

PUBLIC SUBMISSION

As of: 12/7/17 10:32 AM
 Received: November 27, 2017
 Status: Pending_Post
 Tracking No. 1k1-901f-ujq6
 Comments Due: January 08, 2018
 Submission Type: Web

Docket: NRC-2017-0215

Yttrium-90 Microsphere Brachytherapy Sources and Devices Therasphere and SIR-Spheres

Comment On: NRC-2017-0215-0001

Yttrium-90 Microsphere Brachytherapy Sources and Devices TheraSphere and SIR-Spheres;
 Draft Guidance for Comment

Document: NRC-2017-0215-DRAFT-0067

Comment on FR Doc # 2017-24129

11/7/2017
 82 FR 51655

Submitter Information

Name: Siddharth Padia

66

General Comment

I am a practicing Interventional Radiologist at the University of California at Los Angeles. To date, I have performed over 1000 Y90 radioembolization procedures, both at University of Washington and UCLA, since 2010. I am writing to oppose the proposed changes to the "Yttrium-90 Microsphere Brachytherapy Sources and Devices TheraSphere and SIR-Spheres Licensing Guidance" that would eliminate vendor involvement from the Interventional Radiology pathway to Authorized User (AU) status.

I trained to be an authorized user under Pathway 2, which has been in place since the concept evolved that Interventional Radiologists are the natural AU's of these devices. For these procedures, as an Interventional Radiologist, I:

- Perform required dosimetry to deliver appropriate activity to the patient
- Oversee and ensure appropriate Yttrium-90 (Y90) device handling and preparation
- Directly administer the therapy to the patient
- Coordinate the longitudinal care of patients following these procedures

Interventional Radiologists are the only Authorized Users capable of performing all four steps

SUNSI Review Complete

Template = ADM - 013

E-RIDS= ADM-03

Add= L. DUMICK (LDS)

above. The existing collaboration between physicians and industry helps ensure safe and comprehensive training in the use of Y90 devices.

In over 1000 infusions, I have not had a single safety issue with regards to radioembolization. I believe this level of safety was in part due to the rigorous training set forth in Pathway 2. The existing guidelines have been tremendously successful while maintaining impeccable safety. Manufacturer And User facility Device Experience (MAUDE) reports have remained 10/year for both devices since 2013. The majority of the MAUDE reports focus on procedural complication and treatment toxicities seen with all types of hepatic embolization, not specific to the Y90 devices. It seems statistically implausible that reduced vendor involvement will result in a measurable improvement in safety.

Proposed changes to the current arrangement, in which physicians and industry work closely together to ensure the appropriate training of interventional radiologists in the safe use of these devices will make it exceedingly difficult for Interventional Radiologists developing a clinical practice in radioembolization. Without the current direct training provided offsite by the device vendors, physician training will have to be solely performed by direct proctoring. Securing physician proctors is a challenge and can result in the delay in care, impacting cancer outcomes; physicians have limited time and availability away from their own clinical practices. Placing additional responsibilities on physician proctors may also have the untoward effect of limiting access to care, particularly for programs in underserved areas. The unanticipated consequence of the proposed changes is that training Interventional Radiologists in the safe and effective use of these devices will suffer greatly and patient access to care will diminish.

In summary, Interventional Radiologists deliver high quality minimally invasive care via imaging guidance, employing a variety of technologies. Training with other devices, such as aortic stent grafts, spinal augmentation devices, and atherectomy tools frequently involves a combination of vendor and physician collaboration. These relationships are a supplement to core training in hepatic embolization that is accomplished in fellowship. However, fine details regarding all devices may not be included in all programs. The existing NRC guidelines have facilitated training Interventional Radiologists in the safe and effective use of the Y90 devices, benefiting patients, physicians, and the government. There is no evidence of a need for change to the current NRC guidelines.

Regards,
Siddharth Padia, MD