

**REGISTRATION AND LISTING FOR OWNERS
AND OPERATORS OF DOMESTIC TOBACCO
PRODUCT ESTABLISHMENTS**

Form Approved: OMB No. 0910-0650

Expiration Date: xx/xx/20xx

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FAMILY SMOKING PREVENTION AND TOBACCO CONTROL ACT

On June 22, 2009, the President signed the Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act) (Public Law 111-31) into law. The Tobacco Control Act amended the Federal Food, Drug, and Cosmetic Act (the act) by, among other things, adding a new chapter granting FDA important new authority to regulate the manufacture, marketing, and distribution of tobacco products to protect the public health generally and to reduce tobacco use by minors.

Complete the following question and answer form to register your establishment and submit your product listing to FDA's Center for Tobacco Products. For additional information on the legislation and guidance document, access the web links provided on page 11.

STATUTORY REQUIREMENTS

All owners and operators must fulfill the requirements for section 905 of the act, as detailed below. *In order to reduce redundant submissions, FDA strongly encourages owners to register and submit product listing information for themselves and on behalf of their operators.*

Section 905(b) of the act requires that “every person who owns or operates any establishment in any State engaged in the manufacture, preparation, compounding, or processing of a tobacco product or tobacco products shall register with the Secretary the name, places of business, and all such establishments of that person.”

Section 905(i)(1) of the act requires that all registrants “shall, at the time of registration ... file with [FDA] 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, such as all labeling.

Section 905(i)(3) of the act requires that certain changes in the product list be submitted biannually, once during June and once during December.

DEFINITIONS

FDA intends to use the following definitions in implementing the registration and product listing requirements of section 905 of the act.

- 1. Commercial Distribution:** The term “commercial distribution” includes any distribution of a tobacco product to consumers or to another person for future manufacturing through sale or otherwise. As examples, it includes the distribution of a tobacco product as a promotional sample and the delivery of a tobacco product to another manufacturer for further processing via contract without a change in the formal ownership of the product. Commercial distribution does not include internal or interplant transfer of a tobacco product between registered establishments within the same parent, subsidiary, and/or affiliate company and it does not include providing a tobacco product for product testing in cases where such products are not made available for consumption or resale.
- 2. Domestic Establishment:** The term “domestic establishment” means an establishment in any State or Territory or possession of the United States.

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3. **Establishment:** The term “establishment” means a place of business under one ownership at one general physical location. A single building may house more than one distinct establishment if the establishments are under separate ownership.
 4. **Labeling:** The term “labeling,” based on section 201(m) of the act (21 U.S.C. 321(m)), means all labels and other written, printed, or graphic matter (1) upon any tobacco product or any of its containers or wrappers, or (2) accompanying such tobacco product.
 5. **Manufacturing:** The term “manufacturing” means the manufacture, preparation, compounding, or processing of a tobacco product, including repackaging or otherwise changing the container, wrapper, or labeling of any tobacco product package (section 905(a)(1) of the act). This term includes the activities of reconstituting and blending tobacco leaf; testing for quality control and product release; and applying any chemical, additive, or substance to the tobacco leaf other than potable water in the form of steam or mist. This term excludes the activities of de-stemming, drying, or packing tobacco leaf; mechanically removing foreign material from tobacco leaves; and humidifying tobacco leaf with nothing other than potable water in the form of steam or mist. 6.
 6. **Operator:** The term “operator” means a person, as defined in section 201(e) of the act (21 U.S.C. 321(e)) who has management authority over an establishment.
 7. **Owner:** The term “owner” means a person, as defined in section 201(e) of the act (21 U.S.C. 321(e)) who has an ownership interest in an establishment.
 8. **Pouch:** The term “pouch” means a permeable pouch, intended to be filled with pre-portioned tobacco product and placed in the oral cavity with the tobacco product.
 9. **Tobacco Product:** The term “tobacco product” means “any product made or derived from tobacco, or containing nicotine from any source, that is intended for human consumption, including any component, part, or accessory of a tobacco product (except for raw materials other than tobacco used in manufacturing a component, part, or accessory of a tobacco product).” This term does not include an article that is a drug, a device, a combination product, or a food if such article contains no nicotine or no more than trace amounts of naturally occurring nicotine, as defined in the act (section 201(rr) of the FD&C Act (21 U.S.C. 321 (rr); as amended by section 111(a) of the Consolidated Appropriations Act, 2022 (Pub. L. 117-103)).
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**REGISTRATION AND LISTING FOR
OWNERS AND OPERATORS OF DOMESTIC DEEMED
TOBACCO PRODUCT ESTABLISHMENTS**

See page 14 for Instructions

Please type. An item followed by an asterisk () denotes a required field.*

Check one of the following.*
For updates, FDA requests
that you also provide your
FEI number or the DUNS
Number used in your
original submission.

☐ New Submission and
Product Listing (per
905(b) and 905(i)(1))

☐ Update to a Registration
(per 905(b)) (previously
submitted to FDA)

☐ Update to a Product List
(per 905(i)(3)) (previously
submitted to FDA)

Identification Number (if update)

SECTION I - IDENTIFICATION

Please check the appropriate boxes.* (Note that owners and operators may register on behalf of the other party.)

REGISTRATION STATUS

- 1 ☐ Owner registering alone (Complete all sections EXCEPT IIIA and IIIB)
- 2 ☐ Owner registering on behalf of operator (Complete all sections)
- 3 ☐ Owner who is also operator of all establishments (Complete all sections)
- 4 ☐ Operator registering alone (Complete all sections EXCEPT IIA and IIB)
- 5 ☐ Operator registering on behalf of owner (Complete all sections)

SECTION IIA - REGISTRATION

Owner Information

Owner Name (Name of the Corporation/Partnership or Individual Owner)*

Address*

City*

State, Province or Territory*

Country*

ZIP or Postal Code*

Owner Headquarters D&B DUNS Number:

Owner Point of Contact

Title (e.g., Mr., Ms., Dr.):

First/Given Name

Middle Name

Last Name

Position Title

Email Address

Telephone (Include Country Code if applicable)

FAX

SECTION IIB - REGISTRATION
Owner Business Structure

Select the type of business structure (Sole Proprietorship, Partnership, or Corporation) and provide indicated information.* (Continuation sheets may be used if necessary.)

☐ **Sole Proprietorship** (Enter owner name)

☐ **Partnership** (Enter name of each partner)

1.

2.

3.

4.

5.

6.

☐ **Corporation** (Enter the name of each corporate officer and director)

1.

2.

3.

4.

5.

6.

Identify State of incorporation.

Please describe further. (If applicable, give name of country if incorporation made outside U.S.)

If the owner does business by any other name, please list all such names.

1.

2.

3.

4.

SECTION IIIA - REGISTRATION
Operator Information

Multiple copies of this page may be submitted if you are registering on behalf of multiple operators.

Operator Name*

Address*

City*

State*

ZIP Code*

Operator D&B DUNS Number:

Operator Point of Contact

Title (e.g., Mr., Ms., Dr.):

First/Given Name

Middle Name

Last Name

Position Title

Email Address

Telephone (Include Country Code if applicable)

FAX

SECTION IIIB - REGISTRATION
Operator Business Structure

Select the type of business structure (Sole Proprietorship, Partnership, or Corporation) and provide indicated information.* (Continuation sheets may be used if necessary. Multiple copies of this page may be submitted if you are registering on behalf of multiple operators.)

☐ **Sole Proprietorship** (Enter operator name)

☐ **Partnership** (Enter name of each partner)

1.

2.

3.

4.

5.

6.

☐ **Corporation** (Enter the name of each corporate officer and director)

1.

2.

3.

4.

5.

6.

Identify State of incorporation.

Please describe further. (If applicable, give name of country if incorporation made outside U.S.)

If the operator does business by any other name, please list all such names.

1.

2.

3.

4.

SECTION IV - REGISTRATION
Establishment Information

Enter contact and registration information for each establishment being registered. (Multiple copies of this page may be submitted.)

Establishment Name*

Address*

City*

State*

ZIP Code*

Establishment D&B DUNS Number:

Is this establishment an Electronic Nicotine Delivery System (ENDS) Retail Establishment?

☐ Yes ☐ No

Operation (Check all that apply)

☐ Blending	☐ Packaging	☐ Storing
☐ Manufacturing	☐ Labeling	☐ Testing
☐ Reconstituting	☐ Saucing (or casing)	☐ Other (Specify): _____

Establishment Point of Contact

Title (e.g., Mr., Ms., Dr) :

First/Given Name

Middle Name

Last Name

Position Title

Email Address

Telephone (Include Country Code if applicable)

FAX

SECTION V – DEEMED TOBACCO PRODUCT LISTING

Sections V should be completed for each product listed. (Multiple copies of pages 6 through 8 may be submitted.)

1. Product Name* (i.e., brand/sub-brand or other commercial name used in commercial distribution – e.g., Acme E-Cig or Acme Cigar)

2. Product Identification Number (Must be provided if needed to uniquely identify the product)

3. Type of Product Identification Number (Check only one)

- ☐ Item/Catalog Number
☐ SKU Number
☐ UPC Number

4. Intended Use of Product (Check one)*

- ☐ Consumer Use (Go to question 5) ☐ Further Manufacturing Use (Skip to question 6)

5. Consumer Use Product Category (Check applicable, then skip to question 7)*

Cigar

- ☐ Cigar Tobacco
☐ Cigar
☐ Other (Specify below)

Cigar Component or Part

- ☐ Cigar Filter
☐ Cigar Paper
☐ Cigar Tip
☐ Cigar Tipping Paper
☐ Cigar Wrapper
☐ Other (Specify below)

Pipe Tobacco

- ☐ Pipe Tobacco
☐ Pipe Tobacco Kit
☐ Other (Specify below)

Waterpipe Tobacco

- ☐ Waterpipe Tobacco
☐ Other (Specify below)

Waterpipe Tobacco Component or Part

- ☐ Waterpipe Tobacco Base
☐ Waterpipe Tobacco Bowl
☐ Waterpipe Tobacco Cinder

- ☐ Waterpipe Tobacco Diffuser
☐ Waterpipe Tobacco Flavor Enhancer
☐ Waterpipe Tobacco Foil/Screen
☐ Waterpipe Tobacco Gasket
☐ Waterpipe Tobacco Grommet
☐ Waterpipe Tobacco Hose
☐ Waterpipe Tobacco Hose Cooling Attachment
☐ Waterpipe Tobacco Mouthpiece
☐ Waterpipe Tobacco Valve
☐ Waterpipe Tobacco Stem
☐ Waterpipe Tobacco Filtration Base Additives
☐ Other (Specify below)

Electronic Nicotine Delivery System

ENDS Open (Select from list)

ENDS Closed (Select from list)

Electronic Nicotine Delivery System Component or Part

- ☐ E-Liquid
☐ ENDS Adapter
☐ ENDS Atomizer

- ☐ ENDS Battery
☐ ENDS Bridge
☐ ENDS Cartomizer
☐ ENDS Cartridge
☐ ENDS Charger
☐ ENDS Clearomizer
☐ ENDS Coil
☐ ENDS Digital Display/Lights
☐ ENDS Drip Tip
☐ ENDS Drip Well
☐ ENDS Filler Material
☐ ENDS Filter
☐ ENDS Mouthpiece
☐ ENDS Software
☐ ENDS Tank
☐ Other (Specify below)

☐ Other (Specify below)

6. Further Manufacturing Use Product Category (Check applicable)*

Cigar

- ☐ Cigar Tobacco
☐ Cigar
☐ Other (Specify below) _____

Cigar Component or Part

- ☐ Cigar Filter
☐ Cigar Paper
☐ Cigar Tip
☐ Cigar Tipping Paper
☐ Cigar Wrapper
☐ Other (Specify below) _____

Pipe Tobacco

- ☐ Pipe Tobacco
☐ Pipe Tobacco Kit
☐ Other (Specify below) _____

Waterpipe Tobacco

- ☐ Waterpipe Tobacco
☐ Other (Specify below) _____

Waterpipe Tobacco Component or Part

- ☐ Waterpipe Tobacco Base
☐ Waterpipe Tobacco Bowl
☐ Waterpipe Tobacco Cinder

- ☐ Waterpipe Tobacco Diffuser
☐ Waterpipe Tobacco Flavor Enhancer
☐ Waterpipe Tobacco Foil/Screen
☐ Waterpipe Tobacco Gasket
☐ Waterpipe Tobacco Grommet
☐ Waterpipe Tobacco Hose
☐ Waterpipe Tobacco Hose Cooling Attachment
☐ Waterpipe Tobacco Mouthpiece
☐ Waterpipe Tobacco Valve
☐ Waterpipe Tobacco Stem
☐ Waterpipe Tobacco Filtration Base Additives
☐ Other (Specify below) _____

Electronic Nicotine Delivery System

ENDS Open (Select from list) _____

ENDS Closed (Select from list) _____

Electronic Nicotine Delivery System Component or Part

- ☐ E-Liquid
☐ ENDS Adapter
☐ ENDS Atomizer

- ☐ ENDS Battery
☐ ENDS Bridge
☐ ENDS Cartomizer
☐ ENDS Cartridge
☐ ENDS Charger
☐ ENDS Clearomizer
☐ ENDS Coil
☐ ENDS Digital Display/Lights
☐ ENDS Drip Tip
☐ ENDS Drip Well
☐ ENDS Filler Material
☐ ENDS Filter
☐ ENDS Mouthpiece
☐ ENDS Software
☐ ENDS Tank
☐ Other (Specify below) _____

☐ Other (Specify below) _____

7. Flavor (Check applicable)

- ☐ Menthol ☐ None
☐ Other (Specify): _____

8. If submission is an Update to a Product List (per 905(i) (3)) (previously submitted to FDA) (Make applicable entries)

If known, enter the FDA-assigned tracking number (e.g., TP#####) for your tobacco product.

If your product has been introduced to market, discontinued or reintroduced since your last product listing, indicate the most recent change.*

Provide the appropriate date:*

9. Advertising (A representative sampling of advertising may be required. Please see the guidance document, Section IV.C.2. for additional details. Representative samples, appropriately identified, are to be submitted with this form. For each advertisement, we request that you provide the following optional information below. You may use Appendix A as a continuation sheet if needed.)

9a. Type of Advertising Material (e.g., magazine ad)	9b. Title	9c. Unique ID or Internal ID Number	9d. Date First Disseminated (mm/dd/yyyy)

10. Labeling* (All labeling, appropriately identified, is to be submitted with this form. For each item of labeling, we request that you provide the following optional information below. You may use Appendix B as a continuation sheet if needed.)

10a. Universal Product Code(s) (UPC)

10b. Type of Labeling Material (e.g., package label)	10c. Title	10d. Unique ID or Internal ID Number	10e. Date First Disseminated (mm/dd/yyyy)

11. Consumer Information (Consumer information may be required. Please see the guidance document, Section IV.C.2. for additional details. All consumer information, appropriately identified, is to be submitted with this form. For each item, we request that you provide the following optional information below. You may use Appendix C as a continuation sheet if needed.)

11a. Type of Consumer Information (e.g., consumer brochure)	11b. Title	11c. Unique ID or Internal ID Number	11d. Date First Disseminated (mm/dd/yyyy)

SECTION VI - CONFIRMATION STATEMENT

The data and information in this submission have been reviewed and, to the best of my knowledge, are certified to be true and accurate. I agree to report changes to this information as required under section 905(i)(3) of the act.

☐ **Agree****WARNING:**

A willfully false statement is a criminal offense, U.S. Code, Title 18, Section 1001.

Signature of Responsible Person or Agent

Typed Name and Title

Date

Identity of the Signatory

☐ Owner (*Listed in section IIA*)☐ Operator (*Listed in section IIIA*)☐ Authorized Agent (*Complete section below*)**Authorized Agent Contact Information**Title (*e.g., Mr., Ms., Dr.*):

First/Given Name

Middle Name

Last Name

Position Title

Email Address

Telephone (*Include Country Code if applicable*)

FAX

Company Name

Address

City

State, Province or Territory

Country

ZIP or Postal Code

REFERENCES

Reference for the Tobacco Control Act:

<https://www.fda.gov/tobacco-products/rules-regulations-and-guidance/family-smoking-prevention-and-tobacco-control-act-table-contents>

Reference for *Guidance on Registration and Product Listing for Owners and Operators of Domestic Tobacco Prod*

<https://www.fda.gov/regulatory-information/search-fda-guidance-documents/registration-and-product-listing-owners-and-operators-domestic-tobacco-product-establishments>

For regulatory questions regarding sections 904 and 905 of the act,
email TobaccoIndustryQuestions@fda.hhs.gov.

Regulatory Submissions can be mailed to:

Food and Drug Administration
Center for Tobacco Products
Document Control Center
Building 71, Room G335
10903 New Hampshire Avenue
Silver Spring, MD 20993-0002

APPENDIX A - ADVERTISING
Continuation Sheet

Enter information below for a representative sample of advertising. (See Section V for details.)

Product Name	Category	Price	Quantity	Total Price
Product A	Electronics	100	5	500
Product B	Electronics	200	3	600
Product C	Electronics	150	4	600
Product D	Electronics	120	5	600
Product E	Electronics	100	6	600
Product F	Electronics	80	7	560
Product G	Electronics	60	8	480
Product H	Electronics	40	10	400
Product I	Electronics	30	12	360
Product J	Electronics	20	15	300
Product K	Electronics	10	20	200
Product L	Electronics	5	30	150
Product M	Electronics	2	40	80
Product N	Electronics	1	50	50
Product O	Electronics	0.5	60	30
Product P	Electronics	0.2	70	14
Product Q	Electronics	0.1	80	8
Product R	Electronics	0.05	90	4.5
Product S	Electronics	0.02	100	2
Product T	Electronics	0.01	110	1.1
Product U	Electronics	0.005	120	0.6
Product V	Electronics	0.002	130	0.26
Product W	Electronics	0.001	140	0.14
Product X	Electronics	0.0005	150	0.075
Product Y	Electronics	0.0002	160	0.032
Product Z	Electronics	0.0001	170	0.017
Product AA	Electronics	0.00005	180	0.009
Product AB	Electronics	0.00002	190	0.0038
Product AC	Electronics	0.00001	200	0.002
Product AD	Electronics	0.000005	210	0.00105
Product AE	Electronics	0.000002	220	0.00044
Product AF	Electronics	0.000001	230	0.00023
Product AG	Electronics	0.0000005	240	0.00012
Product AH	Electronics	0.0000002	250	0.00005
Product AI	Electronics	0.0000001	260	0.000026
Product AJ	Electronics	0.00000005	270	0.0000135
Product AK	Electronics	0.00000002	280	0.0000056
Product AL	Electronics	0.00000001	290	0.0000029
Product AM	Electronics	0.000000005	300	0.0000015
Product AN	Electronics	0.000000002	310	0.00000062
Product AO	Electronics	0.000000001	320	0.00000032
Product AP	Electronics	0.0000000005	330	0.000000165
Product AQ	Electronics	0.0000000002	340	0.000000068
Product AR	Electronics	0.0000000001	350	0.000000035
Product AS	Electronics	0.00000000005	360	0.000000018
Product AT	Electronics	0.00000000002	370	0.0000000074
Product AU	Electronics	0.00000000001	380	0.0000000038
Product AV	Electronics	0.000000000005	390	0.00000000195
Product AW	Electronics	0.000000000002	400	0.0000000008
Product AX	Electronics	0.000000000001	410	0.00000000041
Product AY	Electronics	0.0000000000005	420	0.00000000021
Product AZ	Electronics	0.0000000000002	430	0.000000000086
Product BA	Electronics	0.0000000000001	440	0.000000000044
Product BB	Electronics	0.00000000000005	450	0.0000000000225
Product BC	Electronics	0.00000000000002	460	0.0000000000092
Product BD	Electronics	0.00000000000001	470	0.0000000000047
Product BE	Electronics	0.000000000000005	480	0.0000000000024
Product BF	Electronics	0.000000000000002	490	0.00000000000098
Product BG	Electronics	0.000000000000001	500	0.0000000000005
Product BH	Electronics	0.0000000000000005	510	0.000000000000255
Product BI	Electronics	0.0000000000000002	520	0.000000000000104
Product BJ	Electronics	0.0000000000000001	530	0.000000000000053
Product BK	Electronics	0.00000000000000005	540	0.000000000000027
Product BL	Electronics	0.00000000000000002	550	0.000000000000011
Product BM	Electronics	0.00000000000000001	560	0.0000000000000056
Product BN	Electronics	0.000000000000000005	570	0.0000000000000028

Product Identification Number

[illegible]

APPENDIX B - LABELING
Continuation Sheet

Enter labeling information below.* (See Section V for details.)

Product Name	Product Identification Number

Universal Product Code(s) (UPC):

[illegible]

APPENDIX C - CONSUMER INFORMATION
Continuation Sheet

Enter consumer information below. (See Section V for details.)

Product Name	Category	Price	Quantity	Total Price
Apple iPhone 12	Electronics	999	1	999
Samsung Galaxy S21	Electronics	899	1	899
MacBook Pro 16	Electronics	2499	1	2499
Microsoft Surface Pro 9	Electronics	1299	1	1299
Adidas Ultraboost 21	Footwear	149	1	149
Nike Air Max 270	Footwear	129	1	129
Levi's 501 Jeans	Clothing	69	1	69
Patagonia Fleece Jacket	Clothing	119	1	119
Instant Pot Duo	Home Appliances	99	1	99
Robot Vacuum Cleaner	Home Appliances	299	1	299
Wireless Earbuds	Electronics	49	1	49
Smart Watch	Electronics	199	1	199
Yoga Mat	Fitness	29	1	29
Protein Powder	Fitness	49	1	49
Blender	Home Appliances	79	1	79
Smart TV 55"	Electronics	499	1	499
Gaming Console	Electronics	399	1	399
Desk Chair	Furniture	149	1	149
Bed Frame	Furniture	199	1	199
Refrigerator	Home Appliances	599	1	599
Washing Machine	Home Appliances	499	1	499
Air Conditioner	Home Appliances	799	1	799
Stove	Home Appliances	149	1	149
Water Filter	Home Appliances	49	1	49
Light Fixture	Home Decor	79	1	79
Plant Pot	Home Decor	29	1	29
Wall Art	Home Decor	59	1	59
Throw Pillow	Home Decor	19	1	19
Curtains	Home Decor	39	1	39
Area Rug	Home Decor	99	1	99
Bedding Set	Home Textiles	49	1	49
Bath Towels	Home Textiles	29	1	29
Shower Curtain	Home Textiles	19	1	19
Bath Mat	Home Textiles	14	1	14
Laundry Basket	Home Textiles	19	1	19
Storage Bins	Home Storage	29	1	29
Shelving Unit	Home Storage	49	1	49
Drawer Organizer	Home Storage	14	1	14
Under Bed Storage	Home Storage	29	1	29
Garage Storage	Home Storage	79	1	79
Tool Box	Home Storage	49	1	49
Workbench	Home Storage	99	1	99
Shed	Home Storage	199	1	199
Greenhouse	Home Storage	299	1	299
Compost Bin	Home Storage	49	1	49
Watering System	Home Storage	79	1	79
Plant Fertilizer	Home Storage	14	1	14
Garden Hose	Home Storage	29	1	29
Lawn Mower	Home Storage	149	1	149
String Trimmer	Home Storage	79	1	79
Leaf Blower	Home Storage	49	1	49
Pruning Shears	Home Storage	19	1	19
Garden Gloves	Home Storage	14	1	14
Shovel	Home Storage	29	1	29
Rake	Home Storage	19	1	19
Wheelbarrow	Home Storage	49	1	49
Tool Bag	Home Storage	29	1	29
Tool Roll	Home Storage	19	1	19
Tool Set	Home Storage	99	1	99
Tool Kit	Home Storage	49	1	49
Tool Chest	Home Storage	79	1	79
Tool Box	Home Storage	49	1	49
Tool Bag	Home Storage	29	1	29
Tool Roll	Home Storage	19	1	19
Tool Set	Home Storage	99	1	99
Tool Kit	Home Storage	49	1	49
Tool Chest	Home Storage	79	1	79
Tool Box	Home Storage	49	1	49
Tool Bag	Home Storage	29	1	29
Tool Roll	Home Storage	19	1	19
Tool Set	Home Storage	99	1	99
Tool Kit	Home Storage	49	1	49
Tool Chest	Home Storage	79	1	79
Tool Box	Home Storage	49	1	49
Tool Bag	Home Storage	29	1	29
Tool Roll	Home Storage	19	1	19
Tool Set	Home Storage	99	1	99
Tool Kit	Home Storage	49	1	49
Tool Chest	Home Storage	79	1	79
Tool Box	Home Storage	49	1	49
Tool Bag	Home Storage	29	1	29
Tool Roll	Home Storage	19	1	19
Tool Set	Home Storage	99	1	99
Tool Kit	Home Storage	49	1	49
Tool Chest	Home Storage	79	1	79
Tool Box	Home Storage			

Product Identification Number

[illegible]

INSTRUCTIONS

Section I

- If you check box 1, complete all sections except IIIA and IIIB.
- If you check box 2, complete all sections.
- If you check box 3, complete all sections.
- If you check box 4, complete all sections except IIA and IIB.
- If you check box 5, complete all sections.

Section IIA

Provide all required information for the Owner. FDA requests that you also provide a point of contact, to facilitate communication between the Owner and FDA. If an Operator is registering alone, the Operator may skip Sections IIA and IIB.

Section IIB

Owners must provide the specified details for their business structure. Only one business structure should be selected. If an Owner uses any trade names to conduct business other than the company name identified in Section IIA, FDA requests that such names be listed here.

Section IIIA

Provide all required information for the Operator. FDA requests that you also provide a point of contact, to facilitate communication between the Operator and FDA. Owners registering alone may skip Sections IIIA and IIIB. If you an Owner or Operator registering on behalf of multiple Operators, you must submit a separate IIIA and IIIB for each Operator.

Section IIIB

Operators must provide the specified details for their business structure. Only one business structure should be selected. If an Operator uses any trade names to conduct business other than the company name identified in Section IIIA, FDA requests that such names be listed here.

Section IV

Provide all required information for each Establishment. FDA requests that you also provide a point of contact, to facilitate communication between the Establishment and FDA. The contact person for a given Establishment does not need to be the Operator, but should be an individual authorized to communicate with FDA. Owners and Operators must register and submit a separate Section IV for each Establishment they own or operate. If you are an Operator registering on behalf of an Owner, you must complete a separate Section IV for each Establishment owned by that Owner, even if you are not the Operator of all of the Establishments.

Section V

This section applies to each product manufactured by the registrant. If an Owner or Operator intends to list multiple products, multiple copies of Section V may be submitted. If additional space is needed for submission of advertising information, registrants may use Appendix A to identify additional items of advertising submitted per product. If additional space is needed for submission of labeling information, registrants may use Appendix B to identify additional items of labeling submitted per product. If additional space is needed for submission of consumer information, registrants may use Appendix C to identify additional items of consumer information submitted per product.

Section VI

Registration and listing information may be submitted only by an owner, operator, or authorized agent thereof. If an agent has been authorized to submit registration and listing information, FDA requests that contact information for that agent be entered in this section.