# **Research Development Workshop**

By Boon-How Chew



This training on the key points and fundamentals of research is for academicians, clinicians and health sciences professionals who want to 'go-now' on learning and conducting a proper and high-quality clinical, biomedical and health sciences research. It is on the whole research process and related areas, see attached topics on the table on the next page. The suggested workshop fee is MYR100 for HPUPM/FPSK staff, MYR200 for UPM staff, MYR500 for non-UPM individuals, and USD500 for non-Malaysians. The fees from the HPUPM/FPSK/UPM staff will be waived as a reward upon satisfactory completion of the training.

#### **PLANNING**

- Reading materials: core and complementary lists (see further below)
- Diary of Progress
- Certificate of attendance or completion
- Workshop evaluation

#### **PREPARATION**

• **Preparatory Work**: Participants to decide on professional interest or areas to pursue within own specialty. Then write 500 words essay to introduce and to argue for the topic on the current challenges and the possible future direction in (own) professional practice.

#### <u>IMPLEMENTATION</u>

- **Duration**: 2 Days Workshops (5 slots of 3 hours) + 1 Day proposal presentation
- **Venue**: a hybrid of online and physical meetings
- **Secretariat**: pen or pencils, and colour papers, diary either as online or paper form. Print out the essay submitted by the participants earlier. This will initiate the research idea and topic during and from the workshop. Invite speakers from outside CRU. Organise pre-training meeting with all speakers. Conduct a survey on participants of their research experience:
  - Involvement in research as PI and co-investigator
  - Confidence in planning and conducting a research
  - Confidence in statistical analysis
  - Publication in journal as CA or co-author
  - Knowledge about journal publication process
  - Writing skills in Likert scale
- **Participant**: Laptop with applications and software essential for the research training. To build own research team for the proposed review article and research project. To hold at least ONE meeting each for the respective review and research before presentation of the study proposal.

#### <u>OUTPUT</u>

**Output 1**: Write a complete study proposal & make a presentation

**Output 2**: Write mini narrative or review article.

Output 3: Completed peer reviews for at least one case-report and two original reports.

#### RECOGNITION

**Certificate of Attendance**: Attended without any output **Certificate of Completion**: Attended with 2 of 3 outputs

Register your interest and post your comment/request: https://forms.gle/mnjrZmdfCgddnuRP8

## **Suggested Structure of the Training Module**

Hour	Talk   Topic	Tent. Speake
0800-0815	REGISTRATION	
0815-0830	Introduction: Quality healthcare, research, KPI & career advancement	CBH & TDPA
0830-0845	Testimony I : Personal sharing by an outstanding researcher	TBD
0845-0915	Interactive talk 1: Understanding the whole research process	CBH
0915-1015	Interactive talk 2: Fundamental concepts of clinical epidemiology	CBH
1015-1030	Interactive talk 3: Classification of epidemiologic research	CBH
BREAK		
1045-1115	Interactive talk 4: An introduction to qualitative study & designs	Invited speaker
1115-1145	Interactive talk 5: Research question, literature review & conceptual framework	СВН
1145-1215	Interactive talk 6: An introduction to databases & search strategies	CBH & an invited speaker
1215-1245	Interactive talk 7: Theoretical design	CBH
1245-1315	Interactive talk 8: Data collection design	СВН
LUNCH		
1400-1430	Interactive talk 9: Sample size estimation	СВН
1430-1500	Interactive talk 10: Statistical design	СВН
1500-1515	Interactive talk 11: Summary: clinical epidemiology & research methodology	CBH
1515-1545	Interactive talk 12: Writing up a study proposal	СВН
1545-1615	Interactive talk 13: Ethics clearance for a clinical study	Invited speaker
1615-1645	Interactive talk 14: Funding opportunities	Invited speaker
BREAK & D.	ISMISS	
Day 2		
0800-0815	REGISTRATION	
0815-0915	Interactive talk 15: Statistical analysis	СВН
0915-1000	Interactive talk 16: Comprehensive reporting, quality writing	СВН
1000-1030	Interactive talk 17: Publication process	CBH
BREAK	Threfactive talk 17.1 ablication process	CDIT
1045-1245	Interactive talk 18: Sistem PRiMS (Putra Research & Innovation Management System) and UPM IP Putra Science Park	Invited speaker
LUNCH		
1400-1500	Interactive talk 19: What is evidence-based practice? Appraise the evidence: primary research and systematic reviews & meta-analysis	СВН
1500-1530	Interactive talk 20: Summary: a suggested roadmap for clinicians to higher quality in research and publication	СВН
1530-1545	Testimony II: Personal sharing by an outstanding researcher	TBD
1545-1630	Closure: Summary & What have you learned?  Q & A	СВН

Day 3 after 2-3 months post-workshop		Facilitator
0800-0815	REGISTRATION & Intro	
0830-1230	Study proposal presentation	СВН

### **Reading Materials**

#### **Core List**

- 1. Chew BH. Planning and Conducting Clinical Research: The Whole Process. *Cureus.* 2019 Feb 20;11(2):e4112. doi: 10.7759/cureus.4112
- 2. Boaz A, Hanney S, Jones T, Soper B: Does the engagement of clinicians and organisations in research improve healthcare performance: a three-stage review. *BMJ Open.* 2015, 5(12).
- 3. Ioannidis JPA: Why Most Published Research Findings Are False. PLoS Med. 2005, 2(8):e124
- 4. Ioannidis JP. How to Make More Published Research True. PLoS Med. 11(10): e1001747
- 5. Macleod MR, Michie S, Roberts I, Dirnagl U, Chalmers I, Ioannidis JP, Al-Shahi Salman R, Chan AW, Glasziou P: Biomedical research: increasing value, reducing waste. *Lancet.* 2014, 383(9912):101-104.
- 6. Chalmers I, Bracken MB, Djulbegovic B, Garattini S, Grant J, Gulmezoglu AM, Howells DW, Ioannidis JP, Oliver S: How to increase value and reduce waste when research priorities are set. *Lancet*. 2014, 383(9912):156-165.
- 7. Munn Z, Stern C, Aromataris E, Lockwood C, Jordan Z: What kind of systematic review should I conduct? A proposed typology and guidance for systematic reviewers in the medical and health sciences. *BMC Med Res Methodol.* 2018, 18(1):5.
- 8. Clarke M: Doing new research? Don't forget the old. *PLoS Med.* 2004, 1(2):e35.
- 9. Roberts I, Ker K: How systematic reviews cause research waste. Lancet. 2015, 386(10003):1536.
- 10. Pautasso M: Ten Simple Rules for Writing a Literature Review. PLoS Comput Biol. 2013, 9(7).
- 11. Bordage G. Conceptual frameworks to illuminate and magnify. *Med Educ.* 2009 Apr;43(4):312-9. doi: 10.1111/j.1365-2923.2009.03295.x.
- 12. Bordage G, Lineberry M, Yudkowsky R. Conceptual Frameworks to Guide Research and Development (R&D) in Health Professions Education. *Acad Med.* 2016 Dec;91(12):e2. doi: 10.1097/ACM.00000000001409.
- 13. Vandenbroucke JP, Pearce N: From ideas to studies: how to get ideas and sharpen them into research questions. *Clin Epidemiol.* 2018, 10:253-264.
- 14. Dine CJ, Shea JA, Kogan JR. Generating Good Research Questions in Health Professions Education. *Acad Med.* 2016 Dec;91(12):e8. doi: 10.1097/ACM.000000000001413.
- 15. Meyer H, Varpio L, Gruppen L, Sandhu G. The Ethics and Etiquette of Research Collaboration. *Acad Med.* 2016 Dec;91(12):e13. doi: 10.1097/ACM.000000000001439.
- 16. Emanuel EJ, Wendler D, Grady C: What makes clinical research ethical? Jama. 2000, 283(20):2701-2711.
- 17. Horner J, Minifie FD: Research ethics III: Publication practices and authorship, conflicts of interest, and research misconduct. *J Speech Lang Hear Res.* 2011, 54(1):S346-362.
- 18. Guyatt G, Jaeschke R, Heddle N, Cook D, Shannon H, Walter S: Basic statistics for clinicians: 1. Hypothesis testing. *CMAJ.* 1995, 152(1):27-32.
- 19. Guyatt G, Jaeschke R, Heddle N, Cook D, Shannon H, Walter S: Basic statistics for clinicians: 2. Interpreting study results: confidence intervals. *CMAJ.* 1995, 152(2):169-173.
- 20. Jaeschke R, Guyatt G, Shannon H, Walter S, Cook D, Heddle N: Basic statistics for clinicians: 3. Assessing the effects of treatment: measures of association. *CMAJ*. 1995, 152(3):351-357.
- 21. Guyatt G, Walter S, Shannon H, Cook D, Jaeschke R, Heddle N: Basic statistics for clinicians: 4. Correlation and regression. *CMAJ.* 1995, 152(4):497-504.
- 22. Katz MH: Multivariable analysis: a primer for readers of medical research. Ann Intern Med 2003, 138(8):644-650.
- 23. Peduzzi P, Concato J, Kemper E, Holford TR, Feinstein AR: A simulation study of the number of events per variable in logistic regression analysis. *J Clin Epidemiol.* 1996, 49(12):1373-1379.
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- 26. Boynton PM, Greenhalgh T: Selecting, designing, and developing your questionnaire. *BMJ.* 2004, 328(7451):1312-1315.
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- 28. Lingard L, Watling C. It's a Story, Not a Study: Writing an Effective Research Paper. *Acad Med.* 2016 Dec;91(12):e12. doi: 10.1097/ACM.000000000001389
- 29. Meyer HS, Carline J, Durning SJ. Ten Tips to Move From "Revisions Needed" to Resubmission. Acad Med. 2016 Dec;91(12):e15. doi: 10.1097/ACM.000000000001391
- 30. Research Toolkits & Resources on HPUPM website: <a href="https://hpupm.upm.edu.my/research">https://hpupm.upm.edu.my/research</a> hpupm/research toolkits resources-12873

#### **Complementary List**

- 1. Altman DG. The scandal of poor medical research. BMJ. 1994 Jan 29;308(6924):283-4
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- 6. Chew BH: Understanding and conducting clinical research a clinical epidemiology approach by a clinician for clinicians: Serdang UPM, Malaysia; 2019
- 7. Moher D, Altman DG. Four Proposals to Help Improve the Medical Research Literature. *PLoS Med.* 2015 Sep 22;12(9):e1001864. doi: 10.1371/journal.pmed.1001864. eCollection 2015 Sep
- 8. Kleinert S, Horton R. How should medical science change? Lancet. 2014 Jan 18;383(9913):197-8.
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- 10. Goodman SN, Fanelli D, Ioannidis JP: What does research reproducibility mean? *Science Translational Medicine*. 2016, 8(341):341ps12.
- 11. Florey CD: Sample size for beginners. *BMJ.* 1993, 306(6886):1181-1184.
- 12. Campbell MJ, Julious SA, Altman DG: Estimating sample sizes for binary, ordered categorical, and continuous outcomes in two group comparisons. *BMJ*. 1995, 311(7013):1145-1148.
- 13. Walters SJ: Consultants' forum: should post hoc sample size calculations be done? *Pharm Stat.* 2009, 8(2):163-169.
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- 15. Friedman LM, Furberg CD, DeMets D: Fundamentals of Clinical Trials: Springer; 2010.
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- 20. Tarnow-Mordi WO, Healy MJ. Distinguishing between "no evidence of effect" and "evidence of no effect" in randomised controlled trials and other comparisons. *Arch Dis Child*. 1999 Mar;80(3):210-1. https://adc.bmj.com/content/80/3/210
- 21. Meisel ZF, Gollust SE, Grande D: Translating Research for Health Policy Decisions: Is It Time for Researchers to Join Social Media? *Acad Med.* 2016, 91(10):1341-1343.
- 22. Moser A, Korstjens I. Series: Practical guidance to qualitative research. Part 1: Introduction. *Eur J Gen Pract*. 2017 Dec;23(1):271-273. doi: 10.1080/13814788.2017.1375093
- 23. Korstjens I, Moser A. Series: Practical guidance to qualitative research. Part 2: Context, research questions and designs. *Eur J Gen Pract*. 2017;23(1):274–279. doi:10.1080/13814788.2017.1375090
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