REVOLUTIONISING CLINICAL TRIAL RECRUITMENT: UNLEASHING SOCIAL MEDIA'S POWER! DUNCAN COLYER

SENIOR MANAGER: CLINICAL RESEARCH

29 November 2023





STRATEGIC PROGRAM PLAN (2021-2024)



https://www.viccompcancerctr.org/ab out-us/strategy/strategic-programplan-2021-24/





Overcoming cancer together

FOR ALL

VISION

To save lives through the integration of cancer research, education and patient care. Through innovation and collaboration, the VCCC will drive the next generation of improvements in prevention, detection and cancer treatment.

ASPIRATION

Our alliance will become a global leader in the transformation of cancer outcomes.

MISSION

Harness the capabilities of our partners to position Victoria as a trailblazer in research-led, consumer-engaged, cancer education, prevention, detection, treatment and care.

MOBILISE

Mobilise and extend the collective and diverse strengths of VCCC members

ADVOCATE

Advocate for practice and policy improvements

TRANSLATE

Be a cutting-edge platform for translating new evidence into practice

- 1 Link clinical, biological, genomic and patient experience data to enable discoveries
- Drive the translation of cancer biology to 21st century personalised cancer care, prioritising new technologies and low survival cancers
- Accelerate the development of novel therapies

4 Drive an innovative. high-performance clinical trial sector through capacity building

BOLD

- Implement data-driven value-based cancer
- 6 Extend a distributed model of leadership to drive impact on patient outcomes

outcomes for all Victorians affected by cancer

- 7 Fast track innovations in regions with poorest cancer outcomes
- 8 Build on initiatives to meet the needs of vulnerable groups

- 9 Develop leadership skills and empowerment of researchers, clinicians and consumers
 - 10 Build a state-wide oncology learning hub leveraging Victoria's content and knowledge translation expertise

Supported by

Catalyse innovative concept generation to grow research quality and impact Extend a distributed model of leadership to drive impact on patient outcomes Embed the lived experience of cancer into research and its translation Transform educational programs into digital platforms

FOUNDATION ELEMENTS









COLLABORATING PARTNERS













































Australia





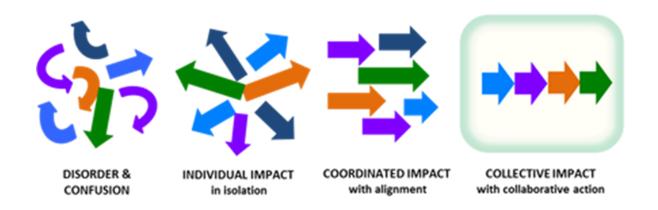


NETWORKED / COLLECTIVE IMPACT APPROACH

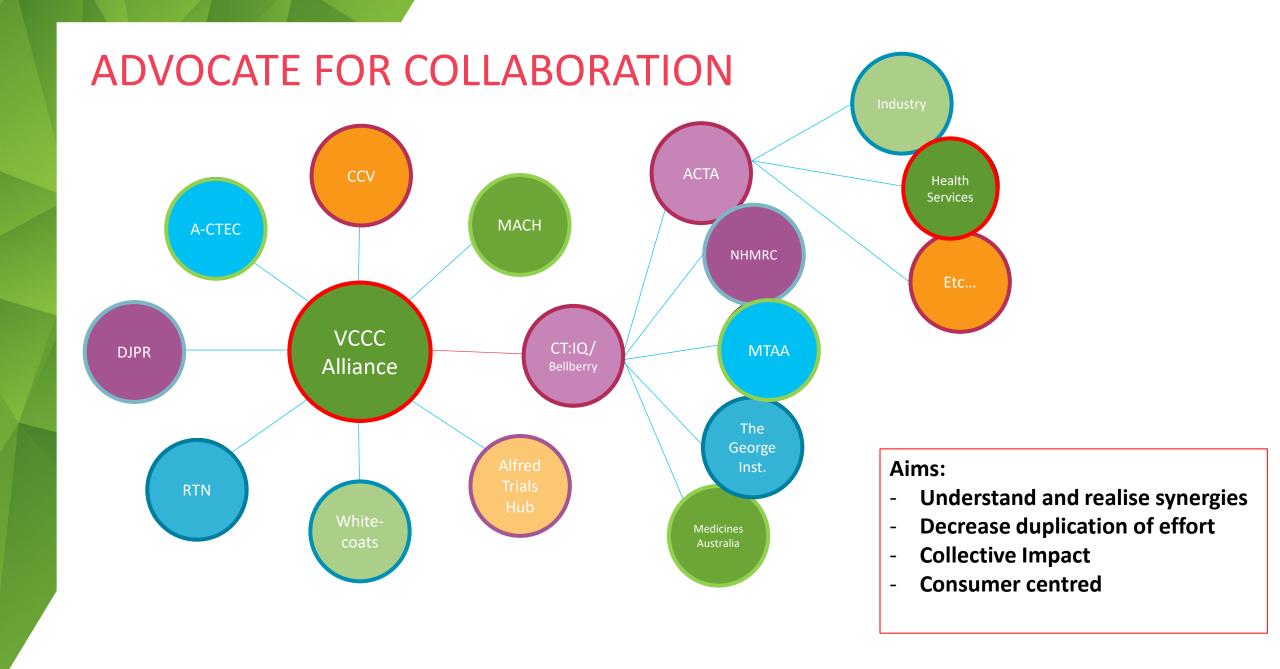
VCCC Alliance's role in Australian cancer eco-system

Facilitator, enabler, connector

- Strong connections with:
 - cancer research institutions and hospitals across Australia
 - o state and national governments
 - advocacy organisations
 - o consumer groups
 - Integrated Cancer Services
 - Cancer Councils







CLINICAL TRIAL INNOVATION OVERVIEW



Program goal

Program 4

Drive an innovative, high-performance clinical trials sector through capacity building

Short title Innovative clinical trials



Objective 1:

Strengthen Clinical Trial Participation through the adoption and conduct of new clinical trial methods

Objective 2:

Support high-performance clinical units through harmonising performance



Expand Registry Trials to new areas

Improve Clinical Trial Participation

Enhance Workforce Capability

Enhance Business Capability



- Project Officers and research fellows
- Expand registry trials to include other disciplines and less established trial groups
 - e.g.
 - other craft groups
 - regional sites
- · Education initiatives for registry trials

- · Consumer led project to explore barriers to participation
- Pilot e-consent/remote consent
- Explore prospective analysis on eligible vs recruited patients
- · Explore central approval for multisite trials

pharmacists

Community of practice for directors Mentor program to support

Education initiatives recommended by CTU community of practice to steering group

groups

process

- Expand competency frameworks e.g.
- development of trials in emerging trial groups

· Initiatives to improve engagement with industry for business sustainability

· Business development for emerging trial

Define metrics for success & collection

Synergies

Program 1, 6 & 7

Program 3 & 5

Program 3 & 10

PROJECT 4.2 – EXPAND OR DEVELOP NEW METHODOLOGIES TO IMPROVE CLINICAL TRIAL PARTICIPATION

Sub-project 4.2.1: New methods of consent

Explore the use of two way communication in clinical research and clinical trials – with CT:IQ

Sub-project 4.2.2: Barriers to clinical trial participation

Patient's view on clinical trial matching websites

Clinician's perceptions on registry trials

Consumer awareness campaign (boosting awareness, debunking myths)

Social media as a recruitment tool

WHY SOCIAL MEDIA MATTERS... IN CLINICAL TRIALS

Patient recruitment remains a (huge) problem

Around 20% close for poor recruitment but many more are affected

Lack of or slow recruitment can lead to issues with:

- Unnecessary cost to sites
- Obsolescence
- Ethical issues (unnecessary risk to patients), confidence by researchers and patients and
- Progression of the science!



ADVERTISING TO SOLVE POOR RECRUITMENT

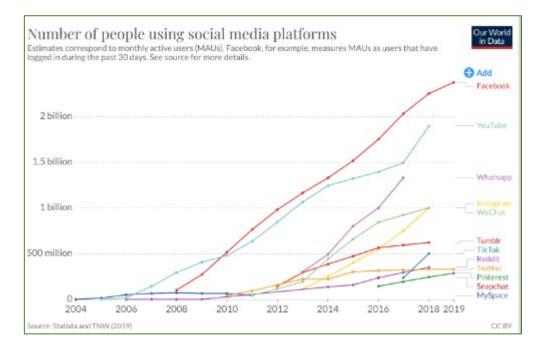
Traditional:

- Flyers
- Posters
- Signs

Digital:

Social Media





PROMISES, PROMISES

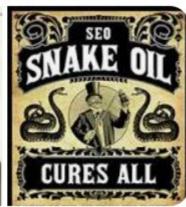
Cheap Effective Targeted

But...

Lots of work
Unclear how/who to moderate
False positives in screening







OUR HYPOTHESIS

 Improving the understanding of the use of social media can improve clinical trial recruitment

Steps

- Scoping work
- Contacting stakeholders
- Understand the landscape and limitations of current resources



Bellberry Applications

BA G9

Advertising and social media

WHAT WE FOUND

- Confusion!
- Information was out of date
- Updates were needed
- Information was often brief
- Emphasis was on 'effective' social media use
- The enigma of a 'social media plan'

National Statement on Ethical Conduct in Human Research

5.1.19 For many research projects, researchers should provide reviewers with proposed recruitment materials (e.g. notices, flyers, advertisements, and social media posts) prior to use, including those materials that are developed subsequent to the initial review of the research proposal. However, for some research designs or where recruitment material needs to be ad lib, adapted or tailored to the context (such as some social media, radio or other oral communication) a description of the strategy and broad messages is sufficient.



Social Media: Use in Research

Social Media can be used as a tool for research in a number of ways, such as advertising, recruiting and communicating with participants. As well as taking the time to understand the advantages and limitations of the different social media platforms (to be used in your research project), please also ensure you consider the relevant ethical and governance issues when considering the use of social media in your research study.

Governance

- All use of Social Media e.g. Facebook, for a research study must be approved by the RCH Human Research Ethics Committee (HREC) and must comply with RCH and MCRI policies and guidelines. Please ensure you have read the relevant Institution policy:
 - RCH Social Media Policy
 - RCH Media Protocol
 - Media Policy & Procedure (MCRI7001)
 - MCRI Social Media Policy
- Researchers must not contravene the relevant terms and conditions of the Social Media sites they use e.g. Facebook Policy.

KNOWLEDGE GAP

What was available



What was needed

THE GOOGLY!

January 12, 2022

Meta ban on health-targeting ads will soon restrict clinical trial recruiters





A new ban on "Detailed Targeting" options for advertisers on Facebook and other Meta-owned social media platforms – which includes restricting marketing based on "health causes" – takes effect January 19, and may make it more challenging for health care and life sciences companies to use these platforms to



Hogan Lovell

Messenger, and WhatsApp.



its new policy that bans "Detailed Targeting" options for advertisers, Meta

cited "Lung cancer awareness," "World Diabetes Day," and "LGBT culture."

However, Facebook's announcement of its new policy also includes recommendations for advertisers trying to find the right audience, including broad targeting via gender and age. Facebook advertisers will still be able to employ location based-targeting, including finding individuals based on ZIP code. Without the corresponding disease information, these are seemingly of less value.



WHAT SUPPORT AND GUIDANCE WAS IN OUR SCOPE?

IN

Support
Awareness
= FAQ and
Procedures

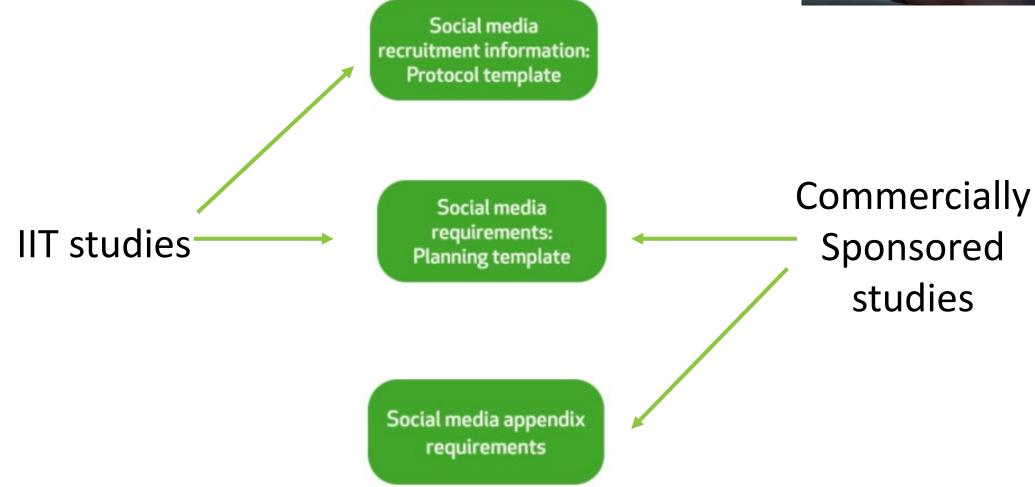


OUT

Policies SOPs

SUPPORTING RESEARCHERS FROM THE HREC'S LENS





Social media requirements for clinical trials

The information gathered here can be used to communicate with HREC.

	Planning
Target audience	E.g. 18-30 year olds
Social media platforms to	Twitter
run ads	LinkedIn
1	Facebook
	Instagram
	Google Ads □
Why social media will be	E.g. clinical recruitment is limited by xxx and we need to
used	reach a wider audience because of xxx
Clinical trial dates	[Insert dates]
Cililical trial dates	[moert dates]
Period ads will be live	[Insert dates]
/recruitment phase	[]
What assets/text will be	E.g. Image on tile/text on tile/post copy/ad copy
used (if available)	
Budget	Can include a breakdown of spend per platform
Mn	
Who is paying for the advertisement?	Include the funder of the post
advertisement?	
What measures are in	E.g. turning off ability to respond to the post of outline
place to protect the	monitoring
privacy of those that	E.g. including a warning such as "Depending on your
respond to the post?	Facebook privacy settings, posts that you follow, like or
	comment on may be able to be seen by your Facebook
	Friends or others".

Social media requirements: Planning template



Directions for researchers: please amend the blue sections below.

Appendix:

Social media will be used to recruit participants. Social media is going to be used because [e.g it will allow for recruitment of a wider audience that we are unable to recruitment from the hospital alone]. We intend to use [Twitter, LinkedIn, Facebook, Instagram, Google Ads etc] to advertise to [insert target audience e.g. adults 18 years and older]. Advertising is expected to last for [insert duration e.g. 6 months] and begin [specific date or as soon as all necessary approvals have been provided]. Please see below for an outline of the planned advertisements.

Advertisement 1:

Link	Insert URL to recruitment page
Key message	What is the one thing people need to know about this
	clinical trial?
Headline	25 characters max
Description including any	100 characters max
images	

Advertisement 2:

Link	Insert URL to recruitment page
Key message	What is the one thing people need to know about this
, ,	clinical trial?
Headline	25 characters max
Description including any	100 characters max
images	

Non-Interventional Protocol Description



Project Title		
Version Date	DD/MM/YYYY	
This document is a protocol for a research project. This study will be conducted in compliance with the NHMRC		
National Statement on ethical Conduct in Human Research (2007), the Note for Guidance on Good Clinical		
Practice (CPMP/ICH-135/95) a	nd any stipulations as outlined by the reviewing Human Research Ethics Committee.	
Project Ethics Number	HREC/XXXXX/Austin-202X	
(Office Use Only)		

Section 1 - Administrative Information

1.1 - Project Classification

Please tick the correct classification for your project	☐ Intention to publish in scientific journal and any samples taken are part of standard of care. This is classified as research, therefore requiring approval from Ethics Committee or their delegate.
	☐ No intention to publish, part of Organisation's "Quality and Safety" continuous improvement processes and to be registered in the Projects and Improvements Database. This means you are exempt from Ethical Review but you cannot publish in a scientific forum. Register your QI project via Quality & Safety on the Projects and Improvements Database.

1.2- Site Specific Investigators

Name	Site Department	Role e.g. Associate Investigator	Email
		Principal Investigator	
		Associate	

Social media recruitment information: Protocol template

FAQ

Designed as a quick reference

Covers the basics

Includes a comparison table

Audience is researchers and other inquisitive people!



Social Media Advertising

The final social media post, including any

What is the

needed?

'advertisement'?

What approvals are

SOCIAL MEDIA FREQUENTLY ASKED QUESTIONS (FAQS)

s a trial is considered

advertising, examples include the paid advert for a specific study, or a post on the institution's social media web page.

How are social media recruitment strategies different to traditional advertising?

Social media and traditional advertising are actually very similar. However, because social media has a wider reach and people may be able to publicly comment on the advertisement/post, there are additional privacy concerns to consider.

What are these additional privacy concerns?

When adverts are noted on social media, people can leave comments and are identifiable. There need to

The printed advertising material (e.g.

Traditional Advertising

mments or 'liking' the ot collect personal,

video needs to be submitted to HREC and RGO for approval

HREC needs to approve the material

The institution's social media page needs to agree that it can be used (as appropriate)

Social media company guidelines need to be

What happens when the advertising material is updated?

Each update needs to be approved as above. Each update needs to be approved as above. be advertising material is updated?

Each update needs to be approved as above. Each update needs to be approved as above.

that where the material needs to be 'ad lib', tailored or adapted, "a description of the strategy and broad messages is sufficient"

(National Statement 3.1.20).

What are the specific Privacy concerns of the audience need to be privacy issues? Privacy communicated with HREC a social media site.

GUIDANCE PROCEDURE

More content than the FAQ

Audience is researchers

Can support an institution's SOPs/Policy



SOCIAL MEDIA GUIDANCE PROCEDURE FOR RESEARCHERS

Background

Social media has emerged as a widely used and powerful communication tool. It is therefore no surprise that it has become a major platform for advertising to the public, and most social media applications derive their income from advertising. Given its widespread usage and reach, it is increasingly being applied to reach potential participants to inform them about clinical research and to encourage participation.

From a regulatory perspective, use of social media as an advertising platform does not differ in any material way from the use of other media – that is, the content of any written or spoken material must abide by ethical and legal requirements. However, there are some additional privacy-related ethical and legal concerns raised due to the ability to engage with people in open forums where individuals may not understand the implications and reach of the personal information on their profiles.

When considering use of social media, it is important that researchers understand the advantages and limitations of these platforms to ensure they have considered and addressed the relevant ethical and research governance issues.

Purpose

To describe the specific compliance requirements for advertising research projects using social media.

Responsibilities

This document covers the responsibilities for researchers involved with using social media for advertising and patient recruitment.

Procedure

Ethical approval

NEXT STEPS

Website launch

Conferences

Abstracts

Journal paper...

Effective use of social media as a clinical trial recruitment tool



Authors: Duncan Colyer³. Peter Gibb³. Nik Zeps³. Heidi Gaulke⁴. Eleonora Kay³

Institutions: VCCC Alliance, Melbourne, VIC, Australia Walter and Eliza Hall Institute, Melbourne, VIC, Australia Monash Partners, Melbourne, VIC, Australia 'Austin Health, Melbourne, VIC, Australia

Project aims

The successful recruitment of patients is widely regarded as one of the most challenging aspects of conducting clinical trials. Poor participant recruitment is the most frequent reason for premature discontinuation of clinical trials which has a significant impact on conclusive results, confidence of trials staff, and ethical consideration for participants. Social media has emerged as a commonly used and powerful communication tool, however its use in reaching potential participants in clinical trials remains underrealised. The improve Participation project at the VCCC All rendeavoured to develop reliable guidance resources to support researchers to understand both the opportunities and the additional considerat in using social media for clinical trials recruitment, with a particular focus on its regulatory and ethical requirements.

Methodology

Advertising clinical trials is not new, from posters in waiting rooms to the sides of busses. The introduction of social media as a medium has not provided the solution to the recruitment problems noted above. Investigating this phenomenon, an initial scoping exercise was undertaken by the VCCC Alliance looking at current advice on the use

Guidance was noted to be:

- Outdated, often by a decade
- Focused on making social media 'effective' > Centred on a 'social media plan', without an example
- > Related to the policies and guidelines of an institution

In conclusion, the current guidance was failing to address the needs of researchers, and prevented social media

recruitment content that was either appropriate for potential participants or acceptable to Human Research Ethics Committees (HRECs) for approval.

Working with subject matter experts, the requirements of resources were developed. Alongside the needs for general guidance, it was acknowledged that differences between Sponsored and investigator initiated Trials (IITs), related to

Project output and conclusions

It is anticipated that such resources can assist researchers in developing suitable approach to social media recruitment and communicating these to appropriately to HRECs. An evalu of the resources has been established within the webpage.

























RIAL INNOVATIONS

HOME | OUR WORK | RESEARCH AND TRANSLATION | CLINICAL TRIAL INNOVATIONS | SOCIAL MEDIA RECRUITMENT

sator-Initiated Trials d Toolkit

y Based Trials

ig Consumer Awareness

Media Recruitment

g Capability

cents and Young Adults

t/Feedback

al Innovations a.kay@unimelb.edu.au

edia guidance procedure for

edia frequently asked

edia requirements: Planning

edia recruitment information:

edia appendix requirements

Social Media Recruitment

Social media has emerged as a widely used and powerful communication tool. It is therefore no surprise that it has become a major platform for advertising to the public. Given its widespread usage and reach, it is increasingly being applied to reach potential participants to inform them about clinical research and to encourage participation.

This is a new resource. We'd love to hear how you're finding it:

To assist researchers and highlighting the specific compliance requirements, the VCCC Alliance, with input from The Royal Children's Hospital, Murdoch Childrens' Research Institute, and Austin Health have collaborated on a quick-reference page, resources, and answers to Frequently Asked Questions

In this section you'll find:

- What do I need to know about social media advertising first?

How to proceed

What additional considerations do I need to think about?

Resources

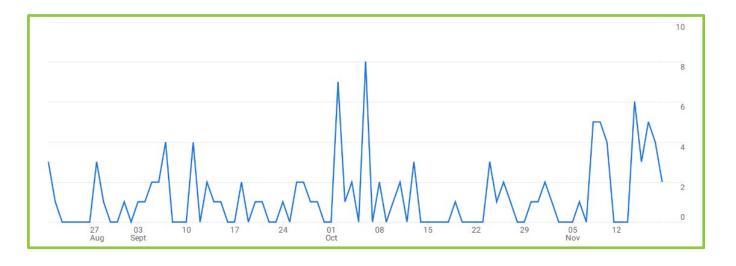
For a procedural overview, please download the below Guidance Document.

Background Information

What is considered advertising on social media?

low are social media recruitment strategies different to traditional advertising

SO FAR...



Average session duration	Views per user	Users	↓ Views
2m 08s	1.51	74	112
Avg +<0.01%	Avg 0%	100% of total	100% of total
2m 08s	1.51	74	112
	2m 08s Avg +<0.01%	1.51 2m 08s Avg 0% Avg +<0.01%	74 1.51 2m 08s 100% of total Avg 0% Avg +<0.01%

Evaluate the resources!

Happy to work with Research Offices too!

	What kind of study are you involved with that you are seeking information on social media recruitment?
	Phase 1
	Phase 2
	Phase 3
	Non-treatment intervention trial
	Other /comments:
2.	Have you ever used social media as a recruitment tool for a clinical trial?
	Yes
	No
3.	Did you consider your social media strategy before or after contacting HREC?
	Before
	After
4.	Did the information on the web page provide you with new insights or considerations for using social media as a recruitment tool for a clinical trial
	Yes
	No
5.	On a scale of 1 to 5, how informative did you find the information on the web page?
	1 - Not informative at all
	2 - Somewhat informative
	3 - Neutral
	4 - Informative
	5 - Extremely informative
6.	How likely are you to recommend this web page to others involved in clinical trial recruitment?
	1 - Not likely at all
	2 - Somewhat likely
	3 - Neutral
	4 - Likely
	5 - Very likely
7.	Any Comments: (Free text response)

ANY QUESTIONS?



More information here:

