QOL for participants, thereby providing novel insights which may have not occurred with reliance on either the QUALID or the EQ-5D-5. These data provide rich information on HRQOL issues that will help to improve endof-life care and experiences for people with advanced dementia, and will contribute to the discussion on how best to measure health-related quality of life in people with significantly reduced cognitive abilities. EQ-5D-5L is an extension of a commonly used HRQOL measure EO-5D-3L which, at the time this study was conducted, remained to be validated in a number of clinical settings such as advanced dementia, and therefore, provides a reliable point for comparison of a disease-specific measure such as QUALID.

It is difficult to establish whether the current study is subject to proxy bias. Because this study utilised nursing home workers as proxies, and not informal carers or relatives, it is possible that the bias was avoided or, at least, minimised. It is also possible that nurses may introduce the bias of perceived benefit of care delivered by them [44].

Another concern is the risk of inconsistencies in proxy-HRQOL measures, as these have been completed by different nurses at different study timelines. As the same nurse completed both instruments at the same time point for each participant, the consistency between the two instruments should not be affected. The longitudinal validity of the study may be reduced by the fact that different raters completed the instruments at different time-points, potentially limiting the comparability of the instrument findings over the study period. However, the experience of nurses conducting the rating, and their familiarity with each participant may have had an effect on the measurements/weights reported [9]. While the validity of the QUALID has been demonstrated [26, 45], it remains a subjective measure (e.g. having to judge whether or not a subject enjoys eating food) and the final scores may be rater-biased.

Conclusions

One of the ways to develop and evaluate interventions is to assess these concepts using valid and reliable dementia-specific QOL and HRQOL scales. This sub-study employed the OUALID and the EO-5D-5L to obtain proxy ratings of HRQOL in people with advanced dementia living in residential care homes over 12 months. The study showed that QUALID is a suitable and reliable instrument for proxy measurement of HRQOL in people with severe dementia, compared to EQ-5D-5L, as it is more sensitive to the particular features of HRQOL in dementia. The EQ-5D-5L fails to identify aspects of HRQOL that are obtained with the QUALID. The main limitation of QUALID is that it is not preference-based and cannot easily be used in economic evaluations, where QALYs are the main outcome. Further research should focus on further comparisons of QUALID with other generic and disease-specific HRQOL measures and development of a preference-based measure based on QUALID.

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Compliance with ethical standards

Conflict of interest Elizaveta Sopina declares that she has no conflict of interest. Lynn Chenoweth declares that she has no conflict of interest. Tim Luckett declares that he has no conflict of interest. Meera Agar declares that she has no conflict of interest. Georgina M Luscombe declares that she has no conflict of interest. Patricia M. Davidson declares that she has no conflict of interest. Patricia M. Davidson declares that she has no conflict of interest. Constance D Pond declares that she has been on an advisory committee on dementia for Elli-Lilly, has received speaker fees for speaking about dementia to Australian General Practitioners, Primary Health Networks and Alzheimer's Australia, as



well as airfares and accommodation to prepare educational material on dementia for General Practitioners. Jane Phillips declares that she has no conflict of interest. Stephen Goodall declares that he has no conflict of interest.

Ethical approval All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Human Research Ethics Committees of the University of NSW, University of Technology Sydney, Queensland University of Technology approved the trial. The trial was registered under trial ID no ACTRN12612001164886 https://www.anzctr.org.au/Trial/Registration/TrialReview.aspx?id=362912.

Human and animal rights This article does not contain any studies with animals performed by any of the authors.

Informed consent Informed consent was obtained from family members with legal authority on behalf of all study participants.

Appendix

See Fig. 1.

References

- de Wit, M., & Hajos, T. (2013). Health-related quality of life. In M. D. Gellman & J. R. Turner (Eds.), Encyclopedia of behavioral medicine (pp. 929–931). New York: Springer.
- Chenoweth, L., et al. (2009). Caring for Aged Dementia Care Resident Study (CADRES) of person-centred care, dementia-care mapping, and usual care in dementia: a cluster-randomised trial. *The Lancet Neurology*, 8(4), 317–325.
- Chenoweth, L., et al. (2014). PerCEN: A cluster randomized controlled trial of person-centered residential care and environment for people with dementia. *International Psychogeriatrics*, 26(7), 1147–1160.
- Smith, S., et al. (2005) Measurement of health-related quality of life for people with dementia: Development of a new instrument (DEMQOL) and an evaluation of current methodology. Health Technology Assessment (Winchester, England). 9(10): 1–93.
- 5. Moyle, W., et al. (2015). Influencers on quality of life as reported by people living with dementia in long-term care: A descriptive exploratory approach. *BMC Geriatrics*, 15(1), 50.
- Kind, P. (1996). The EuroQoL instrument: an index of healthrelated quality of life. Quality of Life and Pharmacoeconomics in Clinical Trials, 2, 191–201.
- Reisberg, B. (1988). Functional assessment staging (FAST). Psychopharmacology Bulletin, 24(4), 653–659.
- 8. Tombaugh, T. N., & McIntyre, N. J. (1992). The mini-mental state examination: A comprehensive review. *Journal of the American Geriatrics Society*, 40(9), 922–935.

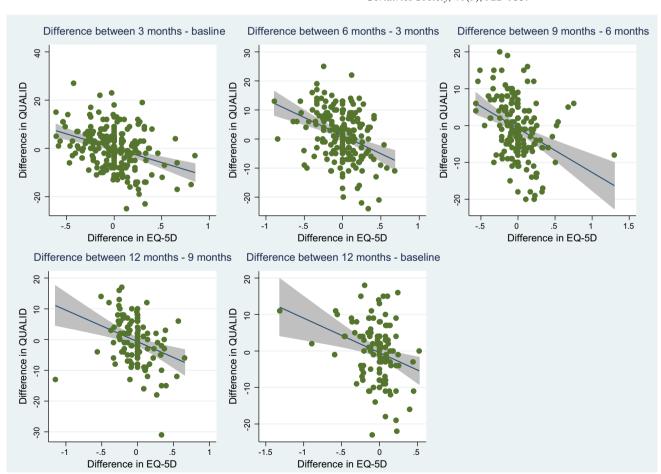


Fig. 1 Correlation between changes in QUALID and changes in EQ5D over different follow-up time-points



- Robertson, S., et al. (2017). Proxy rated quality of life of care home residents with dementia: a systematic review. *International Psychogeriatrics*, 29(4), 569–581.
- 10. Ettema, T. P., et al. (2005). A review of quality of life instruments used in dementia. *Quality of Life Research*, 14(3), 675–686.
- Moniz-Cook, E., et al. (2008). A European consensus on outcome measures for psychosocial intervention research in dementia care. *Aging & Mental Health*, 12(1), 14–29.
- Agar, M., et al. (2017). Effects of facilitated family case conferencing for advanced dementia: A cluster randomised clinical trial. PLoS ONE, 12(8), e0181020.
- Herdman, M., et al. (2011). Development and preliminary testing of the new five-level version of EQ-5D (EQ-5D-5L). Quality of Life Research, 20(10), 1727–1736.
- 14. Wolfs, C. A., et al. (2007). Performance of the EQ-5D and the EQ-5D + C in elderly patients with cognitive impairments. Health and Quality of Life Outcomes, 5(1), 33.
- Ready, R. E., (2011) Measuring quality of life in dementia. In C. Jenkinson (Ed.) Quality of Life measurement in neurodegenerative related conditions, (pp. 82–94), Cambridge: Cambridge University Press.
- Bowling, A., et al. (2015). Quality of life in dementia: A systematically conducted narrative review of dementia-specific measurement scales. Aging & Mental Health, 19(1), 13–31.
- 17. Banerjee, S., et al. (2009). What do we know about quality of life in dementia? A review of the emerging evidence on the predictive and explanatory value of disease specific measures of health related quality of life in people with dementia. *International Journal of Geriatric Psychiatry*, 24(1), 15–24.
- 18. Logsdon, R. G., et al. (1999). Quality of life in Alzheimer's disease: Patient and caregiver reports. *Journal of Mental Health and Aging*, *5*, 21–32.
- 19. Agar, M., et al. (2015). Pragmatic cluster randomised controlled trial of facilitated family case conferencing compared with usual care for improving end of life care and outcomes in nursing home residents with advanced dementia and their families: The IDEAL study protocol. BMC Palliative Care, 14(1), 63.
- Abernethy, A. P., Shelby-James, T., Fazekas, B. S., Woods, D., & Currow, D. C. (2005). The Australia-modified Karnofsky Performance Status (AKPS) scale: a revised scale for contemporary palliative care clinical practice. *BMC Palliative Care*, 4(1), 7.
- Coventry, P. A., et al. (2005). Prediction of appropriate timing of palliative care for older adults with non-malignant life-threatening disease: A systematic review. Age & Ageing, 34(3), 218–227.
- Volandes, A. E., et al. (2009). Video decision support tool for advance care planning in dementia: Randomised controlled trial. BMJ: British Medical Journal, 338, b2159.
- Candy, B., Jones, L., & Sampson, E. L. (2009). Enteral tube feeding in older people with advanced dementia: Findings from a Cochrane systematic review. *International Journal of Palliative Nursing*, 15(8), 396–404.
- Volicer, L., Hurley, A. C., & Blasi, Z. V. (2001). Scales for evaluation of end-of-life care in dementia. *Alzheimer Disease & Associated Disorders*. 15(4): 194–200.
- Viney, R., et al. (2011). Time trade-off derived EQ-5D weights for Australia. Value in Health, 14(6), 928–936.
- Weiner, M. F., et al. (2000). The quality of life in late-stage dementia (QUALID) scale. *Journal of the American Medical Directors Association*, 1(3), 114–116.
- 27. Cohen, J., et al. (2013). Applied multiple regression/correlation analysis for the behavioral sciences. Abingdon: Routledge.

- StataCorp (2015). Stata statistical software: Release 14. College Station: StataCorp LP.
- Brazier, J. E., et al. (2010). A review of studies mapping (or cross walking) non-preference based measures of health to generic preference-based measures. The European Journal of Health Economics, 11(2), 215–225.
- Kunz, S. (2010). Psychometric properties of the EQ-5D in a study of people with mild to moderate dementia. *Quality of Life Research*, 19(3), 425–434.
- 31. Aguirre, E., et al. (2016). How does the EQ-5D perform when measuring quality of life in dementia against two other dementia-specific outcome measures? *Quality of Life Research*, 25(1), 45–49.
- 32. Rokstad, A. M. M., et al. (2013). The effect of person-centred dementia care to prevent agitation and other neuropsychiatric symptoms and enhance quality of life in nursing home patients: A 10-month randomized controlled trial. *Dementia and Geriatric Cognitive Disorders*, 36(5–6), 340–353.
- Sheehan, B. D., et al. (2012). Patient and proxy measurement of quality of life among general hospital in-patients with dementia. Aging & Mental Health, 16(5), 603–607.
- Chua, K.-C., et al. (2016). Quality-of-life assessment in dementia: The use of DEMQOL and DEMQOL-Proxy total scores. *Quality of Life Research*, 25(12), 3107–3118.
- Sopina, E., et al. (2017). Cost-effectiveness of a randomised trial
 of physical activity in Alzheimer's disease: A secondary analysis exploring patient and proxy-reported health-related quality of
 life measures in Denmark. *British Medical Journal Open*, 7(6),
 e015217.
- Bhattacharya, S., et al. (2010). Generic and disease-specific measures of quality of life in patients with mild Alzheimer's disease.
 Dementia and Geriatric Cognitive Disorders, 30(4), 327–333.
- 37. Hounsome, N., Orrell, M., & Edwards, R. T. (2011). EQ-5D as a quality of life measure in people with dementia and their carers: Evidence and key issues. *Value in Health*, *14*(2), 390–399.
- Arons, A. M., et al. (2013). Quality of life in dementia: A study on proxy bias. BMC Medical Research Methodology, 13(1), 110.
- 39. Meyer, J., et al. (2006). Moving from victim blaming to an appreciative inquiry: Exploring quality of life in care homes. *Quality in Ageing and Older Adults*, 7(4), 27–36.
- 40. Schermer, M. (2003). In search of the good life' for demented elderly. *Medicine, Health Care and Philosophy*, 6(1), 35–44.
- 41. Koo, T. K., & Li, M. Y. (2016). A guideline of selecting and reporting intraclass correlation coefficients for reliability research. *Journal of Chiropractic Medicine*, *15*(2), 155–163.
- Rowen, D., et al. (2012). Estimating preference-based single index measures for dementia using DEMQOL and DEMQOL-proxy. Value in Health, 15(2), 346–356.
- Ratcliffe, J., et al. (2016) An empirical comparison of the EQ-5D-5L, DEMQOL-U and DEMQOL-Proxy-U in a post-hospitalisation population of frail older people living in residential aged care. Applied Health Economics and Health Policy, 15(3): 399–412.
- Clare, L., et al. (2014). Care staff and family member perspectives on quality of life in people with very severe dementia in long-term care: A cross-sectional study. *Health and Quality of Life Outcomes*, 12(1), 175.
- Falk, H., Persson, L.-O., & Wijk, H. (2007). A psychometric evaluation of a Swedish version of the Quality of Life in Late-Stage Dementia (QUALID) scale. *International Psychogeriatrics*, 19(6), 1040–1050.

