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For your reference, see the Guidance: [Registration and Product Listing for Owners and Operators of Domestic Tobacco Product Establishments](#).

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this information collection is 0910-0650 (expires 10/31/2015)

Please Note: The system will automatically time out if there is no activity for 30 minutes.

Section 905(i)(1) of the Federal Food, Drug, and Cosmetic Act requires that every person who registers with FDA also submit a list of all tobacco products which are being manufactured, prepared, compounded, or processed by that person for commercial distribution, along with certain accompanying information, including all labeling. In addition, section 905(i)(3) of the Act requires that certain changes in the product list be submitted biannually. Please update your product listings with these changes in June and December of each year as necessary, if you have not done so already.