

### **Consent Form for Research Participants**

**Participating in Interview** 

**SPHS hotline:** 6478 9608

Protocol title: Singapore Population Health Studies (SPHS) – First Follow-up

**Principal Investigator (PI):** Professor Rob Martinus <u>van Dam</u>, Saw Swee Hock School of Public Health (SSHSPH), National University of Singapore (NUS), Tahir Foundation Building, 12 Science Drive 2, #10-01, Singapore 117549

Part A. I hereby acknowledge that:			**This research has been explained to me in			
۱.	I have received a copy of the Participant Information	ation Sheet	t (state language),			
	that explains the use of my data in this study.		which I understand, by			
2.	By signing this consent form, I declare that I und	derstand its	5			
	contents and agree to:	of their	(name of translator as per NRIC).			
	(i) provide my NRIC number for the purpose of study; and	or unis				
	(ii) undergo an interview that will be audio-reco	orded and				
	if needed, be contacted to verify my survey	<mark>//interview</mark>				
	data for quality control purposes.		Name (as per NRIC) of participant			
3.	I can withdraw from the study at any point of time by		Name (as per NNO) of participant			
	informing the PI. I am aware that the withdrawal consent does not affect the research information					
	before the consent is withdrawn and such inform		,			
	still be retained and used for research.					
oar	t B. I hereby further agree to:		Signature/ thumb print (participant)			
		YES NO				
١.	Allow SSHSPH to obtain information about my					
	health from the national registries or my health		NRIC (participant)			
	records for the purposes of this study.		Date of Consent			
2.	Allow SSHSPH to obtain information about my		Day (DD) Month (MM) Year (YYYY)			
	health from the national registries or my health records to be used by NUS and/or NUS'					
	collaborators for future Public Health					
	Research.					
3.	Be re-contacted for further consent under the		Name (as per NRIC) of consent taker			
	circumstances identified in the Participant		<b>-</b>			
	Information Sheet.					
1.	Be re-contacted for invitation to future follow- ups related to this study in 3-5 years. I		]			
	understand that future follow-ups will be		Signatura (consent taker)			
	subject to an Institutional Review Board's		Signature (consent taker)  Date of			
	(IRB) approval and SSHSPH may request the		Consent			
	relevant ministry for my updated contact information.		Day (DD) Month (MM) Year (YYYY)			
<u>.</u>	Be re-contacted for invitation to future Public		** This research has been explained to the participant			
<i>,</i> .	Health Research. I understand that future					
	studies will be subject to an IRB's approval.		in (state language),			
ar	t C. Donation of data for use in future Public Hea	alth	-			
Res	<mark>earch.</mark>					
		YES NO	Name (as per NRIC) of <mark>translator</mark>			
_	ree to donate my data collected for this study to used in future Public Health Research. I		Ivalie tas bei ivivor of translator			
	erstand that the donated data will be coded and					
	dentified for research use by NUS researchers					
and/or NUS' collaborators including overseas collaborators. The future research will be subject to			Signature (translator)			
	RB's approval. SPHS operations team may re-					
der	tify the data in order to contact me for the		Date of Consent			
on	ditions I consent to in Part B above.		Day (DD) Month (MM) Year (YYYY)			



# **Consent Form for Research Participants**

Participating in Health Screening

SPHS hotline: 6478 9608

Protocol title: Singapore Population Health Studies - First Follow-up Principal Investigator (PI): Professor Rob Martinus van Dam

Saw Swee Hock School of Public Health (SSHSPH), National University of Singapore (NUS), Tahir Foundation Building, 12 Science Drive 2, #10-01, Singapore 117549

#### Part A. I hereby acknowledge that:

- 1. I have received a copy of the Participant Information Sheet that explains the use of my data and samples in this study.
- 2. By signing this consent form, I understand its contents and agree to a health screening. At the health screening, I may choose to donate up to 29mls (~3 tablespoons) of blood and 20mls of urine for research.
- 3. I can withdraw from the study at any point of time by informing the PI although if I have donated samples for the study, I can only withdraw my consent to discontinue the use of any unused portion of samples that are re-identifiable. I am aware that the withdrawal of consent does not affect the research information obtained before the consent is

withdrawn and Such information may sti	iii be retained and used for research.							
Part B. I hereby further agree to:  1. Be contacted if a medical issue is found during the course of this study or in future research, in order to facilitate my seeking proper medical advice. I understand that my participation in the study is not considered medical treatment, and that the researchers are not qualified medical practitioners able to provide diagnoses.								
Part C. Donation of data and samples for use in future Public Health Research.  I understand that the donated data and samples will be coded and used for research by NUS researchers and/or NUS' collaborators including overseas collaborators. The future research will be subject to an IRB's approval. If I refuse the donation of samples for future research, no samples will be stored for future research. The SPHS operations team may reidentify the data in order to contact me for the condition I consent to in Part B above. I hereby agree to:  Yes No								
1. Donate 6mls of my urine sample to be u	used in future research.							
2. Donate 20mls (~2 tablespoons) of my blood sample to be used in future research.								
3. The use of my data and samples in future genetic research.								
<ol> <li>The use of my data and samples in f commercial firms. I will not have any fir the research.</li> </ol>								
** This study has been explained to me in (name of translator).  Name (as per NRIC) of participant Signature / the	humbprint of participant (state language), which		Day 1	Month Year e of consent				
I, the witness, certify to the following:  a) I am 21 years of age or older. b) I have taken reasonable steps to ascertain the identity of the participant. c) To the best of my knowledge, the participant had the study fully explained in a language understood by him/ her and clearly understands the nature, risks and benefits of his/ her participation in the study. d) I have taken steps to ascertain that the consent has been given voluntarily without any coercion or intimidation.								
Name (as per NRIC) of witness	Signature of witness	Day	Month Date of cons	Year ent				
Name (as per NRIC) of consent taker	Signature of consent taker	Day	Month Date of cons	Year cent				
**The study has been explained to the participa	ant in (state language							
Name (as per NRIC) of translator	Signature of translator	Day	Month Date of cons	Year eent				

<sup>\*\*</sup> Fill in these sections if the participant is unable to read participant information sheet and consent or any of the translated versions.

NUS. May I speak to [participant name]?							
If participant is not available	Is there a better day and time to reach [Mr/Mrs/Ms] [participant name]? Thank you for your assistance. I will try to call back then.						
When participant is on the phone	This is a follow up about a research project that you've previously taken part on [state year] at [state site].						
Introduction	This follow up is for us to learn how has your lifestyle and health changes since the last time we interviewed you. As this is a cohort study, your participation is valuable to us to study how lifestyle factors contribute to our health.						
	There are 2 parts to this study. First is telephone interview which will take about 20 minutes. Second is an optional health screening at one of our health screening sites.						
Brief explanation	The interview will cover questions related to your health and lifestyle. If there are questions that you are not comfortable with, just let me know and we will skip those. Also our conversation will be audio recorded for quality control and training purposes.						
Pre-consent	May I proceed to describe this study in detail? At the same time, I will ask for your consent about taking part in this study. As a proof of your consent or refusal, I will audio-record this conversation, is that ok?						
	[If YES, proceed.] [If NO, reiterate the reason. If participant still refuse, end the recruitment]						
<u>Verbal Consent</u>							
We are inviting you to take part in Singapore Population Health Studies – First Follow-up study. This study is conducted by Saw Swee Hock School of Public Health, National University of Singapore. The principal investigator (person in charge of this research) is Professor Rob Martinus van Dam. The purpose of this revisit is to find out how factors like diet, exercise and smoking etc. are related to heart disease, diabetes, stroke and other common diseases in Singapore and how these important factors have changed since the last survey and health screening.							
We need your NRIC/FIN to be sure that we are contacting the correct person for this study and for future contact purposes. May we use your NRIC for the use of this research?							
YES, ve	rify NRIC/FIN NO, do not proceed						
The interview will will take about 20	bu over the phone about your health, exercise, use of tobacco, alcohol and medicines? be audio-recorded for quality control and training purposes. This telephone interview min. During the interview, if you are not willing to answer a particular question, let me ove on to the next question YES NO, do not proceed						
We may need to access to relevant national registries or your medical records to obtain the relevant information about your health status and medical condition for this study or future study. If you agree to this, we will be able to obtain the health information useful for this study. Researchers will only be given the health data without the names and NRICs attached. However, if you disagree to allow us to access the use of your relevant health status and medical condition for this study or future study, you can still participate in this study.							

records that are useful for this study? \_\_\_\_\_ YES \_\_\_\_ NO

Do you agree to allow us to gather your health data from the relevant national registries or medical

## F1 CATI PS Annex J PHONE SCRIPT

Do you agree to allow us to gather your health data from the relevant national registries or medical records that are useful for future research? YES NO
As we are studying long term diseases, we will need to follow up on your health again later and to collect information about your lifestyle and health changes after 3-5 years. We will ask you for your consent each time we invite you to continue this study.
Do you agree to be re-contacted in 3-5 years' time to conduct further follow-up or related studies of major health conditions? YES NO
Besides this study, there are various public health research also conducted by the school. Do you allow us to contact you to provide news about our research or invite you to participate in other research studies by the school? YES NO
It is your choice whether you want to participate or not. You may also withdraw your participation at any time by calling the SPHS hotline at 6478 9608 during office hours or send an email to <a href="mailto:sphs@nus.edu.sg">sphs@nus.edu.sg</a> . You can also contact us to ask about the study. If you wish to ask about your rights and welfare as a participant of this study, you can contact the NUS Institutional Review Board (IRB) at 6516 1234.
I certify that the participant consented verbally to participate in this study.
Name and signature of staff obtaining verbal consent  Date

#### Interview

Now let us start the interview.

[End of interview]

#### Offer health screening

If you want to go for the health screening, I will arrange an appointment for you. The health screening will take around 1 ½ to 2 hours to complete. During the health screening there will be blood and urine tests, measurement of your height, weight and blood pressure, ECG, foot assessment and a hand grip test.

#### [For SCCS/CR/AR/SH2012 participants]

You will also receive a vision test and a test of your ability to walk a short distance. You may also be selected for a test of your lung function.

#### [For MECT participants]

You may also be selected for a survey about your well-being.

#### [For SH2 participants]

You will also receive a vision test and a test of your ability to walk a short distance. You may also be selected for a survey about the environment in your community.

After completing the health screening, you will be reimbursed another \$50 in cash for your transportation and as a token of appreciation for your time taken to go for the health screening. A copy of the results will be mailed to you around 4 weeks after the health screening.

Would you want to go for the health screening?

[If Yes, offer location and appointment slots and book]

At the health screening centre, our staff will explain to you the procedures and ask you to sign a consent form before your health screening.

# F1 CATI PS Annex J PHONE SCRIPT

In appreciation for your time, we would like to reimburse you with a token \$5. This could be transferred to you by PayNow if your personal mobile number is registered to receive payment via PayNow.

[If participant does not have this service set up] Alternatively, I could send to you a \$5 NTUC voucher via mail.

[Verify again mobile number/address is correct]

Please note that your personal information such as your name and contact details may be disclosed to third parties authorised by NUS to enable them to perform processes or services related to this project. Let me know if you need more details. [The processes/services include mail, telecommunications, messaging, IT and financial services]

Thank you for your support and participation in our research. Have a nice day.

[THE END]

F1 CF-S Annex N

## CONSENT FORM (FOR ONLINE SURVEY)

I have read the Participant Information Sheet that explains the use of my data in this study.

By submitting this consent form, I declare that I understand its contents and agree to provide my NRIC number for the purpose of this study and complete a survey about my social background, health and lifestyle, and if needed, be contacted to verify my survey data for quality control purposes.

I understand that I can withdraw from the study at any point of time by informing the Principal Investigator. I am aware that the withdrawal of consent does not affect the research information obtained before the consent is withdrawn and such information may still be retained and used for research.

		res	NO
1.	I agree to allow the Saw Swee Hock School of Public Health (SSHSPH) to obtain information about my health from the national registries or my health records for the purposes of this study.		
<mark>2.</mark>	I agree to allow SSHSPH to obtain information about my health from the national registries or my health records to be used by NUS and/or NUS' collaborators for future Public Health Research.		
<mark>3.</mark>	I agree to be re-contacted for further consent under the circumstances identified in the Participant Information Sheet.		
<b>4</b> .	I agree to be re-contacted for invitation to future follow-ups related to this study in 3-5 years. I understand that future follow-ups will be subject to an Institutional Review Board's (IRB) approval and SSHSPH may request the relevant ministry for my updated contact information.		
<u>5.</u>	I agree to be re-contacted for invitation to future Public Health Research. I understand that future studies will be subject to an IRB's approval.		
6.	I agree to donate my data collected for this study to be used in future Public Health Research. I understand that the donated data will be coded and de-identified for research use by NUS researchers and/or NUS' collaborators including overseas collaborators. The future research will be subject to an IRB's approval. SPHS operations team may re-identify the data in order to contact me for the conditions I consent to above.  I agree to I do not wish to		
	participate participate		

Participant who fills this form online prior to the online survey and chooses "agree" will be directed to a page where the PIS and CF can be printed out and where he/she can start the survey. Participant who chooses otherwise will be directed to a page where the following options will be available:

- Print PIS and CF can be printed out and where he/she can.
- Register interest to be given an appointment for health screening
- Exit the application