



Scale Development and Model Validation for the Process of Exercise Engagement for People with Prediabetes

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Purpose: This study had two objectives: 1) to develop a scale for the process of exercise engagement (SPEE) for prediabetic individuals (PDIs); 2) to validate a structural model for the process of exercise engagement for PDIs. **Methods:** A cross-sectional survey with simple random sampling was conducted from September 2013 to December 2015 (in Taiwan). A total of 310 PDIs were enrolled for scale development and model validation via item analysis, factor analyses, and structural equation modeling. The Kuo model was used as the basis for developing the Chinese version of the SPEE for PDIs. **Results:** The SPEE contains five subscales with a total of twenty-one items that account for 54.9% to 65.9% of the total variance explained for assessing participants' process of engagement during exercise. For Kuo model validation, the model measures indicated goodness of fit between the Kuo model and sample data. Analysis further revealed a direct effect between the creating health blueprints (CHB) stage and the spontaneous regular exercise (SRE) stage ($\beta=.60$). **Conclusion:** The SPEE includes five subscales for assessing the psychological transition and behavioral expression at each stage of the process of exercise engagement for PDIs. The SPEE for people with prediabetes provides deeper insights into the factors of behavioral change stages that are required to initiate long-term health care outcomes and avoid developing diabetes. These insights are significant as they allow for patient-specific mapping and behavior modification to effect exercise.

Key words: Prediabetes; Exercises; Prevention; Instruments; Structural Models

INTRODUCTION

The global population with prediabetes is increasing annually. The International Diabetes Federation has stated that 318 million people worldwide were estimated to have prediabetes in 2015, and this number is expected to reach 482 million by 2040 [1]. Prediabetes is a condition in which an individual's blood glucose exceeds the normal range, but he or she has not been diagnosed with diabetes. The criteria for a prediabetes diagnosis are fasting blood glucose levels between 100 and 125 mg/dl (known as im-

paired fasting glucose), 2-h postprandial plasma glucose between 140 and 199 mg/dl (known as impaired glucose tolerance), or hemoglobin A1c (HbA1c) between 5.7% and 6.5% [1]. Prediabetes is associated with a high risk of developing diabetes. If no active preventive approach is adopted during the prediabetic stage, prediabetes usually progresses to type 2 diabetes within an average of 3.6 years [2]. In one study, it was shown that annually in Taiwan, 3.2% of prediabetic individuals (PDIs) develop type 2 diabetes [3]. Prediabetes is also associated with increased risk of cardiovascular disease (14.8%), early stage nephropathy

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(17.7%), and retinopathy (7.9%) [4–7]. Diabetes has a negative impact on patients and requires complicated, lifelong medical care. Diabetes not only impacts the patients but also their families and their associations within society.

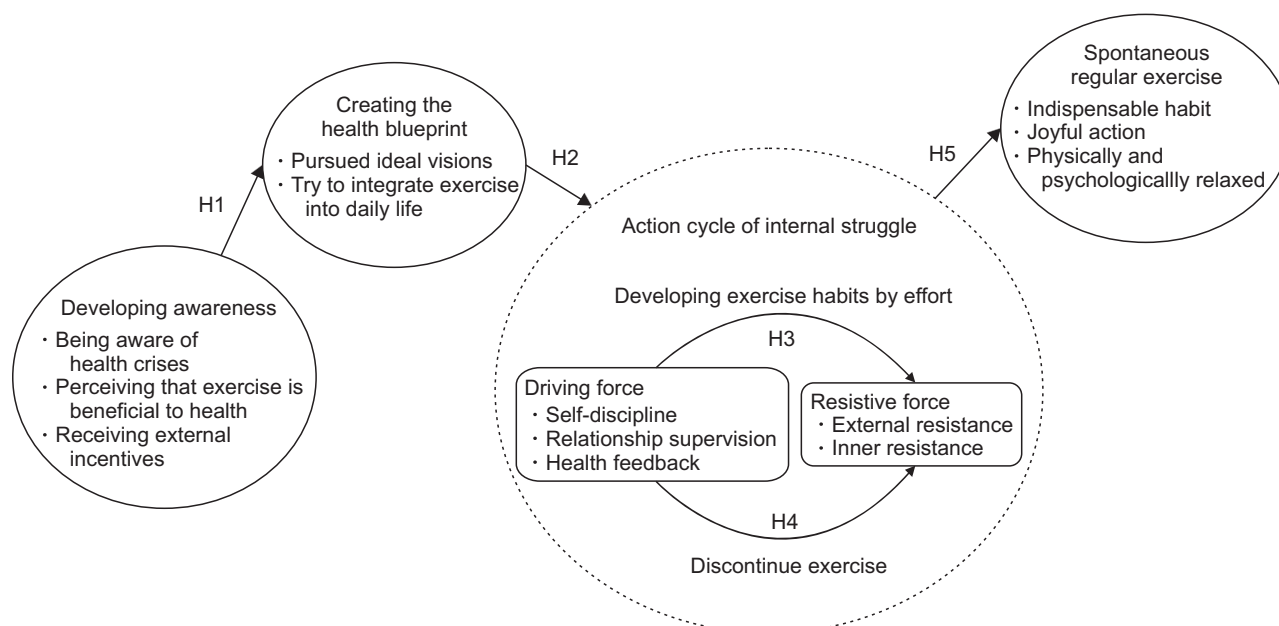
For PDIs, lifestyle modification is the easiest way to prevent or delay the onset of diabetes. Studies have shown that healthy eating habits, regular exercise, and weight control lower the incidence of developing diabetes among prediabetics [8–11]. Specifically, engaging in regular exercise is the most effective intervention for preventing diabetes. One study showed that while adopting a healthy diet can reduce diabetes incidence by 31.0%, regular exercise further reduces the incidence by 46.0% [12]. Another study indicated that PDIs engaging in regular exercise can reduce the risk of developing diabetes by between 30.0 and 69.0% compared to those who do not exercise [9–10,12]. Therefore, increasing physical activity among individuals with prediabetes should be prioritized to prevent the onset of diabetes.

Physical activity is associated with significant reductions in fasting glucose and HbA1c [13]. Physical activity is noted for: 1) reduced blood pressure, 2) reduced overall body fat, 3) reduced low-density lipoprotein cholesterol, 4) increased insulin sensitivity, 5) improved muscle strength, and 6) increased high-density

lipoprotein cholesterol [14,15]. Unfortunately, exercise is suboptimal among individuals with prediabetes in Taiwan for cultural reasons. Fewer than one-third of PDIs exercise routinely, and most of those who exercise do not do so for 150 minutes per week, which is the goal for effective diabetes prevention [16]. Furthermore, most of those who begin exercising after they recognize the importance of physical activity maintain their exercise routines for no more than six weeks [17]. Therefore, it is a major challenge for individuals with prediabetes to maintain consistent, lifelong exercise habits.

Based on previous qualitative research using grounded theory, the psychological status of PDIs' willingness to engage in exercise was modeled [18]. The Kuo model contains four stages: 1) developing awareness (DA) stage, 2) creating the health blueprint (CHB) stage, 3) action cycle of internal struggle (ACIS) stage, and 4) spontaneous regular exercise (SRE) stage. Furthermore, the ACIS stage contains two opposing forces: driving force (DF) and resistive force (RF) (Figure 1). Correctly assessing these two conflicting forces is critical for determining whether PDIs will enter the SRE stage.

According to the Kuo model, the DA stage is the key point in time at which PDIs may be strongly influenced to begin exercis-



H1=hypothesis 1; H2=hypothesis 2; H3=hypothesis 3; H4=hypothesis 4; H5=hypothesis 5.

Figure 1. Kuo model of exercise-engagement process for pre-diabetic individuals [18].

ing. After progressing through the DA stage, PDIs enter the CHB stage and focus on self-determination with regard to their exercise goals. However, in the process, all such individuals must progress through the ACIS stage. When the self-discipline, relationship supervision, and health feedback of the DF are more powerful than the RF, the exercise behavior is maintained. In contrast, if the internal and external resistance of the RF are stronger than the DF, exercise behavior is interrupted. The RF and DF work against each other until the individual breaks the action cycle to enter the SRE stage. In this model, the stages of PDIs' exercise-engagement process are closely interlinked and interrelated.

The objective of the present study was to develop a scale and validate the model for the process of exercise engagement in PDIs from a quantitative perspective, as seen in Figure 1 [18]. Currently, several exercise behavior assessment scales such as the Exercise Self-Efficacy Scale (ESES) [19], the Sport Motivation Scale (SMS) [20], and the Trans-theoretical Model (TTM) [21] are available. Using the mentioned scales and theories to validate the model of the process of exercise engagement for PDIs has several constraints: 1) none of these theories or scales conduct investigations from the PDIs' perspective; 2) none of the scales cover the complete exercise-engagement process; and 3) none of the scales evaluate PDIs' psychological or emotional status at each stage of the exercise-engagement process.

The TTM and the Kuo model are fundamentally similar, while differences remain to be understood within the context of the disease as the region of interest and control from the perspective of the PDI. Behavioral modification from the individual's perspective to benefit from making specific changes is similar but slightly different between the TTM and the Kuo model. The Kuo model is specific for understanding and developing a predictive methodology to assess PDIs and their capacity to alter long-term health outcomes. Presently, scales that are suitable for investigating PDIs' exercise-engagement process remain lacking. To effectively resolve this shortcoming, this study developed a scale consisting of five subscales based on the five subcategories of the Kuo model (Figure 1). The scale was subsequently used to collect responses from PDIs to evaluate the interrelationships between DA, CHB, DF, RF, and SRE based on the following hypotheses:

Hypothesis 1 (H1): A relationship exists between DA and CHB,

where DA will have a positive effect on CHB.

Hypothesis 2 (H2): A relationship exists between CHB and DF, where CHB will have a positive effect on DF.

Hypothesis 3 (H3) and Hypothesis 4 (H4): An inner conflict-based relationship exists between DF and RF, where the magnitudes of DF and RF may positively or negatively affect each other.

Hypothesis 5 (H5): A direct relationship exists between ACIS and SRE.

This study had two objectives: 1) to develop a scale for assessing the process of exercise engagement (SPEE) for PDIs, and 2) to validate the Kuo model for the process of exercise engagement for PDIs.

METHODS

1. Design and participants

A cross-sectional correlational survey with simple random sampling was conducted to validate the measurement scales and structural model. The participants were selected from a list of health checkups at a medical center in Hualien County in eastern Taiwan. The inclusion criteria consisted of: 1) elevated fasting blood glucose levels between 100 and 125 mg/dl, and 2) over 18 years of age. Exclusion criteria included: 1) previous diagnosis of type 1 or type 2 diabetes, 2) history of psychological illness, and 3) other serious illnesses (such as disability after stroke, liver disease, kidney disease, or cancer).

The study sample size was determined according to the following two criteria: (1) Stevens [22] suggested that five subjects should be allocated for each item of scale. (2) Gorsuch [23] and Thompson [24] suggested that at least 100 and 200 subjects are required for exploratory factor analysis (EFA) and structural equation modeling (SEM), respectively. As the initial version of the SPEE included forty-seven items (Appendix 1), a total of 235 ($5 \times 47 = 235$) subjects were required according to Criterion 1. In order to fulfill Criterion 2 (100 subjects for EFA and 200 subjects for SEM) simultaneously, a total of 300 subjects were required. A simple random sampling technique was employed for randomly selecting PDIs from the list of health checkups over the previous three years. Based on an estimated participation rate of 65%, 465 PDIs were randomly selected for invitation to engage in the

study. A total of 310 participants were recruited for the study where (n=110) for EFA and (n=200) for SEM analysis. The study was conducted from September 2013 to December 2015.

2. Data collection

After the research approval was granted by the institutional review board, the research institution made available a list of people with fasting blood glucose levels between 100 and 125 mg/dl as measured in health examinations over the past three years. Those who did not meet the inclusion criteria or whose phone number and address on the list were incorrect were eliminated from consideration, and subsequently, sampled randomly an estimated number of eligible individuals. A fasting blood glucose tracking notification was mailed to 450 eligible people, and 310 chose to participate (final participation rate of 68.9%). Potential participants were contacted via telephone and home visit and were sent multiple notifications inviting them to join this study. Participants were invited to the hospital for free blood glucose testing and a diabetes prevention health education seminar. In some instances, the researcher provided health education on an individual basis at the participants' homes. Participant recruitment continued until the required sample size was obtained.

3. Measures and variables

The Chinese version of the SPEE developed for use in this study is based on the Kuo model for the process of exercise engagement in PDIs in Taiwan (Figure 1) [18]. The scale includes five subscales for quantifying the PDIs' statuses in each of the five stages during the process. The subscales include: 1) the DA stage, 2) CHB stage, 3) DF stage, 4) RF stage, and 5) SRE stage. The scale's content validity was assessed by five experts to ensure its appropriateness. The resulting content validity index [25] of the five subscales ranged from 0.78 to 0.79, demonstrating the SPEE's acceptable content validity.

Preliminary testing was conducted for assessing the suitability of the SPEE. A total of thirty PDIs were asked about their understanding of the contents of the scale. Cronbach's α of .63~.94 for the five subscales showed good internal consistency of the initial version of the SPEE. Minor modifications were made to the scale based on their feedback. Appendix 2 summarizes the number of items in the SPEE at each stage of development. The final

version of the scale with a total of twenty-one items is provided in Appendix 3.

The score ranges were 4~20, 3~15, 6~30, 4~20, and 4~20 for the DA, CHB, DF, RF, and SRE subscales, respectively. Each item was scored from 1 (*never* for the DA, CHB, and DF subscales; *disagree* for the RF and SRE subscales) to 5 (*almost always* for the DA, CHB, and DF subscales; *strongly agree* for the RF and SRE subscales). Higher scores on the DA, CHB, DF, and SRE subscales represented higher levels of motivation for those stages, whereas a higher score on the RF subscale represented a greater resistance to exercise.

Questionnaires were administered either at the hospital or the participant's home. Researchers were made available to answer any questions that the participant may have had. The average time for completing the questionnaire was approximately thirty minutes. Participants could opt for either location at their discretion and convenience. The questionnaires were in Traditional Chinese, and all participants were literate. The completed questionnaires were managed by the research team and remain confidential.

4. Ethical considerations

This study was approved by the institutional research board (Number IRB 101-69) and the research institute. The research objectives and procedure were explained to each participant before the start of the study. Participants were enrolled in the study after providing informed, written consent and right of withdrawal at any time was open to their discretion. Participants' identifying information, i.e., name, phone number, and address, were kept private and confidential in a secure data base that could only be accessed by the interviewing researchers. The research team assured confidentiality with all data accessed.

5. Data analysis

The Chi-square test was conducted to compare the demographic variables of the study sample for item analysis/EFA with the study sample for confirmatory factor analysis (CFA)/SEM. A $p < .05$ was considered statistically significant.

The following steps were performed for the development of the SPEE based on the Kuo model (Figure 1):

- Step 1: Item analysis was implemented to exclude unquali-

fied items with low discrimination ability and internal consistency. Within each subscale, the mean, variance, skewness, corrected item total correlation [26], Cronbach's α coefficient after item deletion, and difference between lower 27% and upper 27% groups were evaluated.

- Step 2: EFA with maximum likelihood estimation was adopted for exploring latent structures with further item reduction in each subscale. First, the Kaiser–Meyer–Olkin (KMO) measure and Bartlett's test of sphericity were calculated and performed to evaluate sample adequacy for EFA. Parallel analysis [27] method was then used to decide the number of factors in each subscale. Only the factors with a derived eigenvalue by EFA greater than the corresponding values by parallel method were retained. Considering the sample size of 110 for EFA, the items with a factor loading of less than 0.55 [28] were also deleted due to low correlation with explored factors.

- Step 3: The SEM technique combining CFA and path analy-

sis was conducted to validate measurement specificity of the subscales as well as the Kuo model (Figure 1). Through CFA, the scales' reliability and validity obtained from EFA were confirmed. *Reliability and convergent validity of the factors were acceptable only if the corresponding composite reliability (CR) and average variance extracted (AVE) were greater than .70 and .50, respectively.* Items were deleted due to lack of convergent validity if the derived factor loading by CFA was less than 0.55. The path coefficients were estimated via path analysis to evaluate the relationships of the DA, CHB, DF, RF, and SRE stages. *The model's goodness of fit was evaluated based on χ^2/df , goodness-of-fit index (GFI), comparative fit index (CFI), root mean square error of approximation (RMSEA), parsimonious normed fit index (PNFI), normed fit index (NFI), incremental fit index (IFI), and Tucker-Lewis index (TLI) [29].*

The parameters with the corresponding criteria for the scale and model are summarized in Table 1. The scale with items and

Table 1. Evaluation Criteria for Reliability and Validity of Scales and Fit of Model

Analysis	Parameter	Criterion
Item analysis	Mean	≥ 1.50 and ≤ 4.50
	Variance	> 1.00
	Skewness	> -1.00 and < 1.00
	Corrected item total correlation	$\geq .30$
	Cronbach α after item deleted	Increase
	Upper 27% vs. lower 27% groups	Reach statistical significant difference ($p < .05$)
	An item was deleted if:	
	(1) Any two of the mean, variance, and skewness did not meet the criteria.	
	(2) Any one of the corrected item total correlation, Cronbach's α after item deletion, or difference between the upper and lower 27th percentiles did not meet the criteria.	
EFA	Kaiser-Meyer-Olkin (KMO) measure	$> .80$
	Bartlett's test of sphericity	$p < .05$
	Factor loading	$> .55$
SEM	Average variance explained (AVE)	$> .50$
	Composite reliability (CR)	$> .70$
	Standardized factor loading	$> .55$
	The modification index	≤ 3.84
	χ^2/df	< 3.00
	GFI	$> .90$
	RMSEA	$< .08$
	PNFI	$> .50$
	NFI	$> .90$
	CFI	$> .90$
	IFI	$> .90$
TLI	$> .90$	

CFI=Comparative fit index; GFI=Goodness-of-fit index; IFI=Incremental fit index; NFI=Normed fit index; PNFI=Parsimonious normed fit index; RMSEA=Root mean square error of approximation; TLI=Tucker-Lewis index.

the Kuo model were considered suitable if they met the corresponding criteria. All analyses were performed using IBM SPSS Statistics 21.0 and AMOS 21.0 for Windows (SPSS, Chicago, IL) software.

RESULTS

1. Demographics of the sampled participants

Of the total 310 valid questionnaires, 110 were collected using the initial version of the scale (Appendix 1) and analyzed using item analysis and EFA. The remaining 200 were subsequently collected for scale and model validation with SEM by using the revised and updated version based on the results of item analysis and EFA. The two respondent groups had similar demographic characteristics and did not differ significantly (all $p > .05$), as shown in Table 2. Slightly over half of the respondents were men, and most were aged 50 or older (>70%), well educated (>75% had >9 years of educational experience), married (>80%), and living with

a partner (>89%). In addition, more than half of the entire PDI group had family histories of diabetes.

2. Reliability and exploration of the scale

The items omitted by item analysis due to low discrimination ability and internal consistency based on the criteria in Table 1 are shown in Table 3. An item was deleted if: 1) any two of the mean, variance, and skewness did not meet the criteria; 2) any one of the corrected item total correlation, Cronbach's α after item deletion, or difference between the upper and lower 27th percentiles did not meet the deletion criteria. Consequently, items Q1~Q4 (item numbers in the initial version of the scale, Appendix 1) in the DA subscale and items Q31~Q33 in the RF subscale were omitted. However, the EFA data in Table 3 shows the items retained based on the EFA results and the corresponding criteria in Table 1. The values of the KMO measure (.80~.90) and $p < .001$ of Bartlett's test revealed that the questionnaire data was adequate for EFA. Items Q12, Q25~Q27, and Q34 were omitted, as their factor loading was lower than .55. After implementing EFA, one factor was retained in each of the five SPEE subscales. The corresponding percentages of total variance explained ranged from 50.0% to 65.9%, demonstrating the subscales' acceptable ability to assess variability in the participants' responses after reduction and extraction following EFA [30].

3. Validation of the scale and conceptual model

SEM was conducted to evaluate the validity of the measurement model of the subscales and structure of the Kuo model (Figure 1). The final SEM model with the path coefficients (standardized regression coefficients) presented in Figure 2. As shown in Figure 2, the final version of the five subscales was further reduced to 4, 3, 6, 4, and 4 items for the DA, CHB, DF, RF, and SRE subscales, respectively. Specific items were eliminated based on the modification index of SEM (Table 1). A CR of .73~.94, standardized factor loading of .55~.90, and AVE of 0.57~0.78 demonstrated acceptable convergent validity in each subscale. Only the AVE of .47 for the CHB subscale was slightly lower than .50.

The analysis of the Kuo model included the measure of model fit $\chi^2/df=2.08$, GFI=.85, RMSEA=.07, PNFI=.77, NFI=.88, CFI=.93, IFI=.93, and TLI=.92, which indicated good fit between

Table 2. Participant Demographics

Variables	Item analysis/ EFA (n=110) n (%)	CFA/SEM (n=200) n (%)	ρ^{\dagger}
Gender			.570
Man	59 (53.6)	101 (50.5)	
Woman	51 (46.4)	99 (49.5)	
Age (yr)			.452
<39	6 (5.5)	9 (4.5)	
40~49	24 (21.8)	33 (16.5)	
>50	80 (72.7)	158 (79.0)	
Educational attainment (yr)			.199
≤9	18 (16.4)	45 (22.5)	
>9	92 (83.6)	155 (77.5)	
Marital status			.273
Single	11 (10.0)	13 (6.5)	
Married	90 (81.8)	161 (80.5)	
Divorced or widowed	9 (8.2)	26 (13.0)	
Living arrangement			.350
Living alone	8 (7.3)	21 (10.5)	
With a partner	102 (92.7)	179 (89.5)	
Family diabetic history			.830
Yes	58 (52.7)	108 (54.0)	
No	52 (47.3)	92 (46.0)	

CFA=Confirmatory factor analysis; EFA=Exploratory factor analysis; SEM=Structural equation modeling.

[†]Chi-square test.

Table 3. Item Analysis and Exploratory Factor Analysis

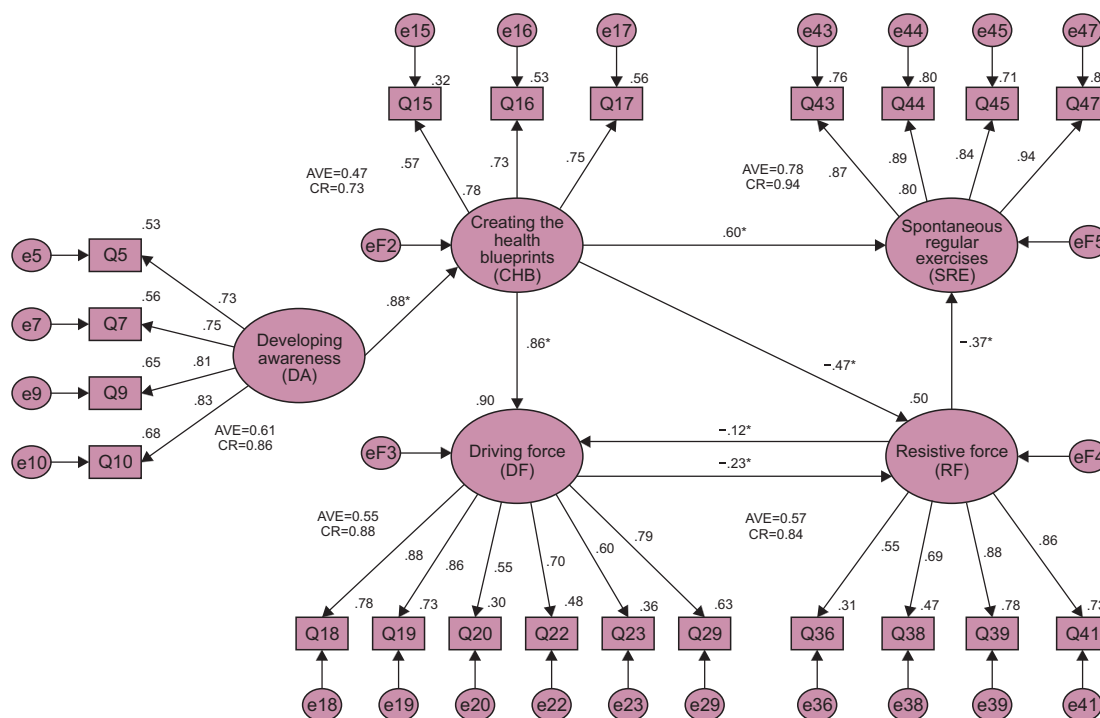
Item analysis								EFA							
Sub-scale	Item omitted [†]	M	V	SK	Overall α	Item del. α	CITC.	Lower 27% vs. upper 27%		KMO	Bartlett's test		% TVE.	Item retained [†]	Factor loading
								t	ρ		χ^2	ρ			
DA	Q1	4.85	.37	-5.13	.81	.82	.22	-2.18	.033	.80	498.58	<.001	50.2	Q5	.80
	Q2	4.84	.36	-5.16		.82	.24	-2.36	.022		Q6	.81			
	Q3	4.86	.34	-5.66		.81	.26	-2.36	.022		Q7	.68			
	Q4	3.49	1.72	-0.48		.84	.10	-2.21	.031		Q8	.69			
													Q9	.74	
													Q10	.80	
													Q11	.65	
CHB	None of items were deleted									.81	269.50	<.001	56.4	Q13	.92
													Q14	.80	
													Q15	.57	
													Q16	.63	
													Q17	.78	
DF	None of items were deleted									.89	914.59	<.001	50.0	Q18	.88
													Q19	.78	
													Q20	.76	
													Q21	.79	
													Q22	.63	
													Q23	.64	
													Q24	.81	
													Q28	.64	
													Q29	.72	
													Q30	.82	
RF	Q31	3.45	1.96	-.31	.88	.90	-.12	.00	>.999	.90	884.89	<.001	65.9	Q35	.73
	Q32	3.19	1.75	-.09		.89	.17	2.82	<.001		Q36	.85			
	Q33	3.15	1.60	-.02		.89	.26	3.53	<.001		Q37	.78			
											Q38	.86			
													Q39	.89	
													Q40	.84	
													Q41	.90	
													Q42	.89	
SRE	None of items were deleted									.88	312.30	<.001	63.7	Q43	.69
													Q44	.77	
													Q45	.81	
													Q46	.87	
													Q47	.83	

M=Mean; V=Variance; SK=Skewness; Overall α =Total score Cronbach α ; Item del. α =Cronbach α after item deleted; CITC=Corrected item total correlation; TVE=Total variance explained; DA=Developing awareness; CHB=Creating the health blueprint; DF=Driving force; RF=Resistive force; SRE=Spontaneous regular exercise.

[†]Item numbers of the items in the initial version of the scale (Appendix 1).

the Kuo model and the sample data (Figure 2). The exceptions were GFI and NFI, which were both slightly lower than .90. The standardized regression coefficients of DA on CHB, CHB on DF, DF on RF, and RF on DF were all statistically significant and supported H1, H2, H3, and H4 of the Kuo model (Figure 1). The

standardized regression coefficients of DA to CHB ($\beta=.88$) and CHB to DF ($\beta=.86$) exhibited strong positive effects, whereas the coefficients of DF to RF ($\beta=-.23$) and RF to DF ($\beta=-.12$) exhibited negative effects. H5 for progression from the ACIS stage to the SRE stage was supported by the significant direct



$\chi^2/df=2.08$, GFI=.85, RMSEA=.07, PNFI=.77, NFI=.88, CFI=.93, IFI=.93, TLI=.92.

SEM=Structural equation modeling; DA=Developing awareness; CHB=Creating the health blueprint; DF=Driving force; RF=Resistive force; SRE=Spontaneous regular exercise; AVE=Average variance explained; CR=Composite reliability; GFI=Goodness-of-fit index; RMSEA=Root mean square error of approximation; PNFI=Parsimonious normed fit index; NFI=Normed fit index; CFI=Comparative fit index; IFI=Incremental fit index; TLI=Tucker-Lewis index.

* $p < .05$.

[†]Item numbers in the figure are the item numbers of the items in the initial version of the scale (Appendix 1).

Figure 2. Structural equation modeling model.

effect of RF on SRE ($\beta = -.37$). This revealed that SRE was directly motivated by a decline in RF. However, the DF effected SRE actions indirectly through the mediation effect of RF ($\beta = -.23$ of DF on RF). The conceptual process outlined by Kuo et al. [18] did not mention the effect of the CHB stage on the SRE stage. However, the results of this study revealed that the CHB stage also directly affected the SRE stage ($\beta = .60$).

DISCUSSION

The development of the SPEE for PDIs provides the medical profession with a convenient instrument to assess the psychological probability that a PDI will implement exercise. The SPEE is a 5-point scale containing 21 items, comprising DA (4 items), CHB (3 items), DF (6 items), RF (4 items), and SRE (4 items) (Appendix 3). This scale is the first exercise-engagement scale de-

signed for PDIs and the first to assess the psychological milieu and behavioral outcome at each stage of the exercise-engagement process. The development process of the SPEE differs from other scales in several key manners: The hypoglycemia problem-solving scale developed by Wu et al. [31] for people with diabetes mellitus was constructed based on the original SPSI and its Chinese version, unlike the SPEE, which relied on the Kuo model foundation. Ebrahimi et al. [32] developed the Diabetes Adjustment Assessment Scale (DAAS) based on their qualitative study but without a conceptual model infrastructure. The DAAS is focused on patients diagnosed with type 2 diabetes and their adjustment to disease onset, not PDIs.

The DA and CHB subscales can be used to explore the levels of PDIs' self-awareness in terms of the importance of physical activity and psychological barriers to exercise. The concept is similar to Lin and Chi's SMS [20], which explores the factors

that motivate an individual to engage in exercise. The main focus is interest in exercise, desire to learn a new sport, improved health, and sense of accomplishment. However, the SMS is designed for general public assessment and not for individuals with specific potential chronic health care needs. The DA subscale of the present study is designed to assess the perspective from the PDIs' point of view. Recognition of health risks is the cause or motivation, and the choice to engage in exercise is the effect, which leads to improved health and a reduction diabetes outcomes. Comparatively, the SMS adopted a broader perspective while covering more dimensions to determine the reasons that individuals engage in exercise.

The DF and RF subscales are similar to the ESES [19] in that both explore the factors that promote or hinder exercise habits among individuals. As with the SMS, the ESES focuses on the general population and not a specific at-risk subgroup. Both scales explore the counteracting effect between positive and negative factors that influence the choice to engage in exercise behavior. Further distinguishing the ESES from the SPEE, the DF and RF subscales are more effective at exploring PDIs' psychological and emotional status during the internal struggle stage. The psychological and emotional status in DF assesses willpower, self-blame, self-motivation, and consistency with regard to exercise. The SPEE is different from the ESES, where the focus explored positive motivations: 1) sense of happiness, 2) improved health, 3) enhanced muscle strength, and 4) bone density. However, the ESES failed to examine motivations to continue exercise over the long term. By comparison, the RF subscale explored: 1) denial, 2) difficulty, 3) sedentary passiveness, and 4) laziness, which are not assessed in the ESES.

The validation of the SPEE is consistent with the Kuo model of the process of exercise engagement in PDIs (Figure 1). The DA stage and its impact on PDIs is of greatest importance and strongly effects the CHB stage. Data showed a direct effect of the DA stage on the CHB stage. This is highly predictive of PDIs' potential to achieve exercise goals that will mediate or eliminate the onset of diabetes, as PDIs may choose to further educate themselves on the dangers of diabetes. The CHB stage motivates the SRE stage not only directly but also indirectly through the DF and RF stages. Furthermore, the effect of the SRE stage on the DF stage is positive, while it is negative on the RF stage.

PDIs are able to empower themselves through education via the hospital or other means, which explains the effects on the DF and RF. The DF and RF stages are negatively and significantly correlated. For PDIs, the more acute the DA stage, the greater their motivation is for engaging in exercise and the smoother their entrance is into the CHB stage. DA stage acuteness does not influence the DF; as such, it may be proposed that during the DA stage, the type of education is extremely important. Once PDIs have crossed the CHB threshold and engaged in exercise, they enter the internal struggle cycle between DF and RF. This influences their exercise behaviors, which enhance or exacerbate their probability for developing diabetes. During the exercise-engagement process, DF and RF counteract each other; however, when a higher DF is present, PDIs more readily enter the SRE stage. In essence, PDIs must first reduce their RF to successfully enter the SRE stage. Therefore, although the RF is a mediating variable in this model, it is critical for determining whether PDIs will develop long-lasting exercise habits.

The findings of this study suggest that whether PDIs initially engage in exercise depends on the impact of the DA stage and, subsequently, whether exercise habits are established. Attaining the SRE stage is dependent to the extent that the RF is reduced, as increasing the DF is not useful. Providing PDIs with physical activity-related health education with the purpose of motivating regular exercise behaviors is crucial. The priority is to increase their awareness of the importance of exercise, followed with solutions and supporting strategies to eliminate factors that prevent regular exercise. In doing so, it is possible to reduce the effect of the RF.

The single difference between the study results and the Kuo model (Figure 1) is the direct and significant ($\beta=.60$) effect of the CHB stage on the SRE stage. As PDIs enter the CHB stage, they may enter the SRE stage directly, bypassing the DF and RF stages. The findings of this study strongly support the Kuo model and are consistent with Self-Determination Theory. This theory states that human behaviors are generated from intrinsic motivation, and the expression of these behaviors is subject to the influence of extrinsic factors [33]. Therefore, when providing relevant clinical health education, medical staff will best serve PDIs by developing awareness, specifically by helping PDIs to plan health blueprints. If PDIs perceive a positive outcome for behav-

ioral change, the exercise intervention may be more effective.

Several factors may influence the transferability of the SPEE and the findings of this study. Due to time constraints, this study could not assess the long-term benefits of exercise and the predictive capacity of the SPEE. As a result of limited research budget and time, the research was confined to one geographic location. Another confounding factor is that the level of participants' education was consistently high, as the government of Taiwan has an enforced state education policy that has been in existence for the last 50 years. It is known that level of education influences behavioral modification.

In order to enhance the transferability of the SPEE, further studies should focus on: 1) psychological status of exercise for PDIs, 2) behavioral expression at various stages of the exercise-engagement process, and 3) contributing factors to the formation of long-term exercise habits. This research would have benefited greatly with the inclusion of a quantitative approach to assess changes in glucose levels as participants became engaged in exercise to reduce the long-term liability of developing diabetes.

Although the sample size of 110 in EFA of the current study met the minimum requirement suggested by Gorsuch [23] and Thompson [24]. However, the sample size of 235 based on the rule of five subjects for each item is the more preferable guideline for scale development [22]. Lack of enough power due to less sample size might be a limitation of the current study. Therefore, a further study with larger sample is suggested to conduct for providing stronger evidence to validate the developed scale and model.

CONCLUSION

This study is the first to develop a scale that includes the five subscales of DA, CHB, DF, RF, and SRE for assessing the psychological transition and behavioral expression at each stage of the process of exercise engagement for PDIs. The Kuo model for the process of exercise engagement of PDIs is valid and has a goodness of fit. Higher scores on the DA, CHB, DF, and SRE subscales represents higher levels for psychological and behavioral transitions, whereas a higher score on the RF subscale represents lower psychological and behavioral transitions. The SPEE

for people with prediabetes provides deeper insights into the behavioral change stages required to initiate extended long-term health care outcomes in order to avoid developing diabetes. These insights are significant as they allow for patient-specific mapping of their DF and RF and the factors affecting their transition to the SRE stage. The use of SPEE within the clinical setting may prove useful in developing tailored exercise strategies that may lead to a reduction in the rate of diabetes onset in PDIs.

CONFLICTS OF INTEREST

The authors declared no conflict of interest.

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REFERENCES

1. International Diabetes Federation (IDF). IDF diabetes atlas (seventh edition) [Internet]. Brussels: IDF; c2015 [cited 2019 Jan 13]. Available from: <https://diabetesatlas.org/upload/resources/previous/files/7/IDF%20Diabetes%20Atlas%207th.pdf>.
2. Li G, Zhang P, Wang J, Gregg EW, Yang W, Gong Q, et al. The long-term effect of lifestyle interventions to prevent diabetes in the China Da Qing diabetes prevention study: A 20-year follow-up study. *The Lancet*. 2008;371(9626):1783–1789. [https://doi.org/10.1016/S0140-6736\(08\)60766-7](https://doi.org/10.1016/S0140-6736(08)60766-7)
3. Chen KT, Chen CJ, Gregg EW, Imperatore G, Narayan KMV.

- Impaired fasting glucose and risk of diabetes in Taiwan: Follow-up over 3 years. *Diabetes Research and Clinical Practice*. 2003;60(3):177-182.
[https://doi.org/10.1016/s0168-8227\(03\)00037-8](https://doi.org/10.1016/s0168-8227(03)00037-8)
4. Tabák AG, Herder C, Rathmann W, Brunner EJ, Kivimäki M. Prediabetes: A high-risk state for diabetes development. *Lancet*. 2012;379(9833):2279-2290.
[https://doi.org/10.1016/S0140-6736\(12\)60283-9](https://doi.org/10.1016/S0140-6736(12)60283-9)
 5. Shaye K, Amir T, Shlomo S, Yechezkel S. Fasting glucose levels within the high normal range predict cardiovascular outcome. *American Heart Journal*. 2012;164(1):111-116.
<https://doi.org/10.1016/j.ahj.2012.03.023>
 6. Plantinga LC, Crews DC, Coresh J, Miller ER 3rd, Saran R, Yee J, et al. Prevalence of chronic kidney disease in US adults with undiagnosed diabetes or prediabetes. *Clinical Journal of American Society of Nephrology*. 2010;5(4):673-682.
<https://doi.org/10.2215/CJN.07891109>
 7. Diabetes Prevention Program Research Group. The prevalence of retinopathy in impaired glucose tolerance and recent-onset diabetes in the diabetes prevention program. *Diabetic Medicine*. 2007;24(2):137-144.
<https://doi.org/10.1111/j.1464-5491.2007.02043.x>
 8. Lindström J, Ilanne-Parikka P, Peltonen M, Aunola S, Eriksson JG, Hemiö K, et al. Sustained reduction in the incidence of type 2 diabetes by lifestyle intervention: Follow-up of the Finnish diabetes prevention study. *The Lancet*. 2006;368(9548):1673-1679.
[https://doi.org/10.1016/S0140-6736\(06\)69701-8](https://doi.org/10.1016/S0140-6736(06)69701-8)
 9. Lindström J, Louheranta A, Mannelin M, Rastas M, Salminen V, Eriksson J, et al. The Finnish diabetes prevention study (DPS): Lifestyle intervention and 3-year results on diet and physical activity. *Diabetes Care*. 2003;26(12):3230-3236.
<https://doi.org/10.2337/diacare.26.12.3230>
 10. The Diabetes Prevention Program (DPP) Research Group. The diabetes prevention program (DPP): Description of lifestyle intervention. *Diabetes Care*. 2002;25(12):2165-2171.
<https://doi.org/10.2337/diacare.25.12.2165>
 11. Pedley CF, Case LD, Blackwell CS, Katula JA, Vitolins MZ. The 24-month metabolic benefits of the healthy living partnerships to prevent diabetes: A community-based translational study. *Diabetes & Metabolic Syndrome*. 2018;12(3):215-220.
<https://doi.org/10.1016/j.dsx.2017.09.011>
 12. Pan XR, Li GW, Hu YH, Wang JX, Yang WY, An ZX, et al. Effects of diet and exercise in preventing NIDDM in people with impaired glucose tolerance: The Da Qing IGT and diabetes study. *Diabetes Care*. 1997;20(4):537-544.
<https://doi.org/10.2337/diacare.20.4.537>
 13. Boniol M, Dragomir M, Autier P, Boyle P. Physical activity and change in fasting glucose and HbA1c: A quantitative meta-analysis of randomized trials. *Acta Diabetologica*. 2017;54(11):983-991.
<https://doi.org/10.1007/s00592-017-1037-3>
 14. Sato Y, Nagasaki M, Kubota M, Uno T, Nakai N. Clinical aspects of physical exercise for diabetes/metabolic syndrome. *Diabetes Research and Clinical Practice*. 2007;77(3 Suppl):S87-S91. <https://doi.org/10.1016/j.diabres.2007.01.039>
 15. Bai YL, Wang LM, Hung SY, Chiou CP. [Exercise benefits for health of elderly diabetics]. *Journal of Taiwan Nephrology Nurses Association*. 2012;11(3):1-7. Chinese.
 16. Kuo YL, Chang SC, Chang M, Wang YW, Yeh SC. [The effects of multi-approach health education on people with pre-diabetes]. *Journal of Evidence-Based Nursing*. 2008;4(4):297-306. Chinese. <https://doi.org/10.6225/JEBN.4.4.297>
 17. Chang SC, Hayter M, Yeh HC, Hsieh TC, Kuo YL. The effectiveness of different health education strategies in people with pre-diabetes: A randomized controlled trial. *Journal of Health Science*. 2016;6(2):22-29.
<https://doi.org/10.5923/j.health.20160602.02>
 18. Kuo YL, Wu SC, Hayter M, Hsu WL, Chang M, Huang SF, et al. Exercise engagement in people with prediabetes—a qualitative study. *Journal of Clinical Nursing*. 2014;23(13-14):1916-1926. <https://doi.org/10.1111/jocn.12424>
 19. McAuley E. Self-efficacy and the maintenance of exercise participation in older adults. *Journal of Behavioral Medicine*. 1993;16(1):103-113. <https://doi.org/10.1007/bf00844757>
 20. Lin CY, Chi LK. [Development of the sport motivation scale: Analysis of reliability and validity]. *Bulletin of Sport and Exercise Psychology of Taiwan*. 2003;(2):15-32. Chinese.
 21. Prochaska JO, Velicer WF. The transtheoretical model of health behavior change. *American Journal of Health Promotion*. 1997;12(1):38-48.
 22. Stevens J. *Applied multivariate statistics for the social sciences*. 4th ed. Mahwah (NJ): Lawrence Erlbaum Associates; 2002. p. 203-204.
 23. Gorsuch RL. *Factor analysis*. 2nd ed. Hillsdale (NJ): Lawrence Erlbaum Associates; 1983. p. 109-110.
 24. Thompson B. *Exploratory and confirmatory factor analysis: Understanding concepts and applications*. Washington, D.C.: American Psychological Association; 2004. p. 273-275.
 25. Davis LL. Instrument review: Getting the most from a panel of experts. *Applied Nursing Research*. 1992;5(4):194-197. [https://doi.org/10.1016/S0897-1897\(05\)80008-4](https://doi.org/10.1016/S0897-1897(05)80008-4)
 26. Ferketich S. Focus on psychometrics. Aspects of item analysis. *Research in Nursing & Health*. 1991;14(2):165-168. <https://doi.org/10.1002/nur.4770140211>
 27. Lautenschlager GJ, Lance CE, Flaherty VL. Parallel analysis criteria: Revised equations for estimating the latent roots of random data correlation matrices. *Educational and Psychological Measurement*. 1989;49(2):339-345.
<https://doi.org/10.1177/0013164489492006>

28. Hair JF, Black WC, Babin BJ, Anderson RE. *Multivariate data analysis*. Harlow: Pearson Education Limited; 2014. p. 115.
29. Hooper D, Coughlan J, Mullen M. Structural equation modeling: Guidelines for determining model fit. *Electronic Journal of Business Research Methods*. 2008;6(1):53–60.
<https://doi.org/10.21427/D7CF7R>
30. Lattin JM, Carroll JD, Green PE. *Analyzing multivariate data*. Pacific Grove (CA): Thomson Brooks/Cole; 2003. p. 351–352.
31. Wu FL, Juang JH, Lin CH. Development and validation of the hypoglycaemia problem-solving scale for people with diabetes mellitus. *Journal of International Medical Research*. 2016;44(3):592–604.
<https://doi.org/10.1177/0300060516636752>
32. Ebrahimi H, Karimi Moonaghi H, Asghari Jafarabadi M, Namdar Areshtanab H, Jouybari L. Development and preliminary validation of diabetes adjustment assessment scale (DAAS): A new measure of adjustment with type 2 diabetes. *Journal of Caring Sciences*. 2016;5(2):145–152.
<https://doi.org/10.15171/jcs.2016.015>
33. Deci EL, Ryan RM. The “what” and “why” of goal pursuits: Human needs and the self-determination of behavior. *Psychological Inquiry*. 2000;11(4):227–268.
https://doi.org/10.1207/S15327965PLI1104_01

Appendix 1. Initial Version of the Scale for Process of Exercise Engagement (SPEE)

Part 1: Developing awareness (DA)

- 1 In my opinion, we are each responsible for our physical health.
- 2 In my opinion, exercise is of great importance to my health.
- 3 In my opinion, exercise is beneficial to me.
(e.g., it can promote normal blood glucose levels, weight loss, improve physical agility and promote sleep.)
- 4 In my opinion, I belong to the diabetes high-risk group.
- 5 When I detect changes in my physical condition (e.g., back pain, aging, weight gain, depleted physical fitness), I am more motivated to exercise.
- 6 When I realize that I am symptomatic of a high risk disease (i.e. rise in blood glucose or blood pressure, steatohepatitis, heart disease, etc.), I am motivated to exercise.
- 7 I remind myself to exercise because I worry about or am afraid of developing diabetes.
- 8 When I see my family members or friends suffering from diabetes, I remind myself to be responsible and exercise.
- 9 If I have fitness equipment at home or a convenient venue for physical activity, I am motivated to exercise.
- 10 When I am reminded that according to the advocacy of health education lectures, media, newspapers, magazines and health professionals, that regular sports can prevent diabetes, I am motivated to exercise.
- 11 If I see someone I admire (e.g., a colleague or friend) acquiring exercise habits, I am motivated to exercise.
- 12 When exercise is directly rewarded or a monetary bonus accompanies it, I am motivated to exercise.

Part 2: Creating the health blueprint (CHB)

- 13 I exercise for the purpose of physical health.
- 14 I exercise so that my children will not worry about my health and I do not want to place a burden on my family members.
- 15 I exercise to stay healthy and maintain a positive image (e.g., my figure or level of attractiveness).
- 16 When I feel that I need to engage in exercise, I first evaluate the feasibility of exercise (e.g., When is the best time to exercise? Which type of exercise suits me best?).
- 17 When I feel that I should engage in exercise, I try to incorporate exercise into my daily routine (e.g., I perform exercise casually when I like, arrange time for exercise, or perform exercise according to my work schedule).

Part 3: Driving force (DF)

- 18 I attempt to exercise every day.
- 19 Even if I feel lazy, through willpower I will force myself to exercise.
- 20 I am annoyed if I feel I should exercise but fail to do so.
- 21 I discipline myself to exercise at a fixed time.
- 22 I have methods of motivating myself to exercise (e.g., establishing goals, joining group activities or classes, posting reminders or signs, writing an exercise journal, scheduling time for exercise).
- 23 If I miss my workout, I perform other forms of exercise to increase my physical activity level (e.g., I do simple exercise routines, stretch at home, or perform arm swings during my break at work).
- 24 After I exercise, I feel calm and empowered about my health.
- 25 My family members or friends urge me to exercise.
- 26 My family members or friends invite me to exercise together with them.
- 27 I push myself to exercise by means of team sports, association or courses (Such as ball team, country dance, shadowboxing, aerobics class, sport competition held by workplace or other units, etc.).
- 28 Prior to my next physical examination (i.e. blood glucose or physical fitness testing), I do my best to exercise regularly.
- 29 If I feel my physical condition has improved (i.e., smoother bowel movement, lower blood glucose, alleviated back pain, able to walk up and down stairs more easily, alleviated allergies) because of exercise, I push myself to exercise consistently.
- 30 When I feel that exercise can bring me a sense of achievement, I push myself to exercise continuously.

Appendix 1. Continued

Part 4: Resistive force (RF)

- 31 When I feel uncomfortable or have restricted movement function (heart trouble, catch a cold, feel pain, having no physical strength, knee pain or foot injury), I stop exercising.
- 32 When it is bad weather (i.e. rainy days, too cold or hot weather), I stop exercising.
- 33 When I am very busy in life (i.e. heavy work, study pressure, traveling, etc.), I stop exercising.
- 34 When I have no field nor space for exercise, I stop exercising.
- 35 In spite of not exercising, I do not fall ill.
- 36 In my opinion, exercise is neither necessary nor crucial.
- 37 Exercise makes me feel stressful and I cannot relax.
- 38 Exercise is physically demanding and makes me feel exhausted.
- 39 I require external encouragement to exercise (e.g., I do not exercise without supervision).
- 40 When I have reached my goal (such as recovery of blood glucose or lose of weight), I stop exercising.
- 41 I am too lazy to exercise.
- 42 I have no interest in exercise.

Part 5: Spontaneous regular exercise (SRE)

- 43 Exercise is indispensable in my everyday life.
- 44 Routine exercise helps me to relax.
- 45 I like to exercise consistently and I do it for no specific purpose.
- 46 Regular exercise helps me feel happy and joyful.
- 47 I do not need anyone to accompany me or supervise me to exercise consistently.

CHB=creating the health blueprint; DA=developing awareness; DF=driving force; RF=resistive force; SPEE=scale for process of exercise engagement; SRE=spontaneous regular exercise.

Appendix 2. Number of Items of the SPEE Scale at Each Stage of Analysis

Subscale	Initial version	After item analysis	After EFA	Final version (after CFA)
DA	12	8	7	4
CHB	5	5	5	3
DF	13	13	10	6
RF	12	9	8	4
SRE	5	5	5	4
Total	47	40	35	21

DA=Developing awareness; CHB=Creating the health blueprint; DF=Driving force; RF=Resistive force; SRE=Spontaneous regular exercise; EFA=Exploratory factor analysis; CFA=Confirmatory factor analysis.

Appendix 3. Final Version of Scale for Process of Exercise Engagement (SPEE)

Scale/item [†]	Option
Part 1. Developing awareness (DA)	
• (Q5) When I detect changes in my physical condition (e.g., back pain, aging, weight gain, depleted physical fitness), I am more motivated to exercise.	5 - Almost always 4 - Often
• (Q7) I remind myself to exercise because I worry about or am afraid of developing diabetes.	3 - Sometimes
• (Q9) If I have fitness equipment at home or a convenient venue for physical activity, I am motivated to exercise.	2 - Seldom
• (Q10) When I am reminded that according to the advocacy of health education lectures, media, newspapers, magazines and health professionals, that regular sports can prevent diabetes, I am motivated to exercise.	1 - Never
Part 2. Creating the health blueprint (CHB)	
• (Q15) I exercise to stay healthy and maintain a positive image (e.g., my figure or level of attractiveness).	5 - Almost always
• (Q16) When I feel that I need to engage in exercise, I first evaluate the feasibility of exercise (e.g., when is the best time to exercise? which type of exercise suits me best?).	4 - Often 3 - Sometimes
• (Q17) When I feel that I should engage in exercise, I try to incorporate exercise into my daily routine (e.g., I perform exercise casually when I like, arrange time for exercise, or perform exercise according to my work schedule).	2 - Seldom 1 - Never
Part 3. Driving force (DF)	
• (Q18) I attempt to exercise every day.	5 - Almost always
• (Q19) Even if I feel lazy, through willpower I will force myself to exercise.	4 - Often
• (Q20) I am annoyed if I feel I should exercise but fail to do so.	3 - Sometimes
• (Q22) I have methods of motivating myself to exercise (e.g., establishing goals, joining group activities or classes, posting reminders or signs, writing an exercise journal, scheduling time for exercise).	2 - Seldom 1 - Never
• (Q23) If I miss my workout, I perform other forms of exercise to increase my physical activity level (e.g., I do simple exercise routines, stretch at home, or perform arm swings during my break at work).	
• (Q29) If I feel my physical condition has improved (i.e., smoother bowel movement, lower blood glucose, alleviated back pain, able to walk up and down stairs more easily, alleviated allergies) because of exercise, I push myself to exercise consistently.	
Part 4. Resistive force (RF)	
• (Q36) In my opinion, exercise is neither necessary nor crucial.	5 - Strongly Agree
• (Q38) Exercise is physically demanding and makes me feel exhausted.	4 - Agree
• (Q39) I require external encouragement to exercise (e.g., I do not exercise without supervision).	3 - Undecided
• (Q41) I am too lazy to exercise.	2 - Disagree Strongly 1 - Disagree
Part 5. Spontaneous regular exercise (SRE)	
• (Q43) Exercise is indispensable in my everyday life.	5 - Strongly Agree
• (Q44) Routine exercise helps me to relax.	4 - Agree
• (Q45) I like to exercise consistently and I do it for no specific purpose.	3 - Undecided
• (Q47) I do not need anyone to accompany me or supervise me to exercise consistently.	2 - Disagree Strongly 1 - Disagree

CHB=creating the health blueprint; DA=developing awareness; DF=driving force; RF=resistive force; SPEE=scale for process of exercise engagement; SRE=spontaneous regular exercise.

[†]Item numbers in the parentheses are the item numbers of the items in the initial version of the scale (Appendix 1).