

Application for Use of Radiation Producing Devices in Clinical Research

Complete this form and submit it to: 1) the Radiation Safety Committee with the iStar application, protocol and consent forms to cpickering@chla.usc.edu (and cc radsafetyoffice@chla.usc.edu) via e-mail.

Principal Investigator:	CHLA IRB Number:	
Position/Title	Date Submitted to IRB:	
CHLA Mailstop:	Attach Protocol	
Telephone Number:	Begin Date:	End Date:
Project Title		
Primary Purpose of the Proposed Radiologic Procedure		
<input type="checkbox"/>	Obtain Basic Research Information	
<input type="checkbox"/>	Benefit to Health of Participating Subjects	
<input type="checkbox"/>	Other (define)	
Brief description of the project, radiation requirements, and how the radiation procedure differs from standard of care.		
Type of Procedure	Location of Radiation Producing Device	
<input type="checkbox"/>	Radiographic (X-ray, portable, dental)	
<input type="checkbox"/>	CT studies	
<input type="checkbox"/>	Fluoroscopic (C-arm, Fluoroscanner)	
<input type="checkbox"/>	Nuclear medicine (PET, SPECT)	
<input type="checkbox"/>	DEXA	
<input type="checkbox"/>	Accelerator	
<input type="checkbox"/>	Other:	
Personnel		
Name of Device Operator(s):		
<input type="checkbox"/>	Board Certified Radiologist	
<input type="checkbox"/>	Certified Radiological Technologist	
<input type="checkbox"/>	Other: (attach qualifications)	

Personnel Required to Be in the Room While Radiation Device is On or Energized		
	Role/Duties:	
	Role/Duties:	
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	Role/Duties:	
Subject Radiation Dosimetry		
Total Effective Dose Equivalent (TEDE)		
Total CTDI for CT Scan Dose		
Total Entrance Skin Exposure per Procedure		
Primary Critical Organ		
Dose to Primary Critical Organ per Procedure		
Number of Procedures per Subject per Year		
Number of Years each Subject will be Receiving Radiation		
Number of Subjects per Study		
Human Subjects Information		
Number of Subjects With Manifest Disease:	Age Range:	Gender:
Describe pathology of subjects with manifest disease:		
Number of Subjects Without Manifest Disease:	Age Range:	Gender:
<input type="checkbox"/> Patients Will Be Hospitalized	<input type="checkbox"/> Normals Will Be Hospitalized	
Attachments		
Attach list of pertinent references including dosimetry references		
Attach copy of complete iStar application and the Informed Consent Form(s)		
Authorization		
Principal Investigator Signature:		Date: