

paragraphs (d)(5)(vi)(A) and (d)(5)(vi)(B);

- f. In newly redesignated paragraph (d)(5)(vi)(A), removing the word “animal” in paragraph (a) and adding in its place the word “nonclinical”; and
- g. In newly redesignated paragraph (d)(5)(vi)(B), removing the word “animal” in paragraph (b) and adding in its place the word “nonclinical”.

§ 314.81 [Amended]

- 13. Amend § 314.81 by:

- a. In paragraph (b)(2)(v), removing the phrase “toxicological findings in animal studies and in vitro studies (e.g., mutagenicity)” and adding in its place the phrase “nonclinical toxicological findings, including, for example, from mutagenicity studies”; and
- b. In paragraph (b)(2)(vii)(a)(7), removing the phrase “an animal” and adding in its place the phrase “a nonclinical”.

§ 314.93 [Amended]

- 14. In § 314.93, amend paragraph (e)(2) by removing the word “animal” and adding in its place the word “nonclinical”.

§ 314.200 [Amended]

- 15. In § 314.200, in the analysis format in paragraph (d)(3), amend the heading for item I.A. by removing the word “Animal” and adding in its place the word “Nonclinical.”

§ 314.430 [Amended]

- 16. In § 314.430, amend paragraph (a) by removing the phrase “all studies and tests of a drug on animals and humans” and adding in its place the phrase “all nonclinical and clinical studies and tests of a drug”.

PART 315—DIAGNOSTIC RADIOPHARMACEUTICALS

- 17. The authority citation for part 315 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 355, 371, 374, 379e; sec. 122, Pub. L. 105–115, 111 Stat. 2322 (21 U.S.C. 355 note).

- 18. Revise § 315.2 to read as follows:

§ 315.2 Definitions.

(a) For purposes of this part, *diagnostic radiopharmaceutical* means:

(1) An article that is intended for use in the diagnosis or monitoring of a disease or a manifestation of a disease in humans and that exhibits spontaneous disintegration of unstable nuclei with the emission of nuclear particles or photons; or

(2) Any nonradioactive reagent kit or nuclide generator that is intended to be used in the preparation of such article

as defined in paragraph (a)(1) of this section.

(b) For purposes of this part, *nonclinical study* means a test or study conducted in vitro, in silico, or in chemico, or a nonhuman in vivo test or study. Such test or study may include the following:

- (1) Cell-based assays.
- (2) Organ chips and microphysiological systems.
- (3) Computer modeling.
- (4) Other nonhuman or human biology-based test methods, such as bioprinting.
- (5) Animal tests or studies.

§ 315.6 [Amended]

- 19. Amend § 315.6 by:

- a. In paragraph (c)(2), removing the word “preclinical” and adding in its place the word “nonclinical”; and
- b. In paragraph (d), removing the word “animal” and adding in its place the word “nonclinical”.

PART 361—PRESCRIPTION DRUGS FOR HUMAN USE GENERALLY RECOGNIZED AS SAFE AND EFFECTIVE AND NOT MISBRANDED: DRUGS USED IN RESEARCH

- 20. The authority citation for part 361 continues to read as follows:

Authority: 21 U.S.C. 321, 351, 352, 353, 355, 371; 42 U.S.C. 262.

§ 361.1 [Amended]

- 21. In § 361.1, amend paragraph (d)(7) by removing the phrase “animal studies” and adding in its place the phrase “nonclinical studies, as defined in § 312.3(b) of this chapter”.

PART 601—LICENSING

- 22. The authority citation for part 601 continues to read as follows:

Authority: 15 U.S.C. 1451–1561; 21 U.S.C. 321, 351, 352, 353, 355, 356b, 360, 360c–360f, 360h–360j, 371, 374, 379e, 381; 42 U.S.C. 216, 241, 262, 263, 264; sec. 122, Pub. L. 105–115, 111 Stat. 2322 (21 U.S.C. 355 note), sec. 7002(e), Pub. L. 111–148, 124 Stat. 817, as amended by sec. 607, Division N, Pub. L. 116–94, 133 Stat. 3127.

- 23. Revise § 601.31 is amended to read as follows:

§ 601.31 Definitions.

(a) For purposes of this part, *diagnostic radiopharmaceutical* means:

(1) An article that is intended for use in the diagnosis or monitoring of a disease or a manifestation of a disease in humans and that exhibits spontaneous disintegration of unstable nuclei with the emission of nuclear particles or photons; or

(2) Any nonradioactive reagent kit or nuclide generator that is intended to be

used in the preparation of such article as defined in paragraph (a)(1) of this section.

(b) For purposes of this part, *nonclinical study* means a test or study conducted in vitro, in silico, or in chemico, or a nonhuman in vivo test or study. Such test or study may include the following:

- (1) Cell-based assays.
- (2) Organ chips and microphysiological systems.
- (3) Computer modeling.
- (4) Other nonhuman or human biology-based test methods, such as bioprinting.
- (5) Animal tests or studies.

§ 601.35 [Amended]

- 24. Amend § 601.35 by:

- a. In paragraph (c)(2), removing the word “preclinical” and adding in its place the word “nonclinical”; and
- b. In paragraph (d), removing the word “animal” and adding in its place the word “nonclinical”.

§ 601.70 [Amended]

- 25. In § 601.70, amend paragraph (b)(7) by removing the phrase “an animal” and adding in its place the phrase “a nonclinical”.

Robert F. Kennedy, Jr.,

Secretary, Department of Health and Human Services.

[FR Doc. 2026–19350 Filed 9–21–26; 8:45 am]

BILLING CODE 4164–01–P

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 73

[MB Docket Nos. 19–310, 17–105; FCC 24–66; FR ID 368194]

Reinstatement of Radio Non-Duplication Rule for Commercial FM Stations; Correction

AGENCY: Federal Communications Commission.

ACTION: Correcting amendment.

SUMMARY: This document corrects the final rule portion of a **Federal Register** document published on July 3, 2024. That document inadvertently included an error in a section heading of regulatory text. This document corrects the final regulations.

DATES: Effective on September 22, 2026.
FOR FURTHER INFORMATION CONTACT: John Bat, Media Bureau, Industry Analysis Division, John.Bat@fcc.gov, (202) 418–7921.

SUPPLEMENTARY INFORMATION: This document corrects the final rule document published at 89 FR 55078,

July 3, 2024. It corrects the section heading for the radio non-duplication rule in 47 CFR 73.3556 to “Duplication of programming on commonly owned or time brokered stations” from “Sponsorship identification; list retention; related requirements.”

List of Subjects in 47 CFR Part 73

Radio, Reporting and recordkeeping requirements.

Federal Communications Commission.

Aleta Bowers,

Federal Register Liaison Officer, Office of the Secretary.

Accordingly, 47 CFR part 73 is corrected by making the following correcting amendment:

PART 73—RADIO BROADCAST SERVICE

■ 1. The authority citation for part 73 continues to read as follows:

Authority: 47 U.S.C. 154, 155, 301, 303, 307, 309, 310, 334, 336, 339.

■ 2. Amend § 73.3556 by revising the section heading to read as follows:

§ 73.3556 Duplication of programming on commonly owned or time brokered stations.

* * * * *

[FR Doc. 2026–19311 Filed 9–21–26; 8:45 am]

BILLING CODE 6712–01–P

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 73

[DA 26–986; FR ID 368505]

Radio Broadcasting Services; Various Locations

AGENCY: Federal Communications Commission.

ACTION: Final rule.

SUMMARY: This document amends the Table of FM Allotments, of the Federal Communications Commission’s (Commission) rules, by reinstating certain channels as a vacant FM allotment in various communities. The FM allotments were previously removed from the FM Table because a construction permit and/or license was granted. These FM allotments are now considered vacant because of the cancellation of the associated FM authorizations. A staff engineering analysis confirms that all of the vacant FM allotments complies with the minimum distance separation requirements and principle community coverage requirements of the Commission’s rules. The window period for filing applications for these vacant

FM allotments will not be opened at this time. Instead, the issue of opening these allotments for filing will be addressed by the Commission in subsequent order.

DATES: Effective September 22, 2026.

FOR FURTHER INFORMATION CONTACT:

Rolanda F. Smith, Media Bureau, (202) 418–2054, Rolanda-Faye.Smith@fcc.gov.

SUPPLEMENTARY INFORMATION: This is a synopsis of the Commission’s *Order*, adopted September 15, 2026, and released September 16, 2026. The full text of this Commission decision is available online at <https://apps.fcc.gov/ecfs/>. The full text of this document can also be downloaded in Word or Portable Document Format (PDF) at <https://www.fcc.gov/edocs>. This document does not contain information collection requirements subject to the Paperwork Reduction Act of 1995, Public Law 104–13. The Commission will not send a copy of the *Order* in a report to be sent to Congress and the Government Accountability Office pursuant to the Congressional Review Act, *see* 5 U.S.C. 801(a)(1)(A), because these allotments were previously reported.

List of Subjects in 47 CFR Part 73

Radio, Radio broadcasting.

Federal Communications Commission.

Nazifa Sawez,

Assistant Chief, Audio Division, Media Bureau.

Final Rules

For the reasons discussed in the preamble, the Federal Communications Commission amends 47 CFR part 73 as follows:

PART 73—RADIO BROADCAST SERVICES

■ 1. The authority citation for part 73 continues to read as follows:

Authority: 47 U.S.C. 154, 155, 301, 303, 307, 309, 310, 334, 336, 339.

■ 2. Amend § 73.202(b) in table 1 (Table of FM Allotments) by;

■ a. Adding the entries for “Fowler” and “Parachute” in alphabetical order under Colorado;

■ b. Revising the entry for “Horseshoe Beach” in alphabetical order under Florida;

■ c. Adding the entry for “Coushatta” in alphabetical order under Louisiana;

■ d. Adding the entries for “Arcadia” and “Ellington” in alphabetical order under Missouri;

■ e. Adding the entry for “Ellsworth AFB” in alphabetical order under South Dakota; and

■ f. Adding the entry for “Huntingdon” in alphabetical order under Tennessee.

The additions and revision reads as follows:

§ 73.202 Table of Allotments.

* * * * *

(b) * * *

TABLE 1 TO PARAGRAPH (b)
[U.S. States]

	Chan- nel No.
Colorado	
* * * * *	
Fowler	257C1
* * * * *	
Parachute	266A
* * * * *	
Florida	
* * * * *	
Horseshoe Beach	234C3
* * * * *	
Louisiana	
* * * * *	
Coushatta	235C2
* * * * *	
Missouri	
Arcadia	280A
* * * * *	
Ellington	294C2
* * * * *	
South Dakota	
* * * * *	
Ellsworth AFB	285C
* * * * *	
Tennessee	
Huntingdon	265C3
* * * * *	

[FR Doc. 2026–19329 Filed 9–21–26; 8:45 am]

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