

safety in air commerce. This regulation is within the scope of that authority because it addresses an unsafe condition that is likely to exist or develop on products identified in this rulemaking action.

### Regulatory Findings

We determined that this proposed AD would not have federalism implications under Executive Order 13132. This proposed AD would not have a substantial direct effect 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.

For the reasons discussed above, I certify this proposed regulation:

1. Is not a “significant regulatory action” under Executive Order 12866;
2. Is not a “significant rule” under the DOT Regulatory Policies and Procedures (44 FR 11034, February 26, 1979);
3. Will not affect intrastate aviation in Alaska; and
4. Will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

### List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Incorporation by reference, Safety.

### The Proposed Amendment

Accordingly, under the authority delegated to me by the Administrator, the FAA proposes to amend 14 CFR part 39 as follows:

### PART 39—AIRWORTHINESS DIRECTIVES

- 1. The authority citation for part 39 continues to read as follows:

**Authority:** 49 U.S.C. 106(g), 40113, 44701.

#### § 39.13 [Amended]

- 2. The FAA amends § 39.13 by adding the following new airworthiness directive (AD):

**Dassault Aviation:** Docket No. FAA–2017–0502; Directorate Identifier 2016–NM–120–AD.

#### (a) Comments Due Date

We must receive comments by July 17, 2017.

#### (b) Affected ADs

None.

#### (c) Applicability

This AD applies to Dassault Aviation Model FALCON 7X airplanes, certificated in any category, manufacturer serial numbers 15 through 89 inclusive, 92 through 94

inclusive, 97 through 101 inclusive, 105, and 106.

#### (d) Subject

Air Transport Association (ATA) of America Code 51, Structures.

#### (e) Reason

This AD was prompted by a discovery of noncompliant rivets in the flight deck upper skin. We are issuing this AD to prevent interference between the rivet shank and the flight deck mounted overhead panel when the flight deck upper skin deforms due to impact (e.g., bird strike); a condition, that if not corrected, could affect the functioning of essential flight control systems, and result in reduced control of the airplane.

#### (f) Compliance

Comply with this AD within the compliance times specified, unless already done.

#### (g) Modification

Before exceeding 99 months or 4,100 flight cycles, whichever occurs first since the date of issuance of the original airworthiness certificate or the date of issuance of the original export certificate of airworthiness, modify the airplane by replacing certain MGPL type rivets installed on the flight deck skin panel with solid type-rivets, in accordance with the Accomplishment Instructions of Dassault Aviation Service Bulletin 7X–176, dated February 3, 2016.

#### (h) Other FAA AD Provisions

The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs):* The Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA, has the authority to approve AMOCs for this AD, if requested using the procedures found in 14 CFR 39.19. In accordance with 14 CFR 39.19, send your request to your principal inspector or local Flight Standards District Office, as appropriate. If sending information directly to the International Branch, send it to the attention of the person identified in paragraph (i)(2) of this AD. Information may be emailed to: [9-ANM-116-AMOC-REQUESTS@faa.gov](mailto:9-ANM-116-AMOC-REQUESTS@faa.gov). Before using any approved AMOC, notify your appropriate principal inspector, or lacking a principal inspector, the manager of the local flight standards district office/certificate holding district office.

(2) *Contacting the Manufacturer:* For any requirement in this AD to obtain corrective actions from a manufacturer, the action must be accomplished using a method approved by the Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA; or EASA; or Dassault Aviation’s EASA Design Organization Approval (DOA). If approved by the DOA, the approval must include the DOA-authorized signature.

#### (i) Related Information

(1) Refer to Mandatory Continuing Airworthiness Information (MCAI) EASA AD 2016–0124, dated June 22, 2016, for related information. This MCAI may be found on the Internet at <http://www.regulations.gov> by

searching for and locating Docket No. FAA–2017–0502.

(2) For more information about this AD, contact Tom Rodriguez, Aerospace Engineer, International Branch, ANM–116, Transport Airplane Directorate, FAA, 1601 Lind Avenue SW., Renton, WA 98057–3356; telephone 425–227–1137; fax 425–227–1149.

(3) For service information identified in this AD, contact Dassault Falcon Jet Corporation, Teterboro Airport, P.O. Box 2000, South Hackensack, NJ 07606; telephone: 201–440–6700; Internet: <http://www.dassaultfalcon.com>. You may view this service information at the FAA, Transport Airplane Directorate, 1601 Lind Avenue SW., Renton, WA. For information on the availability of this material at the FAA, call 425–227–1221.

Issued in Renton, Washington, on May 18, 2017.

**Michael Kaszycki,**

*Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.*

[FR Doc. 2017–10977 Filed 5–30–17; 8:45 am]

**BILLING CODE 4910–13–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Part 74

[Docket No. FDA–2017–C–2902]

#### Glo Eyes, LLC; Filing of Color Additive Petition

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of petition.

**SUMMARY:** The Food and Drug Administration (FDA or we) is announcing that we have filed a petition, submitted by Glo Eyes, LLC, proposing that the color additive regulations be amended to provide for the safe use of D&C Yellow No. 8 as a color additive in contact lens solution.

**DATES:** The color additive petition was filed on April 18, 2017.

**FOR FURTHER INFORMATION CONTACT:** Molly A. Harry, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1075.

**SUPPLEMENTARY INFORMATION:** Under section 721(d)(1) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379e(d)(1)), we are giving notice that we have filed a color additive petition (CAP 7C0311), submitted by Glo Eyes, LLC, 5501 Highway 199, suite 202, Fort Worth, TX 76114. The petition proposes to amend the color additive regulations in 21 CFR part 74, *Listing of Color Additives Subject To Certification*, to provide for the safe use of D&C Yellow

No. 8 (principally the disodium salt of fluorescein) as a color additive in contact lens solution.

We have determined under 21 CFR 25.32(l) 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.

Dated: May 24, 2017.

**Anna K. Abram,**

*Deputy Commissioner for Policy, Planning, Legislation, and Analysis.*

[FR Doc. 2017-11165 Filed 5-30-17; 8:45 am]

**BILLING CODE 4164-01-P**

## POSTAL SERVICE

### 39 CFR Part 20

#### International Mailing Services: Mailing Services Rules Changes

**AGENCY:** Postal Service™.

**ACTION:** Advance notice of proposed rulemaking; request for comments.

**SUMMARY:** The Postal Service is considering limiting First-Class Mail International® service to documents only.

**DATES:** Comments on this advance notice are due July 17, 2017.

**ADDRESSES:** Mail or deliver written comments to the manager, Product Classification, U.S. Postal Service, 475 L'Enfant Plaza SW., Room 4446, Washington, DC 20260-5015. You may inspect and photocopy all written comments at USPS® Headquarters Library, 475 L'Enfant Plaza SW., 11th Floor North, Washington, DC, by appointment only between the hours of 9 a.m. and 4 p.m., Monday through Friday. Call 1-202-268-2906 in advance for an appointment. Email comments, containing the name and address of the commenter, may be sent to: [ProductClassification@usps.gov](mailto:ProductClassification@usps.gov), with a subject line of "First-Class Mail International." Faxed comments are not accepted.

**FOR FURTHER INFORMATION CONTACT:** Mico Milanovic at (202) 268-5348 or Sylvia Baylis at (202) 268-6464.

**SUPPLEMENTARY INFORMATION:** This is an advance notice of the Postal Service's intent to modify some of its International Mailing rules to conform with the new Universal Postal Union (UPU) requirements for certain Letter Post mail, effective January 1, 2018. After lengthy deliberations, UPU member countries voted to identify and separate items by content as documents

versus goods, for purposes of adopting a universal, world-wide standard.

In order for the Postal Service to meet this new standard, the contents of First-Class Mail International postcard, letter, and large envelope (flat) mail; International Priority Airmail (IPA) postcard, letter, and large envelope (flat) mail; and International Surface Air Lift (ISAL) postcard, letter, and large envelope (flat) mail will be limited to documents.

Effective January 1, 2018, mailers who wish to mail any type of goods, regardless of shape, must use First Class Package International Service or another available service. Technical details and proposed IMM changes will be published at a later date and before implementation.

**Stanley F. Mires,**

*Attorney, Federal Compliance.*

[FR Doc. 2017-11120 Filed 5-30-17; 8:45 am]

**BILLING CODE 7710-12-P**

## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 52

[EPA-R03-OAR-2017-0064; FRL-9962-76-Region 3]

#### Approval and Promulgation of Air Quality Implementation Plans; Pennsylvania; Revisions to Allegheny County Health Department Rules

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Proposed rule.

**SUMMARY:** The Environmental Protection Agency (EPA) proposes to approve the state implementation plan (SIP) revision submitted by the Commonwealth of Pennsylvania for the purpose of updating the SIP to include administrative and definition amendments made to Allegheny County Health Department's (ACHD) Rules and Regulations Article XXI, Air Pollution Control. The amendments update the name of the Bureau of Environmental Quality to the Bureau of Environmental Health and revise the definition of "County Executive" to agree with the definition contained in the Allegheny County Home Rule Charter. In the Final Rules section of this **Federal Register**, EPA is approving the Commonwealth's SIP submittal as a direct final rule without prior proposal because the Agency views this as a noncontroversial submittal and anticipates no adverse comments. A detailed rationale for the approval is set forth in the direct final rule. If no adverse comments are

received in response to this action, no further activity is contemplated. If EPA receives adverse comments, the direct final rule will be withdrawn and all public comments received will be addressed in a subsequent final rule based on this proposed rule. EPA will not institute a second comment period. Any parties interested in commenting on this action should do so at this time.

**DATES:** Comments must be received in writing by June 30, 2017.

**ADDRESSES:** Submit your comments, identified by Docket ID No. EPA-R03-OAR-2017-0064 at <https://www.regulations.gov>, or via email to [rehn.brian@epa.gov](mailto:rehn.brian@epa.gov). For comments submitted at [Regulations.gov](https://www.regulations.gov), follow the online instructions for submitting comments. Once submitted, comments cannot be edited or removed from [Regulations.gov](https://www.regulations.gov). For either manner of submission, the EPA may publish any comment received to its public docket. Do not submit electronically any information you consider to be confidential business information (CBI) or other information whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.* on the web, cloud, or other file sharing system). For additional submission methods, please contact the person identified in the **FOR FURTHER INFORMATION CONTACT** section. For the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit <https://www2.epa.gov/dockets/commenting-epa-dockets>.

**FOR FURTHER INFORMATION CONTACT:** Sara Calcinore, (215) 814-2043, or by email at [calcinore.sara@epa.gov](mailto:calcinore.sara@epa.gov).

**SUPPLEMENTARY INFORMATION:** For further information, please see the information provided in the direct final action, with the same title, Revisions to Allegheny County Health Department Rules, that is located in the "Rules and Regulations" section of this **Federal Register** publication. Please note that if EPA receives adverse comment on an amendment, paragraph, or section of this rule and if that provision may be severed from the remainder of the rule, EPA may adopt as final those provisions of the rule that are not the subject of an adverse comment.