Appendix D: Access to Data and Aggregate Reporting

ReCKord Request Form for Use of Aggregate Data:

Eligibility Criteria for Participating Centers

 Enter > 50 patients into ReCKord annually (The Radiosurgery Society will not be held to this criterion)

Submission Requirements

Please attach each of the following with your Data Request Form:

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

General Information

Submitted Data Request Forms (DRF) are posted on the Radiosurgery Society homepage under the Clinician Resource dropdown. Applications are reviewed twice/year (April & September). You will receive a letter from the ReCKord Registry Review Committee regarding the status of your request within 30 days. Accepted DRFs will be forwarded to Adverteksm by the RSS to generate the requested custom data report. Custom reports will require a minimum of 2 weeks to generate. Submit DRF to Nalani Brown at nbrown@therss.org. Applications are posted on the Radiosurgery Society website at www.theradiosurgerysociety.org.

Administrative Information

Date of Submission (mm/dd/yy):	May 15, 2013		
Name of Organization:	The Radiosurgery Society		
Project Title:	The ReCKord Registry: Patterns of Care and Outcomes Research Through Multi-institutional User Database		
Principal Investigator:	Anand Mahadevan MD		
Co-Investigators:	Buddy Medbery MD, Joanne Davis PhD		
Corresponding Contact Name:	Joanne Davis		
Contact Title:	Consultant		
Contact Telephone Number:	248-719-2998		
Contact E-mail Address:	Davis.joanne@hotmail.com		
Contact Address: City, State, Zip:	40791 Kingsley Lane Novi, MI 48377		

Project Description

Project Start Date (mm/dd/yy):	February, 1 2013
Project End Date (mm/dd/yy):	June 1, 2013

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Type of Research Project:	Prospective Study		
	□ Retrospective Analysis		
	□ Technical Study		
	x Other		
If "Other," please explain nature of	The purpose of this project is to describe the role and use		
project:	of Multi-institutional prospective user databases to		
	perform outcomes research. We describe the design,		
	rationale, procedures for data collection, the clinical,		
	treatment and outcome data variables that are tracked in		
	The ReCKord Registry and present the patient		
	demographics, tumor/lesion characteristics, SRS/SBRT		
	treatment plan and delivery information of the first 11,457		
NAME of the state	patients enrolled in ReCKord database.		
What is the research question being	The objectives of the analysis are to describe the patient		
asked?	characteristics/demographics and lesion characteristics,		
	and to determine the treatment patterns of SRS/SBRT in a		
	clinical setting. The purpose of this project is to demonstrate that collective patterns of care and outcomes		
	_		
	research can be performed from data entered prospectively by users in a common database.		
What is the background or rationale			
for the research question? (if			
needed, please attach as a separate	technology and clinical outcomes from single institutions,		
page to application)	information on physician practice patterns of SRS and		
,	SBRT in the daily clinical practice is limited. A registry		
	provides a systematic and inclusive database of		
	information which can reveal and evaluate the		
	effectiveness of management practices. In contrast to a		
	clinical trial, where patient enrollment is defined by		
	specific inclusion and exclusion criteria, treatment is		
	dictated by protocol guidelines, and treatment evaluation		
	is measured at specific follow-up time intervals, a registry		
	documents actual care, representing a broad spectrum of		
	patients where treatments are not specified by protocol		
	guidelines and patient follow-up schedules are conducted		
	in a real-life setting. A registry can provide information		
	as to whether clinicians are adhering to practice		
	guidelines, may complement randomized clinical trials and/or identify new clinical applications and treatment		
	benefits. They would also generate powerful hypothesis to		
	test in future clinical trials to advance the science of		
	SRS/SBRT		
	The ReCKord Registry was conceptualized and		
	designed by a board of ReCKord Clinical Advisors in		
	2005. The ReCKord Clinical Advisory Committee is		
	comprised of radiation oncologists, neurosurgeons,		
	surgeons, medical oncologists, and medical physicists to		
	create and oversee the scientific conduct of the registry.		
	The goals and objectives of the registry are to provide a		
	method to collect standardized data on the use of		
	SRS/SBRT treatment practices and outcomes to help		
	determine the most effective clinical use of SRS/SBRT in		

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	management of patients with life threatening tumors and other diseases. The purpose of this request is to analyze the aggregate screening and treatment data of the first 11,457 patients enrolled in ReCKord. By analyzing the aggregate data we intend to publish the value of such registries and also explore specific areas (e.g lung cancer, prostate cancer) of outcomes research.
Patient Inclusion/Exclusion Criteria:	All patients entered into ReCKord between January 2005 and February, 2013 will be included in the analysis.
Data Requested	
Description of patient population to be analyzed:	All patients entered into ReCKord between January 2005 and February, 2013 will be included in the analysis.
Time frame to be studied:	January 2005 – February 2013
List exact data variables requested	All screening variables and treatment variables for all
(i.e. pathology, treatment planning	patients. Outcome and follow up data may also be
information, outcome,	explored for future projects.
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).	
Deadline for receipt of data	May, 2013
(mm/dd/yy):	
Data Use	
Are these data for internal research	No
purposes only? (yes/no)	
If requesting party will seek to	All data to be included and reviewed for this study is
share data with persons not already	anonymsed and HIPAA compliant. Other than the
listed on this request, list the	investigators listed, aggregate data may be shared with the
organizations with which data	scientific review committee of the Radiosurgery Society
would be shared and in what	for scrutiny, monitoring and approval. The data may be
capacity? (e.g., FDA for a clinical	made available for the IRB's of applicable investigators.
trial, NIH for a grant proposal,	
consultant for project development)	
Peer-reviewed publications to	To be determined once analysis is complete.

Additional Submission Requirements

presentation is anticipated (if any)

which submission is anticipated (if

Please attach each of the following:

• Copy of IRB approval letter for use of ReCKord at your institution

National meetings at which abstract | RSS Meeting 2014, ASTRO 2014

• 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;

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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)	anonymsed and HIPAA compliant. Other than the investigators listed, aggregate data may be shared with the scientific review committee of the Radiosurgery Society for scrutiny, monitoring and approval. The data may be made available for the	
Peer-reviewed publications to which submission is anticipated (if any)	To be determined once analysis is complete.	
National meetings at which abstract presentation is anticipated (if any)	RSS Meeting 2014, ASTRO 2014	

Additional Submission Requirements

Please attach each of the following:

- Copy of IRB approval letter for use of ReCKord 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	Lam Davis
Print Name	Joanne Davis, Phi
Title	Consultant
Date	5/25/13

Please submit Request for Use of Collaborative Data to nalani@theradiosurgerysociety.org

For Internal Use Only:

Date application received:	May 28, 2013	
ReCKord Registry Request #	2013-0528	

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