

**Salish Kootenai College
Institutional Review Board (IRB)**

**Application for Continuing Review
Protocol Final Report Form**

Check Applicable Box:

IRB APPLICATION for CONTINUING REVIEW The Institutional Review Board (IRB) is required by Title 21, Code of Federal Regulations (Part 56.109) and Title 45, Code of Federal Regulations (Part 46.109) to conduct continuing review of ongoing projects not less than once per year.	
PROTOCOL FINAL REPORT The purpose of this form is to make a final report of conclusions of your study.	

Section I

IRB#		Title of Protocol	
Principal Investigator		Secondary Investigator	
Contact Phone		Contact Phone	
Mailing Address		Mailing address	
Email address		Email Address	

Project Personnel List: Only complete if project is continuing. Please list the names of all people working on this project. This would include the principal and secondary investigators, research assistants, graduate students and other people who have contact with human participants or have access to any identifiable data. Please include the person's role in the project (investigator, assistant, etc). All people listed will have to complete human subjects and cultural property protection training as specified in the Salish Kootenai College IRB policies.

Name of Individual:	Role in the Project:

Status of Study. Mark the status of the study (1-5).

1.	No subjects recruited for study, therefore termination requested.
2.	Inactive with no subjects recruited for study to date, study will become active.
3.	Active with ongoing recruitment of subjects.
4.	Active with subject recruitment completed.
5.	Completed (data collection & follow-up complete).

Section 2.

When did the study actually begin?		
What is the estimated completion date for the study?		
How many subjects have completed the study?		
Will new subjects be enrolled in the study?	Yes	No
Did any subjects voluntarily withdraw from the study?	Yes	No
Provide any known reasons for which subjects withdrew from the study:		
Were there any non-medical problems or complications in the that affected the subject or others?	Yes	No
If yes, a description of any problems or complications must be provided.		
Did any subject suffer an unanticipated problem or adverse event which was reported to the IRB since the last IRB review?	Yes	No
If yes, specify the number of reported events and describe briefly their nature and significance.		
Are there any other expected changes in data collection, data analysis, or reporting procedures?	Yes	No
If yes, please describe the changes.		
What is the funding source for this project? What is the anticipated length of funding?		

Section III.

Cont_rev_final_report
 C:\Documents and Settings\nursas.NUR59-105-1\My Documents\irb
 website\IRB\IRB website\continuing_review.doc

Provide a brief summary of any results (preliminary or final) obtained in the study. If the project is still active and no results are appropriate to report to the IRB at this time, this should be stated and explained.

--

Has anything occurred during the conduct of the study that may have altered the risk/benefit relationship? If the answer is yes, provide a current assessment of the risk/benefit relationship of the research based upon the results, adverse events and other factors.

--

Section IV.

If you are requesting continuation of your study, your signature indicates agreement that the study will be performed in accordance with the Salish Kootenai College Institutional Review Board policies. If the study is completed the information provided on this form reports the final status of the study.

<hr/> <p>Signature of Principal Investigator</p>	<hr/> <p>Date</p>
--	-------------------