SMC IRB Approval Request Form

IRB Approval

1. Investigator Name *
2. Position Title (if applicable)
3. Department (if applicable)
4. Organization (Institution or Company Name) *
Show/hide trigger exists. 5. Are you an SMC employee? Yes No

Hidden unless: #5 Question "Are you an SMC employee?" is one of the following answers ("No") 6. Are you collaborating with a staff member, program, or department at SMC? Yes (Name person, program, dept) No
7. Email *
8. Contact Phone Number *
9. Today's date *
10. If applicable, provide your co-PI name(s):
IRB Approval from Other Organization 11. Project Title *

12. Have you received any IRB approval from another organization? *
O Yes
O No
Hidden unless: #12 Question "Have you received any IRB approval from another organization?" is one of the following answers ("No") 13. Does your study falls in one of the exempt categories in the Common Rule? (Check ALL that apply). Click here for more information on Exempt Categories. *
☐ Education research
 Surveys, interviews, educational tests, public observations (that do not involve children)
☐ Benign behavioral interventions
☐ Analysis of previously-collected, identifiable info/specimens
☐ Federal research/demonstration projects
☐ Taste and food evaluation studies
□ None of the above
Acceptance Review
Page entry logic: This page will show when: #12 Question "Have you received any IRB approval from another organization?" is one of the following answers ("Yes")
14. Please upload and submit a copy of the IRB application you submitted to your original organization/institution. * Browse

15. Please upload and submit documentation indicating that your IRB application at the other organization/institution has been approved. *
Browse
Exempt for Further Review
Page entry logic: This page will show when: #13 Question "Does your study falls in one of the exempt categories in the Common Rule? (Check ALL that apply). Click here for more information on Exempt Categories. " is one of the following answers ("Education research", "Surveys, interviews, educational tests, public observations (that do not involve children)", "Benign behavioral interventions", "Analysis of previously-collected, identifiable info/specimens", "Federal research/demonstration projects", "Taste and food evaluation studies")
40 D
16. Provide a brief narrative of the proposed study, including purpose.*
17. Describe the study population.*

18. Will all study participants be 18 years of age or older? C Yes No
19. If applicable, include information on your plan for recruiting participants and sampling procedures. *
20. If applicable, describe the design, the methods, procedures, and tools that will be used to collect data. *
21. Describe your plan to ensure privacy and confidentiality of study participants including the ways identifiable information will be collected, stored, and shared. *

22. How will you ensure the confidentiality of Santa Monica Collège?				
23. Please confirm that, at a minimum, the followand enforced:	wing m	neasur	es will be taken	
	Yes	No	Does not apply	

nd enforced:					
	Yes	No	Does not apply		
Information maintained physically will be stored with appropriate physical safeguards, such as in locked cabinets and/or in restricted areas limited to authorized study personnel.	o	О	O		
Electronic data will be stored with appropriate electronic safeguards, such as unique usernames/passwords, and limited to authorized study personnel.	0	O	С		
Copying and use of study related materials will be restricted.	С	0	O		
Audio and/or recordings will be transcribed and then will be destroyed.	C	0	О		
Audio and/or recordings and/or photos or images will be modified to eliminate the possibility that study participants could be identified.	0	0	С		
Direct identifiers and/or the key to the codes will be destroyed upon completion of the research (all data will be stripped of identifying information and/or the key to codes destroyed, paper documents shredded, electronic files purged, electronic media securely erased)	0	0	O		
Study data and records (signed consent forms, etc.) will be retained for at least the minimum amount of time required by federal regulations and policies (minimum three years)	o	О	С		

24. How will the findings of your study be used (for example, dissertation, publish in journal, present at conference)? *
25. Please upload and submit a copy of the informed consent to be used with human subjects at Santa Monica College. * Browse
26. Please upload and submit a copy of the instruments to be used to collect
data from human subjects at Santa Monica College. If you are administering more than one instrument in your study, please attach them in a single document. *
Browse
27. Please upload and submit a copy of the invitation to participate in research (if applicable).
Browse
Minimal Risk
Dago ontry logica
Page entry logic: This page will show when: #13 Question "Does your study falls in one of the exempt categories in the Common Rule? (Check ALL that apply). Click here for more information on Exempt Categories. " is one of the following answers ("None of the above")

28. Minimal risk : the probability and magnitude of harm or discomfort anticipated in the research are not greater than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or ; tests or involves administration or use of drugs or devices.
Does your study involves no more than minimal risk?
C Yes
O No
Expedited Review
Page entry logic: This page will show when: #28 Question "Minimal risk: the probability and magnitude of harm or discomfort anticipated in the research are not greater than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations or; tests or involves administration or use of drugs or devices. Does your study involves no more than minimal risk?" is one of the following answers ("Yes")
29. Provide a brief narrative of the proposed study, including purpose.*

30. Describe the study population. *
31. Will all study participants be 18 years of age or older?
Yes
O No
32. If applicable, include information on your plan for recruiting participants and
sampling procedures. *
33. Describe the design, the methods, procedures, and tools that will be used to collect data. *

34. Describe your plan to ensure privacy and confidentiality of study participants including the ways identifiable information will be collected, stored, and shared. *
35. How will you ensure the confidentiality of Santa Monica Collège?

36.	Please confirm that,	at a minimum,	the following	measures	will be	taken
anc	l enforced:					

	Yes	No	Does not apply
Information maintained physically will be stored with appropriate physical safeguards, such as in locked cabinets and/or in restricted areas limited to authorized study personnel.	0	O	О
Electronic data will be stored with appropriate electronic safeguards, such as unique usernames/passwords, and limited to authorized study personnel.	0	O	С
Copying and use of study related materials will be restricted.	О	0	О
Audio and/or recordings will be transcribed and then will be destroyed.	О	0	О
Audio and/or recordings and/or photos or images will be modified to eliminate the possibility that study participants could be identified.	O	0	О
Direct identifiers and/or the key to the codes will be destroyed upon completion of the research (all data will be stripped of identifying information and/or the key to codes destroyed, paper documents shredded, electronic files purged, electronic media securely erased)	0	O	O
Study data and records (signed consent forms, etc.) will be retained for at least the minimum amount of time required by federal regulations and policies (minimum three years)	0	O	О

37. How will the findin	gs of your study be used	d (for example,	dissertation,
publish in journal, pres	sent at conference)? *		

38. Please upload and submit a copy of the informed consent to be used with
human subjects at Santa Monica College. *

Browse...

39. Please upload and submit a copy of the instruments to be used to collect data from human subjects at Santa Monica College. If you are administering more than one instrument in your study, please attach them in a single document. *

Browse...

40. Please upload and submit a copy of the invitation to participate in research (if applicable).

Browse...

Full IRB

Page entry logic:

This page will show when: #28 Question "**Minimal risk**: the probability and magnitude of harm or discomfort anticipated in the research are not greater than those ordinarily encountered in daily life or during the performance of routine physical or psychological examinations **or**; tests or involves administration or use of drugs or devices.

Does your study involves no more than minimal risk?" is one of the following answers ("No")

41. Provide a brief narrative of the proposed study, including purpose.*
42. Describe the study population. *
43. Will all study participants be 18 years of age or older?YesNo
44. If applicable, include information on your plan for recruiting participants and sampling procedures. *

45. Describe the design, the methods, procedures, and tools that will be used to collect data. *
46. Describe your plan to ensure privacy and confidentiality of study participants including the ways identifiable information will be collected, stored, and shared. *
47. How will you ensure the confidentiality of Santa Monica Collège?

48. Please confirm that, at a minimum,	the following	measures	will be	taken
and enforced:				

	Yes	No	Does not apply
Information maintained physically will be stored with appropriate physical safeguards, such as in locked cabinets and/or in restricted areas limited to authorized study personnel.	0	O	О
Electronic data will be stored with appropriate electronic safeguards, such as unique usernames/passwords, and limited to authorized study personnel.	0	O	О
Copying and use of study related materials will be restricted.	O	O	О
Audio and/or recordings will be transcribed and then will be destroyed.	С	0	O
Audio and/or recordings and/or photos or images will be modified to eliminate the possibility that study participants could be identified.	O	0	O
Direct identifiers and/or the key to the codes will be destroyed upon completion of the research (all data will be stripped of identifying information and/or the key to codes destroyed, paper documents shredded, electronic files purged, electronic media securely erased)	O	O	0
Study data and records (signed consent forms, etc.) will be retained for at least the minimum amount of time required by federal regulations and policies (minimum three years)	0	O	O

49. How will the findings of your study be used publish in journal, present at conference)? *	d (for example, dissertation,

50. Please upload and submit a copy of the informed consent to be used with human subjects at Santa Monica College. *

Browse...

51. Please upload and submit a copy of the instruments to be used to collect data from human subjects at Santa Monica College. If you are administering more than one instrument in your study, please attach them in a single document. *

Browse...

52. Please upload and submit a copy of the invitation to participate in research (if applicable).

Browse...

Thank You!

The Santa Monica College IRB will review your request and get in contact with you shortly. Most reviews take approximately two weeks.

Approval to conduct original research at SMC does not imply endorsement of the research by the Institutional Review Board or Santa Monica College.

If you have any questions, please contact the IRB Co-Chairs, Dr. Hannah Lawler (lawler_hannah@smc.edu) or Dr. Luis Andrade (andrade_luis@smc.edu).