PART 420—BASIN REGULATIONS—WATER SUPPLY CHARGES

3. The authority citation for part 420 continues to read as follows:

Authority: Delaware River Basin Compact, 75 Stat. 688.

4. Revise § 420.41 to read as follows:

§ 420.41 Schedule of water charges.

The schedule of water charges established in accordance with § 420.22 shall be as follows:

(a) $80 per million gallons for consumptive use, subject to paragraph (c) of this section; and

(b) $0.80 per million gallons for non-consumptive use, subject to paragraph (c) of this section.

(c) On July 1 of every year, beginning July 1, 2017, the rates established by this section will increase commensurate with any increase in the annual April 12-month Consumer Price Index (CPI) for Philadelphia, published by the U.S. Bureau of Labor Statistics during that year. In any year in which the April 12-month CPI for Philadelphia declines or shows no change, the water charges rates will remain unchanged. Following any indexed adjustment made under this paragraph (c), revised consumptive and non-consumptive use rates will be published in the Federal Register by July 1 and posted on the Commission’s Web site. Interested parties may also obtain the rates by contacting the Commission directly during business hours.

Dated: December 20, 2016.

Pamela M. Bush,
Commission Secretary.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1105

[Docket No. FDA–2016–N–1555]

Refuse To Accept Procedures for Premarket Tobacco Product Submissions

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is issuing a final rule describing when FDA will refuse to accept a tobacco product submission (or application) because the application has not met a minimum threshold for acceptability for FDA review. Under the rule, FDA will refuse to accept a tobacco product submission, for example, that is not in English, does not pertain to a tobacco product, or does not identify the type of submission. By refusing to accept submissions that have the deficiencies identified in the proposed rule, FDA will be able to focus our review resources on submissions that meet a threshold of acceptability and encourage quality submissions.

DATES: This rule is effective January 30, 2017.

FOR FURTHER INFORMATION CONTACT: Annette Marthaler or Paul Hart, Office of Regulations, Center for Tobacco Products (CTP), Food and Drug Administration, Document Control Center, Bldg. 71, Rm. G335, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 877–287–1373, CTPRegulations@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Executive Summary

A. Purpose of the Rule

FDA is issuing this refuse to accept rule to identify deficiencies that will result in FDA’s refusal to accept certain tobacco product submissions under sections 905, 910, and 911 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act), as amended by the Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act) (21 U.S.C. 387e, 387f, and 387k). Because these submissions will be refused before they enter FDA’s review queue, more resources will be available for submissions that are ready for further review. This rule establishes a refuse to accept process for premarket tobacco product submissions, including premarket tobacco product applications (PMTAs), modified risk tobacco product applications (MRTPAs), substantial equivalence (SE) applications (also called SE reports), and exemption requests (including subsequent abbreviated reports).

B. Summary of the Major Provisions of the Regulatory Action

The rule explains when FDA will refuse to accept a premarket submission, including PMTAs, MRTPAs, SE applications, and exemption requests (including subsequent abbreviated reports). The rule is based on FDA’s experience in reviewing these submissions. Under the rule, FDA will refuse to accept a premarket submission that: (1) Does not pertain to a tobacco product; (2) is not in English (or does not include a complete translation); (3) is submitted in an electronic format that FDA cannot process, read, review, or archive; (4) does not include the applicant’s contact information; (5) is from a foreign applicant and does not include the name and contact information of an authorized U.S. agent (authorized to act on behalf of the applicant for the submission); (6) does not include a required form(s); (7) does not identify the tobacco product; (8) does not identify the type of submission; (9) does not include the signature of a responsible official authorized to represent the applicant; or (10) does not include an environmental assessment or claim of a categorical exclusion, if applicable. Under the rule, if FDA refuses to accept the submission, FDA will send the contact (if available) a notification. If the submission is accepted for further review, FDA will send an acknowledgement letter.

II. Background

FDA published two rulemaking documents concerning refuse to accept procedures in the Federal Register of August 8, 2016: A direct final rule (81 FR 52329) and a companion proposed rule (81 FR 52371). We published the direct final rule because we believed that the rule was noncontroversial, and we did not anticipate that it would receive any significant adverse comments. As a companion to the direct final rule, we published a proposed rule with the same codified language published in the proposed rules section of the Federal Register. The companion proposed rule provides a procedural framework to finalize the rule in the event that the direct final rule receives any adverse comment and is withdrawn. We received adverse comment on the direct final rule and withdrew the direct final rule by issuing a notice in the Federal Register of November 16, 2016 (81 FR 80567). We are now finalizing the proposed rule and responding to the comments we received.
III. Purpose and Legal Authority

A. Purpose

FDA is issuing this refuse to accept rule to efficiently handle submissions that do not meet a threshold of acceptability for FDA review (e.g., the submission lacks certain information FDA needs for substantive review of the submission). Currently, FDA often expends extensive time and resources in attempts to obtain information and resolve the deficiencies identified in the rule simply to begin substantively processing the submission. FDA expects that this rule will enhance the quality of the submissions and that submissions will move expeditiously through the review process. In addition, this rule will help submitters better understand the common hurdles FDA encounters in conducting a substantive review of submissions.

The rule identifies deficiencies that FDA has seen across types of premarket submissions and will result in FDA refusing to accept the submission. This rule applies to all tobacco product applications; we note that there are additional deficiencies that are not covered in this rule that may arise for specific types of premarket submissions that would also result in FDA’s refusal to accept that specific type of premarket submission (e.g., omission of labeling for a PMTA that is required under section 910(b)(1)(F) of the FD&C Act).

FDA’s refusal to accept a tobacco product submission does not preclude an applicant from resubmitting a new submission that addresses the deficiencies. In addition, acceptance of a submission does not mean that FDA has determined that the submission is complete, rather only that the submission meets the basic, minimum threshold for acceptance. Substantive review of the submission will begin once FDA accepts the submission, and for submissions with filing requirements (i.e., PMTAs and MRPTAs), once filed. This rule establishes a general process for refusing to accept submissions for premarket tobacco review, including PMTAs, MRPTAs, SE applications, and exemption requests (including subsequent abbreviated reports) for the reasons listed in paragraph (a)(1) through (a)(10), if applicable.

B. Legal Authority

Section 701(a) of the FD&C Act (21 U.S.C. 371(a)) provides FDA with the authority to issue regulations for the efficient enforcement of the FD&C Act. This rule will allow FDA to more efficiently use our resources to review premarket submissions under sections 905, 910, and 911 of the FD&C Act. FDA has processed and reviewed many submissions since the enactment of the Tobacco Control Act, and submissions with the deficiencies identified in the rule have been repeatedly identified by FDA as reflecting submissions that are incomplete and not prepared for further review.

IV. Overview of the Final Rule

We are finalizing the proposed rule with only editorial changes. The rule adds part 1105 (21 CFR part 1105) to title 21, specifically § 1105.10. Section 1105.10 provides that FDA will refuse to accept, as soon as practicable, PMTAs, MRPTAs, SE applications, and exemption requests (including subsequent abbreviated reports) for the reasons listed in paragraphs (a)(1) through (a)(10), if applicable.

V. Comments on the Proposed Rule

We consider any comments that were submitted on the direct final rule to have been submitted on the proposed rule. We received two sets of comments on the proposed rule, one from a tobacco product manufacturer and another from a public health group. In general, one of the commenters expressed strong support for this rule, asking that it be applied to a broader set of applications, while the other commenter identified concerns with the rulemaking, including that “promulgating a direct final rule was procedurally improper.”

One commenter suggested that FDA withdraw the rule in its entirety and issue any future rule only after engaging in notice and comment rulemaking. This rulemaking, however, did provide both notice and an opportunity for comments. As previously noted, FDA withdrew the direct final rule and is proceeding with the rulemaking under the procedural framework of the proposed rule. FDA has considered the comments submitted to the docket for the rulemaking and responds to the comments in the following paragraphs.

To make it easier to identify comments and our responses, the word “Comment,” in parentheses, will appear before each comment, and the word “Response,” in parentheses, will appear before each response. We have numbered the comments to make it easier to distinguish between comments; the numbers are for organizational purposes only and do not reflect the order in which we received the comments or any value associated with the comment. We have combined similar comments under one numbered comment.

(Comment 1) One commenter suggested that FDA apply the rule to provisional substantial equivalence applications submitted by manufacturers under section 910(a)(2)(B) of the FD&C Act for new tobacco products that were first introduced or delivered for introduction into interstate commerce between February 15, 2007, and March 22, 2011.

(Comment 2) One commenter suggested that FDA apply this “commonsense regulation” to premarket submissions for newly designed tobacco products submitted during the compliance period announced in the Deeming rule.

(Comment 3) One commenter requested that FDA refine and expand the requirements of this rule to allow FDA to refuse to accept substantial equivalence applications that fail to...
comply with certain criteria that relate to the substantial equivalence pathway, such as creating product-identifying information requirements for predicate products.

(Response) FDA disagrees with this comment. The rule creates a minimum threshold of acceptability for all premarket submissions, regardless of the type of submission, and is not intended to address content specific to only one type of premarket submission. FDA plans to consider including refuse to accept criteria that are specific to a particular premarket pathway as part of future rulemakings. For example, FDA has already issued one such rule, “Tobacco Products, Exemptions From Substantial Equivalence Requirements,” which contains refuse to accept criteria relating specifically to exemption requests (July 5, 2011, 76 FR 38961).

(Comment 4) One commenter argued that FDA lacks the legal authority to implement the rule. The commenter stated that because the Tobacco Control Act does not set forth content requirements for substantial equivalence applications or exemption requests, FDA has no statutory justification for pre-review of those submissions. The commenter further stated that while the Tobacco Control Act does set forth content requirements for premarket tobacco product applications and modified risk tobacco product applications that grant FDA authority to conduct filing reviews of those submissions, FDA lacks the statutory authority to conduct separate reviews for content part of the pre-review of an application. In sum, the commenter argued that FDA does not have the statutory authority, either explicit or implicit, to refuse to accept tobacco product submissions.

(Response) FDA disagrees with this comment. As described in section III.B of the rule, section 701(a) grants FDA the authority to issue regulations for the efficient enforcement of the FD&C Act. As also discussed in the proposed rule, this rule will allow FDA to efficiently enforce the premarket review requirements of sections 905, 910, and 911 of the FD&C Act by allowing FDA to refuse to accept submissions that do not meet basic criteria and focus its resources on those submissions that are ready for review.

(Comment 5) One commenter argued that unless FDA establishes a time by which FDA will refuse to accept a premarket submission, the rule is legally problematic for a number of reasons. While two of the specific reasons are discussed in separate comments and responses, overall, the commenter suggested that FDA should, similar to its approach for new drug applications and premarket approval applications for medical devices, create a limit of 15 days in which to determine whether it will refuse to accept a premarket submission.

(Response) FDA declines the suggestion that FDA adopt a 15-day time limit similar to the refusal to accept review periods for reference to its promotional claims. Courts have upheld FDA premarket reviews in other product areas based on a similar scheme. See, e.g., 12375 (7th Cir. 2016); Whitaker v. Thompson, 353 F.3d 947 (D.C. Cir. 2004); United States v. Cole, 84 F. Supp. 3d 1159, 1166 (D. Or. 2015). Third, there is a split in authority regarding whether the prior restraint doctrine applies to commercial speech; the Sixth Circuit in Discount Tobacco found that the doctrine did not apply to evaluation of the MRTP provisions and defines such product in part by reference to its promotional claims.

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Thus, as of August 8, 2016, marketing these products without FDA authorization is prohibited by statute. However, FDA is affording staggered compliance periods during which FDA does not intend to enforce the premarket review requirements. These compliance periods are general statements of policy that do not establish any rights for any person, and are not binding on FDA or the public. (See e.g., Professionals and Patients for Customized Care v. Shalala, 56 F.3d 592 (5th Cir. 1995).) The commenter gives a vague reference to the rule depriving manufacturers of a “substantive right” to market with no hearing or substantive review, but without citing any authority for such a right. Irrespective of the rule, a manufacturer does not have a right to market a product that is in violation of the FD&C Act because it does not have a required premarket authorization.

(Comment 8) One commenter stated that FDA should allow manufacturers to amend applications that FDA finds to be deficient and consider the amended applications to be received as of their original submission dates. The commenter explained that this approach would not tie up Agency resources because FDA could simply notify an applicant of any deficiencies and consider the amended application content requirements to apply to all premarket submissions. Detailed criteria that are specific to each premarket pathway are not necessary to implementing a rule that applies to all types of premarket submissions generally without any consideration of content specific to each premarket pathway. Any additional grounds for which FDA may refuse a premarket submission exist independently from this rulemaking; therefore, the vagueness of such grounds, if any, is not attributable to the rule and does not cause it to violate the Due Process clause of the 5th Amendment or the APA. A law is impermissibly vague if it does not give “a person of ordinary intelligence a reasonable opportunity to know what is prohibited.” Grayned v. City of Rockford, 408 U.S. 104, 108 (1972). To the extent that the commenter identifies concerns with specific requirements of the rule, we address them in the responses to comments 10–14; however, FDA believes that the requirements of this rule are sufficiently clear to give submitters a reasonable opportunity to be aware of what information must be included with a tobacco product application.

(Comment 10) One commenter argued that FDA must edit the rule so that it comprehensively states all potential refuse to accept criteria for each premarket pathway and commit to accepting all submissions that meet those specific criteria because granting FDA discretion to refuse to accept submissions on the basis of criteria not specified in this rule violates the principles of fair notice embodied in the Constitution and the APA.

(Response) FDA disagrees. Under § 1105.10(b), FDA “may accept the submission” if it “finds that none of the reasons in paragraph (a) of this section exists for refusing to accept a premarket submission.” The use of the word
“may” in this section reflects the fact that this rule addresses the basic threshold of acceptability that all premarket submissions must meet; however it does not address other grounds on which FDA could refuse to accept a specific type of premarket submission, such as the omission of labeling from a PMTA that is required by section 910(b)(1)(F) of the FD&C Act. Any additional grounds on which FDA may refuse to accept a premarket submission exist independently from this rulemaking and are outside of its scope.

(Comment 11) One commenter argues that FDA’s discussion in the preamble of the proposed rule regarding “other information” that FDA recommends be included as part of the product-identifying information submitted under §1105.10(a)(7) should either be deleted or modified to provide a full and complete description of what “other information” applicants should provide. The commenter also suggests that FDA must state whether failure to provide such information would be grounds for FDA to refuse to accept a submission.

(Response) FDA disagrees with this comment. Section 1105.10(a)(7) specifically lists the product-identifying information that is required under the rule: The manufacturer of the tobacco product; the product name, including the brand and subbrand; the product category and subcategory; package type and package quantity; and characterizing flavor. The preamble of the proposed rule notes that other information may be needed to identify the product, such as product descriptors that are not a part of the product name (e.g., premium), but it merely requests such information be submitted to facilitate FDA’s review. Failure to include additional product-identifying information beyond those specifically listed in §1105.10(a)(7) is not grounds for FDA to refuse to accept a submission under the rule.

(Comment 12) One commenter argued that FDA must either remove the requirement in §1105.10(a)(7) that applicants specify the category and subcategory of the tobacco product or provide a list of all potential categories and subcategories. The commenter further noted that FDA could require a uniform system of product identification under 21 U.S.C. 387(e)(section 905(e) of the FD&C Act), but it has not yet issued a regulation doing so.

(Response) FDA disagrees with this comment. The rule requires applicants to describe the category and subcategory of the tobacco product that is the subject of the premarket submission. This is a requirement to provide basic product-identifying information, such as describing the product category as “Smokeless Tobacco Product” and the subcategory as “Dissolvable,” which in no way creates a rigid system of product identification with which an applicant must comply. Creating an exhaustive product categorization system is not necessary for applicants to describe the product’s category and subcategory and in some cases may not allow applicants to accurately describe new tobacco products that fall into novel categories or subcategories. Table 1 in the preamble of the proposed rule provides some recommendations on how an applicant may satisfy this requirement, but it is not intended to be an exhaustive list (for example, although recommendations for waterpipes were not included in table 1, submissions on waterpipes should include similar information). While the table is not an exhaustive list of every tobacco product category and subcategory that exists, manufacturers have enough information to reasonably understand how to comply with the requirement and can provide information based on internal classifications. Applicants unable to identify the category or subcategory of the tobacco product that will be the subject of a premarket submission are encouraged to contact FDA prior to submission.

(Comment 13) One commenter argued that FDA should not require an applicant to identify the submission type as part of a premarket submission because the list of submission types provided in §1105.10(a)(8) is incomplete. To support this statement, the commenter notes that the list in the preamble of the proposed rule does not mention Product Quantity Change SE Reports as a potential premarket submission type.

(Response) FDA disagrees with the suggestion that manufacturers should not be required to identify the type of application they are submitting and that the list of submission types described in the preamble of the proposed rule is incomplete. The type of submission is necessary for FDA to review a premarket submission because it enables FDA to determine the appropriate decisional standard to apply to a submission (e.g., whether it is a PMTA subject to the requirements of section 910 of the FD&C Act or an MRTPA subject to the requirements of section 911 of the FD&C Act). The commenter is also incorrect in its assertion that the proposed rule’s discussion of the types of premarket submissions is incomplete. The only example the commenter provides to support this assertion is the Product Quantity Change SE Reports, which are SE applications. The preamble of the proposed rule described the types of premarket submissions, which are PMTAs, MRTPAs, SE applications, and exemption requests (and subsequent abbreviated reports). Applicants are welcome to provide additional information regarding their submission type, such as specifying that their SE application is being submitted for a product quantity change, provided that the basic submission type remains clear. Applicants unsure of how to identify the type of application that they are submitting are encouraged to contact FDA prior to submission.

(Comment 14) One commenter argued that FDA should remove the requirement that a premarket submission be accompanied by required forms because FDA has yet to require any forms and it is unclear what those forms may eventually require. The commenter stated that if and when FDA creates required forms, it can issue regulations providing how and when the forms must be submitted.

(Response) We disagree with the suggestion that this requirement should be removed from the rule. As described in section IV of the proposed rule, if and when FDA issues any forms it would need to do so in accordance with applicable requirements, e.g., notice and opportunity to comment on what forms in accordance with rulemaking procedures and the Paperwork Reduction Act of 1995 and rulemaking under the APA. We have chosen to include the form submission requirement in this rule to provide notice that the failure to submit any required forms, if and whenever they are issued, will be grounds for refusing to accept a premarket submission.

(Comment 15) One commenter argued that FDA should not require applicants to identify whether a product has a characterizing flavor until FDA has issued a full explanation of what it considers to be a characterizing flavor and how it expects manufacturers to determine what the characterizing flavor of a tobacco product is. The commenter also argued that the requirement to identify a characterizing flavor has no statutory basis and is not necessary to identify a product in light of all other information FDA is requiring, such as the product name, brand, subbrand, category, and subcategory.

(Response) FDA disagrees with this comment. This requirement, along with
the other product-identifying information in § 1105.10(a)(7), will identify to FDA the specific tobacco product that is the intended subject of the application. As explained in the preamble to the proposed rule, FDA is requiring this product-identifying information under section 701 of the FD&C Act to efficiently enforce premarket review requirements for tobacco requirements. For example, FDA needs to be able to distinguish between products that have the same brand and subbrand, but different flavors (e.g., brand X menthol or brand X cinnamon). This also helps ensure that FDA ultimately issues an order that addresses the intended tobacco product. For the purposes of the refuse to accept process and to appropriately identify the specific product that is the subject of the submission, FDA is solely looking to see how the applicant identifies the tobacco product as having no characterizing flavor or having a particular characterizing flavor. Thus, for example, a firm would give “menthol” as the characterizing flavor of a tobacco product it identifies as “Brand A menthol”. At the acceptance stage, FDA would not review beyond how the product is identified, such as to determine whether the product contains a different or additional characterizing flavor. Applicants that have questions regarding how to describe their product’s characterizing flavor are encouraged to contact FDA prior to submission.

Comment 16) One commenter argued that FDA should either modify the rule so that it contains procedures to resolve disputes regarding whether FDA should have refused to accept an application, or it should specify whether the procedures for internal Agency review of decisions specified in § 10.75 (21 CFR 10.75) applies.

Response The procedures for internal Agency review of decisions in § 10.75 apply to a decision of an FDA employee, other than commissioner, on a matter. Applicants seeking review of a refuse to accept decision may use this mechanism or consider other mechanisms set out in part 10. FDA expects, however, that most applicants will find that addressing any deficiencies in the application will quickly resolve issues.

VI. Paperwork Reduction Act of 1995

FDA concludes that this rule contains no collection of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

VII. Federalism

We have analyzed this rule in accordance with the principles set forth in Executive Order 13132. We have determined that the rule does not contain policies that have substantial direct effects on the States, on the relationship between the National Government and the States, or on the distribution of power and responsibilities among the various levels of government. Accordingly, we conclude that the rule does not contain policies that have federalism implications as defined in the Executive Order and, consequently, a federalism summary impact statement is not required.

VIII. Tribal Consultation

We have analyzed this rule in accordance with the principles set forth in Executive Order 13175. We have determined that the rule does not contain policies that would have substantial direct effects on one or more Indian tribes, on the relationship between the Federal Government and Indian tribes, or on the distribution of power and responsibilities between the Federal Government and Indian tribes. Accordingly, we conclude that the rule does not contain policies that have tribal implications as defined in the Executive Order; consequently, a tribal summary impact statement is not required.

IX. Analysis of Environmental Impact

We have determined under 21 CFR 25.30(h) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

X. Economic Analysis of Impacts

We have examined the impacts of the rule under Executive Order 12866, Executive Order 13563, the Regulatory Flexibility Act (5 U.S.C. 601–612), and the Unfunded Mandates Reform Act of 1995 (Pub. L. 104–4). Executive Orders 12866 and 13563 direct us to assess all costs and benefits of available regulatory alternatives and, when regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety, and other advantages; distributive impacts; and equity). We believe that this rule is not a significant regulatory action as defined by Executive Order 12866. The Regulatory Flexibility Act requires us to analyze regulatory options that would minimize any significant impact of a rule on small entities. Because this rule establishes a procedure that FDA is responsible for implementing and has the effect of providing all entities useful feedback on the readiness of a submission, we certify that the rule will not have a significant economic impact on a substantial number of small entities.

The Unfunded Mandates Reform Act of 1995 (section 202(a)) requires us to prepare a written statement, which includes an assessment of anticipated costs and benefits, before proposing “any rule that includes any Federal mandate that may result in the expenditure by State, local, and tribal governments, in the aggregate, or by the private sector, of $100,000,000 or more (adjusted annually for inflation) in any one year.” The current threshold after adjustment for inflation is $146 million, using the most current (2015) Implicit Price Deflator for the Gross Domestic Product. This rule does not result in expenditure in any year that meets or exceeds this amount.

This rule identifies 10 significant and common deficiencies in premarket tobacco submissions that will cause FDA to refuse to accept them. Encouraging submissions that are free of the deficiencies listed in this rule does not represent a change in Agency expectations. One of the 10 deficiencies is required by statute (i.e., must be a tobacco product). One of the deficiencies is required by another regulation (i.e., must comply with requirements related to environmental assessments or exclusions from such assessments). The remaining eight deficiencies are basic expectations for an application to enter the review process. Therefore, this rule clarifies these expectations. This clarification will result in cost savings for both the applicant and FDA as less time is spent by FDA working with applicants to address these significant deficiencies. Applicants have clarity about basic expectations regarding requirements for acceptance of premarket applications. In addition, refusing to accept submissions with these deficiencies will allow Agency staff to more efficiently process submissions and quickly move those submissions without these deficiencies into review of substantial scientific issues.

List of Subjects in 21 CFR Part 1105

Administrative practices and procedures, Tobacco, Tobacco products.
of Food and Drugs. 21 CFR chapter I is amended by adding part 1105, consisting of § 1105.10, to read as follows:

PART 1105—GENERAL

Authority: 21 U.S.C. 371(a), 387e, 387j, and 387k.

Subpart A—General Submission Requirements

§ 1105.10 Refusal to accept a premarket submission.

(a) FDA will refuse to accept for review, as soon as practicable, a premarket tobacco product application, modified risk tobacco product application, substantial equivalence application, or exemption request or subsequent abbreviated report for the following reasons, if applicable:

1. The submission does not pertain to a tobacco product as defined in 21 U.S.C. 321(rr).

2. The submission is not in English or does not contain complete English translations of any information submitted within.

3. If submitted in an electronic format, the submission is in a format that FDA cannot process, read, review, and archive.

4. The submission does not contain contact information, including the applicant’s name and address.

5. The submission is from a foreign applicant and does not identify an authorized U.S. agent, including the agent’s name and address, for the submission.

6. The submission does not contain a required FDA form(s).

7. The submission does not contain the following product-identifying information: The manufacturer of the tobacco product; the product name, including the brand and subbrand; the product category and subcategory; package type and package quantity; and characterizing flavor.

8. The type of submission is not specified.

9. The submission does not contain a signature of a responsible official, authorized to represent the applicant, who either resides in or has a place of business in the United States.

10. For premarket tobacco applications, modified risk tobacco product applications, substantial equivalence applications, and exemption requests only: The submission does not include a valid claim of categorical exclusion in accordance with part 25 of this chapter, or an environmental assessment.

(b) If FDA finds that none of the reasons in paragraph (a) of this section exists for refusing to accept a premarket submission, FDA may accept the submission for processing and further review. FDA will send to the submitter an acknowledgement letter stating the submission has been accepted for processing and further review and will provide the premarket submission tracking number.

(c) If FDA finds that any of the reasons in paragraph (a) of this section exist for refusing to accept the submission, FDA will notify the submitter in writing of the reason(s) and that the submission has not been accepted, unless insufficient contact information was provided.


Leslie Kux, Associate Commissioner for Policy.

[FR Doc. 2016–31370 Filed 12–28–16; 8:45 am]

BILLING CODE 4164–01–P

EQUAL EMPLOYMENT OPPORTUNITY COMMISSION

29 CFR Part 1610

RIN 3046–AB05

Availability of Records


ACTION: Interim final rule.

SUMMARY: The Equal Employment Opportunity Commission (EEOC or Commission) proposes to revise its Freedom of Information Act (FOIA) regulations in order to implement the substantive and procedural changes to the FOIA identified in the FOIA Improvement Act of 2016 and update two district offices addresses and the Office of Legal Counsel’s fax number.

DATES: This interim final rule is effective on December 29, 2016. Comments must be received on or before January 30, 2017.

ADDRESSES: Written comments should be submitted to Executive Secretariat, Equal Employment Opportunity Commission, 131 M Street NE., Suite 6NE03F, Washington, DC 20507. As a convenience to commenters, the Executive Secretariat will accept comments by facsimile (“FAX”) machine. The telephone number of the FAX receiver is (202) 663–4114. (This is not a toll-free FAX number). Only comments of six or fewer pages will be accepted via FAX transmital to ensure access to the equipment. Receipt of FAX transmittals will not be acknowledged, except that the sender may request confirmation of receipt by calling the Executive Secretariat staff at (202) 663–4070 (voice) or (202) 663–4074 (TTY). (These are not toll-free telephone numbers.) You may also submit comments and attachments electronically at https://www.regulations.gov, which is the Federal eRulemaking Portal. Follow the instructions online for submitting comments. Copies of comments submitted by the public will be available for review by prior appointment at the Commission’s Library, 131 M Street NE., Suite 4NW08R, Washington, DC 20507, or can be reviewed anytime at https://www.regulations.gov.

FOR FURTHER INFORMATION CONTACT:

Stephanie D. Garner, Assistant Legal Counsel (202) 663–4642 or Draga G. Anthony, Senior Attorney Advisor, Office of Legal Counsel (216) 522–7452 (voice) or (202) 663–7026 (TTY). (These are not toll free numbers.)

Requests for this document in an alternative format should be made to the Office of Communications and Legislative Affairs at (202) 663–4191 (voice) or (202) 663–4494 (TTY).

SUPPLEMENTARY INFORMATION:

Executive Summary

The interim final rule, as directed by the FOIA Improvement Act of 2016, Public Law 114–183, updates the Commission’s FOIA regulations to reflect substantive and procedural changes to the FOIA and updates the addresses of two district offices and the Office of Legal Counsel’s fax number.

Background

On June 30, 2016, President Obama signed the FOIA Improvement Act of 2016 (“Act”). The Act requires agencies to update FOIA regulations to conform to the Act by:

• Requiring federal agencies to make available their disclosable records and documents for public inspection in an electronic format;
• Making available for inspection in an electronic format records that have been requested three or more times (frequently requested records);
• Requiring that the Annual FOIA data be downloadable;
• Prohibiting agencies from charging a fee for providing records if the agency misses a deadline for complying with a FOIA request unless unusual circumstances apply and more than 5,000 pages are necessary to respond to the request;
• Prohibiting agencies from withholding information requested under FOIA Exemption (b)(5) unless the agency reasonably foresees that disclosure would harm an interest