Instrumental Activities of Daily Living Scales to Detect Cognitive Impairment and Dementia in Low- and Middle-Income Countries: A Systematic Review

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16 Abstract.

- Background: The largest proportion of people with dementia worldwide live in low- and middle- income countries (LMICs),
 with dementia prevalence continuing to rise. Assessment and diagnosis of dementia involves identifying the impact of
- 19 cognitive decline on function, usually measured by instrumental activities of daily living (IADLs).
- Objective: This review aimed to identify IADL measures which are specifically developed, validated, or adapted for use in
 LMICs to guide selection of such tools.
- 22 Methods: A systematic search was conducted (fourteen databases) up to April 2020. Only studies reporting on development,
- validation, or adaptation of IADL measures for dementia or cognitive impairment among older adults (aged over 50) in
 LMICs were included. The QUADAS 2 was used to assess quality of diagnostic accuracy studies.
- Results: 22 papers met inclusion criteria; identifying 19 discrete IADL tools across 11 LMICs. These were either translated
- from IADL measures used in high-income countries (n = 6), translated and adapted for cultural differences (n = 6), or newly
- developed for target LMIC populations (n=7). Seven measures were investigated in multiple studies; overall quality of diagnostic accuracy was moderate to good.
- Conclusion: Reliability, validity, and accuracy of IADL measures for supporting dementia diagnosis within LMICs was reported. Key components to consider when selecting an IADL tool for such settings were highlighted, including choosing
- culturally appropriate, time-efficient tools that account for gender- and literacy-bias, and can be conducted by any volunteer
- with appropriate training. There is a need for greater technical and external validation of IADL tools across different regions,
- countries, populations, and cultures.
- Keywords: Activities of daily living, cognitive dysfunction, cross-cultural comparison, dementia, developing countries,
 diagnosis, functional status

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INTRODUCTION 36

It is estimated that 54 million people are living 37 with dementia globally [1], with numbers set to rise 38 to 75 million by 2030 [2]. Two-thirds of demen-39 tia cases occur in low- and middle-income countries 40 (LMICs) [1, 3], yet less than 10% of people with 41 dementia in LMICS receive a diagnosis [1]. This 42 highlights the difficulty in accurately assessing preva-43 lence of dementia globally and leads to difficulties 44 in accessing appropriate care in LMICs. Dementia 45 is a progressive neurodegenerative condition charac-46 terized by decline of cognitive performance across 47 multiple cognitive domains, which impacts an indi-48 vidual's ability to carry out activities of daily living 49 (ADLs) [4]. There are a number of reasons for the 50 low rates of dementia diagnosis in LMICs, includ-51 ing stigmatization, lack of funding and resources for 52 health and social care, variations in assessment and 53 characterization of dementia, and cultural differences 54 regarding the expectation of older adults within soci-55 ety which contributes to low recognition of dementia 56 by family members and society as a whole [2, 3]. 57 Accurate and timely diagnosis of dementia is vital to 58 appropriately treat and manage the disease, educate 59 carers about the condition, and to ensure that people 60 with dementia from LMICs are represented within 61 global dementia research. As such, it is recommended 62 that valid and accurate tools are developed to support 63 dementia screening in LMICs, which are appropriate 64 for variations in culture, education, and language [3]. 65

Subtle cognitive impairments occur years before 66 formal diagnosis of dementia and can manifest thr-67 ough increasing impairments in ADLs [5]. ADLs 68 refer to everyday activities which are associated with 69 functional independence and are a fundamental part 70 of dementia diagnosis [4]. Clinically, they can be sep-71 arated into more cognitively-driven activities known 72 as instrumental ADLs (IADLs; e.g., shopping, finan-73 cial management), and more procedural activities 74 known as basic ADLs (BADLs; e.g., eating, bathing) 75 [5]. While difficulties in BADLs tend to occur in 76 later stages of dementia, impairments in IADLs may 77 become increasingly apparent early in the disease 78 course prior to formal diagnosis and reflect the onset 79 of cognitive decline [6]. As such, IADL assessments 80 are recommended as simple and effective screening 81 tools for dementia in LMICs [3]. 82

Multiple questionnaires have been developed to 83 assess IADLs in dementia [7]; however, most are 84 targeted at high-income Western countries and may 85 be culturally-inappropriate for use in LMICs due to 86

different age- and gender-roles, literacy rates and geographical variations [3]. For example, in certain countries there are cultural expectations that younger family members will manage household and financial matters while older adults play a more social role within the community [8]. Therefore, IADL tools with a significant focus on financial management or household chores may not be suitable, while tools which are weighted to social activities, such as presiding over ceremonies or following local affairs, could better reflect cognitive decline. Additionally, some LMICs have unique activities that reflect discrete cultural practices, and which would be considered IADLs (e.g., tying a sari) while their equivalent in 100 Western culture would be characterized as BADLs 101 (e.g., getting dressed). When choosing an IADL ass-102 essment to support dementia screening in LMICs, 103 it is important to consider if the tool is culturally-104 appropriate for the target population in order to 105 maximize the efficacy and accuracy of its use for 106 dementia diagnosis [3]. Therefore, this review aims to 107 support researchers and clinicians in selecting cultur-108 ally appropriate IADL tools by 1) identifying IADL 109 tools that have been developed or adapted for use 110 in LMICs and 2) reporting how reliable, valid, and 111 accurate these tools are for identifying dementia. 112

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METHODS

Identification of studies

Search terms and databases

Searches were conducted across fourteen data-116 bases, including databases of LMIC-based litera-117 ture to maximize the opportunity to locate studies 118 involving LMIC populations. The databases searched 119 were: 3ie, AIM, African Journals Online, CINAHL, 120 Eldis, Embase, KCI, LILACS, MedCarib, MED-121 LINE, PsycInfo, RSCI, SciELO, and World Bank. 122 Search results were limited to studies conducted 123 prior to April 2020 (the date searching commenced) 124 with no lower date limit. To identify studies from 125 LMICs, the Cochrane filter for LMICs was used 126 in databases where this was possible. A list of all 127 countries listed as low-, lower middle-, or upper 128 middle-income as of April 2020 was also obtained 129 from the World Bank Database. Combinations of 130 the search terms described in the Supplementary 131 Material were searched across the databases. This 132 review was pre-registered on PROSPERO (Refer-133 ence: CRD42018107882).

Inclusion criteria 134

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Inclusion criteria were as follows: 135

1. The study assessed IADLs in older adults aged 136 50 years or older who had been given a diagno-137 sis of, or were being assessed for, dementia or 138 cognitive impairment. 139

- 2. The study was conducted in an LMIC setting, 140 as defined by the World Bank country classification by income database as of April 2020. 142
 - 3. The study reported at least one of the following:
 - a. The validity and reliability of the IADL measure
 - b. The sensitivity and specificity of the IADL measure
 - c. Positive and/or negative predictive value of the IADL measure
 - d. Comparison with a previously validated IADL measure

Exclusion criteria 152

Studies were excluded if they focused on IADL 153 assessments in populations other than those living 154 with dementia or cognitive impairment, as were 155



Fig. 1. Flowchart of the screening and eligibility evaluation for studies included in the review.

studies which only involved populations from highincome countries. Studies which did not report any statistical assessments of the diagnostic accuracy or validity of the IADL measure were also excluded. Finally, studies which were not available in English language were excluded due to a lack of resources available for translation.

Selection process

Results from all searches were imported into Microsoft Excel to assist with screening against the inclusion and exclusion criteria. All titles and abstracts were screened by four reviewers (RMA, HY, MG, AN) according to the inclusion criteria. Any discrepancies were referred to an adjudicator to obtain a consensus view. Full text versions of articles that met the inclusion criteria were obtained and each were assessed for final inclusion by two reviewers (from RMA (all texts; n = 44), HY (n = 5), MP (n = 10), MG (n=10), SMP (n=9), AN (n=10)) with discrepancies referred to an adjudicator who had not previously reviewed the specific text (CD (n=12)). Eligibility of identified articles was recorded at every stage to document the review process. Duplicates were identified and removed prior to commencing the screening process. A hand search of reference lists of included studies was also conducted to identify any studies which had not been detected in the search process (HY, CD; see Fig. 1 for further details).

Data analysis

Data extraction

Data were extracted from all eligible articles, with key measures of interest as follows: 1) LMIC country involved; 2) setting (urban/rural, clinic/ community/care); 3) type of IADL tools (translated, translated, and adapted, newly developed for target population); 4) criteria used to characterize cognitive impairment/dementia; 5) domains included in the IADL tool (basic, instrumental, advanced); 6) scoring of IADL tool; and 7) clinometric properties of IADL tool (i.e., reliability, validity, accuracy).

Interpretation of data

Data was synthesized according to the type of IADL tool each study employed, i.e., translated, translated and adapted, and newly developed for a target population. This approach was determined after reviewing all studies included in this review. Translated tools refer to IADL tools which were used

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and/or validated in another country and language, 203 and which were directly translated into a target lan-204 guage (e.g., English to Portuguese). Translated and 205 adapted tools refer to IADL tools which were used 206 and/or validated in another country and language, and 207 which were translated into a target language using a 208 cross-cultural approach, such as making adaptions for 200 terminology or changing items to ensure appropriate-210 ness for the target culture. Tools which were newly 211 developed for a target population refers to IADL tools 212 which were developed specifically for the population 213 being studied, usually through consensus processes 214 to ensure that items included in the IADL tool were 215 appropriate and relevant to the culture (e.g., inclusion 216 of "tying lower garments appropriately" in Indian 217 populations). 218

All studies included in this review reported relia-219 bility (internal consistency (e.g., Cronbach's alpha), 220 test-retest, inter-rater (e.g., ICCs, Pearson/Spearman 221 correlations)), validity (concurrent (e.g., correlati-222 ons), construct (e.g., correlations), convergent (e.g., 223 correlations), discriminative (e.g., between-group 224 comparisons)), and diagnostic accuracy (criterion 225 validity, sensitivity, specificity, positive/negative pre-226 dictive values, area under the curve (AUC)). The-227 refore, the current review examined these three types 228 of reliability, four types of validity, and the range 229 of diagnostic accuracy measures. IADL tools which 230 were assessed in multiple studies were highlighted 231

in the results and data were synthesized to provide a comprehensive overview of the evidence.

Quality assessment

The Quality Assessment of Diagnostic Accuracy Studies version 2 (QUADAS-2) tool [9] was used to evaluate the quality of included studies. This measure assesses four key domains: 1) method of participant selection; 2) index test use and interpretation; 3) reference standard use and interpretation; and 4) flow and timing of tests. Some of the included articles were not diagnostic accuracy studies and so it was not possible to use the QUADAS-2 to fully assess these as certain domains were not covered. Two reviewers (RMA and SMP) determined quality of all diagnostic accuracy studies in a blinded assessment. Disagreements were settled through consensus.

RESULTS

Search yield

The search yielded 4,247 articles, of which 1,741 250 were duplicates and removed. Following title and 251 abstract search, 47 full texts were obtained and 252 (Fig. 1). An additional four articles were identified via 254



Fig. 2. Heat map of locations for research into the development, adaption, and validation of assessments for instrumental activities of daily living to support dementia diagnosis in low-middle income countries.

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a hand search of reference lists of included studies.In total, 22 studies were included in this review.

The characteristics of the 22 eligible studies are 258 summarized in Table 1. All articles were published 259 between 1999-2019. Only one study carried out 260 longitudinal analysis [10]. In order of quantity, coun-261 tries represented by this review include Brazil (41%; 262 n=9), India (13%; n=3), Turkey (9%; n=2), Tan-263 zania (9%; n=2), Argentina (5%; n=1), Nigeria 264 (5%; n=1), Republic of Congo (5%; n=1), Cen-265 tral African Republic (5%; n = 1), Iran (5%; n = 1), 266 Sri Lanka (5%; n=1), and Thailand (5%; n=1) 267 (Fig. 2). The sample size ranged from 40-632 par-268 ticipants across studies. Additionally, 82% of studies 269 reported > 50% of participants as female. Prevalence 270 of cognitive impairment in the sample ranged from 271 1-100% across studies. Studies were conducted in 272 clinical (59%; n = 13), community (36%; n = 8) and 273 care (5%; n=1) settings, and in urban (50%; n=11), 274 rural (23%; n = 5), both urban and rural (9%; n = 2), 275 and unspecified (23%; n=4) environments. 276

Nineteen IADL tools were identified and catego-277 rized into three types: translated (n=6), translated 278 and adapted (n = 6), and newly developed for the tar-279 get population (n = 7). Results relating to reliability, 280 validity and diagnostic accuracy for all tools can be 281 found in Table 3. Seven discrete IADL tools were 282 assessed by multiple studies and synthesized data for 283 these will be presented below. 284

285 Quality assessment

Eleven of the studies included diagnostic accuracy 286 measures and where therefore assessed for quality 287 using the OUADAS 2. Most studies demonstrated 288 some risk of bias; scores are presented in Table 2. 289 All studies were included in the review regardless of 290 the assessed quality to demonstrate the full available 291 data related to the IADL tools assessed within the 292 current literature. 293

Translated high-income country developed IADL
 tools in LMICs

296 Activities of daily living questionnaire (ADL-Q)

The ADL-Q assesses both BADLs and IADLs, evaluating 28 items across six domains: social interaction, social participation, planning/organizing, intellectual activities, feeding, and self-care [11]. This scale is based on an observer's report, whereby the observer rates the individual's abilities on a scale of 0-3; higher scores reflect greater impairment. A response option "don't know/has never done" is also available, and if selected, the item is excluded from the total score. Scores from discrete items are summed to form subdomain scores, and then transformed into a percentage score. No/mild impairment is classified as 0-33%, moderate impairment is 34-66%, and severe impairment is 67-100%.

Two studies assessed the use of the ADL-Q, translated into Spanish and Portuguese and conducted in Argentina [12] and Brazil [13], respectively. Both studies took place in clinical settings and urban environments. For Gleichgerrcht et al. [12], 100% of participants had a diagnosis of dementia, and for Fransen et al. [13], 31% had Alzheimer's disease and 39% had mild cognitive impairment (MCI). On average, people with dementia had 12–13 years of education in Gleichgerrcht et al. [12], while they had 6.7 years in Fransen et al. [13]'s study. Reliability and validity findings are described in Table 3.

Fransen et al. [13] examined diagnostic accuracy of the ADL-Q for detecting MCI compared to normal aging, and for distinguishing Alzheimer's disease from MCI. With a cut-off of 1%, MCI could be distinguished from controls with 66% sensitivity and 69% specificity (AUC: 0.653; based on Winblad et al. [14]), and with a cut-off of 21%, MCI could be differentiated from Alzheimer's disease with 93% sensitivity and 91% specificity (AUC: 0.977; based on Frota et al. [15]).

Disability assessment for dementia scale (DADS)

The DADS is an informant-based scale which assesses both BADLs and IADLs, evaluating 40 items (17 basic, 23 instrumental) across ten domains. BADL domains include hygiene, dressing, continence, and eating, while IADL domains involve meal preparation, telephoning, going on an outing, finance, and correspondence, taking medication, leisure activities, and housework. Response to each item is yes (1 point) or no (0 points), with the total score ranging from 0–100. Total scores are calculated by summing the score of each item and a percentage is calculated by excluding not applicable answers (e.g., does not do this activity). Lower scores reflect greater impairments in ADLs.

Two studies assessed the use of DADS, translated into Turkish and Portuguese and conducted in Turkey [16] and Brazil [17], respectively. Both studies took place in clinical settings with Bahia et al. [17] 301

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IADL Tool	Study	Country	Setting	Language of IADL tool	Participant No.	Mean Age	% Female	Education
Thai ADL Scale	Senanarong et al. [50]	Thailand	Clinic, urban	Thai	181	Dementia: 69.51 ± 9.16 Controls: 67.73 ± 9.35	Dementia: 64.8% Controls: 72.7%	Dementia: 0-4 y: 50.28% > 12 y: 11.9% Controls: 0-4 y: 31.82% > 12 y: 26.4%
FAQ- BR/PFAQ	Jomar et al. [20]	Brazil	Community, urban	Portuguese	265	Elderly: 74–84: 44.2% Informants: 75+: 36.6%	Elderly: 74% Informants: 82.1%	\geq 9 y Elderly: 45.7% Informants: 85.7%
	Aprahamian et al. [22]	Brazil	Clinic, urban	Portuguese	106	AD: 80.28 Controls: 77.95	71.70%	100% illiterate
	Sanchez et al. [21]	Brazil	Community, Urban	Portuguese	68	58 ± 12.9	79.40%	>9 y: 75%
ADLQ-SV	Gleichgerrcht et al. [12]	Argentina	Clinic, urban	Spanish	40	AD: 79 ± 5.9 bvFTD: 75.4 ± 11 Other: 76.6 ± 8.9	AD: 66% bvFTD: 60% Other: 76%	AD: 12.2 ± 4.7 y FTD: 12.9 ± 3.7 y Other: 12.6 ± 4.1 y
ADLQ-BR	Fransen et al. [13]	Brazil	Clinic, urban	Portuguese	90	Controls: 68.07 ± 5.57 MCI: 69.34 ± 7.04 AD: 75.07 ± 6.65	Controls: 74.1% MCI: 71.4% AD: 78.6%	Controls: 14.19 ± 5.57 y MCI: 10.26 ± 4.60 y AD: 6.71 ± 5.16 y
EASI	Pandav et al. [27]	India	Community, rural	Not specified	632	66.5 ± 7.6	46.90%	73.3% illiterate
	Fillenbaum et al. [26]	India	Community, rural	Not specified	387	55–64: 123 participants 65–74: 145 participants 75+: 119 participants	47%	78% illiterate
CSADL	Noroozian et al. [32]	Iran	Clinic, unspeci- fied	Persian	277	Not stated	55%	Male: 9 y Female: 5 y
DADS- Turkish	Tozlu et al. [16]	Turkey	Clinic, unspeci- fied	Turkish	157	77.7±6.8	63.70%	31.8% illiterate
DADS-BR	Bahia et al. [17]	Brazil	Clinic, urban	Portuguese	129	AD: 76.4 ± 6.9 Controls: 74.5 ± 7.3	AD: 64% Controls: 57.5	AD: 6.4 ± 5.1 y Controls: 6.5 ± 4.9 y
IADL-E	Mathuranath et al. [31]	India	Clinic, urban, rural	Not specified	240	67.8 ± 10.5	45%	Dementia: 9.9 ± 4.9 y Controls: 8.9 ± 5.8 y

 Table 1

 Demographic and geographical characteristics of all instrumental activities of daily living tools (n = 19) included in the review

CHIF	Hendrie et al. [30]	Nigeria/USA	Community, rural	Yoruba/ English	Nigeria: 295 USA: 155	Nigeria: Dementia: 82.9 ± 10.7 Without Dementia: 78.2 ± 6.6 USA: Dementia: 83.4 ± 6.8 Without Dementia: 80.7 ± 6.4	Nigeria: Dementia: 86.8% Without Dementia: 73.9% USA: Dementia: 75% Without Dementia: 70.4%	Nigeria Dementia 0% Without dementia: 13.6% USA Dementia: 8.9 ± 2.5 Without dementia: 9.4 ± 3.0
CA-DFI	Edjolo et al. [29]	Central African Republic/ Republic of	Community, urban, rural	"local lan- guages"	301	76.1 ± 7.4	94%	99.7% Low educational level
IDEA- IADL	Collingwood et al. [8]	Tanzania	Community, rural	Swahili	449 Grouped by IDEA Cognitive Scale scores: $\leq 7: 40$ 8–9: 57 $\geq 10: 352$	IDEA Cognitive score levels: ≤7: 80 (IQR: 73.75-85.5) 8–9: 76(IQR: 70-81.25) ≥10: 72 (IQR: 67-79)	IDEA Cognitive score levels: ≤7: 85% 8–9: 71.9% ≥10: 50.6%	Not specified
	Stone et al. [10]	Tanzania	Community, rural	Swahili	Baseline: 153 Follow-up: 98	Baseline: 21.6% 65–69 22.9% 70–74 20.9% 75–79 20.3% 80–84 14.4% 85+ Follow up 15.3% 65–69 17.3% 70–74 15.3% 75–79 28.6% 80–84 23.5% 85+	Baseline: 67.3% female Follow up: 66.3% female	Without formal education: Baseline: 33.3% Follow up: 29.6%
IDEA- IADL Short	Stone et al. [10]	Tanzania	Community, rural	Swahili	As previous	As previous	As previous	As previous
ADCDS- ADL Turkish	Aysun et al. [24]	Turkey	Clinic, unspeci- fied	Turkish	73	AD: 72.56 ± 10.55 Controls: 68.38 ± 8.82	AD: 56.3% Controls: 58.1%	5.16 ± 3.83 y
ADCDS- ADL Brazil	Cintra et al. [25]	Brazil	Clinic, urban	Portuguese	95	75.9±7.6	60%	Controls: 5.7 ± 4.4 y MCI: 5.2 ± 3.9 y AD: 3.6 ± 3.3 y

(Continued)

					Table 1 (<i>Continued</i>)			
IADL Tool	Study	Country	Setting	Language of IADL tool	Participant No.	Mean Age	% Female	Education
GADLS	Paula et al. [34]	Brazil	Clinic, urban	Portuguese	178	MCI <75: 67.04 ± 4.53 MCI 75+: 81.17 ± 5.1 AD <75: 68.97 ± 4.13 AD 75+: 79.47 ± 3.40	Not specified	MCI <75: $5.15 \pm 4.29 \text{ y}$ MCI 75: $3.92 \pm 3.40 \text{ y}$ Dementia <75: $4.68 \pm 3.92 \text{ y}$ Dementia 75: $5.26 \pm 3.61 \text{ y}$
DAFS-R	Pereira et al. [23]	Brazil	Clinic, urban	Portuguese	89	73.8±6.7	AD: 58% MCI: 74% Controls: 75%	10.3 ± 6 y
	Fransen et al. [13]	Brazil	Clinic, urban	Portuguese	As previous	As previous	As previous	As previous
LBI	Marra et al. [33]	Brazil	Clinic, urban	Portuguese	90	75.46 ± 7.66	75.50%	No education: 24.4% 1–7 y: 56.6% 8 + y: 18.8%
PI	Marra et al. [33]	Brazil	Clinic, urban	Portuguese	As previous	As previous	As previous	As previous
Bristol ADL	Umayal et al. [44]	Sri Lanka	Care	Sinhalese	70	>75: 47.1%	74.30%	≤5 y: 70%
Blessed ADL	Umayal et al. [44]	Sri Lanka	Care	Sinhalese	As previous	As previous	As previous	As previous

ADL, activities of daily living; FAQ, Functional activities questionnaire; BR, Brazil; PFAQ, Portuguese Functional Activities Questionnaire; ADLQ, Activities of daily living questionnaire; SV, Spanish Version; EASI, Everyday Activities Scale – India; CSADL, Cleveland Scale of Activities of Daily Living; DADS, Disability Assessment for Dementia; IADL, Instrumental activities of daily living for elderly people; CHIF, Clinician Home-based Interview to assess Function; CA-DFI, Central Africa Daily Functioning Interference Scale; IDEA-IADL, IDEA study Instrumental Activities of Daily Living Questionnaire; ADCDS-ADL, Alzheimer's Disease Co-operative Study – Activities of Daily Living Scale; GADLS, General Activities of Daily Living Scale; DAFS-R, Revised Direct Assessment of Functional Status; LBI, Lawton Brody Index; PI, Pfeffer Index; AD, Alzheimer's disease; MCI, mild cognitive impairment.

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Table 2
Consensus scores for the QUADAS-2 demonstrating quality of all diagnostic
accuracy studies $(n = 11)$ included in this review

		Ris	k of Bias		Applica	ability C	oncerns	
	Patient	Index	Reference	Patient	Index	Reference	1	
	Selection	Test	Standard	timing	Selection	Test	Standard	
Jomar et al. [20]	+	+	-	+	+	-	-	
Pandav et al. [27]	+	-	-	?	+	-	-	
Noroozian et al. [32]	+	+	+	+	+	?	+	
Edjolo et al. [29]	+	+	-	-	+	-	-	
Stone et al. [10]	?	-	-	?	-	-	-	
Collingwood et al. [8]	-	+	+	-	+	+	+	
Cintra et al. [25]	-	-	-	?	-	-	-	
Paula et al. [34]	+	-	-	-	-	-	-	
Pereira et al. [23]	-	-	-	-	+	-	-	
Umayal et al. [44]	+	+	-	+	+	-	-	
Bahia et al. [17]	+	-	+	+	-	-	-	
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+, high risk; -, low risk; ?, unclear risk

reporting an urban environment. Tozlu et al. [16] included 100% of participants with dementia, whereby
31.8% were illiterate. Bahia et al. [17] reported 69%
of participants to have dementia, with a mean of
6.4–6.5 years of education.

Diagnostic accuracy was only investigated for DAD-Brazilian version (AUC: 0.993 [17]). With a cut off of 94.6%, dementia could be distinguished from controls with a sensitivity of 94.6%, specificity of 100%, a positive predictive value of 100%, and a negative predictive value of 93% (based on [18, 19]; alternative cut-offs provided in Table 3).

364 Translated and adapted IADL tools in LMICs

365 *Functional activities questionnaire (FAQ)*

The FAO is an informant based IADL scale 366 with 10 items: finances, handling documents, shop-367 ping, games/hobbies, making tea/coffee, preparing 368 a balanced meal, paying attention/understanding/ 369 discussing a tv program/book/magazine, keeping 370 track of current affairs, remembering appointments/ 371 occasions/medication, and travelling. Every item is 372 rated between 0-3, with higher scores reflecting 373 greater impairment. If activities are not usually com-374 pleted by the individual, informants specify whether 375 the individual would be able to carry out the activity. 376 The maximum score is 30. 377

Three studies examined the FAQ [20–22]. All studies were based in Brazil and used Portuguese versions of the scale. Transcultural adaptions of the FAQ for Brazil were designed, which included reviewing and adapting items and expressions to increase relevance to Brazilian culture. All studies took place in urban environments, with two in community settings [20, 21] and one in clinic [22]. Within each sample, dementia accounted for 43% [20] and 62% [22] of participants. Sanchez et al. [21] did not characterize people with dementia, but all those included had a MMSE score of < 27. For Sanchez et al. [21] and Jomar et al. [20], 75% and 85.7% of informants had 9 + years of education, while the sample in Aprahamian et al. [22] was 100% illiterate.

Both Jomar et al. [20] (AUC: 0.797) and [22] (AUC: 0.864) provided diagnostic accuracy measures. Jomar et al. [20] reported a sensitivity of 80%, specificity of 72%, positive predictive value of 68.7%, and negative predictive value of 82.4% with a cut-off score of 14. Aprahamian et al. [22] used a cut-off of 11.5, showing a sensitivity of 85.3% and specificity of 76.5%.

Direct assessment of function scale (DAFS)

The DAFS is an observation-based scale which includes BADLs and IADLs. It requires approximately 25 minutes to administer and involves simulating 23 daily tasks across seven domains: time orientation, communication, transportation, finance, shopping, grooming, and eating. The maximum score is 106, with lower scores reflecting greater impairments in ADLs.

Two studies examined DAFS in clinical urban settings in Brazil [13, 23]. The scale was translated into Portuguese and revised to improve relevance for Brazilian culture. For example, currency and stimulus cards with phone numbers and addresses were

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Table 3
Key results relating to reliability, validity, and diagnostic accuracy of instrumental activities of daily living tools $(n = 19)$ in low to middle income countries

IADL Tool	Study	Dementia Criteria	% Demen- tia/CI	No of items	Total Score	Type of IADL tool	Method	Reliability	Validity	Diagnostic Accuracy/ Criterion Validity
Thai ADL Scale	Senanarong et al. [50]	DSM-IV	88%	13	26	Newly developed for target population	Collected from infor- mants	Inter-rater (<i>n</i> = 30): Evaluation 1 ICC: 0.96 (95%CI: 0.91–0.98) Evaluation 2 ICC: 0.93 Test-retest: Rater 1 ICC: 0.92 (95%CI: 0.83–0.96) Rater 2 ICC: 0.89 (95%CI:	Discriminative: Scores: CDR 2 > CDR 1 > CDR 0.5 > CDR 0 Construct: Significant association between each item and the Thai MSE (r=0.69) and CDR (r=0.81) Convergent: Controlling for cognition, correlations between Thai ADL and Bathel Index $(r=0.64)$	
FAQ- BR/PFAQ	Jomar et al. [20]	DSM-IV	43%	10	30	Translated and adapted	Collected from infor- mants	0.78-0.95)	and FAQ (r = 0.30) remain. Concurrent: FAQ BR negatively correlated with MMSE (r = 0.624, p < 0.001) and positively with IQCODE-BR (r = 0.755, p < 0.001).	Cut off:≥14/30 Sensitivity: 80% (CI: 71.5–86.9) Specificity: 72 (CI: 64.1–79.0) AUC: 79.7% (IC: 74.3%–84.4) PPV: 68.7% (CI: 60.1–76.4) – 96/115 people NPV: 82.4% (CI:
	Aprahamian et al. [22]	DMS-IV, NINCDS- ADRDA	62%				97	hor	Discriminative: PFAQ significantly different between AD and controls $(p < 0.001)$.	74.8–88.5) – 49/150 Cut off: 11.5 Sensitivity: 85.3 Specificity: 76.5 AUC: 86.4% (SE: 4.3%; 95%CI: 78.94.96%)
	Sanchez et al. [21]	Not used	100% with MMSE<27, dementia not specified					Cronbach's alpha: 0.95 Test-retest: ICC: 0.97	roo,	(

ADLQ-SV	Gleichgerrcht et al. [12]	NINCDS- ADRDA: AD McKeith: LBD Lund and Manch- ester: bvFTD NINDS- AIREN: VaD Benson et al: PCA	100%	28	100	Translated	Collected from infor- mants – based on observa- tion	Cronbach's alpha for all factors: 0.82-0.96 Inter-rater: Cohen's K: 0.90 Test-Retest: r=0.95, $p < 0.001$	Concurrent Validity: Correlation with FAQ total $(r=0.67, p<0.001)$ and CDR $(r=0.54, p<0.001)$.	
ADLQ-BR	Fransen et al. [13]	AD: Frota et al., 2011 MCI: Winblad et al., 2004	Dementia: 31% MCI: 39%	28 900	100	Translated	Based on observa- tion	Cronbachs alpha = 0.759	Construct: Correlation between ADLQ-BR and DAFS-R (rho = 0.743).	Controls versus MCI Cut-off 1/100 Sensitivity: 66% Specificity: 69% AUC: 65.3% MCI versus AD Cut off: 21/100 Sensitivity: 93% Specificity: 91% AUC: 97.7%
EASI	Pandav et al. [27]	DSM-III	1%	11	11	Newly developed for target population	Collected from infor- mants	5		Cut off $\geq 3/11$ Dementia versus Controls Sensitivity: 62.5% Specificity: 89.7% AUC: 88.4% PPV: 24.4% NPV: 97.8%
Fillenbaum et al. [26]	Based on Hindi Mental State Examina- tion Scores	Not specified					•	Cronbach's alpha: 0.82 Inter-rater reliability: 100% agreement Test-retest: 82–100% agreement	Discriminative and Construct: Differences between Hindi Mental State Examination Stages for EASI (p < 0.001).	
									0/	(Continued)

						Table 3 (Continu	3 ed)			
IADL Tool	Study	Dementia Criteria	% Demen- tia/CI	No of items	Total Score	Type of IADL tool	Method	Reliability	Validity	Diagnostic Accuracy/ Criterion Validity
CSADL	Noroozian et al. [32]	Expert opinion	85%	48	138	Translated	Collected from infor- mants		Discriminative: CSADL Scores: Dementia + AD > MCI	Cognitive impairment versus controls Full scale Cut off: 20 Sensitivity: 90% Specificity: 93% Cut off: 26 Sensitivity: 87% Specificity: 100% IADL Scale Cut off: 20 Sensitivity: 91% Specificity: 100%
DADS- Turkish	Tozlu et al. [16]	DSM-IV, NINCDS- ADRDA		40	100	Translated	Collected from infor- mants	Cronbach's alpha: 0.942 Inter-rater: ICC: 0.994 (95%CI: 0.987-0.997) Test-retest: ICC: 0.996 (95%CI: 0.991-0.998)	Discriminative: Significant differences for DAD scores between GDS stages: Stage 4 > Stage $5 >$ Stage $6 + 7$. No difference between stages 6 and 7 Construct: Correlation between DAD and Lawton IADL Scale ($r = 0.928$, p < 0.001). Convergent: Correlation between MMSE and DADS ($r = 0.812$, p < 0.001), DADS and CDS ($r = 0.920$ and $2 < 0.001$)	Specificity: 10078
DADS-BR	Bahia et al. [17]		69%	40	100	Translated	Collected from infor- mants	Cronbach's alpha: 0.77	Convergent: Correlation between DADS and MMSE scores ($r=0.044$, p < 0.001) Scores lower in AD than controls ($p < 0.01$)	AUC: 99.3% Cut-off: 94.6 Sensitivity: 96.6% Specificity: 100 PPV: 100 NPV: 93 Cut-off: 90 Sensitivity: 90% Specificity: 100 PPV: 100 NPV: 81.6 Cut-off: 85 Sensitivity: 81.8% Specificity: 100 PPV: 100 NPV: 71.4

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IADL-E	Mathuranath et al. [31]	DSM-IV. AD: NINCDS- ADRDA VaD: NINDS- AIREN	44%	11	22	Newly developed for target population	Collected from infor- mants	Ibadan Results: Cronbach's alpha: 0.83 Inter-rater: r=0.87, p<0.001	Convergent: IADL-CDI correlated with MMSE (co-efficient: 0.31) – increasing when MMSE increased and vice versa. Construct: IADL-E correlated with DSM-IV (r = 0.89), CDR (r = 0.82), MMSE (r = 0.74) and ACE (r = 0.60)	Only cognitive sub score used. Cut off: 16/22 Dementia versus Controls Sensitivity: 91% Specificity: 99% AUC: 97% (94–99) PPV: 0.76%
CHIF	Hendrie et al. [30]	ICD-10, DSM-III AD: NINCDS- ADRDA	Nigeria: 13% USA: 26%	10	20	Newly developed for target population	Clinician interview		Discriminative: Participants without dementia performed better on CHIF than with dementia (p < 0.001) Construct: Correlation between CHIF and Blessed Dementia Scale $(r = 0.56, p < 0.001)$ and MMSE (r = 0.44, p < 0.001)	Dementia versus Controls AUC: 92.5% Cut off: 18/20 Sensitivity: 89.5% Specificity: 68.5% Cut off: 17/20 Sensitivity: 68.4% Specificity: 82.5%
CA-DFI	Edjolo et al. [29]	DSM-IV AD: NINCDS- ADRDA MCI: Peterson's Criteria	Dementia: 26.6% MCI: 20.3%		Unknown	Newly developed for target population	Collected from infor- mants	Cronbach's alpha: 0.92	Convergent: 10 item CADFI correlated with walking speed (<i>r</i> =0.431) and CDR (<i>r</i> =0.62) Construct: Item response theory showed <ask STELLA></ask 	Cognitive Impairment versus Controls Based only on laundry score. Cut off: 0.35 Sensitivity: 96% Specificity: 69% AUC: 87.8% (83 0-91 6)
IDEA-IADL	Collingwood et al. [8]	DSM-IV	26.90%	11	33	Newly developed for target population	Collected from infor- mants	Cronbach's alpha: 0.959	Criterion: Dementia diagnosis a significant predictor of IADL score Construct: Factor analysis revealed only one factor with eigenvalue > 1, explaining 71.6% of variance.	Controls AUC: 89.6% (CI: 84.2–95.1)
	Stone et al. [10]	DMS-IV	Baseline: 25% Follow- up: 36.7%					Cronbach's alpha: 0.956	100×	Dementia versus controls Baseline AUC: 90.3% (CI: 85.2–95.3) Follow-up AUC: 62.5% (CI: 50.8–74.2)

						Table 3 (Continue)	3 ed)			
IADL Tool	Study	Dementia Criteria	% Demen- tia/CI	No of items	Total Score	Type of IADL tool	Method	Reliability	Validity	Diagnostic Accuracy/ Criterion Validity
IDEA-IADL Short	Stone et al. [10]	As previous	As previous	3	6	Newly developed for target population	Collected from infor- mants		Construct: Factor analysis revealed 2 factors as most strongly predicting dementia.	Baseline AUC: 99.5% ($82.0-94.9$) Follow up AUC: 62.1% ($50.2-73.9$) Criterion: Significantly predicted dementia with regression co-efficient: 0.868 ($p < 0.001$)
ADCDS- ADL Turkish	Aysun et al. [24]	NINCDS- ADRDA		23	78	Translated	Collected from infor- mants	Cronbach's alpha: 0.938 Test-Retest: ICC: 0.998 (95%CI: 0.997–0.999)	Discriminative: ADCS-ADL Scores for CDR Stages 0.5 > 1>2 > 3 Construct: ADSC-ADL highly correlated with BADL (rho = 0.826) and IADL scores (rho = 0.826) on the Modified OARS Convergent: ADCDS-ADL scores are highly correlated with CDR ($r = 0.828$), GDS ($r = 0.743$), but not ADAS Cog ($r = 0.191$)	
ADCDS- ADL Brazil	Cintra et al. [25]	AD: NINCDS- ADRDA MCI: Albert and Peterson Criteria	Dementia: 35% MCI: 34%	23	79	Translated and adapted	Collected from infor- mants	Cronbach's alpha: 0.89	Discriminative: Controls had better ADCDS = ADL scores than MCI and AD (p < 0.001). Subitem scores were also better in controls for advanced (p < 0.001), IADL (p < 0.001), IADL (p = 0.004). Convergent: Association between ADCS-ADL and clinical/ neuropsychological diagnosis (ROC = 0.89, p < 0.001).	Full scale Cut off: 71/79 Cognitive Impairment versus Controls Sensitivity: 86.2% Specificity: 70% AUC: 81.1% PPV: 86.2% NPV: 70% AD versus Controls Sensitivity: 97% Specificity: 70% AUC: 84.1% PPV: 78% NPV: 95.4% MCI versus Controls Sensitivity: 75% Specificity: 70% AUC: 72.6% PPV: 72.7%

		nc	0	0	576	9	1.			Sensitivity: 97% Specificity: 25% AUC: 61.5% PPV: 42.9% NPV: 88.9% IADL Scale Cut-off: 32 Cognitive Impairment versus Controls Sensitivity: 81.5% Specificity: 76.7% AUC: 80% PPV: 88.3% NPV: 65.7% AD versus Controls Sensitivity: 93.9% Specificity: 76.7% AUC: 85.7% PPV: 81.6% NPV: 92% MCI versus Controls Sensitivity: 68.8% Specificity: 76.7% AUC: 72.6% PPV: 75.9% NPV: 69.7% AD versus MCI Sensitivity: 93.9%
GADLS F e	Paula t al. [34]	AD: NINCDS- ADRDA MCI: Peterson Criteria	Dementia: 52% MCI: 48%	13	28	Translated and adapted	Collected from infor- mants	Cronbach's alpha: 0.849	D _r	AUC: 63.1% PPV: 41.5% NPV: 83.3% Young MCI versus Young AD (≤74) Sensitivity: 69% Specificity: 62% AUC: 72.5% (CI: 59.9–81.8)

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(Continued)

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IADL Tool	Study	Dementia Criteria	% Demen- tia/CI	No of items	Total Score	Type of IADL tool	Method	Reliability	Validity	Diagnostic Accuracy/ Criterion Validity	
DAFS-R	Pereira et al. [23]	DSM-IV AD: NINCDS- ADRDA MCI: Peterson's Criteria	Dementia: 29% MCI: 35%	23	##	Translated and adapted	Simulation observed by clinicians	Cronbach's alpha: 0.78 Inter-rater: ICC: 1–0.918 for all items Test-Retest: ICC: 1–0.915 for all items	Discriminative: Subitems Time Orientation and Communication Scores: MCI + Controls > AD. Subitems Finances and Shopping scores: Controls > MCI > AD. Convergent: Correlation between DAFS and IQCODE (r =0.65, p < 0.001). No correlation between DAFS and IQ-CODE when groups examined separately. Construct: Correlation between ADLQ-BR and DAFS-R (rho = 0.743).	AD versus Controls:Cut-off: 86 Sensitivity: 100% Specificity: 93.7% AUC: 99.8% MCI versus Controls: Cut-off: 93 Sensitivity: 80.60% Specificity: 84.4% AUC: 86.8%	
	Fransen et al. [13]	As previous	As previous	0	ie.	9	1			Controls versus MCI Cut off: 91/105 Sensitivity: 68% Specificity: 63% AUC: 72.6% MCI versus AD Cut off: 70/105 Sensitivity: 89% Specificity: 83% AUC: 90.5%	
LBI	Marra et al. [33]	DSM-IV	100%	8	8 for women 5 for men	Translated	Collected from infor- mants	hor	Construct: Negative correlation found between PI and LBI for full sample (p < 0.0001, rho = 0.818) - when looking in each severity - mild $(p = 0.007$, rho = 0.530), severe (p < 0.0001, r = 0.0723) - in moderate dementia, the questionnaires were not correlated. Discriminative: All dementia severity groups different for LBI scores (p < 0.001)		

Table 3 (Continued)

PI	Marra	As previous	As previous	10	30	Translated	Collected	Construct: Negative	
	et al. [33]						from	correlation found between	
							infor-	PI and LBI for full sample	
							mants	(p < 0.0001, rho = 0.818) -	
								when looking in each	
								severity - mild ($p = 0.007$,	
								rho = 0.530), severe	
								(p < 0.0001, r = 0.0723) - in	
								moderate dementia, the	
								questionnaires were not	
			Orn					correlated.	
								Discriminative: All	
								dementia severity groups	
								(r < 0.001)	
Bristol ADL	Umayal et al. [44]	ICD-10NA	44%	14	42 .	Translated and	Collected	(<i>p</i> < 0.001)	Cut off: 20
					42	adapted	from		Sensitivity: 100%
						adapted	infor-		Specificity: 74.2%
					~ I K		mants		AUC: 93 3% (95%CI:
							mants		87 1–99 5%)
Blessed	Umaval	As previous	As previous	13	19	Translated and	Collected		Cut-off: 10.5
CERAD	et al. [44]	no previous				adapted	from		Sensitivity: 100%
						1	infor-		Specificity: 89.2%
							mants		AUC: 89.2% (95%CI:
							· <i>(</i> / / /		81.6-96.7%)

ADL, activities of daily living; FAQ, Functional activities questionnaire; BR, Brazil, PFAQ, Portuguese Functional Activities Questionnaire; ADLQ, Activities of daily living questionnaire; SV, Spanish Version; EASI, Everyday Activities Scale – India; CSADL, Cleveland Scale of Activities of Daily Living; DADS, Disability Assessment for Dementia; IADL, Instrumental activities of daily living for elderly people; CHIF, Clinician Home-based Interview to assess Function; CA-DFI, Central Africa Daily Functioning Interference Scale; IDEA-IADL, IDEA study Instrumental Activities of Daily Living Questionnaire; ADCDS-ADL, Alzheimer's Disease Co-operative Study – Activities of Daily Living Scale; GADLS, General Activities of Daily Living Scale; DAFS-R, Revised Direct Assessment of Functional Status; LBI, Lawton Brody Index; PI, Pfeffer Index; AD, Alzheimer's disease; MCI, mild cognitive impairment; AUC, area under the curve; PPV, positive predictive value; NPV, negative predictive value.

adapted to Brazilian standards. In Fransen et al. [13],
31% of participants had Alzheimer's disease and 39%
had MCI, while Pereira et al. [23] included 29% of
participants with dementia and 35% with MCI. On
average, people with dementia had 6.7 years of education in Fransen et al. [13], and 10.3 years in Pereira
et al. [23].

Only Pereira et al. [23] reported diagnostic accuracy between dementia and controls (AUC: 0.998, based on [15]). Using a cut-off of 86, DAFS showed
a sensitivity of 100% and specificity of 93.7%. Diagnostic accuracy for discriminating MCI from controls and Alzheimer's disease can be found in Table 3.

Alzheimer's disease cooperative study–activities of daily living scale (ADCS-ADLS)

The ADCS-ADLS is a 23-item informant-based 431 scale which includes assessments of BADLs (6 432 items), IADLs (10 items), and advanced ADLs (7 433 items). Each item is rated as either dependent, par-434 tially independent, or totally independent, with a 435 maximum score of 79 points, where lower scores 436 reflect greater impairments. It requires approximately 437 12 minutes to administer. 438

Two studies assessed ADCS-ADLs, translating it 439 into Turkish and Portuguese and conducted in Turkey 440 [24] and Brazil [25], respectively. For the Turkish 441 version, only minor adjustments to wording were 442 made. For the Brazilian version, an expert committee 443 applied changes to the format of questions, cultural 444 expressions, and vocabulary, and added one sub-445 item. This adapted ADCS-ADLS Brazilian version 446 was tested in community dwellers with and with-447 out cognitive impairment, which led to the removal 448 of "selecting/choosing clothes" and modification of 449 "eating with knives and forks" to "eating indepen-450 dently". People with dementia encompassed 44% of 451 Aysun et al. [24]'s sample, and 35% of Cintra et al. 452 [25]'s sample with an additional 34% MCI. Mean 453 education ranged from 3.6-5.7 years across the sam-454 ples. 455

Cintra et al. [25] reported diagnostic accuracy 456 measures for the Brazilian ADCS-ADLS. Using a 457 cut-off score of 71, dementia could be distinguished 458 from controls with 97% sensitivity, 70% specificity, 459 78% positive predictive value, and 95.4% nega-460 tive predictive value (AUC: 0.841, based on [19]). 461 Table 3 provides values for distinguishing controls 462 from overall cognitive impairment and MCI, and for 463 differentiating MCI from dementia.

Newly developed IADL tools in LMICs

Everyday abilities scale for india (EASI)

The EASI is an 11-item informant-based scale involving BADLs and IADLs across four domains: personal care, mobility, social interaction, and cognitive function. A point is scored for each item where impairments are reported, with higher scores reflecting greater impairments. The EASI was developed for a largely illiterate rural Indian population, involving consolation with professional experts, village leaders, and field workers familiar with the community. Items were selected based on activities older adults are culturally expected to carry out, regardless of social status (e.g., wrap/tie lower garments appropriately, express opinions in important family matters).

Two studies assessed EASI in community-based rural settings in India [26, 27]. In Pandav et al. [27], 1% of participants had a dementia diagnosis, while this information was not specified in Fillenbaum et al. [26]. In both studies, there were high levels of illiteracy (73–78%).

Pandav et al. [27] reported diagnostic accuracy measures (AUC: 0.884, based on DSM-III criteria) for distinguishing dementia from controls. Using a cut-off of 3, sensitivity was 62.5%, specificity 89.7%, positive predictive value 24.4%, and negative predictive value 97.8%.

IDEA-instrumental activities of daily living scale (IDEA-IADL)

The IDEA-IADL is an 11-item informant-based scale assessing IADLs. It can be administered by local healthcare workers to caregivers or relevant informants. It was developed through consultation with district enumerators and local healthcare workers who had extensive training on dementia. Activities that would be expected of an older person, regardless of gender or physical/sensory impairments, were identified (e.g., settle conflicts, preside over ceremonies), resulting 12 relevant activities heavily weighted toward social functions. Following pilot work, one activity was removed ("They make their will and testament and make decisions about their property when they are gone") as administrators felt uncomfortable asking this. Each item had a four-point scale (0-3) with higher scores reflecting greater impairments. The maximum score is 33.

Two studies examined the IDEA-IADL in community-based rural Tanzania [10, 28]. Paddick et al. [28] reported 26.9% of participants with a diagnosis of dementia, while in the longitudinal study by 465

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Stone et al. [10] had 25% with dementia at baseline (n = 153), and 36.7% at follow-up (n = 98). Additionally, 33.3% of participants at baseline and 29.6% at follow-up had no formal education; education and literacy rates were not specified in Paddick et al. [28]. Both studies reported area under the curve scores

for accuracy of distinguishing dementia from controls, based on American Psychiatric Association [18] criteria, ranging from 0.625 (follow-up, [10]), 0.896 [28], and 0.903 (baseline, [10]).

524 DISCUSSION

In terms of reliability, validity, and accuracy, we 525 highlighted the seven IADL tools which were rep-526 orted by multiple studies, giving them stronger 527 evidence bases to potentially identify dementia in 528 LMICs, describing their key characteristics (dom-529 ains, time commitment, scoring process); how they 530 have been developed, translated or adapted; and their 531 accuracy at distinguishing cognitive impairment from 532 normal ageing. We now critically discuss the practi-533 cal implications of this review in terms of clinical 534 practice and future research. 535

536 Choosing an IADL tool: key considerations

Our findings demonstrate three different cate-537 gories of IADL tools validated in LMICs. These 538 include translated, translated and adapted, and those 539 newly developed for target populations (i.e., national 540 or regional populations within specific countries). 541 In addition, there were geographical trends in the 542 selection of IADL tools assessed. In African and 543 South Asian LMICs, bespoke culturally-specific 544 tools were predominately investigated [10, 26-31], 545 while translated and adapted tools were mainly used 546 in South America and West Asian LMICs [12, 13, 547 16, 20-25, 32-34]. This made synthesis of results 548 difficult. Diagnostic accuracy appeared highest in 549 translated/translated and adapted tools, but these find-550 ings cannot be readily generalized to African and 551 South Asian LMICs due to cultural differences. For 552 example, while most included LMICs have cultural 553 expectations whereby younger family members assist 554 older members with BADLs when significant disabil-555 ity is present [35], studies based in Africa and South 556 Asia placed significant emphasis on social IADLs 557 (e.g., presiding over ceremonies, keeping up with 558 local affairs/festivals) as younger family members 559 have responsibility over more traditional IADLs mea-560 sured in Western scales (e.g., financial management) 561

[10, 29]. It is difficult to compare the efficacy of tools which used directly translations of IADL scales used in high-income countries (i.e., translated) and tools which used a cross-cultural adaption process (i.e., translated and adapted). These tools were generally used in Brazil and Turkey, which may hold similarities with the cultures that the tools were originally developed for. This highlights the necessity of first understanding cultural expectations of the target population when choosing an IADL tool, as it should include relevant activities for older adults within that culture to ensure sensitivity for detecting dementiarelated impairments [3].

The influence of gender norms and literacy rates are another key consideration when selecting an IADL tool. Most included studies had a predominantly female sample. While this likely reflects the higher prevalence of dementia in women compared to men [36], this limits our understanding of the suitability of IADL tools for men within LMICs. For example, IADL tools with a significant weighting on household activities may not reflect subtle impairments in men within LMICs, as traditional gender roles within most societies dictate that older women predominately carry out household activities (e.g., cooking, cleaning), while men may mainly perform management activities (e.g., keeping financial records) [37]. To account for this, the Lawton Brody Index provided discrete scoring systems for men and women [33] and the IADL-E has an equal number of male- and female-dominant items [31]. An alternative way to negate gender bias is to focus on social IADLs, which both older men and women within the community commonly carry out, such as giving advice [10].

Additionally, low literacy and education rates significantly impact dementia screening and may introduce performance differences across the spectrum of literacy [22]. Articles included in this review similarly highlight significant rates of illiteracy and low educational levels [22, 26, 27, 29, 30]. These illiteracy and education rates can be considered barriers to comprehensive cognitive assessment, and as such, brief cognitive assessments and IADL tools are recommended to reduce bias [38]. Both translated and bespoke IADL questionnaires assessed in populations with high illiteracy and low education demonstrated excellent diagnostic accuracy scores [22, 27, 29], showing that evaluation of the sensitivity and specificity of cut-off IADL scores have been established for illiterate populations in LMICs. Furthermore, Hendrie et al. [30] reported the use of 562

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an observational IADL tool (i.e., CHIF) in a Nigerian population with less than four years of education
which reported excellent accuracy for identifying
cognitive impairment. Ensuring selected IADL tools
accommodate for gender or literacy bias is vital to
capture cognitively driven impairments early in the
disease course.

A final consideration for the selection of IADL 621 tools is the time and expertise required to conduct 622 the assessment. This review describes tools which 623 utilize data collection through informant report, infor-624 mant interview and direct observation. Informants 625 may provide inaccurate answers to IADL questions 626 due to their perception of the "normal" aging pro-627 cess or the stigma surrounding cognitive impairment 628 [10]. Direct observation is generally considered the 629 gold standard of IADL assessment, demonstrated 630 by the excellent diagnostic accuracy scores reported 631 [12, 13, 23]. However, such tools require signifi-632 cant staff training, time, and resources which may 633 be inappropriate for wide-scale dementia screening 634 in LMICs. The WHO mhGAP (Mental Health Gap 635 Action Programme) proposes that community health 636 workers could deliver interventions and diagnostic 637 services, with basic training. Thus the most appro-638 priate tool for dementia screening in LMICs may be 639 short, simple to score IADL questionnaires, based on 640 informant report, tailored for use in community set-641 tings [3]. In four studies, where IADL assessments 642 were carried out by community/field workers, good 643 diagnostic accuracy and inter-rater reliability were 644 found [10, 26-28]. However, Stone et al. [10] found 645 significant discrepancy in diagnostic accuracy val-646 ues in a longitudinal follow up, with baseline scores 647 showing excellent accuracy for identifying dementia 648 (AUC: 0.99-0.90) and one year follow-up indicating 649 poor accuracy (AUC: 0.625). Baseline assessment 650 was conducted by a skilled health-care team while 651 longitudinal follow-up was carried out by village 652 enumerators. It is proposed that discrepancies were 653 due to subjectivity in interpreting answers provided 654 to the questions introduced by village enumerators. 655 This highlights the importance of appropriate asses-656 sor training and selecting IADL tools which do not 657 require a high dependency on individual judgement 658 in the grading process, such as dichotomous scales 659 (e.g., "yes/no"). 660

661 Strengths and limitations of this review

A significant strength of this review was our comprehensive and rigorous search strategy (see

Supplementary Material) and use of multiple elec-664 tronic databases to identify potential articles for 665 inclusion. We also hand-searched reference lists of all 666 included articles to maximize the scope of our search. 667 We carried out independent title, abstract, and full-668 text screening and all disagreements were adjudicated 660 by a third reviewer. Our quality assessment indi-670 cated that, although most diagnostic accuracy studies 671 included demonstrated some risk of bias, overall, 672 they showed moderate-good quality. However, we 673 only included articles available in English due to lim-674 ited resources and may not have captured all relevant 675 IADL tools for LMICs. For example, we have limited 676 representation of Asian countries despite significant 677 work reported on cognitive assessments in Asia [39]. 678 Additionally, we excluded studies which combined 679 IADL questions with cognitive assessments within 680 one tool (e.g., Everyday Cognition Scale [40]) as 681 they did not fall within the strict remit of our review 682 question. These tools could also be considered within 683 the diagnostic process in LMICs, and further inves-684 tigation should determine how useful they may be. 685 A variety of IADL tools were assessed within this 686 review across a diverse range of populations. As such, 687 a meta-analysis was inappropriate to conduct at this 688 time but may be useful in the future when greater evi-689 dence bases are built for discrete measures. At this 690 time, the evidence for any tool is limited by incon-691 sistencies in validation methods, and lack of external 692 validation across all scales. As such, we do not rec-693 ommend any particular IADL tool as a diagnostic aid 694 for dementia in LMICs but do provide suggestions to 695 bridge this gap. 696

Recommendations for future research

A significant gap identified by this review is the lack of research around the generalizability of IADL tools, both across LMICs and within LMICs, as illustrated by the seven newly developed tools across six LMICs included in this review. Their item domains are similar; for example, both the EASI and the IDEA-IADL consider variations in ability to be involved in family matters and to take part in festivals and ceremonies [10, 26-28]. However, there has been no investigation into the feasibility of using bespoke IADL tools created for a specific LMIC in LMICs that hold similar cultural ideals. In contrast, there is significant evidence that tools which have been translated and adapted from Western high-income countries are feasible and acceptable to use in South America. For example, the FAQ

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shows acceptable-excellent diagnostic accuracy in 714 Brazil [20-22] and is one of the most commonly used 715 IADL scales worldwide [41]. This lends more con-716 fidence to the generalizability of translated scales on 717 a global scale, but these tools have not been investi-718 gated in Africa or South Asian countries which may 710 have unique cultural expectations, as discussed previ-720 ously. Therefore, we recommend that already existing 721 tools-either translated from Western high-income 722 countries or bespoke tools developed for LMICs (e.g., 723 EASI, IDEA-IADL) be considered and evaluated for 724 use before new scales are developed for specific target 725 populations. 726

Additionally, within LMICs there is limited under-727 standing of how transferable IADL tools of all 728 types are between urban and rural communities. For 729 example, most translated tools investigated in South 730 America were applied in clinical urban environments 731 and required skilled professionals to conduct the 732 assessments, which may not be applicable or feasible 733 for rural communities. In contrast, Edjolo et al. [29] 734 highlights that items included in the CA-DFI may 735 not be applicable to urban settings, such as assess-736 ing one's ability to work in fields. As such, suitable 737 urban alternatives need to be validated. Only two 738 studies explicitly included both urban and rural com-739 munities, highlighting a significant gap that should 740 be addressed through future studies [29, 31]. 741

A major limitation to the current state of research 742 is the lack of external validation of IADL tools 743 within LMICs. Most studies included in this review 744 involved scale development or initial validation. For 745 the majority, reliability and technical validity were 746 established, whereby IADL tools showed accept-747 able internal consistency, inter/intra-rater reliability, 748 and associations with other measures of cognitive 749 impairment (e.g., cognitive scales). However, without 750 external validity, findings of each IADL tool cannot 751 be generalized to communities beyond those investi-752 gated or to individuals who present in a different way 753 (e.g., prodromal dementia). This is particularly rele-754 vant to newly developed tools for target populations 755 as translated tools have generally demonstrated good 756 validity in populations from different backgrounds 757 and cultures, such as the FAQ [20-22, 41-43]. Several 758 studies also excluded people with physical impair-759 ments or other neurological conditions [12, 13, 16, 760 17, 22-25, 34, 44], limiting our understanding of how 761 IADL tools might distinguish dementia from other 762 disorders in a population-level cohort. The validity 763 of IADL tools could also be strengthened by estab-764 lishing their relationship with recognized objective 765

gold-standard biomarkers, such as blood tests and neuroimaging [45]. While this may not be standard clinical practice in LMICs due to the expensive nature and resource-intensity of these biomarkers, it would improve confidence for clinicians to apply these simple IADL tools as diagnostic benchmarks. Ideally, further technical, and external validity within a population sample should be established before wide-scale adoption of an IADL tool within a LMIC.

Implications for practice

Due to limited financial and healthcare resources within LMICs, it is vital to establish simple, sensitive dementia screening and diagnostic tools to promote early detection [3]. Timely diagnosis allows individuals and their families to better understand the diagnosis, consider appropriate care and treatment plans and avail of non-pharmacological interventions and drug therapies early in the disease [46]. Beyond clinical use, early and accurate diagnosis is important for researchers and policymakers to identify the true prevalence of dementia in LMICs and develop appropriate action plans for global dementia strategies. Additionally, IADL tools could support both clinicians and researchers by identifying changes in function due to disease progression and determining care needs of an individual. This review has indicated that IADL tools which are culturally appropriate and applicable to settings of different language, education and healthcare resources can be implemented in LMIC settings with good-excellent accuracy for distinguishing dementia from normal ageing. It is important to acknowledge, however, that there is no "perfect" measure; diagnostic practice generally requires a variety of tools to support clinical decisionmaking. It is recommended that IADL tools are used in combination with at least one brief global cognitive assessment [3], such as translated versions of the Mini-Mental State Examination or culturallytailored assessments such as the IDEA Cognitive screen [10, 39]. This combination can strengthen the accuracy of the diagnostic battery. For example, Pandav et al. [27] reported the highest paired sensitivity (90.6%) and specificity (68.2%) when the EASI was coupled with a comprehensive cognitive battery. Similarly, Paddick et al. [28] found that the combination of both the IDEA-IADL and the IDEA cognitive screen showed the highest accuracy for distinguishing cognitive impairment from normal aging (AUC: 0.93) compared to single measures (AUC: 0.84-0.89). These measures could be supported by

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inexpensive digital markers, such as measures col-816 lected from wearable technology (e.g., gait, sleep), 817 which are culturally-naïve [47]. Such devices have 818 been found to be acceptable and feasible to use in 819 older adults in LMICs, as conducted by community 820 field workers [48] and are considered useful support-821 ive markers for dementia diagnosis in high-income 822 settings [49]. Further work is needed to 1) validate 823 their utility in the LMIC diagnostic pathway and 824 2) identify which combination of diagnostic tools 825 provides the greatest sensitivity and specificity for 826 identifying dementia in culturally-diverse LMIC set-827 tings. 828

829 CONCLUSION

This review synthesized the current literature on 830 the reliability, validity, and accuracy of IADL tools 831 for identifying dementia in LMICs. From our find-832 ings, we present the seven IADL tools with the 833 strongest evidence base. We also highlight key con-834 siderations for choosing an IADL tool for use in 835 an LMIC, such as selecting tools that are cultur-836 ally appropriate, account for bias introduced by 837 gender-roles and literacy rates, easy and quick to use 838 and which can be conducted by any volunteer with 839 the right training. There are significant gaps in the 840 research which must be addressed, including greater 841 technical validity against established gold-standard 842 biomarkers of dementia and external validation of 843 IADL tools within different regions, populations, cul-844 tures and across LMICs. Future work should consider 845 combinations of diagnostic markers, such as IADL 846 tools, brief cognitive assessments, and novel mea-847 sures such as those derived from digital technology, 848 to establish the most appropriate and sensitive diag-849 nostic toolkit for dementia in LMICs. 850

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SUPPLEMENTARY MATERIAL

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