

## **INVESTIGATOR INITIATED TRIAL PROPOSAL REQUEST FORM**

For consideration for review by the Olympus Corporation of the Americas (OCA) IIT Committee, please complete this proposal request form with as much information as possible. Please attach a protocol synopsis, a current signed and dated CV, a copy of your medical license, and any other relevant material. Please include a CV and medical license for sub-investigators, if applicable. A completed, signed and dated Financial Disclosure Form will also be required. Please send this completed form and any accompanying documents to: ocaiitsupport@olympus.com

Full Study Title:						
1 Submitter Informati	on					
Name:						
Institution / Department:						
Email:						
Phone Number:						
2 Sponsor-Investigator / Study Site Information						
Name / Title:						
Email:						
Phone Number:						
Street Address, City, State, Zip	:					
Institution Affiliation:						
Institution Address, City, State	, Zip:					
Study Coordinator Name or	N/A:					
Study Coordinator Phone Num	ber:					
Sub-Investigator(s) or $\square$ N/A:						
How many years of experience in clinical research:						
How many publications in the l	ast two	o years:				
How many studies have you participated in during last 12 months:						
Have you ever worked with Olympus:		☐ Yes		No	If <b>YES</b> , what was name of the study and approximate date of participation:	



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## 3 Study Methodology and Design

Study Design: (check all that apply)	☐ Randomized ☐ Blinded ☐ Multicenter ☐ Controlled ☐ Non-Randomized ☐ Non-Blinded ☐ Single Center ☐ Observational					
Name of Institution where Study will be conducted:						
Institution Street Address, City, State, Zi	ip:					
If Multicenter Study, list other sites or $\square$	□ N/A:					
Study Objective / Hypothesis:						
OLYMPUS Product(s) to be used in the	study:					
What is the Study Duration (e.g., months) for a Prospective Study or □ N/A: (Includes IRB approval process, subject recruitment, enrollment and follow-up)						
What is the Study Duration for a Retrospective Study or □ N/A: (Includes IRB approval process and length of time required to review medical records, etc.)						
Primary Endpoint:						
Secondary Endpoint:						
Study Population (including inclusion and exclusion criteria):						
Anticipated Start Date (first patient in):						
Anticipated End Date (last patient out):						
Sample Size:						



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Number of Study Visits or □ N/A if Re	trospective Study:	
Type of Follow-Up Visits or □ N/A:		
Anticipated Output:	☐ Manuscript ☐ Abstract ☐ Poster ☐ Other:	☐ Podium Presentation
Schedule of Anticipated Output (e.g., pla	anned date for submission):	
4 Support Requested		
☐ Monetary Support Requested ( <i>Please</i>	include currency in fields held	ow and give hest estimate for hudge
Total Estimated Budget:		on unu gere eest estimate jer enuge.
Overhead %:		
List any Study-specific fees:		
What are you utilizing Olympus funds		
☐ In-Kind Support Requested		
Product Requested:		
Quantity:		
5 Sponsor-Investigator Sign	ature	
I hereby certify that the information prov of my knowledge; that this request for fur guaranteed and; that any amount awarded a written clinical study agreement.	nding is unsolicited and appro	val by OCA IIT Committee is not
Printed Name of Sponsor-Investigator	Date	
Sponsor-Investigator Signature		