NOTICE OF INTENT TO TERMINATE RECEIVERSHIPS—Continued

Fund	Receivership name	City	State	Date of appointment of receiver
10216	Riverside National Bank of Florida	Fort Pierce	FL	04/16/2010
10534	City National Bank New Jersey	Newark	NJ	11/01/2019

The liquidation of the assets for each receivership has been completed. To the extent permitted by available funds and in accordance with law, the Receiver will be making a final dividend payment to proven creditors. Based upon the foregoing, the Receiver has determined that the continued existence of the receiverships will serve no useful purpose. Consequently, notice is given that the receiverships shall be terminated, to be effective no sooner than thirty days after the date of this notice. If any person wishes to comment concerning the termination of any of the receiverships, such comment must be made in writing, identify the receivership to which the comment pertains, and be sent within thirty days of the date of this notice to: Federal Deposit Insurance Corporation, Division of Resolutions and Receiverships, Attention: Receivership Oversight Section, 600 North Pearl, Suite 700, Dallas, TX 75201. No comments concerning the termination of the above-mentioned receiverships will be considered which are not sent within this timeframe.

(Authority: 12 U.S.C. 1819.)

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on September 1, 2026.

Jennifer M. Jones,

Deputy Executive Secretary.

[FR Doc. 2026–18050 Filed 9–2–26; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Benjamin W. McDonough, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than October 5, 2026.

A. Federal Reserve Bank of Richmond (Brent B. Hassell, Assistant Vice President) P.O. Box 27622, Richmond, Virginia 23261. Comments can also be sent electronically to *Comments.applications@rich.frb.org:*

1. *First Bancorp, Southern Pines, North Carolina;* to acquire First Carolina Bancshares Corporation, Florence, South Carolina, and thereby indirectly acquire Carolina Bank & Trust Company, Lamar, South Carolina.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–18058 Filed 9–2–26; 8:45 am]

BILLING CODE;P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–P–1372]

Determination That PHENAPHEN WITH CODEINE NO. 3 (Acetaminophen; Codeine Phosphate) Oral Capsules, 325 Milligrams and 30 Milligrams, and PHENAPHEN WITH CODEINE NO. 4 (Acetaminophen; Codeine Phosphate) Oral Capsules, 325 Milligrams and 60 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) has determined that PHENAPHEN WITH CODEINE NO. 3 (acetaminophen; codeine phosphate) oral capsules, 325 milligrams (mg) and 30 mg, and PHENAPHEN WITH CODEINE NO. 4 (acetaminophen; codeine phosphate) oral capsules, 325 mg and 60 mg, were not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDAs) for PHENAPHEN WITH CODEINE NO. 3 (acetaminophen; codeine phosphate) oral capsules, 325 mg and 30 mg, and PHENAPHEN WITH CODEINE NO. 4 (acetaminophen; codeine phosphate) oral capsules, 325 mg and 60 mg, if all other legal and regulatory requirements are met.

FOR FURTHER INFORMATION CONTACT: Stacy Kane, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6236, Silver Spring, MD 20993–0002, 301–796–8363, *Stacy.Kane@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Section 505(j) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 355(j)) allows the submission of an ANDA to market a generic version of a previously approved drug product. To obtain approval, the ANDA applicant must show, among other things, that the generic drug product: (1) has the same active ingredient(s), dosage form, route