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Modifiable Risk Factors for Prevention of Dementia in Midlife, Late Life and the Oldest-Old: Validation of the LIBRA Index

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Abstract.

Background: Recently, the Lifestyle for BRAin health (LIBRA) index was developed to assess an individual's prevention potential for dementia.

Objective: We investigated the predictive validity of the LIBRA index for incident dementia in midlife, late life, and the oldest-old.

Methods: 9,387 non-demented individuals were recruited from the European population-based DESCRIPA study. An individual's LIBRA index was calculated solely based on modifiable risk factors: depression, diabetes, physical activity, hypertension, obesity, smoking, hypercholesterolemia, coronary heart disease, and mild/moderate alcohol use. Cox regression was used to test the predictive validity of LIBRA for dementia at follow-up (mean 7.2 y, range 1–16).

Results: In midlife (55–69 y, n = 3,256) and late life (70–79 y, n = 4,320), the risk for dementia increased with higher LIBRA scores. Individuals in the intermediate- and high-risk groups had a higher risk of dementia than those in the low-risk group. In the oldest-old (80–97 y, n = 1,811), higher LIBRA scores did not increase the risk for dementia.

Conclusion: LIBRA might be a useful tool to identify individuals for primary prevention interventions of dementia in midlife, and maybe in late life, but not in the oldest-old.

Keywords: Aging, dementia, modifiable risk factors, prevention

INTRODUCTION

Dementia is one of the fastest growing health problems worldwide, without any cure available so far for its most common forms including Alzheimer's disease. This urges the need for prevention. For prevention, early identification of individuals at high risk for dementia is of great importance. In particular, a better understanding of the role of modifiable risk factors in predicting dementia is crucial, as these risk factors constitute promising targets for prevention strategies across a wide age spectrum [1, 2].

Recently, the Lifestyle for BRAin health (LIBRA) index was developed to assess an individual's room for prevention of dementia in midlife [3]. The index is a dementia risk score that is defined based on empirical evidence from the existing literature and expert consensus. It includes 12 easily assessable and solely modifiable health and lifestyle factors that are all within the reach of interventions (e.g., physical activity and obesity).

Previously defined dementia risk indices are based on single cohort studies rather than the total evidence in the existing literature [4–8] and/or included non-modifiable risk factors (e.g., sex, APOE genotype) [4–9]. This makes them less generalizable and less suitable for global prevention strategies. Moreover, some indices were defined in a midlife population [4] whereas others were defined in an older population [6]. Previous research has shown that dementia risk factors may act differently throughout the adult life span [10, 11]. Thus, more knowledge is needed on the validity of modifiable risk factors in midlife, late life, and the oldest-old to introduce timely and appropriate prevention strategies.

The aim of our study was to test the ability of the LIBRA index to assess the prevention potential by investigating the predictive validity of the LIBRA index for incident dementia in midlife, late life, and the oldest-old in a large multicenter European population-based cohort.

METHODS*Participants*

Participants were recruited from the DESCRIPA study, a multicenter study consisting of eight harmonized European population-based cohorts. For the current study, six cohorts fulfilled our inclusion criteria: The Swedish Prospective Population Study of Women (GPPSW) [12] and Gerontological and Geriatric Population Study of 85-year-olds (GH85) [13], the Italian Longitudinal Study of Aging (ILSA), [14] the Dutch Longitudinal Aging Study Amsterdam (LASA) [15] and Maastricht Aging Study (MAAS), [16, 17] and the French Personnes Agées QUID study (PAQUID). [18] The medical ethics committee at each center approved the study. All subjects provided informed consent.

Inclusion criteria for the current study were age ≥ 55 years, good subjective general health, a direct or indirect baseline measure of at least 7 out of 9 modifiable risk factors as described below, information on educational level, and at least one clinical follow-up. Individuals with dementia at baseline were excluded.

Clinical assessment was performed according to each study protocol. Generally, it included a medical history by interview or questionnaire, medical and neurological examination, and cognitive assessment.

For a subgroup, blood sampling was performed. Clinical follow-up assessment was performed at study-specific intervals up to 16 years after baseline. The outcome measure was incident dementia according to the DSM-III-R [19] or for the LASA study defined as impairment in multiple cognitive domains [15].

Risk factors

Modifiable risk factors were selected from the LIBRA index [3]. We had data available on nine of the 12 risk/protective factors: depression, diabetes, physical inactivity, hypertension, obesity, smoking, hypercholesterolemia, coronary heart disease, and mild/moderate alcohol use. No data were available on cognitive activity, renal dysfunction, and adherence to a Mediterranean diet.

To compose the LIBRA index, direct or indirect measures of the nine risk factors were used according to each study-specific protocol (Supplementary Tables 1 and 2). The LIBRA index was calculated following a previously applied approach based on the relative risk (RR) of all risk factors separately [4, 9], as recently reported [3]. Briefly, the natural logarithm (ln) of the RR was calculated for each factor. Next, these were standardized by taking the lowest ln (RR) as a reference value (score 1) and dividing all other values by this value. Finally, summing the scores of the risk factors resulted in the total LIBRA index. A higher LIBRA score indicates a higher risk for dementia.

Since age, sex, and education are well-established risk factors for dementia, albeit non-modifiable, two extended versions of the LIBRA index were calculated by adding standardized scores of age, sex, and education to the LIBRA index based on the beta weights reported by Anstey and colleagues (age for males, age for females, and years of education; see models 2 and 3 below) [9]. Our educational level categories were slightly different from those used by Anstey and colleagues [9] because the data available in this study were not always coded according to their coding system (Supplementary Table 1). The standardized score for each risk factor is listed in Table 1.

Statistical analyses

Baseline differences in proportion of risk factors between subjects with and without dementia at follow-up were analyzed using χ^2 or Fisher's exact

Table 1
Risk factor scores for incident dementia used to calculate the LIBRA index

		Risk score
<i>Demographic factors</i>		
Age for males	<65 years	0
	65–69 years	0.4
	70–74 years	5.2
	75–79 years	6.8
	80–84 years	11.2
	85–89 years	14.1
Age for females	≥90 years	16.4
	<65 years	0
	65–69 years	2.1
	70–74 years	6.2
	75–79 years	9.2
	80–84 years	12.4
Educational level	85–89 years	15.3
	≥90 years	17.6
	High	0
	Medium	1.4
	Low	2.7
<i>Risk factors of LIBRA index</i>		
Depression	No	0
	Yes	2.1
Hypertension	No	0
	Yes	1.6
Obesity	No	0
	Yes	1.6
Smoking	No	0
	Yes	1.5
Hypercholesterolemia	No	0
	Yes	1.4
Diabetes	No	0
	Yes	1.3
Physical inactivity	No	0
	Yes	1.1
Coronary heart disease	No	0
	Yes	1.0
Low/moderate alcohol use	No	0
	Yes	–1.0

Risk scores were calculated based on relative risks (for the LIBRA risk factors)³ and beta coefficients (for the demographic factors)⁹ in the literature, as described in the Methods. The LIBRA index is the sum of the individual risk scores. LIBRA, Lifestyle for BRAin health.

tests. Cox proportional hazards models were used to examine prediction of dementia on a 16-year follow-up using the LIBRA index (model 1). We performed analyses separately by age groups. Due to the increase in life expectancy and in the mean age of retirement, midlife was defined age 55–69 years, late life age 70–79 years, and oldest-old age 80–97 years. First, analyses were performed for the continuous LIBRA index and C-statistics were calculated as a measure of predictive accuracy. Next, the LIBRA index was categorized into three age-specific risk groups based on tertiles (i.e., low-, intermediate-, high-risk). Additionally, the prediction of dementia by two alternate

versions of the LIBRA index was tested: LIBRA + educational level (model 2); and LIBRA + age, sex and educational level (model 3). Survival curves of all Cox proportional hazards models were plotted. All analyses were corrected for center. Statistical analyses were done with the SPSS version 23.0 (Chicago, IL, USA) and R Survival package, function *SurvConcordance*, with significance set at $p < 0.05$.

RESULTS

Sample characteristics

9,387 participants were included with a mean age of 72.9 (SD 7.3, 55–97) years, of whom 5141 (55%) were female. 31% of the cases had data on APOE genotype available. The average LIBRA index was 2.9 (SD 2.0, range –1.0 to 10.5). After an average follow-up of 7.2 years (SD 3.6, range 1 to 16), 1120 (12%) individuals progressed to dementia. The dementia incidence rate was 16.8 (95% CI 16.0–17.6) per 1000 person-years. The availability and prevalence of risk factors is presented by outcome for each age group separately in Table 2. The availability of risk factors in the total cohort is presented in Supplementary Table 3. Overall, APOE genotype ($\epsilon 4$ carrier versus non-carrier) did not influence predictive accuracy of the LIBRA index for progression to dementia (model 1; LIBRA*APOE HR = 0.97, $p = 0.664$).

Modifiable risk factor profiles for dementia

Midlife, age 55–69 years

3,256 individuals were classified in the midlife group (mean age 65.0 (SD 4.0) years; 51% female), of whom 190 (6%) progressed to dementia after an average follow-up of 8.1 (SD 3.5) years. The dementia incidence rate was 7.2 (95% CI 6.3–8.1) per 1000 person-years. The average LIBRA index for individuals in midlife was 2.6 (SD 2.1, range –1.0 to 10.5). The risk for dementia increased on a log-linear scale with higher LIBRA scores (HR = 1.10, 1.02–1.18, $p = 0.020$; C statistic 0.57, SE 0.03; model 1). When educational level (model 2) or age, sex, and educational level (model 3) were added to the LIBRA index, the risk for dementia slightly increased (HR = 1.13, 1.06–1.21, $p < 0.001$ and HR = 1.11, 1.04–1.17, $p = 0.001$ respectively).

Next, individuals were classified based on tertiles in low ($n = 1070$, score –1.0 to 1.5), intermediate ($n = 1132$, score 1.6 to 3.5), and high ($n = 1054$, score 3.6 to 10.5) dementia risk groups. Table 3

describes the proportions of risk factors in the three risk groups. Individuals in the intermediate- and high-risk groups showed a higher risk for dementia than those in the low-risk group (intermediate HR = 1.56, 1.04–2.36, $p = 0.033$; high HR = 1.92, 1.25–2.96, $p = 0.003$; Table 4; Fig. 1). When education (model 2) or age, sex, and education (model 3) were added to the LIBRA index results remained similar (Table 4), except that now the increased risk for dementia in the intermediate risk group compared to the low-risk group did not reach statistical significance in model 2 (HR = 1.40, 0.96–2.03, $p = 0.078$). The high-risk group had an increased dementia risk compared to the intermediate risk group in model 3 (HR = 1.44, 1.04–1.99, $p = 0.029$).

Late life, age 70–79 years

4,320 individuals were classified in the late life group (mean age 74.5 (SD 2.9) years; 56% female), of whom 580 (13%) progressed to dementia after an average follow-up of 7.3 (SD 3.5) years. The dementia incidence rate was 18.8 (95% CI 17.6–20.0) per 1000 person-years. The average LIBRA index for individuals in late life was 3.2 (SD 2.0, range –1.0 to 9.6). The risk for dementia increased on a log-linear scale with higher LIBRA scores (HR = 1.08, 1.03–1.13, $p = 0.002$; C statistic 0.50, SE 0.01; model 1). The predictive accuracy for dementia slightly increased when educational level (model 2) or age, sex, and educational level (model 3) were added to the index (HR = 1.10, 1.06–1.15, $p < 0.001$ and HR = 1.14, 1.10–1.17, $p < 0.001$, respectively).

Individuals in the intermediate ($n = 1,496$, score 2.4 to 4.1) and high ($n = 1,392$, score 4.2 to 9.6) risk groups presented a greater risk of incident dementia than those in the low ($n = 1,432$, score –1.0 to 2.3) risk group (Intermediate HR = 1.25, 1.02–1.52, $p = 0.030$; High HR = 1.38, 1.11–1.72, $p = 0.005$; Table 4; Fig. 1). When education (model 2) or age, sex, and education (model 3) were added to the LIBRA index, results remained similar (Table 4), except that the high-risk group had an increased risk for dementia compared to the intermediate risk group in model 3 (HR = 2.12, 1.73–2.61, $p < 0.001$).

Oldest-old, age 80–97 years

1,811 individuals were classified in the oldest-old group (mean age 83.2 (SD 2.5) years; 58% female), of whom 350 (19%) progressed to dementia after an average follow-up of 5.5 (SD 3.2) years. The dementia incidence rate was 36.4 (95% CI 34.1–38.7) per 1000 person-years. The average LIBRA index for

Table 2
Risk factors by outcome in midlife, late life, and the oldest-old

	Midlife		Late life		Oldest-old		<i>p</i> values
	No dementia at follow-up N = 3,066	Dementia at follow-up N = 190	No dementia at follow-up N = 3,740	Dementia at follow-up N = 580	No dementia at follow-up N = 1,461	Dementia at follow-up N = 350	
<i>Demographic factors</i>							
Age							
<65 years	1,077 (35)	19 (10)	–	–	–	–	
65–69 years	1,989 (65)	171 (90)	–	–	–	–	
70–74 years	–	–	2,036 (54)	238 (41)	–	–	
75–79 years	–	–	1,704 (46)	342 (59)	–	–	
80–84 years	–	–	–	–	1,080 (74)	241 (69)	0.082
85–89 years	–	–	–	–	347 (24)	103 (29)	
≥90 years	–	–	–	–	34 (2)	6 (2)	
Female	1,570 (51)	105 (55)	2,045 (55)	378 (65)	796 (55)	247 (71)	<0.001
Education level							
Low	1,043 (34)	77 (41)	1,875 (50)	304 (52)	861 (59)	202 (58)	0.056
Medium	1,624 (53)	94 (50)	1,524 (41)	238 (41)	492 (34)	133 (38)	
High	399 (13)	19 (10)	341 (9)	38 (7)	108 (7)	15 (4)	
<i>Risk factors of LIBRA index</i>							
Depression	665(3,065 (22)	40/189 (21)	1,202/3,730 (32)	126/579 (22)	424/1,456 (29)	64/350 (18)	<0.001
Hypertension	1,582/3,045 (52)	132/188 (70)	2,749/3,725 (74)	436/573 (76)	1,150/1,452 (79)	258/348 (74)	0.040
Obesity	524/3,008 (17)	30/187 (16)	664/3,617 (18)	73/556 (13)	185/1,357 (14)	27/320 (8)	0.009
Smoking	884/3,061 (29)	32/190 (16)	1,228/3,637 (34)	135/572 (24)	392/1,455 (27)	50/348 (14)	<0.001
Hypercholesterolemia	1,072/3,062 (35)	47/190 (25)	1,275/3,737 (34)	169/579 (29)	378/1,451 (26)	89/348 (26)	0.856
Diabetes	229/2,832 (8)	25/185 (14)	445/3,636 (12)	59/569 (10)	141/1,444 (10)	34/348 (10)	0.997
Physical inactivity	1,098/2,283 (48)	112/160 (70)	1,346/2,169 (62)	347/470 (74)	491/670 (73)	178/228 (78)	0.152
Coronary heart disease	634/3,066 (21)	42/190 (22)	1,122/3,740 (30)	146/580 (25)	515/1,460 (35)	103/350 (29)	0.038
Low/moderate alcohol use	1,684/3,039 (55)	86/189 (46)	2,570/3,713 (69)	325/575 (57)	1,015/1,454 (70)	219/346 (63)	0.019

Results are numbers (%) of individuals with risk factors by outcome. LIBRA, Lifestyle for BRAin health. *P* values are presented for the comparison of individuals with and without dementia at follow-up. Significance was set at *P* < 0.05.

Table 3
Proportions of risk factors stratified by age and by the LIBRA risk score groups for dementia

	Midlife			Late life			Oldest-old		
	Low risk N = 1,028	Intermediate risk N = 1,146	High risk N = 1,082	Low risk N = 1,432	Intermediate risk N = 1,496	High risk N = 1,392	Low risk N = 602	Intermediate risk N = 655	High risk N = 554
Age									
<65 years	53	32	18	—	—	—	—	—	—
65–69 years	47	69	82	—	—	—	—	—	—
70–74 years	—	—	—	52	52	54	—	—	—
75–79 years	—	—	—	48	48	46	—	—	—
80–84 years	—	—	—	—	—	—	73	72	74
85–89 years	—	—	—	—	—	—	25	26	24
≥90 years	—	—	—	—	—	—	2	3	2
Female	55	51	48	58	59	51	58	60	54
Education									
Low	29	33	40	45	48	58	55	59	63
Medium	56	55	48	46	43	34	38	34	31
High	14	12	12	9	9	8	8	6	7
Depression	2	11	52	4	26	63	1	20	63
Hypertension	9	63	84	48	81	94	54	86	96
Obesity	3	16	32	7	10	37	5	14	21
Smoking	13	28	44	14	25	59	9	25	42
Hypercholesterolemia	21	29	52	17	35	49	12	19	49
Diabetes	2	5	18	3	9	25	3	7	21
Physical inactivity	23	55	81	41	75	87	55	86	93
Coronary heart disease	5	17	40	10	34	45	17	37	50
Low/moderate alcohol use	63	56	46	80	60	62	85	62	59
Dementia at follow-up	4	7	7	14	15	11	23	19	16

Results are proportions (%) of individuals with risk factors for the low, medium, and high dementia risk groups based on the tertile classification of the total risk score. Note that total numbers differ between risk factors because of missing data, as described in Table 2 and Supplementary Table 3.

Table 4
Prediction of dementia at follow-up stratified by age groups and by the LIBRA risk score groups for dementia

Index	Risk group	Midlife		Late life		Oldest-old	
		Hazard ratio	p-value	Hazard ratio	p-value	Hazard ratio	p-value
1) LIBRA	Low	Reference		Reference		Reference	
	Intermediate	1.56 (1.04–2.36)	L: p = 0.033 H: p = 0.208	1.25 (1.02–1.52)	L: p = 0.030 H: p = 0.328	0.91 (0.71–1.17)	L: p = 0.461 H: p = 0.457
	High	1.92 (1.25–2.96)	L: p = 0.003	1.38 (1.11–1.72)	L: p = 0.005	0.82 (0.62–1.09)	L: p = 0.164
2) LIBRA + education	Low	Reference		Reference		Reference	
	Intermediate	1.40 (0.96–2.03)	L: p = 0.078 H: p = 0.199	1.36 (1.11–1.66)	L: p = 0.003 H: p = 0.092	1.29 (1.00–1.66)	L: p = 0.051 H: p = 0.111
	High	1.74 (1.18–2.57)	L: p = 0.005	1.62 (1.29–2.03)	L: p < 0.001	1.04 (0.78–1.40)	L: p = 0.788
3) LIBRA + age, sex, education	Low	Reference		Reference		Reference	
	Intermediate	1.65 (1.07–2.52)	L: p = 0.022 H: p = 0.029	1.37 (1.11–1.69)	L: p = 0.003 H: p < 0.001	1.15 (0.88–1.50)	L: p = 0.294 H: p = 0.066
	High	2.36 (1.53–3.64)	L: p < 0.001	2.12 (1.73–2.61)	L: p < 0.001	1.48 (1.13–1.94)	L: p = 0.004

Results are hazard ratios with 95% confidence intervals for the LIBRA index (model 1) and the 2 additional models with demographics added to the index (models 2 and 3), for midlife, late life, and the oldest-old. P-values are comparisons with the other groups: L = p-value compared to the low-risk group. H = p-value compared to the high-risk group. LIBRA model + education: midlife low n = 1,125 (score –1.0 to 3.1), intermediate n = 1,140 (score 3.1 to 5.4), high n = 991 (5.5 to 12.2), late life low n = 1,270 (score –1.0 to 4.0), intermediate n = 1,652 (score 4.1 to 6.2), high n = 1,398 (score 6.3 to 12.3); oldest-old low n = 527 (score –1.0 to 4.0), intermediate n = 687 (score 4.1 to 5.8), high n = 597 (score 5.8 to 12.2); LIBRA model + age, sex, and education: midlife low n = 1,075 (score –1.0 to 3.7), intermediate n = 1,109 (score 3.7 to 6.2), high n = 1,072 (score 6.2 to 13.6), late life low n = 1,438 (score 4.2 to 10.8), intermediate n = 1,481 (score 10.8 to 13.3), high n = 1,401 (score 13.3 to 21.2); oldest-old low n = 623 (score 10.2 to 16.6), intermediate n = 578 (score 16.6 to 18.7), high n = 610 (score 18.7 to 27.0). LIBRA, Lifestyle for BRAin health.

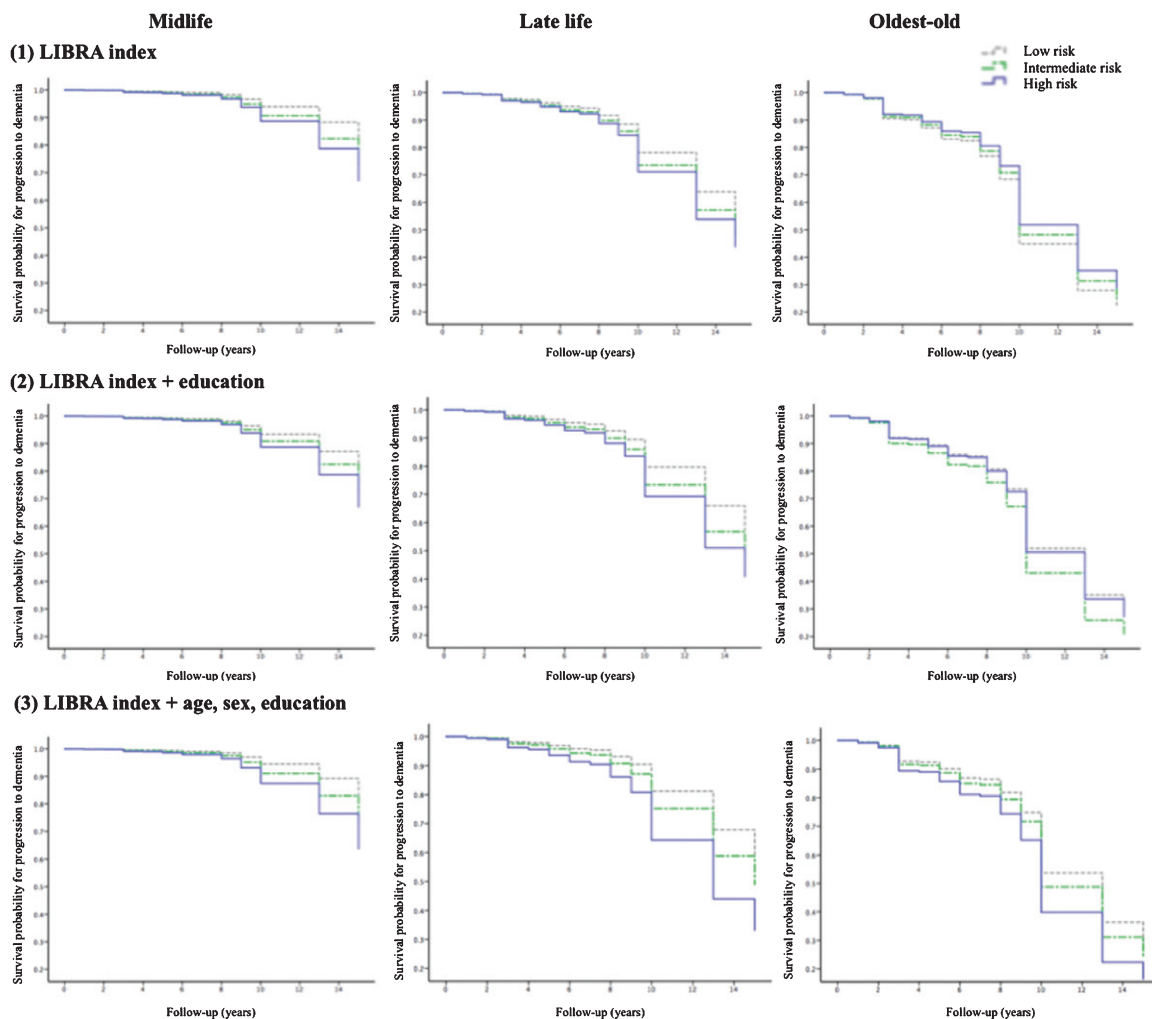


Fig. 1. Survival curves for the LIBRA risk groups. Individuals are classified in risk groups based on tertiles of the LIBRA index score. On the left are survival curves for individuals in midlife, in the middle for individuals in late life, and on the right for the oldest-old. (1) Model 1: Model with the LIBRA index, (2) Model 2: extended model with the LIBRA index + education, and (3) Model 3: extended model with the LIBRA index + age, sex, and education. The dotted grey line represents individuals in the low-risk group, the dotted green line represents individuals in the intermediate-risk group, and the solid blue line represents individuals in the high-risk group.

the oldest-old was 2.9 (SD 1.8, range -1.0 to 9.5). The risk for dementia decreased on a log-linear scale with higher LIBRA scores (HR = 0.93, 0.88–0.99, $p = 0.031$; C statistic 0.54, SE 0.02; model 1). No decreased risk for dementia was found when educational level (model 2) was added to the index (HR = 1.00, 0.94–1.05, $p = 0.870$). When age, sex, and educational level (model 3) were added to the index, the risk for dementia increased with higher LIBRA scores (HR = 1.08, 1.13–1.17, $p < 0.001$).

Individuals in the intermediate ($n = 655$, score 2.1 to 3.7) and high ($n = 554$, score 3.8 to 9.5) risk groups did not present greater risk of incident dementia compared to those in the low ($n = 602$, score

-1.0 to 2.0) risk group (Intermediate HR = 0.91, 0.71–1.17, $p = 0.461$; High HR = 0.82, 0.62–1.09, $p = 0.164$; Table 4; Fig. 1). When education (model 2) or age, sex, and education (model 3) were added to the LIBRA index, results were similar (Table 4) but the high-risk group had an increased risk for dementia compared to the low-risk group in model 3 (HR = 1.48, 1.13–1.94, $p = 0.004$).

Modified LIBRA index

As obesity and hypertension are considered to be major risk factors for later development of dementia only in midlife, we performed an additional analysis

based on model 1 but with exclusion of obesity and hypertension from the LIBRA index for late life and the oldest-old. In late life, higher modified LIBRA scores were associated with an increased risk for dementia (HR = 1.11, 1.05–1.17, $p < 0.001$). After classifying individuals in risk groups based on tertiles of the modified LIBRA index, we found that the intermediate- and high-risk groups had an increased risk for dementia compared to the low-risk group in late life (intermediate HR = 1.41, 1.16–1.72, $p = 0.001$; high HR = 1.47, 1.16–1.86, $p = 0.002$). In the oldest-old, the risk for dementia did not increase with higher modified LIBRA scores (HR = 0.95, 0.89–1.02, $p = 0.179$). The intermediate- and high-risk groups did not have a higher risk for dementia than the low-risk group in the oldest-old (intermediate HR = 0.97, 0.75–1.25, $p = 0.801$; high HR = 0.87, 0.65–1.15, $p = 0.310$).

DISCUSSION

Our study showed that modifiable risk factors based on the LIBRA index could be used to quantify dementia risk in midlife and late life, but not in the oldest-old. Our findings highlight the need for individuals to attain a low LIBRA score, e.g., by adapting a brain-healthy lifestyle and prevention of chronic disease, in order to reduce their risk for dementia.

Alzheimer's disease is the most common form of dementia. Recent evidence suggests that one in three up to one in two Alzheimer's disease cases are potentially attributable to modifiable risk factors. Particularly improved prevention of vascular and metabolic morbidity and depression, and higher levels of education could help to reduce the dementia prevalence worldwide [1, 20]. Our results indicate that the LIBRA index might be a useful tool to identify individuals for primary prevention interventions of dementia and monitor individuals risk-change over time. Moreover, all LIBRA factors can be easily assessed based on interview and routine medical assessment, which makes it feasible and easily applicable, e.g., in daily primary care practice [21]. Also, other risk factors can be added to the LIBRA index to improve its predictive ability as new evidence becomes available.

Our multicenter data support previous findings that modifiable risk factors may act differently in midlife and later life and that this affects the prediction of dementia. The LIBRA index was specifically designed to determine an individual's room for prevention in midlife. We found that the current index is

indeed mainly useful to predict dementia in midlife, and in late life particularly in its modified form, but not in the oldest-old. This corroborates previous reports showing variability in the effects of certain risk factors over the life-course, with some factors having maximum penetrance in midlife rather than in later life, including obesity and hypertension [11, 22]. Moreover, most inconsistent findings on predictive ability of risk factors come from late life studies, probably reflecting multi-comorbidity at the oldest age [10, 11, 23]. Our results show that excluding obesity and hypertension from the LIBRA index would not improve quantifying an individual's prevention potential in the oldest-old. However, we found that adding age, sex, and educational level to the LIBRA index in the oldest-old allowed a better differentiation between individuals with lower and higher risk for dementia, whereas this influence was much lower in midlife and early late life. This could partly be explained by the larger weight that was ascribed to older age but also suggests that demographic factors play a lesser role in midlife over and above modifiable factors. Lifestyle-related risk factors may also be affected by brain aging and ongoing cognitive decline, and therefore play a more important role in midlife and early late life. In addition, at the oldest age, underlying dementia-related pathology might have already accumulated so that the predictive value of risk factors becomes lower. The oldest-old might represent a selected group who have survived into old age because of (unknown) resilience factors that compensate for lifelong exposure to poor health and lifestyle.

The LIBRA index is unique in that it reflects an individual's prevention potential for dementia. The predictive accuracy of the LIBRA index for dementia was somewhat lower compared to that of other prediction indices [24]. This is not unexpected given that previous indices were maximized for risk prediction by including major predictors for dementia such as age, gender, education, and APOE genotype. It seems that most of the variance explained by these indices stem from the inclusion of such deterministic factors that are not amenable to change. Furthermore, previous indices were often developed based on a single cohort, and several lack validation in external datasets, increasing the likelihood of overestimating its predictive accuracy [24].

The strengths of our study included the large sample size, relatively long-term follow-up, and generalizability to a wide age spectrum. The use of a multicenter design allowed for a large sample

size and for testing of age-dependent effects. However, our study had several limitations. Indirect and center-specific measures of risk factors could have introduced heterogeneity in exposure classification. Furthermore, we used only nine out of twelve risk factors of the LIBRA index. The lack of three LIBRA factors (i.e., cognitive activity, renal dysfunction, and adherence to a Mediterranean diet) may have influenced the predictive validity of the LIBRA index. Also, not all individuals had data available for all nine risk factors but when we compared our findings to those for participants with all risk factors available, results remained similar (data not shown). Next, like other indices, LIBRA results in a simple additive score and interactions between risk factors were not taken into account, as this information is not available in the existing literature. Moreover, we used all-cause dementia as outcome measure, given the lack of information on type of dementia. Although Alzheimer's disease was likely the most common diagnosis, investigating the association between risk factors and specific subtypes of dementia would be relevant in future studies. Although our study had a relatively long follow-up (mean 7.2 years, range 1–16) some individuals likely would have progressed to dementia at a later stage, which could increase the predictive validity of the LIBRA index. We did not correct for mortality. Probably, people with higher LIBRA scores present with more multi-morbidity and were more likely to die as a competing risk for dementia. From this perspective, our results might underestimate the true potential of the LIBRA index. Finally, some individuals were already under treatment for a certain risk factor, which may limit the room for prevention.

In sum, our results support the role of modifiable risk factors in the development of dementia and demonstrate the utility and validity of the LIBRA index in midlife and late life up to 79 years. LIBRA can be a useful tool for raising people's awareness and for the identification of individuals who might benefit most from primary prevention strategies through lifestyle change or health management. Further validation of the LIBRA index is needed in external datasets and subgroups such as individuals with mild cognitive impairment.

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SUPPLEMENTARY MATERIAL

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