TENNESSEE VALLEY AUTHORITY

18 CFR Part 1301

Tennessee Valley Authority Procedures for the Protection of National Security Classified Information

AGENCY: Tennessee Valley Authority.

ACTION: Final rule.

SUMMARY: The Tennessee Valley Authority is amending its regulation which contain TVA's procedure for the Protection of National Security Classified Information. These amendments reflect changes in position titles and addresses; conform the references to Protection of National Security Classified Information to the most current publication of TVA's Protection of National Security Classified Information Notices in the Federal Register.

DATES: Effective date: September 25, 2018.

FOR FURTHER INFORMATION CONTACT: Glenn Alan Spencer, Employment & Government Law Attorney, Tennessee Valley Authority, 400 W Summit Hill Dr. (WT6), Knoxville, Tennessee 37902–1401; telephone (865) 632–6255 or by email at gaspenzer@tva.gov.

SUPPLEMENTARY INFORMATION: Section 1301.63(a) currently states that Executive Order 13526 requires that each agency that originates or handles classified information designate a senior agency official to direct and administer its information security program. TVA’s senior agency official is currently the Director, Enterprise Information Security & Policy. TVA is revising §1301.63(a) to align with organizational and personnel changes within the agency.

Section 1301.67(c) currently states that requests shall be in writing, and shall be sent to: Director, Enterprise Information Security & Policy, Tennessee Valley Authority, 1101 Market St., Chattanooga, TN 37402. TVA is revising §1301.67(c) to align with organizational and personnel changes within the agency.

TVA considers this rule to be a procedural rule which is exempt from notice and comment under 5 U.S.C. 533(b)(3)(A). This rule is not a
significant rule for purposes of Executive Order 13526 and has not been reviewed by the Office of Management and Budget. As required by the Regulatory Flexibility Act, TVA certifies that these regulatory amendments will not have a significant impact on small business entities. Since this rule is non-substantive, it is being made effective September 25, 2018.

List of Subjects in 18 CFR Part 1301

Freedom of information, Government in the sunshine, Privacy, Protection of national security classified information.

For the reasons stated in the preamble, TVA amends 18 CFR part 1301 as follows:

PART 1301—PROCEDURES

§ 1301.63 Senior agency official.

1. The authority citation for part 1301 continues to read as follows:


Subpart E—Protection of National Security Classified Information

2. In § 1301.63, revise paragraph (a) to read as follows:

(a) The Executive Order requires that each agency that originates or handles classified information designate a senior agency official to direct and administer its information security program. TVA’s senior agency official is the Director, TVA Police & Emergency Management.

3. In § 1301.67, revise paragraph (c) to read as follows:

(c) Requests shall be in writing, and shall be sent to: Director, TVA Police & Emergency Management, Tennessee Valley Authority, 400 West Summit Hill Drive, Knoxville, TN 37902.

Todd M. Peney,

Director, TVA Police & Emergency Management, Tennessee Valley Authority.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 74

[Docket No. FDA–2017–C–2902]

Listing of Color Additives Subject to Certification; D&C Yellow No. 8

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA or we) is amending the color additive regulations to provide for the expanded safe use of D&C Yellow No. 8 as a color additive in contact lens solution. We are taking this action in response to a color additive petition submitted by Glo Eyes, LLC.

DATES: This rule is effective October 26, 2018. See section VIII for further information on the filing of objections. Submit either electronic or written objections and requests for a hearing on the final rule by October 25, 2018.

ADDRESSES: You may submit objections and requests for a hearing as follows. Please note that late, untimely filed objections will not be considered. Electronic objections must be submitted on or before October 25, 2018. The https://www.regulations.gov electronic filing system will accept comments until midnight Eastern Time at the end of October 25, 2018. Objections received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic objections in the following way:

• Federal eRulemaking Portal: https://www.regulations.gov. Follow the instructions for submitting comments. Objections submitted electronically, including attachments, to https://www.regulations.gov will be posted to the docket unchanged. Because your objection will be made public, you are solely responsible for ensuring that your objection does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your objection, that information will be posted on https://www.regulations.gov.

• If you want to submit an objection with confidential information that you do not wish to be made available to the public, submit the objection as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

• Mail/Hand Delivery/Courier (for written/paper submissions): Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper objections submitted to the Dockets Management Staff, FDA will post your objection, as well as any attachments, except for information submitted, marked and identified, as confidential. If submitted as detailed in “Instructions.”

Instructions: All submissions received must include the Docket No. FDA–2017–C–2902 for “Listing of Color Additives Subject to Certification; D&C Yellow No. 8.” Received objections, those filed in a timely manner (see ADDRESSES), will be placed in the docket and, except for those submitted as “Confidential Submissions,” publicly viewable at https://www.regulations.gov or with the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

• Confidential Submissions—To submit an objection with confidential information that you do not wish to be made publicly available, submit your objections only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states “THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.” The Agency will review this copy, including the claimed confidential information, in our consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on https://www.regulations.gov. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as “confidential.” Any information marked as “confidential” will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA’s posting