

Environmental, Health, and Safety (EHS) Guidance for Distributors and Resellers

Carestream Health values its relationship with its distributors and resellers. These relationships play an important role in extending the ability of customers to obtain Carestream Health Products throughout the world. The obligations in these relationships are defined in individual Distributor Agreements or Reseller Agreements. These Agreements along with supplemental information (e.g., Dealer Policy Guide) establish the framework of the relationship.

As Carestream Health Products (and related systems) are sold into a particular market, the distributor or reseller needs to comply with applicable laws and regulations as stated in the Terms and Conditions of individual Agreements. The following list provides guidance with regards to the types of environmental, health, or safety laws and regulations that may apply in a particular market or Territory. Such laws and regulations may include, but are not limited to,

- Compliance with FDA & regional requirements regarding registration for sales, service and demonstration of X-ray producing equipment and associated certified components.
- Compliance with federal and regional requirements for installing/servicing of X-ray producing equipment as specified in Federal regulations ([Section 21 CFR 1020.30\(d\)](#))
- Compliance with the importation & export of radiation producing equipment (X-ray & lasers) as specified by the CDRH Section 21 CFR 1002.1
- labeling of Products regarding hazard identification during storage, use, service, transportation, and proper disposal
- posting of precautionary signage in the vicinity where the Products are used or stored
- the availability of Material Safety Data Sheets or their equivalent in the local language
- Product Declarations of Conformity, or their equivalent in the local language
- the availability of safety related precautions defined as part of or in addition to the end-user instruction manuals in the local language
- training of local service personnel regarding the identification of and safeguarding against potential hazards
- regulatory notification of imported chemicals into the Territory (e.g. for Japan, Australia, Canada and European Union)
- tracking and reporting of Product volume imported into the Territory as required by the local governmental authorities (chemicals, packaging, electronic products, etc.)
- reporting, accrual, and remittance of any advanced disposal fees for products (requirements vary by region – check with local authorities for applicable product categories)
- collection, treatment, recovery and disposal of the supplied electrical and electronic equipment when it becomes waste (e.g., European Union WEEE Directive)
- reporting, accrual, and remittance of any advanced disposal fees for packaging (e.g., EU Packaging Directive, Polish Packaging and Packaging Waste Law, etc.)

NOTE: If a conflict or discrepancy exists between this EHS Guidance and an individual Distributor or Reseller Agreement, the provisions of the Carestream Health, Inc. Distributor or Carestream Health, Inc. Reseller Agreement will apply.