Request For Use of Aggregate Data Form

Date of Submission (mm/dd/yy):	12/15/2018
Name of Organization:	St.Mary's Medical Center Department of Radiation Oncology
Project Title:	A Retrospective Examination of Clinical Outcomes Following SBRT for Recurrent and Oligometastatic Gynecologic Cancers
Principal Investigator:	Sanjeev Sharma, M.D.
Co-Investigators:	Raj Singh, M.D.; Hayden Ansinelli, M.D.; John Austin Vargo, M.D.
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Project Description

Project Start Date (mm/dd/yy):	1/1/2019
Project End Date (mm/dd/yy):	6/1/2019
Type of Research Project:	□ Retrospective Analysis
If "Other," please explain nature of project:	
What is the research question being asked?	How effective is SBRT in treatment of recurrent and oligomestatic gynecologic malignancies? What are the associated treatment toxicities? What is associated local control, progression-free survival, and overall survival?
What is the background or rationale for the research question? (if needed, please attach as a separate page to application)	There are limited reports on the use of SBRT for recurrent and oligometastatic gynecologic cancers; this will add to the literature with regards to clinical outcomes.
Patient Inclusion/Exclusion Criteria:	Include: All patients in registry treated for recurrent and oligometastatic gynecololologic cancers with SBRT Exclude: Patients treated in the primary setting with SBRT.

Data Requested

Description of patient population to be analyzed:	All patients with recurrent or oligometastatic gynecologic malignancies treated with SBRT.
Time frame to be studied:	1/1/2019-6/1/2019
List exact data variables requested (i.e. pathology, treatment planning information, outcome, reimbursement, etc.): **If the request is not self-evident, write a summary of the request and/or instructions on data output (e.g., table specifications, sample tables)**.	1.Treatment planning (doses,fractions,etc) 2. Outcomes(overall survival,local control, KPS) 3.Toxicity 4. Patient demographics 5. Prior treatments 6. Lesion characteristics (i.e. GTV, histology, location) 7. Adjuvant or concurrent treatments (i.e. chemotherapy)
Deadline for receipt of data (mm/dd/yy):	1/1/2019

Data Use

Are these data for internal research purposes only? (yes/no)	No
If requesting party will seek to share data with persons not already listed on this request, list the organizations with which data would be shared and in what capacity? (e.g., FDA for a clinical trial, NIH for a grant proposal, consultant for project development)	N/A
Peer-reviewed publications to which submission is anticipated (if any)	Possibly Radiotherapy and Oncology
National meetings at which abstract presentation is anticipated (if any)	Possibly ACRO or RSS

Additional Submission Requirements

Please attach each of the following:

- Copy of IRB approval letter for use of RSSearch® at your institution
- Curriculum vitae of the principal investigator

Requestor Certification

In making this request, I certify that:

- All information provided on this form and attachments is accurate and complete;
- I have all requisite institutional authority to submit this Request for Use of Collaborative Data

Signature	
Print Name	Sanjeev Sharma
Title	Principal Investigator

Date (12/15/18)	

Please submit Request for Use of Collaborative Data to jjenkins@therss.org

For Internal Use Only:

Date application received:	12/15/2018
RSSearch Registry Request #	2018-1215