DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration

Potential Tobacco Product Violations Report

Form Approved: OMB No.: 0910-0716 Expiration Date: 08/31/2023 (See page 3 for PRA Statement)

Directions:

Use this form to report potential tobacco-related violations of the Federal Food, Drug, and Cosmetic Act and associated regulations. These submissions are reviewed by FDA's Center for Tobacco Products.

WHO can report? - Any member of the public.

Tell us:

WHEN did you see the potential violation?

WHERE did the potential violation occur?

WHAT is the potential violation?

WHY report? - Information we receive from the public is often very helpful in identifying problems with marketed products and possible violations of the laws that we enforce.

To submit your report, complete the form below:

	Date and State	e Where Violation Occ	cur	red
Date potential violation occurred (mm/	dd/yyyy)	I do not recall the date this potential violation occurred		State in which potential violation occurred
	Desc	ription of Product		
Туре		Tobacco Brand		
Potential violation type	Sales to minors	5		Free samples
(choose all that apply)	Flavored cigare			Self-service display/direct access to cigarette or smokeless tobacco
	_	motion/marketing ne/direct access to cigarette		Sale of cigarettes in packs of less than 20
	or smokeless to	obacco or covered tobacco		Unsure
	Unauthorized to	obacco product		
Type of potentially violative promotional	Newspaper			Price signage
materials <i>(choose all</i>	Magazine			Posters
that apply)	Periodicals			Coupons
	Billboard			Internet
	Direct mail			Unsure
	In-store adverti	sements		
Who potentially violated?	Retailer			Importer
(choose all that apply)	Manufacturer			Distributor

Potentia	I Tobacco Product Violations Repor	t
Description of potential violation		
	cal address of the potential violator	, if known
Retailer, manufacturer, importer, or distributor nan	ne	
Street Address		
Street Address Line 2		
City	State/Province/Region	Postal/Zip Code
,	ű	'
If report is about a website, insert website address		
irreport is about a website, insert website address		
internet policies, please visit: http://	te to the extent allowed by law. For more infor /www.fda.gov/AboutFDA/AboutThisWebsite/W	
	, I want my report to be anonymous. (Please note to A will receive your email address. However, if you	
	s, FDA may contact me. (Please fill in contact infor	
Name	· · · · · · · · · · · · · · · · · · ·	,
Affiliation (such as company, school, or group)		
rumation (cash as sompany, somest, or group)		
Street Address		
Street Address Line 2		
		(continued on next page

	obacco Product Violations Report
City	State/Province/Region
Postal/Zip Code	Phone Number
Email	
Please email me to notify me No	In order to receive a response, please configure your email spam/junk
	filter to allow messages from ctpcompliance@fda.hhs.gov. In most cases
hat FDA got my complaint	
nat FDA got my complaint Yes	this is solved by adding our email address to your address book.
res	this is solved by adding our email address to your address book.
If you would rather subm	
If you would rather subm	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the following address:
If you would rather subm	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the the following address: Food and Drug Administration
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If you would rather subm	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the following address: Food and Drug Administration Center for Tobacco Products Document Control Center
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If you would rather subm	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the following address: Food and Drug Administration Center for Tobacco Products Document Control Center Building 71, Room G335
If you would rather submer plea	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the the following address: Food and Drug Administration Center for Tobacco Products Document Control Center Building 71, Room G335 10903 New Hampshire Avenue Silver Spring, MD 20993-0002
If you would rather subm plea	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the the following address: Food and Drug Administration Center for Tobacco Products Document Control Center Building 71, Room G335 10903 New Hampshire Avenue
If you would rather subm plea	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the the following address: Food and Drug Administration Center for Tobacco Products Document Control Center Building 71, Room G335 10903 New Hampshire Avenue Silver Spring, MD 20993-0002 none, please call 1-877-CTP-1373, and select option 3.
If you would rather subm plea	this is solved by adding our email address to your address book. nit your report to us in writing, along with any attachments, se do so at the the following address: Food and Drug Administration Center for Tobacco Products Document Control Center Building 71, Room G335 10903 New Hampshire Avenue Silver Spring, MD 20993-0002 none, please call 1-877-CTP-1373, and select option 3.

An email message automatically will be produced when you click the SUBMIT BY EMAIL button. In the resulting email message, please don't forget to click the "Send" button or its equivalent when you are ready to send the email.

OMB Paperwork Reduction Act Statement

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 for this collection of information is estimated to average 0.25 hour 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."