

that there is inadequate lead time to permit the development of technology necessary to meet the 2013 HD OBD New or Stricter Requirements that are subject to the waiver request, giving appropriate consideration to the cost of compliance within that time.<sup>39</sup> California's accompanying enforcement procedures would also be inconsistent with section 202(a) if the federal and California test procedures conflicted, *i.e.*, if manufacturers would be unable to meet both the California and federal test requirements with the same test vehicle.<sup>40</sup>

Regarding test procedure conflict, CARB notes that there is no issue of test procedure inconsistency because federal regulations provide that engines certified to California's HD OBD regulation are deemed to comply with federal standards. EPA has received no adverse comment or evidence of test procedure inconsistency. We therefore cannot find that the 2013 HD OBD New or Stricter Requirements are inconsistent with federal test procedures.

EPA also did not receive any comments arguing that the 2013 HD OBD Amendments were technologically infeasible or that the cost of compliance would be excessive, such that California's standards might be inconsistent with section 202(a).<sup>41</sup> In EPA's review of the 2013 HD OBD New or Stricter Requirements, we likewise cannot identify any requirements that appear technologically infeasible or excessively expensive for manufacturers to implement within the timeframes provided. EPA therefore cannot find that the 2013 HD OBD New or Stricter Requirements do not provide adequate lead time or are otherwise not technically feasible.

We therefore cannot find that the 2013 HD OBD New or Stricter Requirements that we analyzed under the waiver criteria are inconsistent with section 202(a).

Having found that the 2013 HD OBD New or Stricter Requirements satisfy each of the criteria for a waiver, and having received no evidence to contradict this finding, we cannot deny a waiver for the amendments.

#### IV. Decision

The Administrator has delegated the authority to grant California section 209(b) waivers to the Assistant Administrator for Air and Radiation.

After evaluating CARB's 2013 HD OBD Amendments and CARB's submissions for EPA review, EPA is hereby confirming that the 2013 HD OBD Amendments, with the exception of the 2013 HD OBD New or Stricter Requirements identified above, are within the scope of EPA's previous waivers for the HD OBD Requirements and HD OBD Enforcement Regulation. In addition, EPA is hereby granting a waiver for the 2013 HD OBD New or Stricter Requirements.

This decision will affect persons in California and those manufacturers and/or owners/operators nationwide who must comply with California's requirements. In addition, because other states may adopt California's standards for which a section 209(b) waiver has been granted under section 177 of the Act if certain criteria are met, this decision would also affect those states and those persons in such states. For these reasons, EPA determines and finds that this is a final action of national applicability, and also a final action of nationwide scope or effect for purposes of section 307(b)(1) of the Act. Pursuant to section 307(b)(1) of the Act, judicial review of this final action may be sought only in the United States Court of Appeals for the District of Columbia Circuit. Petitions for review must be filed by January 6, 2017. Judicial review of this final action may not be obtained in subsequent enforcement proceedings, pursuant to section 307(b)(2) of the Act.

#### V. Statutory and Executive Order Reviews

As with past waiver and authorization decisions, this action is not a rule as defined by Executive Order 12866. Therefore, it is exempt from review by the Office of Management and Budget as required for rules and regulations by Executive Order 12866.

In addition, this action is not a rule as defined in the Regulatory Flexibility Act, 5 U.S.C. 601(2). Therefore, EPA has not prepared a supporting regulatory flexibility analysis addressing the impact of this action on small business entities.

Further, the Congressional Review Act, 5 U.S.C. 801, *et seq.*, as added by the Small Business Regulatory Enforcement Fairness Act of 1996, does not apply because this action is not a rule for purposes of 5 U.S.C. 804(3).

Dated: October 24, 2016.

**Janet G. McCabe,**

*Acting Assistant Administrator, Office of Air and Radiation.*

[FR Doc. 2016-26865 Filed 11-4-16; 8:45 am]

**BILLING CODE 6560-50-P**

#### FEDERAL DEPOSIT INSURANCE CORPORATION

##### Notice of the Termination of the Receivership of 10508, Frontier Bank, FSB Palm Desert, California

The Federal Deposit Insurance Corporation ("FDIC"), as Receiver for 10508 Frontier Bank, FSB, Palm Desert, California ("Receiver") has been authorized to take all actions necessary to terminate the receivership estate of Frontier Bank, FSB ("Receivership Estate"); the Receiver has made all dividend distributions required by law.

The Receiver has further irrevocably authorized and appointed FDIC-Corporate as its attorney-in-fact to execute and file any and all documents that may be required to be executed by the Receiver which FDIC-Corporate, in its sole discretion, deems necessary; including but not limited to releases, discharges, satisfactions, endorsements, assignments and deeds.

Effective November 1, 2016, the Receivership Estate has been terminated, the Receiver discharged, and the Receivership Estate has ceased to exist as a legal entity.

Dated: November 1, 2016.

Federal Deposit Insurance Corporation.

**Valerie J. Best,**

*Assistant Executive Secretary.*

[FR Doc. 2016-26852 Filed 11-4-16; 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 applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of

<sup>39</sup> See, e.g., 38 F.R. 30136 (November 1, 1973) and 40 F.R. 30311 (July 18, 1975).

<sup>40</sup> See, e.g., 43 F.R. 32182 (July 25, 1978).

<sup>41</sup> See, e.g., 78 F.R. 2134 (Jan. 9, 2013), 47 F.R. 7306, 7309 (Feb. 18, 1982), 43 F.R. 25735 (Jun. 17, 1978), and 46 F.R. 26371, 26373 (May 12, 1981).

a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than December 5, 2016.

A. Federal Reserve Bank of Atlanta (Chapelle Davis, Assistant Vice President) 1000 Peachtree Street NE., Atlanta, Georgia 30309. Comments can also be sent electronically to [Applications.Comments@atl.frb.org](mailto:Applications.Comments@atl.frb.org):

1. *Atlantic Coast Financial Corporation*: To become a bank holding company by acquiring 100 percent of the outstanding shares of Atlantic Coast Bank, both of Jacksonville, Florida.

Board of Governors of the Federal Reserve System, November 2, 2016.

**Michele Taylor Fennell,**

*Assistant Secretary of the Board.*

[FR Doc. 2016–26863 Filed 11–4–16; 8:45 am]

BILLING CODE 6210–01–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30Day–17–0950]

#### Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and

clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

#### Proposed Project

The National Health and Nutrition Examination Survey (NHANES), (OMB No. 0920–0950, expires 12/31/2017)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on the extent and nature of illness and disability; environmental, social and other health hazards; and determinants of health of the population of the United States.

The National Health and Nutrition Examination Survey (NHANES) have been conducted periodically between 1970 and 1994, and continuously since 1999 by the National Center for Health Statistics, CDC. Annually, approximately 14,410 respondents participate in some aspect of the full survey. Up to 13,104 additional persons might participate in tests of procedures, special studies, or methodological studies. Participation in NHANES is completely voluntary and confidential. A three-year approval is requested.

The data collected through NHANES allows for the production descriptive statistics which measure the health and nutrition status of the general population. Through the use of physical examinations, laboratory tests, and interviews NHANES studies the relationship between diet, nutrition and health in a representative sample of the civilian noninstitutionalized population of the United States. NHANES monitors

the prevalence of chronic conditions and risk factors. NHANES data are used to produce national reference data on height, weight, and nutrient levels in the blood. NHANES also seeks to be responsive in exploring emerging public health issues and new health related technologies. Results from more recent NHANES can be compared to findings reported from previous surveys to monitor changes in the health of the U.S. population over time. NHANES collects personal identification information. Participant level data items will include basic demographic information, name, address, social security number, Medicare number and participant health information to allow for linkages to other data sources such as the National Death Index (OMB No. 0920–0124, expires 10/31/2016 and data from the Centers for Medicare and Medicaid Services (CMS).

A variety of agencies sponsor data collection components on NHANES. To keep burden down, NCHS cycles in and out various components. Health interviews are conducted in the participants' household. Physical exams are conducted in the Mobile Examination Center (MEC). The 2017–2018 NHANES physical examination includes the following components: Anthropometry (all ages), 24-hour dietary recall (all ages), physician's examination (all ages, blood pressure is collected here), and oral health examination (ages 1 and older, body composition using Dual X-ray Absorptiometry (DXA) exam (ages 8–59) and hearing (ages 6–19 and 70+ (and above)). The hearing age range for 2017–18 is a modification (The 2015–16 hearing age range was 20–69 years).

While at the examination center additional interview questions are asked (6 and older), a second 24-hour dietary recall (all ages) is scheduled to be conducted by phone 3–10 days later. In 2017 we plan to add a liver elastography (ultrasound) exam for participants 12 years and older. A set of alcohol and liver-related questions, and a hip measurement will also be added to complement the liver exam. The age range for liver-related blood test (serum ferritin) already collected in NHANES is being expanded from female participants 12–49 to all participants 12 years and older. The existing collection of serum ferritin in children 1–5 years will continue. We will cycle bone density for hip and spine back into the (DXA) exam for (ages 50+). The Osteoporosis questionnaire will also cycle back into NHANES to complement the changes to the DXA exam. These questions will be asked of participants 50+.