DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

FDA FORM 3779

POTENTIAL TOBACCO PRODUCT VIOLATIONS REPORTING

Form Approved: OMB No. 0910-xxxx
Expiration Date: *xx/xx/xxxx*
(See page 3 for Burden Statement)

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, Office of Compliance and Enforcement.

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, use the form below:

Date and State Where Violation Occurred

Date potential violation occurred

I do not recall the date this potential violation occurred

State in which potential violation occurred

Description of Product

Type

Tobacco brand

Potential violation type (choose all that apply)

- ☐ Sales to Minors
- ☐ Flavored cigarette sales
- ☐ Advertising/promotion/marketing
- ☐ Free samples
- ☐ Vending machine/self-service display/direct access to cigarette or smokeless tobacco
- ☐ Sale of cigarettes in packs of less than 20
- ☐ Unsure

Type of potentially violative promotional materials (choose all that apply)

- ☐ Newspaper
- ☐ Magazine
- ☐ Periodicals
- ☐ Billboard
- ☐ Direct Mail
- ☐ In-store advertisements
- ☐ Price signage
- ☐ Posters
- ☐ Coupons
- ☐ Internet advertising
- ☐ Unsure

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Who potentially violated? (choose all that apply)

- ☐ Retailer
- ☐ Distributor
- ☐ Manufacturer
- ☐ Unsure
- ☐ Importer

Description of potential violation (character limit 1000)

Please provide name and physical address of the potentially violative party/location below by potential violator types, if known

Brand, retailer, manufacturer, importer, or distributor name (character limit 250)

Street Address

Street Address 2

City

State/Province/Region

Postal/ Zip Code

If report is about a Web site, please provide Web site address here:

All reports will remain confidential to the extent allowed by law. For more information about FDA's internet policies please visit: http://www.fda.gov.AboutFDA/AboutThisWebsite/WebsitePolicies/default.htm

May we contact you if we need additional Information?

- ☐ Yes, CTP may contact me. (Please fill in contact information below.)
- ☐ No, I want my report to be anonymous

Name (character limit 250)

Affiliation (such as company, school, or group) (character limit 250)

Street Address

Address Line 2

City

State/Province/Region

Postal/ Zip Code

Phone number

(Continued on next page)

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E-mail

I would like to receive an e-mail to notify me that CTP got my complaint

☐ Yes
☐ No

In order to receive a response, please configure your e-mail spam/junk filter to allow messages from ctpcompliance@fda.hhs.gov. In most cases, this is solved by adding our e-mail address to your address book

If you would rather submit your report to us in writing along with any attachments, please do so at the following address:

Tobacco Product Violation Report, Office of Compliance and Enforcement
FDA Center for Tobacco Products
c/o Document Control Center
9200 Corporate Boulevard
Rockville, MD 20850-3229

To reach us by telephone, please call 1-877-CTP-1373, and select option 3. You may also e-mail us at ctpcompliance@fda.hhs.gov.

Submit by E-mail | Print Form | Reset Form

OMB Paperwork Reduction Act Statement

Public reporting burden for this collection of information is estimated to average 0.25 hours (15 minutes) per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to:

Department of Health and Human Services
Food and Drug Administration
Office of the Chief Information Officer
1350 Piccard Drive, 420A
Rockville, MD 20850

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.