

FEE SCHEDULE FOR EACH VESSEL
SIZE

Vessel size (GRT ¹)	Inspection fee
Extra Small (<3,000 GRT)	US\$1,495
Small (3,001–15,000 GRT) ..	2,990
Medium (15,001–30,000 GRT)	5,980
Large (30,001–60,000 GRT)	8,970
Extra Large (60,001–120,000 GRT)	11,960
Mega (>120,001 GRT)	17,940

¹ Gross register tonnage in cubic feet, as shown in Lloyd's Register of Shipping.

Dated: August 19, 2015.

Pamela J. Cox,

*Director, Division of the Executive Secretariat,
Office of the Chief of Staff, Centers for Disease
Control and Prevention.*

[FR Doc. 2015–21107 Filed 8–25–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND
HUMAN SERVICESAdministration for Children and
FamiliesProposed Information Collection
Activity; Comment Request

Proposed Projects:

Title: State Self-Assessment Review
and Report.

OMB No.: 0970–0223.

Description: Section 454(15)(A) of the
Social Security Act, as amended by the

Personal Responsibility and Work
Opportunity Reconciliation Act of 1996,
requires each State to annually assess
the performance of its child support
enforcement program in accordance
with standards specified by the
Secretary of the Department of Health
and Human Services, and to provide a
report of the findings to the Secretary.
This information is required to
determine if States are complying with
Federal child support mandates and
providing the best services possible. The
report is also intended to be used as a
management tool to help States evaluate
their programs and assess performance.

Respondents: State Child Support
Enforcement Agencies or the
Department/Agency/Bureau responsible
for Child Support Enforcement in each
State.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Self-assessment report	54	1	4	216

Estimated Total Annual Burden
Hours: 216.

In compliance with the requirements
of Section 506(c)(2)(A) of the Paperwork
Reduction Act of 1995, the
Administration for Children and
Families is soliciting public comment
on the specific aspects of the
information collection described above.
Copies of the proposed collection of
information can be obtained and
comments may be forwarded by writing
to the Administration for Children and
Families, Office of Planning, Research
and Evaluation, 370 L'Enfant
Promenade SW., Washington, DC 20447,
Attn: ACF Reports Clearance Officer.
Email address: [infocollection@
acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be
identified by the title of the information
collection.

The Department specifically requests
comments on: (a) Whether the proposed
collection of information is necessary
for the proper performance of the
functions of the agency, including
whether the information shall have
practical utility; (b) the accuracy of the
agency's estimate of the burden of the
proposed collection of information; (c)
the quality, utility, and clarity of the
information to be collected; and (d)
ways to minimize the burden of the
collection of information on
respondents, including through the use
of automated collection techniques or

other forms of information technology.
Consideration will be given to
comments and suggestions submitted
within 60 days of this publication.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–21053 Filed 8–25–15; 8:45 am]

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DEPARTMENT OF HEALTH AND
HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0110]

Agency Information Collection
Activities; Submission for Office of
Management and Budget Review;
Comment Request; Medical Device
Reporting; Manufacturer, Importer,
User Facility, and Distributor Reporting

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 September
25, 2015.

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 emailed to
oira_submission@omb.eop.gov. All
comments should be identified with the
OMB control number 0910–0437. Also
include the FDA docket number found
in brackets in the heading of this
document.

FOR FURTHER INFORMATION CONTACT: FDA
PRA Staff, Office of Operations, Food
and Drug Administration, 8455
Colesville Rd., COLE–14526, Silver
Spring, MD 20993–0002,
PRASaff@fda.hhs.gov.

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.

Medical Device Reporting;
Manufacturer, Importer, User Facility,
and Distributor Reporting (21 CFR Part
803)—(OMB Control Number 0910–
0437)—Revision

Section 519(a)(1) of the Federal Food,
Drug, and Cosmetic Act (the FD&C Act)