RSSearch® Registry Protocol

RSSearch Request Form for Use of Aggregate Data:

Eligibility Criteria for Participating Centers

 Enter > 50 patients into RSSearch annually (The RSS will not be held to this criterion)

Submission Requirements

Administrative Information

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

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

General Information

Submit the request for aggregate data to nbrown@therss.org. You will receive confirmation and a response regarding the status of your request within 30 business days. Requested custom aggregate data report will require a minimum of 2 weeks to generate. Accepted applications are posted on the RSS website at www.therss.org.

Date of Submission (mm/dd/yy): Name of Organization: Project Title: Principal Investigator: Co-Investigators: Corresponding Contact Name: Contact Title: Contact Telephone Number: Contact E-mail Address:

City, State, Zip:

Project Description

Project Start Date (mm/dd/yy):

Project End Date (mm/dd/yy):

Type of Research Project:

Prospective Study
Retrospective Analysis
Technical Study
Other

Contact Address:

If "Other," please explain nature of project:	
What is the research question being asked?	
What is the background or rationale	
for the research question? (if	
needed, please attach as a separate	
page to application)	
Patient Inclusion/Exclusion Criteria:	
Data Requested	
Description of patient population to	
be analyzed:	
Time frame to be studied:	
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).	
Deadline for receipt of data	
(mm/dd/yy):	
Data Use	
Are these data for internal research	
purposes only? (yes/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)	
Peer-reviewed publications to	
which submission is anticipated (if	
any)	
National meetings at which abstract	

Additional Submission Requirements

presentation is anticipated (if any)

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	
Title	
Date	

Please submit Request for Use of aggregate data to nbrown@therss.org

For Internal Use Only:

Date application received:	
RSSearch Registry Request #	