



IMPORTANT REGULATORY REQUIREMENT

The IEC and FDA quality system regulations for our industry require documented evidence of procedures enabling Dunlee to find any item of our manufacture which could be subject to an FDA recall. As a Dunlee customer, we must solicit your help to document the location of every x-ray and CT tube sold.

We understand any concerns for customer confidentiality. Still, we are required to maintain a file that assures us these items can be located if necessary.

Please check the appropriate choice below and provide the information as indicated. Retain a copy for your files and mail or fax the original to Dunlee.

DUNLEE
555 North Commerce Street
Aurora, Illinois 60504
Fax No.: (630) 585-2125
Phone No.: (630) 585-2100

- Our company will retain permanent record of the location of Dunlee products that we sell/install/distribute and will provide such information to Dunlee in the event of a product recall.
- We prefer to forward installation records to Dunlee on all Dunlee products we sell. We will forward a copy of the Product Location report to Dunlee within 30 days of installation. (This form is included in the Rating Chart provided with each tube.)

Company	_____	Signed	_____
Address:	_____	Title	_____
	_____	Date	_____

Phone	_(____)_____		