Manti, UT, Manti-Ephraim, Takeoff Minimums and Obstacle DP, Orig
Manti, UT, Manti-Ephraim, WUXOT ONE, Graphic DP
Manti, UT, Manti-Ephraim, YMONT ONE, Graphic DP
Marion/Wytheville, VA, Mountain Empire, LOC RWY 26, Amdt 3
Marion/Wytheville, VA, Mountain Empire, RNAV (GPS) RWY 26, Amdt 1
Wenatchee, WA, Pangborn Memorial, WENATCHEE TWO, Graphic DP
Black River Falls, WI, Black River Falls Area, RNAV (GPS) RWY 26, Orig-B
Necedah, WI, Necedah, RNAV (GPS) RWY 36, Orig-D
Racine, WI, Batten Intl, Takeoff Minimums and Obstacle DP, Amdt 5A
Stevens Point, WI, Stevens Point Muni, ILS OR LOC RWY 21, Amdt 1
Stevens Point, WI, Stevens Point Muni, RNAV (GPS) RWY 21, Amdt 1
Wausau, WI, Wausau Downtown, RNAV (GPS) RWY 31, Amdt 1
Cheyenne, WY, Cheyenne Rgnl/Jerry Olson Field, ILS OR LOC RWY 27, Amdt 35A

Rescinded: On April 10, 2017 (82 FR 17117), the FAA published an Amendment in Docket No. 31125, Amdt No. 3739 to Part 97 of the Federal Aviation Regulations under section 97.33, the following entries for Majuro Atoll, RM, effective April 27, 2017, and are hereby rescinded in their entirety:

Majuro Atoll, RM, Marshall Islands Intl, RNAV (GPS) RWY 7, Orig-D
Majuro Atoll, RM, Marshall Islands Intl, RNAV (GPS) RWY 25, Orig-D

[FR Doc. 2017–09908 Filed 5–17–17; 8:45 am]

BILLING CODE 4910–13–P

SOCIAL SECURITY ADMINISTRATION

20 CFR Part 421

[Docket No. SSA–2016–0011]

RIN 0960–AH95

Implementation of the NICS Improvement Amendments Act of 2007

AGENCY: Social Security Administration.

ACTION: Final rule; CRA Revocation.

SUMMARY: We are removing from the Code of Federal Regulations the final rules, Implementation of the NICS Improvement Amendments Act of 2007 (NIAA), published on December 19, 2016. We are doing so because Congress passed, and the President signed, a joint resolution of disapproval of the final rules under the Congressional Review Act.

DATES: This rule removal is effective on May 18, 2017.

FOR FURTHER INFORMATION CONTACT: Social Security Administration, 410–965–3735 or *Regulations@ssa.gov.* For information on eligibility or filing for benefits, call our national toll-free number, 1–800–772–1213 or TTY 1–800–325–0778, or visit our Internet site, Social Security Online, at *http://www.socialsecurity.gov.*

SUPPLEMENTARY INFORMATION: On May 5, 2016, we published a notice of proposed rulemaking (NPRM) in the **Federal Register** (81 FR 27059) in which we proposed adding part 421 to our regulations to fulfill responsibilities that we have under the NIAA. On December 19, 2016, we published a final rule (81 FR 91702) for the Implementation of the NICS Improvement Amendments Act of 2007 (NIAA), which had an effective date of January 18, 2017.[1] On February 2, 2017, the United States House of Representatives passed H.J. Res. 40, ''Providing for congressional disapproval under chapter 8 of title 5, United States Code, of the rule submitted by the Social Security Administration relating to Implementation of the NICS Improvement Amendments Act of 2007 (NIAA).''.[2] On February 15, 2017, the United States Senate passed H.J. Res. 40 without amendment,[3] and the President signed H.J. Res. 40 into law on February 28, 2017.[4] Under the terms of Public Law 115–8, the final rules ''shall have no force or effect.'' As a result, we are removing them from the Code of Federal Regulations.

Authority for removal: This document was prepared under the direction of Nancy A. Berryhill, Acting Commissioner of Social Security. We issued it under the authority of section 702 of the Social Security Act (42 U.S.C. 902(a)(5)), and Public Law 115–8, 131 Stat. 15.

List of Subjects in 20 CFR Part 421

Administrative practice and procedure, Freedom of information, Privacy, Reporting and recordkeeping requirements.

Nancy A. Berryhill,

Acting Commissioner of Social Security.

Under the authority of section 702 of the Social Security Act (42 U.S.C. 902(a)(5)), the Congressional Review Act (5 U.S.C. 801 *et seq.*), and Public Law 115–8, 131 Stat. 15, and for the reasons set out in the preamble, we amend title 20, chapter III, of the Code of Federal Regulations as follows:

PART 421—[REMOVED]

■ 1. Remove part 421, consisting of §§ 421.100 through 421.170.

[FR Doc. 2017–10084 Filed 5–17–17; 8:45 am]

BILLING CODE 4191–02–P

[1] Although the final rule had an effective date of January 18, 2017, we delayed the compliance date of the rule until December 19, 2017 (81 FR at 91720). Therefore, we did not report any records to the National Instant Criminal Background Check System (NICS) pursuant to the final rule.

[2] 163 Cong. Rec. H916 (daily ed. Feb. 2, 2017).

[3] 163 Cong. Rec. S1169 (daily ed. Feb. 15, 2017).

[4] Public Law 115–8, 131 Stat. 15.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 201, 801, and 1100

[Docket No. FDA–2015–N–2002]

RIN 0910–AH19

Clarification of When Products Made or Derived From Tobacco Are Regulated as Drugs, Devices, or Combination Products; Amendments to Regulations Regarding ''Intended Uses''; Further Delayed Effective Date; Request for Comments; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; extension of comment period.

SUMMARY: In the **Federal Register** of January 9, 2017, the Food and Drug Administration (FDA or the Agency) issued a final rule entitled ''Clarification of When Products Made or Derived From Tobacco Are Regulated as Drugs, Devices, or Combination Products; Amendments to Regulations Regarding 'Intended Uses' '' (Final Rule). On March 20, 2017, FDA published a document in the **Federal Register** (Final Rule Extension) to delay the effective date of the Final Rule until March 19, 2018, and requested comments on particular issues raised in a petition for reconsideration and stay of action of the Final Rule. The petition for reconsideration raised questions about the amendments to the regulations regarding ''intended uses'' that are set forth in the Final Rule. In the Final Rule Extension FDA also requested comments regarding any aspect of the Final Rule, or with respect to issues relating to ''intended uses'' generally, and on whether the delay in the effective date should be modified or revoked. FDA is now issuing this document to extend the comment period. The Agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.