Dear Healthcare Professional,

The Device Identification Card belongs to the patient. Please follow the instructions below and provide the card to the patient after their implant procedure.

Instructions:
- Start by completing the Patient Surgery Information on the front of the Device Identification Card
- Remove the plastic Device Identification Card from this document
- Take the Patient Record Labels from the thermoform lid inside the packaging
- Place the Patient Record Labels from each product on the back of the plastic Device ID Card
  - TOP for product implanted in the LEFT breast
  - BOTTOM for product implanted in the RIGHT breast
- Please provide the Device Identification Card to your patient and advise her to keep it for her records.

If a Patient Record Label is not available, please use a ballpoint pen to record the product description, serial number, catalog number, and UDI information for each device in the space provided.

Important Information:
- Please retain this card for your records.
- Breast implants have a boxed warning. Please consult the patient information documents at www.allerganlabeling.com.
- The patient information documents may be updated periodically, please go to www.allerganlabeling.com for the most current information, including the patient decision checklist.
- To obtain more information or request paper copies of the patient information documents, contact Allergan at 1-800-678-1605.

Notes:

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