APPENDIX

Appendix 1. Data Management

Appendix 2. FirstCPR Participant Information and Consent form

Appendix 3. FirstCPR evaluation survey questionnaire

Appendix 4. Roles and Responsibilities

Appendix 5. Audit and Monitoring of study

Appendix 1. Data Management

Majority of the data will be collected via surveys via RedCapTM. The study Standard Operating Procedures (SOP) manual and a more specific RedCapTM SOP has been drafted to document all necessary steps in data entry and validation. Any data collected as paper copies will be scanned and uploaded onto the electronic platforms. Data and reports extracted from RedCapTM will be stored in de-identified form on a secure webbased platform and limited to be accessed by authorised study personnel- thus maintaining confidentiality throughout the study period and after the trial. Focus group discussions and in-depth interviews will be recorded on a digital recorder with consenting participants. The recording will be transferred onto the University of Sydney's Research Data Storage (RDS) which is a centralised secure data storage platform. The discussions will then be transcribed following which all the recordings will be destroyed.

Appendix 2. FirstCPR Participant Information and Consent form



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FirstCPR study: Education and training of community members in responding to an Out-of-Hospital Cardiac Arrest (OHCA)



PARTICIPANT INFORMATION STATEMENT

What is this study about?

You are invited to take part in the FirstCPR study, led by Professor Clara Chow at the Westmead Applied Research Centre, University of Sydney. The information in this document will explain what the research is about, why it is being done, and what is involved if you choose to participate. You have been invited to participate in this study because your club/organisation has agreed to work with us to raise awareness and knowledge of what to do when people see a cardiac arrest in the community. This means we are inviting all members (18 years and older) to participate in the educational campaign mainly related to cardiopulmonary resuscitation (CPR). This program is supported by a National Health and Medical Research Council (NHMRC) research grant.

This Participant Information Statement tells you about the research study. Knowing what is involved will help you decide if you want to take part in the research. Please read this sheet carefully and ask questions about anything that you don't understand or want to know more about.

Providing your consent to participate:

Participation in this research study is voluntary. Taking the FirstCPR survey will mean that you consent to take part in this study. This means you are telling us that you:

- ✓ Understand what you read in this sheet.
- ✓ Agree to take part in the research study as outlined below.
- ✓ Agree to the use of your personal information as described (name, email, phone number).
- ✓ Agree that you are 18 years or older
- ✓ Are happy to be contacted in the future about the project

What will happen if I say I want to be in the study?

If you wish to participate there will be a 5-minute questionnaire at the start about cardiac arrest in the community. Then you will be asked if you would like to receive information and reminders about cardiac arrest and CPR via text message and/or email for a period of 10 to 12 months. Some messages will include links to additional information on websites, videos or factsheets which have been selected by the First CPR research team. At the end of the study period, there will be another short questionnaire. Some of you will be invited to continue to receive less frequent reinforcement messages for an additional 12month period and be asked to complete a similar 5-minute questionnaire at the end of this second period. You can opt out of receiving messages at any time by replying 'STOP.'

During the study period you may be contacted about participating in an interview or group discussion about your views on cardiac arrest, CPR education and training. If you are selected for these, you will be sent more information on these closer to the time and can decide if you wish to join at that stage.

Your organisation will also be provided with materials to share with members and will host a face-to-face information session about basic life-saving skills. These sessions will demonstrate how to do CPR, use a defibrillator, and provide you with the opportunity to ask questions of an instructor. Where a need is identified, arrangements will be made for a multilingual interpreter to be present at the information sessions so translation of the information can be relayed in the language of the audience.

How long will the study take?

Your organisation will have access to FirstCPR educational and training material over the next 12 months. However, the time commitment from participants equates to viewing fortnightly short educational and informative messages that we will send via email or text, and these can be viewed in your own time. Participants who enjoy receiving these messages can sign up to receive monthly messages for an additional 12-month period to reinforce learning. You will also be invited to attend a one-hour in-person educational and training session and register to claim one of 30 free vouchers for a 2.5-hour accredited training session that will include hands-on CPR training on a manikin.

Are there any good things about being in the study?

By being a part of this research, you will also be given access to training materials and you will contribute to our research to understand whether programs like this can increase awareness and knowledge about CPR. There will also be up to 30 vouchers available to participants in your organisation to attend a free accredited CPR training course that normally costs \$65. All

participants will also go into a draw to win one of ten Coles/Woolworths vouchers valued at \$30 each, drawn at the end of each annual survey rollout.

Are there any risks or costs associated with being in the study?

Aside from giving up your time to complete the surveys, we do not expect that there will be any risks or costs associated with taking part in this study.

What will happen to information about me that is collected during the study?

Your information will be stored securely, and your identity/information will be kept strictly confidential, except as required by law. Access to this information will only be permitted to authorised researchers directly involved in the study. Study findings may be published, but you will not be individually identifiable in these publications.

Do I have to be in the study? Can I withdraw from the study once I've started?

You can decide if you want to take part in this or not. You don't have to, and it is completely up to you. If you decide you want to be in the study and then you change your mind later, that's ok. All you need to do is tell us by text message, email, or telephone call that you want to withdraw. If you decide to withdraw from the study, we will not collect any more information from you. Any information that we have already collected, however, will be kept in our study records and may be included in the study results.

Will you tell me what you learnt in the study at the end?

Yes, we will if you want us to. There is a question on the consent form that asks you if you want us to tell you what we learnt in the study (study findings). If you select 'Yes', when we finish the study, we will tell you what we learnt. If you have any questions, you can ask us by calling on <u>0412</u> <u>369 519</u> or email us at <u>warc.firstcpr@sydney.edu.au</u>.

What if I am not happy with the study or the people doing the study?

If you are not happy with how we are doing the study, then you can

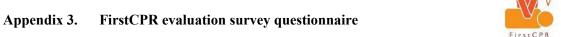
- Call the university on +61 2 8627 8176 or
- Write an email to human.ethics@sydney.edu.au

FirstCPR Participant Consent Form

If you are happy to be in the study, please

- write your name in the space below
- **sign** your **name** at the bottom of the next page
- put the **date** at the bottom of the next page.

You should only say 'yes' to being in the study if you are 18 years or older, understand what it is
about, and you want to be in it.
Yes, I,
this research study.
In saying yes to being in the study you 1) confirm you have read the participant information sheet,
2) Understand what is involved in participation, 3) understand that by providing contact details and
completing the survey you consent to participate in the study and 4) are happy to be contacted in
the future about the project.
Please tick (\$\angle\$) 'Yes' or 'No' if you would like us to email you a summary report at the end
if the study to tell you what we learnt in the study.
ΠVFS
□ YES
□ YES □ NO
□ NO
□ NO
□ NO If, yes please let us know your email so we can send you the report
□ NO If, yes please let us know your email so we can send you the report
□ NO If, yes please let us know your email so we can send you the report Email.
□ NO If, yes please let us know your email so we can send you the report Email.



FirstCPR Baseline survey					
Question	Options				
1. Kindly tell us some information about yourself					
Please enter your age (in years)					
	(Eligible to participate if 18years or older)				
Please select your Gender	☐ Male				
	☐ Female				
	☐ Another term:				
	☐ Prefer not to answer				
What is the highest level of	☐ Primary/Grade School				
schooling you have completed?	☐ Some high school				
	☐ High school graduate				
	☐ Technical college or some University				
	☐ University diploma or degree				
	☐ Postgraduate				
In which country were you born?	☐ Australia				
	☐ China				
	☐ England				
	☐ India				
	☐ Italy ☐ Malaysia				
	☐ New Zealand				
	☐ Philippines				
	☐ South Africa				
	☐ Sri Lanka				
	☐ Vietnam				
	☐ Other, please specify				
(If born outside Australia),					
Approximately how many years have					
	(if less than a year- e.g., 6 months- please enter 0.5)				
Australia					

What language do you mainly speak	П	English					
at home?		☐ Mandarin					
at nome.	☐ Arabic						
		Cantonese					
		Vietnamese	:				
		Italian					
		Greek					
		Hindi					
		Spanish					
		Punjabi					
		Other, pleas	se specify.		•••••		
What is your Postcode?							
Which of the following describes		Working for	r an empl	oyer or cond	lucting a bu	siness	
your current status?		Unpaid wor			8		
		Unemploye	d, looking	g for work			
		Studying	/ 0 /	1			
	Homemaker / Stay-at-home parent						
	☐ Retired ☐ Other, please specify						
(If calcated months of attacks of							
(If selected working / studying / looking forwork): Can you indicate		Medical or Law Enforce					
if your current occupation (or field of		Fitness Inst		nach			
study) fits into any of the following		Social work		oacii			
industry categories		Aged care w		rer			
- if not, please select 'Other' and		Jail or corre					
specify		Transport w	orker				
industry category		Flight attend	dant				
		Firefighter					
		Lifeguard					
		Constructio	n worker				
		Electrician					
		Teacher		ataff			
		Childcare po		stan			
		Other, pleas					
(if category other than medical or		Yes	se speerry				
Health selected above): Have you		No					
ever worked or been trained in a		110					
medical or health-related field?							
In general, would you say that your	Ver	y Poor	Poor	Fair	Good	Excellent	
health is							
2. The next few questions are rela		•	_	-		n). Please	
select theoption that best reflects		_					
I would rate my overall knowledge	Ver	y Poor	Poor	Fair	Good	Excellent	
of CPR as							

Have you heard of Hands-only or Compression-only CPR?	☐ Yes ☐ No					
Standard CPR involves chest con	npressions a	nd mouth-to-n	nouth brea	athing and is		
performed on a personwho is sus	_				pression-	
only CPR involves resuscitation v	with chest co	ompressions or	nly and no	mouth-to-mo	uth	
breathing.						
I feel confident in my ability to perfo						
Standard CPR	Not confident	Somewhat confident		Confident	Very confident	
Hands-only						
CPR						
I would be willing to perform CPR (on not breathing normally if they were a		rd or hands-only) on a perso	on collapsed an	d	
Family member	Definitely not	Probably not □	Maybe	Yes, probably □	Yes, definitely	
Friend		П	П	П	П	
Stranger		П	П			
If Definitely Not, Probably Not or Maybe (for FAMILY MEMBER) which of these statements best describes your reasons for why you would not be prepared to perform CPR on this person? (You can select more than oneresponse)	□ Don't know how to do CPR □ Don't feel confident □ Concerned about hurting the person □ Concerned about being sued □ Concerned about not performing CPR properly □ Physically unable to perform CPR □ Concerned about infection □ Other, please specify					
If Definitely Not, Probably Not or Maybe (for FRIEND) which of these statements best describes your reasonsfor why you would not be prepared to perform CPR on this person? (You can select more than one response)	☐ Other, please specify					

If Definitely Not, Probably Not or Maybe (for STRANGER) which of these statements best describes your reasons for why you would not be prepared to perform CPR on this person? (You canselect more than one response)	□ Don't know how to do CPR □ Don't feel confident □ Concerned about hurting the person □ Concerned about being sued □ Concerned about not performing CPR properly □ Physically unable to perform CPR □ Concerned about infection			
	☐ Other, please specify			
3. Training related questions				
Have you ever been trained in CPR?	☐ Yes ☐ No			
(If Yes), When did you last receivetraining?	☐ Less than 12 months ago			
	☐ 1 to 5 years ago			
	☐ More than 5 years ago			
	☐ Can't recall			
(If Yes), Was your most recent training led by a qualified trainer/instructor?	☐ Yes ☐ No			
(If Yes), Why did you undertake your most recent CPRtraining?	☐ Requirement of my job ☐ Requirement of a community or sporting club ☐ Self-initiated ☐ Other, please specify			
If (No- to ever trained), Which of these statements best describes your reasons for not receiving CPR training? (You can select more than one response)	☐ Never thought about it ☐ Cost ☐ Time ☐ Didn't know where to go to learn ☐ Other, please specify			
known as Defibrillator. An AED i someone having acardiac arrest. I	ut AED (Automatic External Defibrillator) also simply s a portable device that can potentially save the life of it checks the heart's rhythm and sends a shock to the heart easy-to-use and can guide anyone to use it through simple			
Please select the option that best refle	ects your response to the statements below			
I would rate my overall knowledge of a defibrillator (AED) as	 □ Very Poor □ Poor □ Fair □ Good □ Excellent □ Not Applicable as I had never heard of an AED/Defibrillator 			

Please select the option that best refle				d feel to		
I would feel confident to use an AED in an emergency	 □ Not confident □ Somewhat confident □ Confident □ Very confident □ Not Applicable as I have never heard of an AED/Defibrillator 					
If an AED/Defibrillator were availab	ole, I v	would be	willing to use	it in an emerge	ency if they w	vere a
Family member Friend Stranger If Definitely Not, Probably Not or Maybe (for FAMILY MEMBER)		Don't kn	Probably not □ □ □ now how to use a	Maybe	Yes, probably	Yes, definitely
selected, which of these statements best describes your reasons for why you would not be prepared to use an AED on this person? (You can select more than one response)	☐ Concerned about hurting the person ☐ Concerned about being sued ☐ Concerned about not being able to operate it properly ☐ Other, please specify					
If Definitely Not, Probably Not or Maybe (for FRIEND) selected, which of these statements best describes your reasons for why you would not be prepared to use an AED on this person? (You can select more than one response)	 □ Don't know how to use an AED □ Don't feel confident □ Concerned about hurting the person □ Concerned about being sued □ Concerned about not being able to operate it properly □ Other, please specify 					
If Definitely Not, Probably Not or Maybe (for STRANGER) selected, which of these statements best describes your reasons for why you would not be prepared to use an AED on this person? (You can select more than one response)		Don't kn Don't fee Concernd Concernd Concernd properly Other, ple	now how to use a el confident ed about hurting ed about being s ed about not bei	an AED g the person sued ing able to opera	ate it	
Thank you	ı for	comple	ting the surve	y questions.		

Note: Survey items on training, willingness and confidence have been adapted from validated surveys used in Australia [Refs: 10,17]

Appendix 4. Roles and Responsibilities

Committees

Members of the investigating team as well as partner organisations formed the steering committee and continue to participate and provide guidance in various discussions and decisions related with the project. In addition, three subcommittees or working groups have been set up based on members' expertise and interest to oversee and provide regular input and support in three key areas of (i)intervention development, (ii)implementation plan and (iii)scientific/ statistical advice.

Subcommittees:

Intervention(development) committee: oversees the development of educational tools and materials for the intervention. Broadly, key tasks involved discussion on key messages that need to be conveyed in the intervention campaign, and identification, collation and reviewing of all intervention material aligned with these key messages.

Implementation(planning) committee: deliberates on planning the delivery of the intervention and implementation of the project as per the study protocol. Key discussions involved - the approach to selecting and recruiting community groups, planning of study areas, recruitment approaches for urban versus regional areas, examination of feasibility or rolling interventions in organisations within a select timeframe, review, and comment on study timeline, and advise on the process evaluation component.

Scientific(advisory) committee: reviewed and provided advice on the development and amendments to the scientific protocol and included deliberation over the design and methodology, survey items, analysis and evaluation plan, data management and addressing any issues raised by the ethics committee.

Operations committee: The study will be coordinated and managed by the Westmead Applied Research Centre (WARC), The University of Sydney. A core committee of investigators, project officers have been meeting regularly to organise and facilitate the working groups, set up the project, develop key documents and contracts and troubleshoot issues related with intervention development and implementation of the project.

Appendix 5. Audit and Monitoring of study

The research team has established quality control procedures for data collection. Through regular audits of study implementation, data collection procedures will further assure that any issues including will be picked up and attended to in time. All data entry will be completed via RedCapTM, which will allow for real time data query generation for values entered outside of pre-set valid ranges and consistency checking. Only authorised research staff will have access. All entered data forms will be electronically signed (by use of the unique password) by authorised study staff. All changes made following the initial entry will have an electronically dated audit trail. Centralized coding of outcomes will be performed by a trained researcher and reviewed by the team statistician, to confirm accuracy of coding and correct reporting of outcomes by the sites. Data monitoring will thus be conducted throughout the study, but no interim analysis is planned. Monitoring of data will ensure any adverse or unintended events are picked up in time and reported as outlined in the ethics and governance documents. While an independent audit is not planned, an internal audit by the sponsor may occur during the trial.