Report of Tobacco Product Removals Subject to Tax for Tobacco Product User Fee Assessments

(Section 919 of FFDCA; Title 21, Code of Federal Regulations, Part 1150)

The authority for collecting the following information is section 301 et seq. of the Federal Food, Drug, and Cosmetic Act. The information will be used to assess and collect user fees from tobacco product domestic manufacturers and importers. Please be advised that under section 1001 of title 18 anyone who makes a materially false, fictitious or fraudulent statement is subject to criminal penalties.

Return this completed form and copies of supporting documents (TTB Forms 5220.6, 5210.5 and 5000.24 and Customs CBP Form 7501, as appropriate) by mail to:

Food and Drug Administration Center for Tobacco Products Document Control Center Attn: OM, Division of Financial Management, User Fee Team Building 71, Room G335 10903 New Hampshire Avenue Silver Spring, MD 20993-0002

Fax: 301-595-1429 or 301-595-1430

Email: TobaccoUserFees@fda.hhs.gov

1. Company Name/Address (including Zip Code) Alternate	Company Address (if any, for FDA notifications)

2. Contact Person Name

3. TTB Permit Number	4. Telephone Number (including Area Code)	
5. Employer Identification Number	6. Email	
7. Period of Activity	·	
A. Month:	B. Year:	

Privacy Act Notice:

This notice is provided pursuant to the Privacy Act of 1974, 5 U.S.C. 552a. The collection of this information is authorized by 21 U.S.C. 371, 379, 379e, 379h, 379h-1, 379j, 379j-12, 379j-21, 387s and 393(d)(2); 42 U.S.C. 263b(r)(1); 5 U.S.C. 301 and 552; and 42 U.S.C. 3101. FDA will use the information to assess, collect and process user fee payments and to facilitate debt collection under the Debt Collection Improvement Act. FDA may disclose information to courts and the Department of Justice in the context of litigation and requests for legal advice; to other Federal agencies in response to subpoenas issued by such agencies; to HHS and FDA employees and contractors to perform user fee services; to the National Archives and Records Administration and General Services Administration for records management inspections; to banks in order to process payment made by credit card; to Dun and Bradstreet to validate submitter contact information, and to other entities as permitted under the Debt Collection Improvement Act. Furnishing the requested information is mandatory. Failure to supply the information could prevent FDA from processing user fee payments. Additional detail regarding FDA's use of information is available online: http://www.fda.gov/RegulatoryInformation/FOI/PrivacyAct/default.htm, and http://www.fda.gov/AboutFDA/AboutThisWebsite/WebsitePolicies/default.htm

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Domestic Taxable Removals		Imports		
Volume (Number or Pounds)	Taxes (Dollars)	Volume (Number or Pounds)	Taxes (Dollars)	
TTB 5210.5 Monthly Manufacturer Report	TTB 5000.24 Excise Tax Return	Customs CBP 7501 Importer Entry Summary		
8. CIGARETTES				
A. Line 14 Columns C + D	B. Line 13 Column B	C. Box 31 - Monthly Total (Number of Cigarettes)	D. Box 38 - Monthly Total (Taxes on Cigarettes Only)	
9. CIGARS				
A. Line 14 Columns A + B	B. Line 12 Column B	C. Box 31 - Monthly Total (Number of Cigars)	D. Box 38 - Monthly Total (Taxes on Cigars Only)	
10. SNUFF				
A. Line 14 Column F	B. Line 15 Column B (Snuff Only)	C. Box 31 - Monthly Total (Pounds of Snuff)	D. Box 38 - Monthly Total (Taxes on Snuff Only)	
11. CHEWING TOBACCO				
A. Line 14 Column E	B. Line 15 Column B (Chewing Tobacco Only)	C. Box 31 - Monthly Total (Pounds of Chewing Tobacco)	D. Box 38 - Monthly Total (Taxes on Chewing Tobacco Only)	
12. PIPE TOBACCO				
A. Line 14 Column G	B. Line 16 Column B (Pipe Tobacco Only)	C. Box 31 - Monthly Total (Pounds of Pipe Tobacco)	D. Box 38 - Monthly Total (Taxes on Pipe Tobacco Only)	
13. ROLL-YOUR-OWN TOBACCO				
A. Line 14 Column H	B. Line 16 Column B (Roll-Your-Own Tobacco Only)	C. Box 31 - Monthly Total (Pounds of Roll-Your-Own Tobacco Only)	D. Box 38 - Monthly Total (Taxes on Roll-Your-Own Tobacco On	
14. CERTIFICATION				
I hereby certify that the information of	on this form is true and correct. and	d that I am hereby authorized to submit	this form on the company's behali	

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14C. Date Prepared (mm/dd/yyyy)

14B. Title

14A. Signature (Print and Sign)

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

The burden time for this collection of information is estimated to average 4 hours per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to the following address:

Department of Health and Human Services Food and Drug Administration Office of Operations Paperwork Reduction Act (PRA) Staff PRASTAFF@fda.hhs.gov

"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 number."