## **Research Project Summary**

This form must be completed if a study;

(Ver1.0. 20210518)

2. Has been previously approved by the BRE the COVID-19 Pandemic.		rch durinş
Contact Name	Date	
E-mail	Phone	
SECTON (1) STUDY INFORMATION		
1. Study Title		
2. REB#		
3. List Bruyère PI, Investigators & Co-Investigators		
4. Study Status		
5. Briefly describe the study design		
6. Briefly describe the study 's primary objective.		
7. Why is this study important and how will it impact the future	e care of patients at Bruyère.	

8. Briefly describe the study main inclusion and exclusion criteria.

SECTION (2) PARTICIPANTS

9. Where will the research occur (e.g. EBH - Palliative Care, SVH - 5N etc) ?		
11. How many participants will be recruited and over what period of time?		
12. What are the activities and tin	ne commitments required of the participants and how will this fit into the unit workflow?	
13. Are unit staff being asked to help?	13a. If yes, describe what unit staff are being asked to do, what the time commitment will be, when will this be done (during work hours, after hours) and describe any compensation the clinical unit will receive.	
Yes		
No		
N/A		
SECTION (3) COVID AF	PROVAL	
14. Has the research project received IPAC/PPE approval?	14a. If no, please briefly explain why.	
Yes		
No		
15. Does the research project follow the most current version of the Procedure Manual on Conducting Faceto-face research?	15a. If no, please briefly explain why.	
Yes		
No		
Departmental Sign-off (IMPORTANT: please note that sign-off on this form is only required if the study already has REB approval and is seeking to resume research during the COVID pandemic.) All new studies must obtain Departmental Sign-off on Section 25 of the Research Ethics Application.		
Not Applicable		

Date

Name & Title

Signature