Table 1 to Paragraph (g)(1) – Initial Inspection of Affected Part

FCs Accumulated (since new)	Compliance Time
700 FCs or less.	Before exceeding 500 FCs, or within 100 FCs after the effective date of this AD, whichever occurs later
More than 700 FCs up to 1,000 FCs (inclusive).	Within 50 FCs after the effective date of this AD
1,001 FCs or greater.	Within 25 FCs or 30 calendar days, whichever occurs first after the effective date of this AD

(2) An in-shop BSI in accordance with Accomplishment Instructions, paragraph 3.A, of RR Trent 1000 Alert NMSB 72–AK451, Initial Issue, dated November 14, 2019, may be substituted for any on-wing BSI, provided the compliance time specified in Table 1 to paragraph (g)(1) of this AD is not exceeded.

(3) If, during any initial or repetitive BSI of the IPC shaft assembly required by paragraph (g)(1) or (2) of this AD, any crack is detected, before further flight, remove the IPC shaft assembly and replace it with a part eligible for installation.

(h) Definitions

For the purpose of this AD, a ''part eligible for installation'' is:

(1) An IPC shaft assembly that is new (not previously installed on an engine);

(2) An IPC shaft assembly that, before (re)installation, has passed an inspection (no crack detected) in accordance with Accomplishment Instructions, paragraph 3.B., of RR Trent 1000 Alert NMSB 72–AK451, Initial Issue, dated November 14, 2019.

(i) No Reporting Requirement

The reporting requirements in the Accomplishment Instructions, paragraphs 3.A. and 3.B., of RR Trent 1000 Alert NMSB 72–AK451, Initial Issue, dated November 14, 2019, are not required by this AD.

(j) Credit for Previous Actions

You may take credit for the initial BSI of the IPC shaft assembly that is required by paragraph (g)(1) of this AD if you performed the BSI before the effective date of this AD using RR Trent 1000 NMSB 72–K452, Initial Issue, dated October 21, 2019.

(k) Alternative Methods of Compliance (AMOCs)

(1) The Manager, ECO 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 ECO Branch, send it to the attention of the person identified in paragraph (l)(1) of this AD. You may email your request to: *ANE-AD-AMOC@faa.gov.*

(2) 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.

(l) Related Information

(1) For more information about this AD, contact Stephen Elwin, Aerospace Engineer, ECO Branch, FAA, 1200 District Avenue, Burlington, MA, 01803; phone: 781–238–7236; fax: 781–238–7199; email: *stephen.l.elwin@faa.gov.*

(2) Refer to European Union Aviation Safety Agency (EASA) AD 2019–0282, dated November 20, 2019, for more information. You may examine the EASA AD in the AD docket on the internet at *https://www.regulations.gov* by searching for and locating it in Docket No. FAA–2020–0293.

(3) For service information identified in this AD, contact Rolls-Royce Deutschland Ltd & Co KG, Eschenweg 11, 15827 Blankenfelde-Mahlow, Germany; phone: +49 (0) 33 708 6 0; email: *https://www.rolls-royce.com/contact-us.aspx.* You may view this referenced service information at the FAA, Engine and Propeller Standards Branch, 1200 District Avenue, Burlington, MA, 01803. For information on the availability of this material at the FAA, call 781–238–7759.

Issued on March 26, 2020.

Lance T. Gant,

Director, Compliance & Airworthiness Division, Aircraft Certification Service.

[FR Doc. 2020–06736 Filed 4–1–20; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Office of the Secretary

15 CFR Part 4

[Docket No. 200117–0024]

RIN 0605–AA49

Social Security Number Fraud Prevention Act of 2017 Implementation

AGENCY: Office of the Secretary, Department of Commerce.

ACTION: Notice of proposed rulemaking.

SUMMARY: This proposed rule would revise the Department of Commerce (Department) regulations under the Freedom of Information Act (FOIA) and the Privacy Act. The revisions would clarify and update the language of procedural requirements pertaining to the inclusion of Social Security account numbers (SSNs) on documents that the Department sends by mail. These revisions are necessary to implement the Social Security Number Fraud Prevention Act of 2017, which restricts the inclusion of Social Security Numbers (SSNs) on documents sent by mail by the Federal Government.

DATES: Submit comments on or before April 24, 2020. Comments received by mail will be considered timely if they are postmarked on or before that date. The electronic Federal Docket Management System (FDMS) will accept comments until Midnight Eastern Time at the end of that day.

ADDRESSES: You may submit comments, identified by Regulatory Information Number (RIN) 0605–AA49, by any of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Departmental Privacy Act Officer, Office of Privacy and Open

Government, Department of Commerce, 1401 Constitution Ave. NW, Mail Stop 61025, Washington, DC 20230.

Instructions: All submissions received must include the agency name and docket number or RIN for this rulemaking. All comments received will be posted without change to regulations.gov, including any personal information provided. Comments sent by any other method, to any other address or individual, or received after the end of the comment period, may not be considered by the Department.

FOR FURTHER INFORMATION CONTACT: Departmental Privacy Act Officer, Office of Privacy and Open Government, United States Department of Commerce, (202) 482–1190.

SUPPLEMENTARY INFORMATION: The Social Security Number Fraud Prevention Act of 2017 (the Act) (Pub. L. 115–59; 42 U.S.C. 405 note), which was signed on September 15, 2017, restricts Federal agencies from including individuals' SSNs on documents sent by mail, unless the head of the agency determines that the inclusion of the SSN on the document is necessary (section 2(a) of the Act). The Act requires agency heads to issue regulations specifying the circumstances under which inclusion of a SSN on a document sent by mail is necessary. These regulations, which must be issued not later than five years after the date of enactment, shall include instructions for the partial redaction of SSNs where feasible, and shall require that SSNs not be visible on the outside of any package sent by mail (section 2(b) of the Act). This proposed rule would revise the Department regulations under the Freedom of Information Act (FOIA) (subpart A, 15 CFR part 4) and the Privacy Act (subpart B, 15 CFR part 4), consistent with these requirements in the Act. The proposed revisions would clarify the language of procedural requirements pertaining to the inclusion of SSNs on documents that the Department sends by mail. The proposed rule also makes clarifying updates by changing the term ''Privacy Officer'' to ''Privacy Act Officer'' where it occurs in Subpart B of 15 CFR part 4, and by changing the term ''FOI Officer'' to ''FOIA Officer'' in several places in Appendix B, The proposed rule also updates an office name by changing the phrase ''Assistant General Counsel for Employment, Litigation, and Oversight'' to ''Assistant General Counsel for Employment, Litigation, and Information'' where it occurs in part 4.

Classification

This proposed rule has been determined to be significant for purposes of review under Executive Order 12866. This proposed rule is not subject to the requirements of Executive Order 13771 because it is expected to result in no more than *de minimis* costs to citizens and residents of the United States. In accordance with the Regulatory Flexibility Act (5 U.S.C. 605(b)), the Chief Counsel for Regulation has reviewed this rule and certifies that this regulation, if implemented, will not have a significant economic impact on a substantial number of small entities. This rule is largely procedural in nature, and, therefore, will not affect requesters. This regulation does not contain a collection of information as defined by the Paperwork Reduction Act, 44 U.S.C. 3501, *et seq.*

List of Subjects in 15 CFR Part 4

Appeals, Freedom of Information Act, Information, Privacy, Privacy Act.

Catrina D. Purvis,

Chief Privacy Officer, and Director of Open Government.

For the reasons stated in the preamble, the Department of Commerce proposes to amend Subpart B of 15 CFR part 4 as follows:

PART 4—DISCLOSURE OF GOVERNMENT INFORMATION

■ 1. The authority citation for part 4 continues to read as follows:

Authority: 5 U.S.C. 301; 5 U.S.C. 552; 5 U.S.C. 552a; 5 U.SC. 553; 31 U.S.C. 3717; 44 U.S.C. 3101; Reorganization Plan No. 5 of 1950; Pub. L. 115–59, 131 Stat. 1152 (42 U.S.C. 405, note).

Subpart A—Freedom of Information Act

■ 2. In § 4.7, revise paragraph (d) to read as follows:

§ 4.7 Responses to Requests.

* * * * *

(d) All responses shall be made subject to the provisions of § 4.25(b)(2)(iv).

* * * * *

Subpart B [Amended]

■ 3. Amend subpart B by removing the words ''Privacy Officer'' wherever they appear and adding in their place the words ''Privacy Act Officer''.

■ 4. Amend § 4.22:

■ a. In paragraph (b)(7), emoving the words ''Privacy Officer'' and adding, in their place, the words ''Privacy Act Officer''; and

■ b. Adding new paragraph (b)(10).

The addition reads as follows:

§ 4.22 Definitions

* * * * *

(b) * * *

(10) *Un-redacted SSN Mailed Documents Listing (USMDL)* means the Department approved list, as posted at *www.commerce.gov/privacy,* designating those_documents for which the inclusion of the Social Security number (SSN) is determined to be necessary to fulfill a compelling Department business need when the documents are requested by individuals outside the Department or other Federal agencies, as determined jointly by the Senior Agency Official for Privacy and the Departmental Privacy Act Officer.

■ 5. Amend § 4.25 by:

■ a. Adding paragraphs (a)(3) and (4); and

■ b. Revising paragraph (b)(2)(iii), and adding paragraphs (b)(2)(iv) and (v).

The additions and revisions read as follows:

§ 4.25 Disclosure of requested records to individuals [Amended]

(a) * * *

(3) Inclusion of Social Security Numbers (SSNs) on responsive documents.

The Department shall redact SSNs from responsive documents provided to requesters where feasible. Where full redaction is not feasible, partial redaction to create a truncated SSN shall be preferred to no redaction.: The following conditions must be met for the inclusion of an unredacted (full) SSN or partially redacted (truncated) SSN on a responsive document:

(i) The inclusion of the full SSN or truncated SSN of an individual must be required or authorized by law,

(ii) The inclusion of the full SSN or truncated SSN of an individual must be determined by the Senior Agency Official for Privacy and Departmental Privacy Act Officer to be necessary to fulfill a compelling Department business need; and

(iii) The full SSN of an individual may be included only on documents listed on the USMDL.

(4) The following requirements apply when the Department mails or delivers responsive documents containing SSNs or truncated SSNs:

(i) The full SSN of an individual may be included only on documents listed on the USMDL.

(ii) For documents that are listed on the USMDL and that include the full SSN of an individual, the signature of the recipient is required upon delivery.

(iii) For documents that include the truncated form of the SSN of an individual, the signature of the recipient is required upon delivery.

(iv) The full SSN, the truncated SSN, any part of the SSN of an individual must not be visible from the outside of the envelope or package.

(b) * * *

(2) * * *

(iii) Copies of documents may be mailed at the request of the individual, and may be subject to payment of the fees prescribed in §§ 4.25(a)(3) and 4.31. In the event that the Department, at its own initiative, elects to provide a copy by mail, no fee will be charged to the individual.

(iv) Copies of documents listed on the USMDL, include full SSNs, and are requested by an individual are subject to payment of the fees prescribed in § 4.31.

(v) Documents containing SSNs or truncated SSNs that are required to be returned by the individual to the Department will be mailed or delivered along with a prepaid mail or delivery service envelope at the expense of the Department.

* * * * *

Appendix B to Part 4 [Amended]

■ 6. Amend Appendix B to part 4 by adding the word ''Act'' after the phrase ''Departmental Freedom of Information'' wherever it appears, after the phrase ''Executive Secretary; Freedom of Information'', and before the phrase ''Officer for the Office of the Secretary''.

[FR Doc. 2020–06490 Filed 4–1–20; 8:45 am]

BILLING CODE 3510–17–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 866

[Docket No. FDA–2020–N–1088]

Microbiology Devices; Reclassification of Nucleic Acid-Based Hepatitis C Virus Ribonucleic Acid Assay Devices, To Be Renamed Nucleic Acid-Based Hepatitis C Virus Ribonucleic Acid Tests

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed amendment; proposed order; request for comments.

SUMMARY: The Food and Drug Administration (FDA or Agency) is proposing to reclassify nucleic acid-based hepatitis C virus (HCV) ribonucleic acid (RNA) devices intended for the qualitative or quantitative detection or genotyping of HCV RNA, postamendments class III devices (product codes MZP and OBF), into class II (general controls and special controls), subject to premarket notification. FDA is also proposing a new device classification regulation with the name ''nucleic acid-based Hepatitis C virus (HCV) ribonucleic acid tests'' along with the special controls that the Agency believes are necessary to provide a reasonable assurance of safety and effectiveness for these devices. FDA is proposing this reclassification on its own initiative. If finalized, this order will reclassify these types of devices from class III (general controls and premarket approval) to class II (general controls and special controls) and reduce the regulatory burdens associated with these devices, as these types of devices will no longer be required to submit a premarket approval application (PMA), but can instead submit a premarket notification (510(k)) and obtain clearance before marketing their device.

DATES: Submit either electronic or written comments on the proposed order by June 1, 2020. Please see section XI of this document for the proposed effective date when the new requirements apply and for the proposed effective date of a final order based on this proposed order.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before June 1, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of June 1, 2020. Comments received by Mail/Hand Delivery/Courier (for written/paper submissions) will be considered timely.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment 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 comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed below (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 comments submitted to the Dockets Management Staff, FDA will post your comment, 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–2020–N–1088 for ''Reclassification of Nucleic Acid-Based Hepatitis C Virus Ribonucleic Acid Assay Devices, To Be Renamed Nucleic Acid-Based Hepatitis C Virus Ribonucleic Tests.'' Received comments, 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 a comment with confidential information that you do not wish to be made publicly available, submit your comments 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 its 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