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 also will 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 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. Additional information on all bank holding companies may be obtained from the National Information Center website at *www.ffiec.gov/nic/*.

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 June 7, 2010.

A. Federal Reserve Bank of San Francisco (Kenneth Binning, Vice President, Applications and Enforcement) 101 Market Street, San Francisco, California 94105–1579:

1. Carpenter Fund Manager GP, LLC, Carpenter Fund Management, LLC, Carpenter Community Bancfund, L.P., Carpenter Community Bancfund-A, LP, Carpenter Community Bancfund-CA, L.P., SCJ, Inc., and CCFW, Inc., all of Irvine, California; to acquire no more than 35 percent of the voting shares of Bridge Capital Holdings, and thereby indirectly acquire voting shares of Bridge Bank, N.A., both of San Jose, California.

Board of Governors of the Federal Reserve System, May 10, 2010.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2010–11440 Filed 5–12–10; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL TRADE COMMISSION

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Correction

AGENCY: Federal Trade Commission ("FTC" or "Commission").

ACTION: Notice; correction.

SUMMARY: The Federal Trade Commission published a document in the **Federal Register** of April 15, 2010, seeking public comments on its proposal to extend through May 31, 2013, the current Paperwork Reduction Act clearance for information collection requirements associated with the Contact Lens Rule (the Rule), 16 CFR part 315. The document contained an incorrect OMB Control No. for the pre-existing clearance. The correct number is 3084-0127.

FOR FURTHER INFORMATION CONTACT: Requests for additional information or copies of the proposed information requirements should be addressed to Karen Jagielski, Attorney, Division of Advertising Practices, Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue, NW., NJ– 3212, Washington, DC 20580, (202) 326–2509. Correction In the **Federal Register** of April 15, 2010, in FR Doc. 2010–8647, on page 19647, in the third column, under **Supplementary Information: Background**, correct the second paragraph, third sentence, to read: "Pursuant to the OMB regulations, 5 CFR Part 1320, that implement the PRA, the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for the Rule (OMB Control No. 3084-0127)."

By direction of the Commission.

Donald S. Clark

Secretary.

[FR Doc. 2010–11501 Filed 5–12–10; 8:45 am]

BILLING CODE: 6750–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–10–0539]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Estimating the Capacity for National and State-Level Colorectal Cancer Screening through a Survey of Endoscopic Capacity (SECAP II)(OMB No. 0920–0539, exp. 3/31/2003)—Reinstatement with Change—Division of Cancer Prevention and Control, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Colorectal cancer (CRC) is the second leading cause of cancer-related deaths in the United States (U.S.). Most colorectal cancers develop from pre-existing growths, or polyps, which slowly transform into cancers over a period of 10–20 years. As a result, CRC is ideally suited for prevention and early detection through regular screening. Recommended screening procedures include flexible sigmoidoscopy and colonoscopy, which allow qualified medical professionals to identify and remove polyps as well as to detect early cancers. Information regarding the capacity of the U.S. health care system to provide lower GI endoscopic procedures is critical to planning widespread CRC screening programs.

CDC requests OMB approval to reinstate a previously approved data collection, formerly entitled the *National Survey of Endoscopic Capacity (SECAP)* (OMB No. 0920–0539, exp. 3/31/2003), to obtain a current estimate of the number of colorectal cancer screening and follow-up tests being performed, as well as the maximum number of screening and follow-up tests that could be performed in the event of widespread screening. In addition, the reinstatement request describes a plan to conduct state-specific surveys in up to 18 selected states. Similar surveys were conducted in 15 selected states from 2003 to 2005, and provided estimates of endoscopic screening capacity at state and sub-state levels (*State Survey of Endoscopic Capacity,* OMB No. 0920–0590, exp. 6/30/2006). However, in light of recent trends in colorectal cancer screening (*e.g.*, increases in the percentage of public and private insurers that reimburse for screening colonoscopy, increased use of colonoscopy and decreased use of flexible sigmoidoscopy, availability of other colorectal cancer screening procedures), there is a need to update estimates of endoscopic capacity to guide continued screening initiatives.

OMB approval is requested for three years. The proposed national survey will be conducted in 2010–2011 and

will be based on an updated version of the previously fielded paper-and-pencil survey instrument. The target population for the national survey is all facilities in the U.S. that use lower gastrointestinal flexible endoscopic equipment for the detection of colorectal cancer in adults. Information will be collected from a random sample of 1,440 facilities, stratified by U.S. Census region and urban/rural location.

Additional state-level surveys will be conducted from approximately 2010–2012 and will include a census survey of up to 18 selected states, based on methodology employed with the previously fielded state-based survey. An average of 135 facilities will be selected to participate in each state. A total of approximately 1,680 completed state surveys will be collected over the three years of the project.

Facilities will be recruited and screened through a telephone interview. Participation is voluntary. The information collection will inform planning efforts for national and state colorectal cancer screening.

There are no costs to respondents other than their time. The total estimated burden hours are 732.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Medical Facilities that Perform CRC Screening.	National Survey Recruitment Interview	700	1	5/60
	National SECAP Survey	480	1	35/60
	State Survey Recruitment Interview	800	1	5/60
	State SECAP Survey	560	1	35/60

Carol Walker,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010–11413 Filed 5–12–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–D–0043]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Request for a Meeting or Teleconference in Electronic Format to the Center for Veterinary Medicine

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by June 14, 2010.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or e-mailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0452. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., P150–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Guidance for Industry on How to Use E-Mail to Submit a Request for a Meeting or Teleconference in Electronic Format to the Center for Veterinary Medicine—(OMB Control Number 0910–0452)—Extension

CVM holds meetings and/or teleconferences when a sponsor requests a presubmission conference under 21 CFR 514.5, or requests a meeting to discuss general questions. Generally, meeting requests are submitted to CVM on paper. However, CVM now allows registered sponsors to submit information electronically, and to request meetings electronically, if they determine this is more efficient and time saving for them. CVM's guidance on "How to Submit a Request for a Meeting or Teleconference in Electronic Format to CVM," provides sponsors with the option to submit a request for a meeting or teleconference as an e-mail attachment by the Internet. The likely respondents are sponsors for new animal drug applications.

In the **Federal Register** of February 5, 2010 (75 FR 6035), FDA published a 60-day notice requesting public comment on the proposed collection of information.

In response, two comments were received. One comment was completely outside the scope of the notice and the other requested that FDA meet openly with industry rather than closed sessions. Neither comment addressed the paperwork involved in the information collection.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section/ FDA Form 3489	No. of Respondents	Annual Frequency per Response	Total Annual Responses[2]	Hours per Response	Total Hours
10.64	40	2.4	96	.08	7.7

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

[2] Electronic submissions received between January 1, 2008, and December 31, 2008.