

## Front matter

NRC Form XXX <b>U.S. Nuclear Regulatory Commission</b> For each person licensed under 10 CFR 32.11, 32.14, 32.18, 32.22, and 32.26.	Approved by OMB: NO. 3150-XXXX      Expires:    /    /
<b>REPORT OF TRANSFERS TO EXEMPT PERSONS</b>	Estimated burden per response to comply with this voluntary collection request: 20 minutes. This collection is used by NRC to track the distribution of byproduct material. Send comments regarding burden estimate to the Records and FOIA/Privacy Services Branch (T-5-F52), U.S. Nuclear Regulatory Commission, Washington DC 20555-0001, or by e-mail to <a href="mailto:infocollects@nrc.gov">infocollects@nrc.gov</a> .
<p><b>All licensees must comply with the following:</b></p> <ol style="list-style-type: none"> <li>1. All licensees shall file the report, covering the preceding calendar year, on or before January 31<sup>st</sup> of each year.</li> <li>2. Licensees who permanently discontinue activities authorized by the license shall file a report for the current calendar year within 30 days after ceasing distribution.</li> <li>3. If no transfers of byproduct material have been made under 10 CFR 32.11, 32.14, 32.18, 32.22, or 32.26 during the reporting period, the report must so indicate.</li> <li>4. The licensee shall maintain the record of a transfer for a period of one year after the transfer is included in a report to the Commission.</li> </ol>	

## BOX "A"

A. Licensee Information					
1. LICENSEE NAME AND ADDRESS:	2. LICENSE NUMBER:				
	3. DOCKET NUMBER:				
	4. REPORT PERIOD:	FROM:		TO:	
	5. EXEMPTION DESIGNATION — indicate under which NRC regulation the byproduct material is distributed — 10 CFR 30.14, 30.18, 30.19, 30.20, or the appropriate paragraph in 30.15:				

## Box "B"

B. If distributing under 10 CFR 30.14			
1. Quantity of byproduct material introduced into each product or material:	2. Name and address:	3. Type and quantity of radionuclide introduced into each product or material:	4. Initial concentration of the radionuclide:

**BOX "C"**

<b>C. If distributing under 10 CFR 30.15, 30.19, or 30.20</b>				
1. Description or identification of each type of product:	2. Model number, if applicable:	3. Radionuclide in each device:	4. Quantity of the radionuclide in each device:	5. Number of units distributed:

**Box "D"**

<b>D. If distributing under 10 CFR 30.18</b>		
1. Specify each radioisotope distributed under the specific license:	2. Specify each chemical and physical form for each radionuclide:	3. Indicate the total quantity transferred of each isotope:

**Box "E"**

<b>E. Certifying Official: I Certify that the foregoing is true and correct, complete and accurate in all material respects.</b>		
1. Printed Name and Title	2. Signature	3. Date