Tobacco Registration and Listing Module (TRLM)

The FDA has published the revised guidance, Registration and Product Listing for Owners and Operators of Domestic Tobacco Product Establishments.

The agency revised the guidance because the FDA recognizes that product listing for some tobacco products may result in numerous labeling submissions that the manufacturer must prepare and submit via the FDA Unified Registration and Listing System (FURLS). To reduce the amount of labeling to upload, the agency currently does not, at this time, intend to enforce the requirement that owners and operators submit the labeling for each individual listed tobacco product when the registrant submits information that represents the labeling for a selected line of products. The updated guidance includes a new appendix that provides an example of how registrants can provide this information as a single submission.