(d) Subject

Air Transport Association (ATA) of America Code 32, Landing gear.

(e) Reason

This AD was prompted by a report indicating that Belleville washers installed on the shimmy damper of the main landing gear (MLG) may fail due to fatigue. We are issuing this AD to prevent a failed washer segment migrating into the piston cavity and interfering with piston travel. As a result, the shimmy damper performance could be compromised, and an MLG shimmy could occur, potentially leading to an MLG failure and affecting the airplane's safe flight and landing.

(f) Compliance

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

(g) Maintenance or Inspection Program Revision

Within 30 days after the effective date of this AD, revise the airplane maintenance or inspection program, as applicable, by incorporating maintenance review board (MRB) report task number 320100–229, Restoration (Belleville Washer Replacement) of the MLG Shimmy Damper, of the MRB Report of the Bombardier CRJ700/900/1000 Maintenance Requirements Manual (MRM)—Part 1, Volume 1, CSP B–053, Revision 17, dated June 25, 2017. The initial compliance time for MRB report task number 320100–229 is specified in paragraphs (g)(1) and (g)(2) of this AD, as applicable.

(1) For any shimmy damper with 20,000 total accumulated flight cycles or fewer as of the effective date of this AD, the initial compliance time is before the accumulation of 26,000 total flight cycles.

(2) For any shimmy damper with 20,000 total accumulated flight cycles or more as of the effective date of this AD, the initial compliance time is specified in paragraph (g)(2)(i) or (g)(2)(ii), whichever occurs later.

(i) Within 6,000 flight cycles after the effective date of this AD, but prior to the accumulation of 30,000 total flight cycles.

(ii) Within 30 days after effective date of this AD.

(h) Credit for Previous Actions

This paragraph provides credit for actions required by paragraph (g) of this AD, if those actions were performed before the effective date of this AD using Bombardier Temporary Revision MRB–0070, dated October 20, 2015.

(i) No Alternative Actions and/or Intervals

After the airplane maintenance or inspection program has been revised, as required by paragraph (g) of this AD, no alternative actions (*e.g.,* inspections) and/or intervals may be used unless the actions and/or intervals are approved as an alternative method of compliance (AMOC) in accordance with the procedures specified in paragraph (j)(1) of this AD.

(j) Other FAA AD Provisions

The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs):* The Manager, New York ACO Branch, 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 manager of the certification office, send it to ATTN: Program Manager, Continuing Operational Safety, FAA, New York ACO Branch, 1600 Stewart Avenue, Suite 410, Westbury, NY 11590; telephone 516–228–7300; fax 516–794–5531. 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, New York ACO Branch, FAA; or Transport Canada Civil Aviation (TCCA); or Bombardier Inc.'s TCCA Design Approval Organization (DAO). If approved by the DAO, the approval must include the DAO-authorized signature.

(k) Related Information

(1) Refer to Mandatory Continuing Airworthiness Information (MCAI) Canadian Airworthiness Directive CF–2017–14, dated April 21, 2017, for related information. This MCAI may be found in the AD docket on the internet at *http://www.regulations.gov* by searching for and locating Docket No. FAA–2017–1175.

(2) For more information about this AD, contact Cesar Gomez, Aerospace Engineer, Airframe and Mechanical Systems Section, FAA, New York ACO Branch, 1600 Stewart Avenue, Suite 410, Westbury, NY 11590; telephone 516–228–7318; fax 516–794–5531.

(3) Service information identified in this AD that is not incorporated by reference is available at the addresses specified in paragraphs (l)(3) and (l)(4) of this AD.

(l) Material Incorporated by Reference

(1) The Director of the Federal Register approved the incorporation by reference (IBR) of the service information listed in this paragraph under 5 U.S.C. 552(a) and 1 CFR part 51.

(2) You must use this service information as applicable to do the actions required by this AD, unless this AD specifies otherwise.

(i) Maintenance review board (MRB) report task number 320100–229, Restoration (Belleville Washer Replacement) of the MLG Shimmy Damper, of the MRB Report of the Bombardier CRJ700/900/1000 Maintenance Requirements Manual (MRM)—Part 1, Volume 1, CSP B–053, Revision 17, dated June 25, 2017.

(ii) Reserved.

(3) For service information identified in this AD, contact Bombardier, Inc., 400 Côte-Vertu Road West, Dorval, Québec H4S 1Y9, Canada; Widebody Customer Response Center North America toll-free telephone 1–866–538–1247 or direct-dial telephone 1–514–855–2999; fax 514–855–7401; email *ac.yul@aero.bombardier.com;* internet *http://www.bombardier.com.*

(4) You may view this service information at the FAA, Transport Standards Branch, 2200 South 216th St., Des Moines, WA. For information on the availability of this material at the FAA, call 206–231–3195.

(5) You may view this service information that is incorporated by reference at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call 202–741–6030, or go to: *http://www.archives.gov/federal-register/cfr/ibr-locations.html.*

Issued in Des Moines, Washington, on May 21, 2018.

James Cashdollar,

Acting Director, System Oversight Division, Aircraft Certification Service.

[FR Doc. 2018–11827 Filed 6–6–18; 8:45 am]

BILLING CODE 4910–13–P

DELAWARE RIVER BASIN COMMISSION

18 CFR Parts 401 and 420

Regulatory Program Fees and Water Charges Rates

AGENCY: Delaware River Basin Commission.

ACTION: Final rule.

SUMMARY: Notice is provided of the Commission's regulatory program fees and schedule of water charges for the fiscal year beginning July 1, 2018.

DATES: This final rule is effective July 1, 2018.

FOR FURTHER INFORMATION CONTACT: Elba L. Deck, CPA, Director of Administration and Finance, 609–883–9500, ext. 201.

SUPPLEMENTARY INFORMATION: The Delaware River Basin Commission (''DRBC'' or ''Commission'') is a Federal-interstate compact agency charged with managing the water resources of the Delaware River Basin on a regional basis without regard to political boundaries. Its members are the governors of the four basin states—Delaware, New Jersey, New York and Pennsylvania—and on behalf of the federal government, the North Atlantic Division Commander of the U.S. Army Corps of Engineers.

In accordance with 18 CFR 401.43(c), on July 1 of every year beginning July 1, 2017, the Commission's regulatory program fees as set forth in Tables 1, 2 and 3 of that section are subject to an annual adjustment, 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. Pursuant to 18 CFR 420.43(c), the same indexed adjustment applies to the Commission's schedule of water charges for consumptive and non-consumptive withdrawals of surface water within the basin. The referenced April 12-month CPI for 2018 showed an increase of 1.38%. Commensurate adjustments are thus required.

This notification is made in accordance with 18 CFR 401.42(c) and 18 CFR 420.42(c), which provide that a revised fee schedule will be published in the **Federal Register** by July 1. The revised fees also may be obtained by contacting the Commission during business hours or by checking the Commission's website.

List of Subjects

18 CFR Part 401

Administrative practice and procedure, Project review, Water pollution control, Water resources.

18 CFR Part 420

Water supply.

For the reasons set forth in the preamble, the Delaware River Basin Commission amends parts 401 and 420 of title 18 of the Code of Federal Regulations as set forth below:

PART 401—RULES OF PRACTICE AND PROCEDURE

■ 1. The authority citation for part 401 continues to read as follows:

Authority: Delaware River Basin Compact (75 Stat. 688), unless otherwise noted.

Subpart C—Project Review Under Section 3.8 of the Compact

■ 2. In § 401.43, revise Tables 1, 2 and 3 to read as follows:

§ 401.43 Regulatory program fees.

* * * * *

TABLE 1 TO § 401.43—DOCKET APPLICATION FILING FEE

Project type	Docket application fee	Fee maximum
Water Allocation	$411 per million gallons/month of allocation [1], not to exceed $15,401 [1]. Fee is doubled for any portion to be exported from the basin.	Greater of: $15,401 [1] or Alternative Review Fee.
Wastewater Discharge	Private projects: $1,027 [1] Public projects: $513 [1]	Alternative Review Fee.
Other	0.4% of project cost up to $10,000,000 plus 0.12% of project cost above $10,000,000 (if applicable), not to exceed $77,003 [1].	Greater of: $77,003 [1] or Alternative Review Fee.

[1] Subject to annual adjustment in accordance with paragraph (c) of this section.

TABLE 2 TO § 401.43—ANNUAL MONITORING AND COORDINATION FEE

	Annual fee	Allocation
Water Allocation	[1] $308	<4.99 mgm.
	[1] $462	5.00 to 49.99 mgm.
	[1] $667	50.00 to 499.99 mgm.
	[1] $847	500.00 to 9,999.99 mgm.
	[1] $1,027	> or = to 10,000 mgm.
	Annual fee	**Discharge design capacity**
Wastewater Discharge	[1] $308	<0.05 mgd.
	[1] $626	0.05 to 1 mgd.
	[1] $842	1 to 10 mgd.
	[1] $1,027	>10 mgd.

[1] Subject to annual adjustment in accordance with paragraph (c) of this section.

TABLE 3 TO § 401.43—ADDITIONAL FEES

Proposed action	Fee	Fee maximum
Emergency Approval Under 18 CFR 401.40	$5,000	Alternative Review Fee.
Late Filed Renewal Surcharge	$2,000.	
Modification of a DRBC Approval	At Executive Director's discretion, Docket Application Fee for the appropriate project type.	Alternative Review Fee.
Name change	$1,027 [1].	
Change of Ownership	$1,540 [1].	

[1] Subject to annual adjustment in accordance with paragraph (c) of this section.

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. In § 420.41, revise paragraphs (a) and (b) to read as follows:

§ 420.41 Schedule of water charges.

* * * * *

(a) $82.14 per million gallons for consumptive use, subject to paragraph (c) of this section; and

(b) $0.82 per million gallons for non-consumptive use, subject to paragraph (c) of this section.

* * * * *

Dated: June 1, 2018.

Pamela M. Bush,

Commission Secretary.

[FR Doc. 2018–12258 Filed 6–6–18; 8:45 am]

BILLING CODE 6360–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 74

[Docket No. FDA–2017–C–0935]

Listing of Color Additives Subject to Certification; D&C Black No. 4

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 safe use of D&C Black No. 4 for coloring ultra-high molecular weight polyethylene (UHMWPE) non-absorbable sutures for use in general surgery. This action is in response to a color additive petition (CAP) submitted by DSM Biomedical.

DATES: This rule is effective July 10, 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 July 9, 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 July 9, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of July 9, 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–0935 for ''Listing of Color Additives Subject to Certification; D&C Black No. 4.'' 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 at 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.'' We 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 of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf*.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Joseph M. Thomas, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740–3835, 301–796–9465.

SUPPLEMENTARY INFORMATION:

I. Introduction

In the **Federal Register** on March 6, 2017 (82 FR 12531), we announced that we filed a color additive petition (CAP 7C0310) submitted by DSM Biomedical (petitioner), 735 Pennsylvania Dr., Exton, PA 19341. The petition proposed to amend the color additive regulations in part 73 (21 CFR part 73), *Listing of Color Additives Exempt from Certification,* to provide for the safe use of high-purity carbon black for coloring UHMWPE non-absorbable sutures for use in general surgery.[1] After the petition was filed and during our review, we determined that the color additive will require batch certification by FDA. We intend to give each certified batch of the subject color additive the name D&C Black No. 4. Therefore, this color additive will be identified as D&C Black No. 4 and will be listed in part 74 (21 CFR part 74), *Listing of Color Additives Subject to Certification.*

II. Identity and Specifications

D&C Black No. 4 is a high-purity carbon black prepared by the oil furnace process. It is manufactured by injecting

[1] The original petition did not specify that the color additive is to be used in sutures that are non-absorbable. Therefore, our March 6, 2017, notice of filing did not specify that the color additive is intended for use in non-absorbable sutures. However, petitioner's subsequent submissions to FDA indicated that the intended use of the additive is for sutures that are non-absorbable.