# Translation and Validation of the Malay Version of Modified Vaccine Hesitancy Scale (MVHS-M) for Assessment of Parental Vaccine Hesitancy

Mohd Zin ND<sup>a</sup>, Wan Mohammad WMZ<sup>a</sup>, Kueh YC<sup>b</sup>, Mohd Fuzi NMH<sup>c</sup>

<sup>a</sup>Department of Community Medicine, School of Medical Sciences, Health Campus, Universiti Sains Malaysia,Kubang Kerian, Kelantan, Malaysia <sup>b</sup>Unit of Biostatistics and Research Methodology, School of Medical Sciences, Health Campus, Universiti Sains Malaysia, Kubang Kerian, Kelantan, Malaysia <sup>c</sup>Surveillance Unit, CPRC Infectious Disease Section, Kelantan State Health Department, Ministry of Health Malaysia, Kota Bharu, Kelantan, Malaysia

#### ABSTRACT

INTRODUCTION: Parental hesitancy towards routine childhood vaccines has been recognized as one of the public health threats. Since the uptake of child vaccination remains inconsistent, there is a need for a reliable and validated tool to measure this phenomenon. MATERIALS AND METHODS: A cross-sectional study was conducted at government health clinics in Kelantan between April 2023 to July 2023. A permission to use the original version of Modified Vaccine Hesitancy Scale (MVHS) was obtained and translated into the Malay version (MVHS-M) based on established guidelines. Parents who have at least one child aged 7 years or less were recruited by using systematic random sampling to validate the MVHS-M. A confirmatory factor analysis (CFA) was used to confirm the latent domain, while reliability was measured by composite reliability and testretest. The data were analysed using IBM SPSS Version 26 and Mplus version 8. RESULTS: A total of 270 parents who fulfilled the study criteria were selected and completed the survey. The CFA showed a good fit index: RMSEA=0.057 (90% CI 0.031, 0.082), CFI=0.970, TLI=0.957, and SRMR=0.031. The composite reliability for the domain "lack of confidence" was 0.93 (95% CI 0.91, 0.94), while the domain "risk" showed a reliability of 0.74 (95% CI 0.69, 0.79). The test-retest reliability, as measured by the Intra-class Correlation Coefficient (ICC), was 0.77 (95% CI 0.59, 0.87), indicating good stability. CONCLUSION: MVHS-M is a valid and reliable tool that will be useful in identifying parental vaccine hesitancy in Malaysia.

Keywords Validity, Reliability, Vaccine hesitancy, Child vaccination

Corresponding Author Assoc. Prof. Dr Wan Mohd Zahiruddin Wan Mohammad Department of Community Medicine, School of Medical Sciences, Health Campus, Universiti Sains Malaysia, 16150 Kubang Kerian, Kelantan, Malaysia

E-mail : drzahir@usm.my

Received: 4th December 2023; Accepted: 13th Feb 2024

Doi: https://doi.org/10.31436/imjm.v23i02

#### INTRODUCTION

Vaccines saved millions of children's lives every year.<sup>1</sup> The eradication of smallpox and near-elimination of polio are well-known success stories resulting from the worldwide implementation of immunization programs. In Malaysia, since the introduction of the National Immunization Program (NIP) in the 1950s, the program has contributed to a substantial reduction in child mortality due to infectious diseases.<sup>2,3</sup> Despite these successes, there is growing concern about issues related to possible side effects of vaccines, the feeling of unnecessary vaccination since the disease no longer exists, and the emergence of vaccine misinformation, contributing to a phenomenon called vaccine hesitancy.<sup>4–9</sup> This phenomenon had a detrimental effect on vaccination uptake, weakening herd immunity and

causing a resurgence of vaccine-preventable diseases.<sup>10–14</sup>

Vaccine hesitancy refers to uncertainty or indecision towards vaccination. Overall, hesitant individuals are placed on a continuum between complete acceptance and complete refusal of vaccines, despite the availability of immunization services.<sup>15,16</sup> Therefore, it is crucial to identify vaccine hesitant parents before they reach a point of complete refusal, as their viewpoint can still be influence by positive interventions. While extensive research on vaccine hesitancy has been widely studied in Western countries,<sup>17–19</sup> there is limited local evidence on this matter. Hence, a valid measurement tool is deemed necessary to understand this issue in the local context. Based on previous reviews, vaccine hesitancy has been measured using various tools, including heterogeneous assessment questions that vary in cut-off point, the Vaccine Hesitancy Scale (VHS), and the Parent Attitude about Childhood Vaccines (PACV).<sup>20,21</sup> Aiming to standardise the assessment of vaccine hesitancy, the World Health Organization (WHO) has recommended the utilisation of VHS and emphasized the importance of validating it to ensure its applicability across various contexts.<sup>15</sup> The VHS has been validated in numerous countries<sup>18,22,23</sup> and has been translated into Spanish<sup>19</sup>, Arabic,<sup>24,25</sup> Turkish,<sup>26</sup> Korean,<sup>27</sup> and Chinese.<sup>28</sup> Recently, a modified version of VHS has been made available in English and Spanish.<sup>29</sup>

Given the recommendation of WHO for the utilisation of VHS, adapting this questionnaire at a local level would provide a more accurate basis for future analysis. Thus, the Modified Vaccine Hesitancy Scale (MVHS) was selected as a tool to identify vaccine hesitancy among parents in Malaysia. To the best of our knowledge, this MVHS has not been translated or validated in the Malay language. Therefore, our study aimed to translate the MVHS into Malay and subsequently validate the Malay version.

## MATERIALS AND METHOD

#### Study Design and Setting

This was a cross-sectional study conducted over four months starting from April to July 2023. This study took place at government health clinics in ten districts of Kelantan. It consisted of two phases in which were Translation and Cultural Adaptation phase, and Validation and Reliability phase.

#### **Translation and Cultural Adaptation Phase**

A permission to translate the questionnaire into the Malay language from the original author was obtained beforehand. The process of translation and cultural adaptation process followed the established guideline.<sup>30</sup> Forward translation were performed by two independent translators; a medical officer from the Maternal and Child Health Unit at the Ministry of Health (MOH) facility as translator one, and a teacher from the Language,

Literacies and Translation Unit of Universiti Sains Malaysia (USM) as translator two. Both translators were fluent in Malay and English language. Any difficulty or confusing items, words, or sentences were highlighted. Then, a common translation was synthesised from both forward translations during the reconciliation process. In order to ensure the relevancy of the item, content validation was performed by a panel of experts consisting of three public health physicians, two family medicine specialists and one paediatrician. The experts agreed that all items were relevant to Malaysian parents. The reconciled version of the questionnaire was then proceeded for backward translated into English by another two independent translators. It was subsequently harmonized to improve any undue discrepancies. Overall, the translated version demonstrated satisfactory equivalences, and the reconciled version was used as prefinal MVHS-M during face validation.

A face validation or also known as cognitive debriefing was conducted using pre-final MVHS-M with ten respondents. The respondents were selected among the parents through purposive sampling during their visit to Gunong Health Clinics in Bachok district. Their selection criteria were based on the study criteria mentioned below. They were instructed to evaluate the understandability, interpretation, and cultural appropriateness of the translated questionnaire. Necessary amendments were made accordingly. Following this, pilot testing was carried out to identify any further shortcomings in the translated questionnaire and to refine the procedures related to its administration and data preparation for analysis. A total of 30 parents who fulfilled the same inclusion criteria and different respondents were ensured. The time required to answer all the questions ranged 15 to 20 minutes, which was considered acceptable.

## Validation and Reliability Phase

The validation process was conducted with parents who met the same study criteria. A total of 270 parents were recruited from government health clinics in districts other than Bachok. A total of 62 respondents were invited to assess for stability by participating in a retest at seven-day intervals following the initial administration of the questionnaire

#### Sample Size

20% non-response rate,33 the required sample size was Size Calculator Version 2.0.34

#### Sampling Method and Subject Recruitment

Respondents were recruited through a multistage random sampling method from the remaining nine districts. Initially, using simple random sampling, Kota Bharu, Pasir Mas and Pasir Puteh districts were selected. Then, nine government health clinics from the above districts were selected using the same sampling method. Despite the calculated sample size being 250, a total of 270 respondents were recruited after evenly distributing the number of participants for each clinic and rounding the figures. As a result, 30 respondents were recruited at each clinic.

Parents who aged over 18 years, had at least a child who aged  $\leq 7$  years, and provided their consent were included in this study. However, non-Malaysian parents, parents with immunocompromised children, parents who attended the clinics solely for their child immunization and parents with cognitive impairment and intellectual disability were excluded. The sampling frame involved all parents who met the study criteria and attended government health clinics during study period. All participants were provided with a patient information sheet and required to provide written consent before participating. Using systematic random sampling, parents were selected at regular two-unit intervals at the registration counter of the clinics. They were instructed to fill out self-administered 9-item questionnaire of MVHS-M and sociodemographic proforma while waiting for their version 26, followed by descriptive statistics to describe turn in the clinic. The participants were given 15 to 20 the numerical and categorical variables. minutes to complete their answers, after which researcher consistency reliability analyses using Cronbach's alpha and collected the questionnaire and checked for completeness. For test-retest, only participants who had agreed during conducted using the same software. The data was then the initial session were contacted after seven days, and transferred to Mplus version 8 for Confirmatory Factor they were given a similar time frame to complete the Analysis (CFA) and estimation of composite reliability answers again.

#### **Research Tool**

The minimum recommended sample size for the The original Vaccine Hesitancy Scale (VHS) was a selfvalidation phase was 200 samples.31,32 By considering the administered questionnaire, developed by the SAGE Working Group on Vaccine Hesitancy. The development determined to be 250. For test-retest, a sample size involved extensive literature reviews and discussion required of 62 respondents was calculated using Sample among experts.<sup>15</sup> Following the development, the psychometric properties were not initially assessed, and were evaluated by Shapiro et al.23 They identified two subdomains; lack of confidence and risk were identified. However, prior validation studies showed inadequate psychometric properties, leading to development of a modified version of VHS.18,23,29 The Modified Vaccine Hesitancy Scale (MVHS) is a modified version that can be used to predict and identify vaccine hesitancy related to childhood vaccination.29 After some modification, the validated questionnaire was reduced from 10 items to nine items and maintain organized within two subdomains; lack of confidence and risk. The MVHS demonstrated good construct validity (RMSEA=0.09, CFI=0.96, TLI=0.94 and SRMR=0.04) and displayed good internal reliability with Cronbach's alpha coefficients of 0.90 and 0.76, respectively.

> In MVHS, instead of using a 5-point Likert scale as in the original VHS, each statement is measured using a 4-point Likert scale. The response options are "Strongly agree", "Agree", "Disagree" and "Strongly Disagree". The "Neutral" option, scored as 3 in the original VHS was excluded in the MVHS to reduce the potential effect of social desirability bias.35 Numeric scores are still maintained as 1, 2, 4, or 5. Higher scores on MVHS indicate more hesitancy. The respondents were asked to choose the best response for each statement.

#### **Statistical Analysis**

Data entry was performed using IBM SPSS software Internal the Intra-class Correlation Coefficient (ICC) were also Raykov's rho.36 The sociodemographic using

characteristics of the respondents were presented as descriptive statistics.

CFA was performed using a robust maximum likelihood estimator due to the lack of multivariate normality.37 Several fit indices were used to evaluate the model fitness including root mean square error of approximation (RMSEA;  $\leq 0.08$ ), Comparative Fit Index (CFI;  $\geq 0.95$ ), Tucker-Lewis Index (TLI;  $\geq 0.95$ ), and standardized root mean square residual (SRMR  $\leq 0.08$ ), based on established guideline.<sup>31</sup> Items with standardised factor loading of 0.5 and above was as cut-off point in this study.<sup>31</sup>. Besides, the factor loading must be significant at  $\alpha$  level <0.05. The initial model was evaluated using these fit indices. Then, the model specification was further considered by assessing the acceptability of the model, modification indices (MIs) and correlated item's residual based on theoretical justification. To assess reliability, Raykov's rho coefficient and Cronbach's alpha were used to measure internal consistency, with a predetermined cut-off value of  $\geq 0.7.^{31}$  In addition, test-retest reliability was evaluated using the ICC, with values ranging from 0.75 to 0.95 considered indicative of good reliability.38

#### RESULTS

#### Sociodemographic Characteristics of Respondents.

The mean (SD) age of the respondents was 32.3 (SD 6.02). The majority were mothers (81.9%) and were not pregnant (51.6%) at the time of data collection. All participants were Malay, Muslim and had at least one child. Details of sociodemographic characteristics of respondents were displayed in Table I.

# **Confirmatory Factor Analysis of MVHS-M**

The MVHS-M consists of nine items with two subdomains (lack of confidence and risk). In the initial measurement model as in Table II, most of the fit indices were acceptable, except for the TLI (RMSEA=0.07, SRMR=0.033, CFI=0.953, TLI=0.935). Since all items had factor loadings higher than 0.4 (Table III), no items were removed from the model. The initial model was further respecified to improve fit indices. Researcher added a correlation between the residual for item C5 with C6

Variables	N(%)	Mean (SD)
Age of parents (years)		32.29 (6.02)
Ethnicity		
Malay	270 (100.0)	
Age of youngest child (months)		24.00 (39.00) <sup>a</sup>
Relationship to child		
Mother	221 (81.9)	
Father	49 (18.1)	
Number of children		
One	98 (36.3)	
Two	68 (25.2)	
Three	46 (17.0)	
Four and more	58 (21.5)	
Highest formal education		
No formal education	3 (1.1)	
Primary & Secondary school	117 (43.3)	
Certificate/Diploma/STPM	100 (37.0)	
Degree	46 (17.0)	
Postgraduate	4 (1.5)	
Employment Status		
Unemployed	153 (56.7)	
Employed	117 (43.3)	
Household Income		
<rm3030 (b40)<="" td=""><td>225 (83.3)</td><td></td></rm3030>	225 (83.3)	
RM3030 to RM6619 (M40)	38 (14.1)	
≥RM6620(T20)	7 (2.6)	
Current pregnancy status*		
Pregnant	107 (48.4)	
Non-pregnant	114 (51.6)	
Information sources**		
Healthcare providers		
Yes	180 (66.7)	
No	90 (33.3)	
Internet and social media		
Yes	127 (47.0)	
No	143 (53.0)	
Family and friends		
Yes	45 (16.7)	
No	225 (83.3)	

<sup>a</sup>Median(IQR) \*Applicable for female respondents

\*\*Multiple responses

(Model-2) based on adequate theoretical support. Figure 1 shows a path diagram of MVHS-M which indicates present of correlated residual between C5 and C6. Model-2 demonstrated good fits with all fit indices falling within the recommended values.

Composite Reliability (CR) for Model-2 was calculated using Raykov's method<sup>36</sup>. The cut-off point considered in this study was >0.7, which is considered acceptable<sup>31</sup>. The Average Variance Extracted (AVE) values were also provided in Table III. Internal consistency based on Cronbach's alpha ranged from satisfactory to good, with Table III: Standardised item's loading, composite reliability and average values ranging from 0.75 to 0.94. For stability testing,

Table II: Fit indices for the measurement model of MVHS-M (initial and final model)

Model	CFI	TLI	SRMR	RMSEA (90% CI)	CIfit
Model-1 (Initial)	0.953	0.935	0.033	0.070 (0.050, 0.090)	0.071
Model-2	0.970	0.957	0.031	0.057 (0.031, 0.082)	0.291

Notes: CFI=Comparative fit index; TLI=Tucker-Lewis index; SRMR= standardised root mean square residual; RMSEA=root mean square error of approximation; CI=confidence interval

out of 62 distributed questionnaires, 51 respondents completed the questionnaire again on day seven after the initial administration (82.3% response rate). The Intraclass Correlation Coefficient (ICC) based on a two-way mixed effects model with the consistency option and average measures was 0.77 (95% CI 0.59, 0.87), which considered as good reliability over time.38

#### DISCUSSION

Vaccine hesitancy is a global public health concern. Therefore, there is a need for validated scales with international comparability. Additionally, the experts recommended the validation of the VHS in different contexts15 and a modified version of VHS was which produced better psychometric developed, properties.<sup>29</sup> This study aims to translate and validate the MVHS to assess vaccine hesitancy in the Malaysia population, specifically parents with children  $\leq 7$  years old. Hence, the current study contributes as the first report on the validation of MVHS in Malaysia. The findings revealed that the scale provided valid and reliable results for assessing vaccine hesitancy among parents with children  $\leq$ 7 years old in government health clinics.

Based on Confirmatory Factor Analysis (CFA) (Table II), Model-2 exhibit a better fit compared to Model-1, leading to the selection of Model-2 as the best final model. Based on Hair et al<sup>31</sup> as expected when N>250 and observed variables <12, the cut-off values for CFI and TLI should be  $\geq 0.95$  and RMSEA < 0.07, but SRMR may be biased. The fit indices in Model-2 were within the recommended values, indicating good fit for internal validity. Moreover, all fit indices in Model-2 showed better fit compared to the original modified version.29 In this study, Chivariance extracted for the Model-2 of MVHS-M

Domains/Items	Standardised factor loading	CR (95% CI)	AVE
F1: Lack of Confidence			
Childhood vaccines are important for my child's health	0.89		
Getting vaccines is a good way to protect my child from disease	0.81		
Childhood vaccines are effective	0.85		
Having my child vaccinated is important for the health of others in my community	0.88	0.93 (0.91,0.94)	0.71
All childhood vaccines offered by my child's health care provider are beneficial	0.85		
I do what my child's health care provider recommend about vaccines	0.82		
F2: Risk			
New vaccines carry more risks than older vaccines	0.57		
I am concerned about serious side effects of childhood vaccines	0.63	0.74 (0.69,0.79)	0.50
I think childhood vaccines might cause lasting health problems for my child	0.89		



Figure 1: Path diagram for CFA of MVHS-M, which indicates the presence of correlated residual between C5 and C6

square goodness-of-fit ( $\chi^2$ ) was not reported due to its sensitivity to multivariate non-normality and sample size.<sup>39,40</sup> However, normed chi-square (NC= $\chi^2/df$ ) was introduced to reduce the sensitivity of  $\chi^2$  to sample size. Despite, it was also not reported in this study as it does not play a major role in global fit testing.32 The standardised factor loadings for the MVHS-M presented in Table III ranged from 0.572 to 0.899, which were above the recommended cut-off point for factor loading  $(\geq 0.5)$  by previous literature.<sup>31</sup> Thus, no items were removed in the CFA model. In the current study, a One of the limitations of this study, as with previous VHS correlated error item between C5 and C6, which were studies, that, is the mix of positively worded for "lack of within the same factor was added after model re- confidence" items and negatively worded for "risk" items. specification based on theoretical justification as both This mixing of wording could potentially cause differences items seem to measure the same aspect. However, this was in responses based on question-wording rather than not reported in the earlier version.29

factors based on Raykov's rho in this study ranged from worded items can result in confusion and tiredness among 0.74 to 0.93, exceeding the suggested threshold of 0.7. participants, perhaps failing to adequately address Similarly, Cronbach's alpha values ranged from 0.75 to response bias. 0.94. The findings suggested good reliability and accuracy of the MVHS-M in assessing parental vaccine hesitancy Additionally, it should be noted that all respondents in this toward childhood vaccination. Therefore, assessment process was repeated, comparable results the entire Malaysian population. Thus, generalizability would be obtained.<sup>41</sup> Previous version of VHS mostly should be done with caution. Furthermore, since the utilised Cronbach's alpha instead of the CR. The MVHS-M was translated into formal Malay, its usability is Cronbach's alpha was found in the original VHS with values ranging from 0.64 to 0.92 among Canadian samples.<sup>23</sup> This pattern of results aligns with other validation studies.<sup>18,19,22,29</sup> Additionally, Average Variance Extracted (AVE) showed an acceptable value ( $\geq 0.5$ ), indicating adequate convergent validity of the newly translated questionnaire.42 The MVHS-M also demonstrated stability and reliability over time, with a test-retest reliability demonstrated ICC of 0.77 after a 7interval, suggesting stability in assessing vaccine dav hesitancy among parents.

Comprehensive methodologies employed in the translation and cultural adaptation of the MVHS-M contributes positively impact to its clinical utility. These methodologies ensure that the newly translated version maintains the intended meaning of the original MVHS, utilised well-understood concepts and incorporates appropriate language for the Malaysian context. The application of MVHS-M is further enhanced since it was pre-tested, and piloted, and most importantly, the CFA showed a good model fit with good reliability and stability. Notably, it is interesting to note that the questionnaire consists of a concise set of nine items with straightforward and uncomplicated questions, making it suitable for selfadministration. Thus, enhances the clinical utility of the MVHS-M.

genuine underlying constructs within the items. Nevertheless, prior studies indicate that the utilization of Regarding reliability, the composite reliability (CR) of scales that incorporate both positively and negatively

> if the study were Malay ethnicity limiting the generalizability of restricted to those who can read and understand the Malay language. Further study to translate and validate it to Tamil or Mandarin would enable its widespread use across the multi-racial population in Malaysia.

# CONCLUSION

The MVHS-M demonstrates good construct validity and reliability, making it a useful tool for in assessing the magnitude of vaccine hesitancy among the parents regarding their child's vaccinations. This tool enables the identification of high-risk parents, allowing for the development of targeted intervention and more effective strategies to address vaccine hesitancy.

# FUNDING

This research was funded by Universiti Sains Malaysia through a postgraduate development incentive grant (TIPPS, 2023)

## **CONFLICT OF INTEREST**

None

#### INSTITUTIONAL REVIEW

The study was conducted based on principles outlined in the Declaration of Helsinki. Ethical approval was granted from the Ethics Committee, Universiti Sains Malaysia (USM/JEPeM/KK/23010102) and Medical Research &

Ethics Committee (MREC), Ministry of Health Malaysia, [NMRR ID-23-00277-1P7(IIR)].

# ACKNOWLEDGEMENT

We would like to thank the management of Kelantan State Health Department for the permission to conduct this study. Special thanks to all respondents who were involved in this study.

# REFERENCES

- UNICEF. World Immunization Week 2018 [Internet]. UNICEF. 2018 [cited 2022 Dec 20]. Available from: https://www.unicef.org/world-immunization-week-2018
- Faridah K. Immunisation Programme in Malaysia. In: International Symposium for Asia Pacific Experts. Hanoi; 2017.
- Institute for Public Health (IPH). National Health and Morbidity Survey (NHMS) 2016: Maternal and Child Health. Vol II: Findings. 2016;272.
- Wong LP, Wong PF, AbuBakar S. Vaccine hesitancy and the resurgence of vaccine preventable diseases: the way forward for Malaysia, a Southeast Asian country. Hum Vaccines Immunother [Internet]. 2020;16(7):1511–20.Available from: https://doi.org/10.1080/21645515.2019.1706935
- Wan Rohani WT, Noor Ain MY, Tengku Mohamad Ariff, Raja Hussin Aryati A. Issues in vaccine hesitancy in Malaysia. J Biomed Clin Sci. 2017;2 (June):42–6.
- Kalok A, Loh SYE, Chew KT, Abdul Aziz NH, Shah SA, Ahmad S, et al. Vaccine hesitancy towards childhood immunisation amongst urban pregnant mothers in Malaysia. Vaccine [Internet]. 2020 Feb;38 (9):2183–9. Available from: https://doi.org/10.1016/ j.vaccine.2020.01.043
- Joslyn Panting A, Zaikiah MZ, Norrafizah J, Perialathan K, Sheikh Shafizal SI, Muhd Ridwan Z. Potential Factors Contributing to Vaccine Hesitancy among Parents in Malaysia: An Overview. Int J Heal Sci Res [Internet]. 2018;8(7):360. Available from: www.ijhsr.org
- 8. Rumetta J, Abdul-Hadi H, Lee YK. A qualitative study on parents' reasons and recommendations for

childhood vaccination refusal in Malaysia. J Infect Public Health [Internet]. 2020;13(2):199–203. Available from: https://doi.org/10.1016/ j.jiph.2019.07.027

- Vasudevan L, Baumgartner JN, Moses S, Ngadaya E, Mfinanga SG, Ostermann J. Parental concerns and uptake of childhood vaccines in rural Tanzania – a mixed methods study. BMC Public Health [Internet]. 2020 Dec 20;20(1):1573. Available from: https:// bmcpublichealth.biomedcentral.com/ articles/10.1186/s12889-020-09598-1
- Ministry of Health(MOH), Malaysia. Kenyataan akhbar Ketua Pengarah Kesihatan Malaysia:Kes Difteria di Malaysia. 2017;
- Kannan HK. 18 cases of Diphtheria, three deaths so far in Malaysia this year. New Straits Times. 2017 Aug 8;
- Tok PSK, Jilani M, Misnar NF, Bidin NS, Rosli N, Toha HR. A diphtheria outbreak in Johor Bahru, Malaysia: Public health investigation and response. J Infect Dev Ctries [Internet]. 2022 Jul 28;16(07):1159– 65. Available from: https://www.jidc.org/index.php/ journal/article/view/16076
- World Health Organization (WHO). Measles and rubella strategic framework 2021-2030. Geneva; 2020.
- Nuwarda RF, Ramzan I, Weekes L, Kayser V. Vaccine Hesitancy: Contemporary Issues and Historical Background. Vaccines [Internet]. 2022 Sep 22;10(10):1595. Available from: https:// www.mdpi.com/2076-393X/10/10/1595
- World Health Organization (WHO). Report of the Sage Working Group on Vaccine Hesitancy. WHO. 2014.
- MacDonald NE, Eskola J, Liang X, Chaudhuri M, Dube E, Gellin B, et al. Vaccine hesitancy: Definition, scope and determinants. Vaccine. 2015;33 (34):4161–4.
- Kempe A, Daley MF, McCauley MM, Crane LA, Suh CA, Kennedy AM, et al. Prevalence of Parental Concerns About Childhood Vaccines. Am J Prev Med [Internet]. 2011 May;40(5):548–55. Available from: https://linkinghub.elsevier.com/retrieve/pii/ S0749379711000420

- Domek GJ, O'Leary ST, Bull S, Bronsert M, Contreras-Roldan IL, Bolaños Ventura GA, et al. Measuring vaccine hesitancy: Field testing the WHO SAGE Working Group on Vaccine Hesitancy survey tool in Guatemala. Vaccine [Internet]. 2018 Aug;36(35):5273–81. Available from: https:// linkinghub.elsevier.com/retrieve/pii/ S0264410X18310156
- Gentile A, Pacchiotti AC, Giglio N, Nolte MF, Talamona N, Rogers V, et al. Vaccine hesitancy in Argentina: Validation of WHO scale for parents. Vaccine [Internet]. 2021;39(33):4611–9. Available from: https://doi.org/10.1016/j.vaccine.2021.06.080
- Bussink-Voorend D, Hautvast JLAA, Vandeberg L, Visser O, Hulscher MEJLJL. A systematic literature review to clarify the concept of vaccine hesitancy. Nat Hum Behav [Internet]. 2022 Aug 22;6(12):1634–48. Available from: https://www.nature.com/articles/ s41562-022-01431-6
- Oduwole EO, Pienaar ED, Mahomed H, Wiysonge CS. Overview of Tools and Measures Investigating Vaccine Hesitancy in a Ten Year Period: A Scoping Review. Vaccines. 2022;10(8).
- Luyten J, Bruyneel L, van Hoek AJ. Assessing vaccine hesitancy in the UK population using a generalized vaccine hesitancy survey instrument. Vaccine [Internet]. 2019;37(18):2494–501. Available from: https://doi.org/10.1016/j.vaccine.2019.03.041
- 23. Shapiro GK, Tatar O, Dube E, Amsel R, Knauper B, Naz A, et al. The vaccine hesitancy scale: Psychometric properties and validation. Vaccine [Internet]. 2018;36(5):660–7. Available from: https:// doi.org/10.1016/j.vaccine.2017.12.043
- Temsah MH, Alhuzaimi AN, Aljamaan F, Bahkali F, Al-Eyadhy A, Alrabiaah A, et al. Parental Attitudes and Hesitancy About COVID-19 vs. Routine Childhood Vaccinations: A National Survey. Front Public Heal. 2021;9(October):1–11.
- 25. Sabahelzain MM, Dube E, Moukhyer M, Larson HJ, Van Den Borne B, Bosma H. Psychometric properties of the adapted measles vaccine hesitancy scale in Sudan. PLoS One [Internet]. 2020 Aug 1 [cited 2022 Nov 22];15(8 August):1–12. Available from: /pmc/articles/PMC7410231/

- Önal Ö, Eroğlu HN, Evcil FY, Kişioğlu AN, Uskun E. Validity and reliability of turkish version of the vaccine hesitancy scale. Turkish Arch Pediatr. 2021;56(3):230–5.
- Kim J, Han K, Chung SJ, Kim C. Psychometric validation of the Korean versions of the Vaccine Hesitancy Scale and Vaccination Attitudes Examination Scale. Vaccine [Internet]. 2023;41 (32):4685–92. Available from: https:// doi.org/10.1016/j.vaccine.2023.06.046
- Wang Q, Xiu S, Yang L, Han Y, Cui T, Shi N, et al. Validation of the World Health Organization's parental vaccine hesitancy scale in China using child vaccination data. Hum Vaccines Immunother [Internet]. 2022 Jan 31 [cited 2022 Oct 22];18(1). Available from: https:// doi.org/10.1080/21645515.2021.2021060
- Helmkamp LJ, Szilagyi PG, Zimet G, Saville AW, Gurfinkel D, Albertin C, et al. A validated modification of the vaccine hesitancy scale for childhood, influenza and HPV vaccines. Vaccine [Internet]. 2021 Mar 26;39(13):1831–9. Available from: https://doi.org/10.1016/j.vaccine.2021.02.039
- Wild D, Grove A, Martin M, Eremenco S, McElroy S, Verjee-Lorenz A, et al. Principles of Good Practice for the Translation and Cultural Adaptation Process for Patient-Reported Outcomes (PRO) Measures: Report of the ISPOR Task Force for Translation and Cultural Adaptation. Value Heal [Internet]. 2005 Mar;8(2):94–104. Available from: https:// linkinghub.elsevier.com/retrieve/pii/ S1098301510602525
- Hair JF, Black WC, Babin BJ, Anderson RE. Multivariate Data Analysis. 7th ed. United States Of America: Pearson Education Limited; 2014.
- Kline R. Principled and practices of structural equation modelling. 4th ed. New York: The Guiford Press; 2016.
- 33. Arifin WN. Introduction to sample size calculation. Educ Med J [Internet]. 2013 Jun 1;5(2). Available from: http://eduimed.usm.my/EIMJ20130502/ EIMJ20130502\_10.pdf
- Arifin WN. Sample size calculator (Version 2.0) [Internet]. 2017. Available from: http:// wnarifin.github.io

- Chyung SYY, Roberts K, Swanson I, Hankinson A. Evidence-Based Survey Design: The Use of a Midpoint on the Likert Scale. Perform Improv [Internet]. 2017 Nov;56(10):15–23. Available from: https://onlinelibrary.wiley.com/doi/10.1002/ pfi.21727
- 36. Raykov T, Marcoulides GA. Scale Reliability Evaluation Under Multiple Assumption Violations. Struct Equ Model A Multidiscip J [Internet]. 2016 Mar 3;23(2):302–13. Available from: http:// www.tandfonline.com/doi/ full/10.1080/10705511.2014.938597
- 37. Wang J, Wang X. Structural Equation Modeling [Internet]. 2nd ed. Wiley; 2019. (Wiley Series in Probability and Statistics). Available from: https:// onlinelibrary.wiley.com/doi/ book/10.1002/9781119422730
- Koo TK, Li MY. A Guideline of Selecting and Reporting Intraclass Correlation Coefficients for Reliability Research. J Chiropr Med [Internet]. 2016 Jun;15(2):155–63. Available from: https:// linkinghub.elsevier.com/retrieve/pii/ S1556370716000158
- 39. Wang J, Wang X. Structural equation modelling. United Kingdom: Higher Education Press; 2012.
- West SG, Taylor AB, Wu W. Model fit and model selection in structural equation modeling. In: Handbook of structural equation modeling. New York, NY, US: The Guilford Press; 2012. p. 209–31.
- Bolarinwa O. Principles and methods of validity and reliability testing of questionnaires used in social and health science researches. Niger Postgrad Med J [Internet]. 2015;22(4):195. Available from: https:// journals.lww.com/10.4103/1117-1936.173959
- Hair JF, M.Hult GT, Ringle CM, Sarstedt M. A Primer on Partial Least Squares Structural Equation Modeling (PLS-SEM). 2022.