

EPA-APPROVED REGULATIONS IN THE WEST VIRGINIA SIP—Continued

State citation [Chapter 16–20 or 45 CSR]	Title/subject	State effective date	EPA approval date	Additional explanation/citation at 40 CFR § 52.2565
Section 45–40–42	CAIR NO _x Ozone Season Allowance Allocations.	5/1/06	12/18/07 [Insert page number where the document begins].	1. Except for subsections 42.2.d, 42.2.e, 42.3.a.2, and 42.4.b. 2. Only Phase I (2009–2014).

* * * * * (e) * * *

Name of non-regulatory SIP revision	Applicable geographic area	State submittal date	EPA approval date	Additional explanation
Article 3, Chapter 64 of the Code of West Virginia, 1931.	Statewide	5/1/06	12/18/07 [Insert page number where the document begins].	Effective date of March 11, 2006.

PART 97—[AMENDED]

■ 3. The authority citation for 40 CFR part 97 continues to read as follows:

Authority: 42 U.S.C. 7401, 7403, 7410, 7426, 7601, and 7651, *et seq.*

■ 4. Appendix A to Subpart EE is amended by adding the entry for “West Virginia” in alphabetical order under paragraph 1. to read as follows:

Appendix A to Subpart EE of Part 97—States With Approved State Implementation Plan Revisions Concerning Allocations

1. * * *
West Virginia (for control periods 2009–2014)

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■ 5. Appendix A to Subpart EEEE is amended by adding the entry for West Virginia in alphabetical order to read as follows:

Appendix A to Subpart EEEE of Part 97—States With Approved State Implementation Plan Revisions Concerning Allocations

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West Virginia (for control periods 2009–2014)

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[FR Doc. E7–24367 Filed 12–17–07; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Part 488

[CMS–2278–IFC2]

RIN 0938–AP22

Revisit User Fee Program for Medicare Survey and Certification Activities

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Interim final rule with comment period.

SUMMARY: This interim final rule with comment period implements the continuation of the revisit user fee program for Medicare Survey and Certification activities, in accordance with the statutory authority in the Further Continuing Appropriations, 2008 Resolution (“Continuing Resolution”) budget legislation passed by the Congress and signed by the President on November 13, 2007. On September 19, 2007, we published a final rule that established a system of revisit user fees applicable to health care facilities that have been cited for deficiencies during initial certification, recertification or substantiated complaint surveys and require a revisit to confirm that previously-identified deficiencies have been corrected.
DATES: *Effective date:* These regulations are effective December 14, 2007, and applicable beginning November 17, 2007.

Comment date: To be assured consideration, comments must be

received at one of the addresses provided below, no later than 5 p.m. on February 19, 2008.

ADDRESSES: In commenting, please refer to file code CMS–2278–IFC2. Because of staff and resource limitations, we cannot accept comments by facsimile (FAX) transmission.

You may submit comments in one of four ways (no duplicates, please):

1. *Electronically.* You may submit electronic comments on specific issues in this regulation to <http://www.cms.hhs.gov/eRulemaking>. Click on the link “Submit electronic comments on CMS regulations with an open comment period.” (Attachments should be in Microsoft Word, WordPerfect, or Excel; however, we prefer Microsoft Word.)

2. *By regular mail.* You may mail written comments (one original and two copies) to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–2278–IFC2, P.O. Box 8010, Baltimore, MD 21244–8016.

Please allow sufficient time for mailed comments to be received before the close of the comment period.

3. *By express or overnight mail.* You may send written comments (one original and two copies) to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–2278–IFC2, Mail Stop C4–26–05, 7500 Security Boulevard, Baltimore, MD 21244–1850.

4. *By hand or courier.* If you prefer, you may deliver (by hand or courier) your written comments (one original and two copies) before the close of the

comment period to one of the following addresses. If you intend to deliver your comments to the Baltimore address, please call telephone number (410) 786-7195 in advance to schedule your arrival with one of our staff members.

Room 445-G, Hubert H. Humphrey Building, 200 Independence Avenue, SW., Washington, DC 20201; or 7500 Security Boulevard, Baltimore, MD 21244-1850.

(Because access to the interior of the HHH Building is not readily available to persons without Federal Government identification, commenters are encouraged to leave their comments in the CMS drop slots located in the main lobby of the building. A stamp-in clock is available for persons wishing to retain a proof of filing by stamping in and retaining an extra copy of the comments being filed.)

Comments mailed to the addresses indicated as appropriate for hand or courier delivery may be delayed and received after the comment period.

For information on viewing public comments, see the beginning of the **SUPPLEMENTARY INFORMATION** section.

FOR FURTHER INFORMATION CONTACT: Kelley Tinsley, (410) 786-6664.

SUPPLEMENTARY INFORMATION:

Submitting Comments: As the public was provided an opportunity to comment on the substance of the rule during the comment period prior to the publication of the September 19, 2007 final rule, and as the substance of the rule is not changed by this interim final rule with comment period, we are accepting comments only to the extent that they pertain to the applicability of the new authority for the rule. You can assist us by referencing the file code CMS-2278-IFC2.

Inspection of Public Comments: All comments received before the close of the comment period are available for viewing by the public, including any personally identifiable or confidential business information that is included in a comment. We post all comments received before the close of the comment period on the following Web site as soon as possible after they have been received: <http://www.cms.hhs.gov/eRulemaking>. Click on the link "Electronic Comments on CMS Regulations" on that Web site to view public comments.

Comments received timely will be available for public inspection as they are received, generally beginning approximately three weeks after publication of a document, at the headquarters of the Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore, Maryland 21244,

Monday through Friday of each week from 8:30 a.m. to 4 p.m. To schedule an appointment to view public comments, phone 1-800-743-3951.

SUPPLEMENTARY INFORMATION:

I. Background

In the June 29, 2007 **Federal Register** (72 FR 35673), we published the proposed rule entitled, "Establishment of Revisit User Fee Program for Medicare Survey and Certification Activities" and provided for a 60-day comment period. In the September 19, 2007 **Federal Register** (72 FR 53628) we published the Revisit User Fee Program final rule. That final rule set forth final requirements and a final fee schedule for providers and suppliers who require a revisit survey as a result of deficiencies cited during an initial certification, recertification, or substantiated complaint survey.

The Centers for Medicare & Medicaid Services (CMS) has in place an outcome-oriented survey process that is designed to ensure that existing Medicare-certified providers and suppliers or providers and suppliers seeking initial Medicare certification, meet statutory and regulatory requirements, conditions of participation, or conditions for coverage. These health and safety requirements apply to the environments of care and the delivery of services to residents or patients served by these facilities and agencies. The Secretary of the Department of Health and Human Services (HHS) has designated CMS to enforce the conditions of participation/coverage and other requirements of the Medicare program. The revisit user fee will be assessed for revisits conducted in order to determine whether deficiencies cited as a result of failing to satisfy federal quality of care requirements have been corrected.

Pursuant to the requirements of the Continuing Appropriations Resolution budget bill for fiscal year (FY) 2007, the Secretary directed CMS to implement the revisit user fees for FY 2007 for certain providers and suppliers for which a revisit was required to confirm that previously-identified failures to meet federal quality of care requirements had been remedied. The fees recover the costs associated with the Medicare Survey and Certification program's revisit surveys. The primary purpose for implementing the revisit user fees is to ensure the continuance of CMS Survey and Certification quality assurance functions that improve patient care and safety. The fees became effective upon publication September 19, 2007, when the final rule was published.

II. Provisions of the Interim Final Rule

The current Continuing Resolution Pub. L. 110-16 Division B of HR 3222 which amends Pub. L. 110-92 H. J. Res. 52 §§ 101 & 106(2007), authorizes HHS to continue to impose revisit user fees until December 14, 2007, as follows:

* * *

Sec. 101. Such amounts as may be necessary, at a rate for operations as provided in the applicable appropriations Acts for fiscal year 2007 and under the authority and conditions provided in such Acts, for continuing projects or activities (including the costs of direct loans and loan guarantees) that are not otherwise specifically provided for in this joint resolution, that were conducted in fiscal year 2007, and for which appropriations, funds, or other authority were made available in the following appropriations Acts:

* * *

(3) The Continuing Appropriations Resolution, 2007 (division B of Public Law 109-289, as amended by Public Law 110-5). (*H.J.Res.20, § 101(2007)*).

Sec. 106. Unless otherwise provided for in this joint resolution or in the applicable appropriations Act for fiscal year 2008, appropriations and funds made available and authority granted pursuant to this joint resolution shall be available until whichever of the following first occurs:

* * *

(3) December 14, 2007.

As directed by the Secretary, in the September 19, 2007 **Federal Register** (72 FR 53628), we established the revisit user fee program for revisit surveys. We put forth in regulation the relevant definitions, criteria for determining the fees, the fee schedule, procedures for the collection of fees, the reconsideration process, enforcement and regulatory language addressing enrollment and billing privileges, and provider agreements. In the September 19, 2007 final rule, cost projections were based on FY 2006 actual data and were expected to amount to \$37.3 million for FY 2007. These calculations were included in section IV of the final rule (72 FR 53642).

We stated in the final rule that, "if authority for the revisit user fee is continued, we will use the current fee schedule in [the final rule] for the assessment of such fees until such time as a new fee schedule notice is proposed and published in final form." (72 FR 53628). The current Continuing Resolution continues the authority of the FY 2007 Continuing Resolution from November 17, 2007 through December 14, 2007. Accordingly, the revisit fees will continue to be assessed for the entire time period authorized by the current Continuing Resolution.

III. Response to Comments

Because of the large number of public comments we normally receive on **Federal Register** documents, we are not able to acknowledge or respond to them individually. We will consider all comments we receive by the date and time specified in the **DATES** section of this preamble, and, when we proceed with a subsequent document, we will respond to the comments in the preamble to that document.

IV. Waiver of Proposed Rulemaking and Delay in Effective Date

We ordinarily publish a notice of proposed rulemaking in the **Federal Register** and invite public comment on the proposed rule in accordance with 5 U.S.C. 553(b) of the Administrative Procedure Act (APA). The notice of proposed rulemaking includes a reference to the legal authority under which the rule is proposed, and the terms and substance of the proposed rule or a description of the subjects and issues involved. This procedure can be waived, however, if an agency finds good cause that a notice-and-comment procedure is impracticable, unnecessary, or contrary to the public interest and incorporates a statement of the finding and its reasons in the rule issued. We find that the notice-and-comment procedure is unnecessary in this circumstance because providers and suppliers have already been provided notice and an opportunity to comment on the substance of this rule. This interim final rule with comment merely updates the Congressional authority under which the rule operates.

Therefore, we find good cause to waive the notice of proposed rulemaking and to issue this final rule on an interim basis. We are providing a 60-day public comment period.

We ordinarily provide a 30-day delay in the effective date of the provisions of a rule in accordance with the Administrative Procedure Act (APA), 5 U.S.C. 553(d). However, the delay in the effective date may be waived as, in pertinent part, "provided by the agency for good cause found and published with the rule." 5 U.S.C. 553(d)(3). The Secretary finds that good cause exists to waive the 30-day effective date delay.

The good cause exception to the 30 day effective date delay provision of section 553(d) of the APA is read to be broader than the good cause exception to the notice and comment provision of section 553(b) of the APA.

The legislative history of the APA indicates that the purpose for deferring the effectiveness of a rule under section 553(d) was to "afford persons affected a

reasonable time to prepare for the effective date of a rule or rules or to take other action which the issuance may prompt." S. Rep. No. 752, 79th Cong., 1st Sess. 15 (1946); H.R. Rep. No. 1980, 79th Cong. 2d Sess. 25 (1946). In this case, affected parties do not need time to adjust their behavior before this rule takes effect. This rule merely updates the authority under which the revisit fee is assessed and does not provide any additional requirements for the affected parties. Moreover, with or without a revisit fee, a provider or supplier must be found to have corrected significant deficiencies in order to avoid termination. Additionally, the application of a fee for the revisit does not place appreciable administrative burdens on the affected providers or suppliers. We do not expect appreciable cost to State survey agencies because we are undertaking the billing and collection of the revisit user fee.

We identified in the September 19, 2007 final rule the immediacy of this revisit user fee program and the specific statutory requirement contained limited in the Continuing Resolution that required us to implement the revisit user fee program in FY 2007. Accordingly, providers and suppliers have been on notice for some time that these fees will be imposed, and do not need additional time to be prepared to comply with the requirements of this regulation. We believe that given the short timeframe that we have to collect fees before the statutory authority of the current Continuing Resolution expires, there is good cause to waive the 30-day effective date.

V. Collection of Information Requirements

This document does not impose information collection and recordkeeping requirements. Consequently, it need not be reviewed by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995.

VI. Regulatory Impact Analysis

A. Overall Impact

We have examined the impacts of this rule as required by Executive Order 12866 (September 1993, Regulatory Planning and Review), the Regulatory Flexibility Act (RFA) (September 19, 1980, Pub. L. 96-354), section 1102(b) of the Social Security Act, the Unfunded Mandates Reform Act of 1995 (Pub. L. 104-4), and Executive Order 13132.

Executive Order 12866 (as amended by Executive Order 13258, which merely reassigns responsibility of duties) directs agencies to assess all

costs and benefits of available regulatory alternatives and, if regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety effects, distributive impacts, and equity). A regulatory impact analysis (RIA) must be prepared for major rules with economically significant effects (\$100 million or more in any one year). This rule is not a major rule. The aggregate costs will total approximately \$37.3 million in any one year.

The RFA requires agencies to analyze options for regulatory relief of small businesses. For purposes of the RFA, small entities include small businesses, nonprofit organizations, and small governmental jurisdictions. Individuals and States are not included in the definition of a small entity. Small businesses are small entities, either by nonprofit status or by having revenues of \$6.5 million to \$31.9 million or less in any one year for purposes of the RFA. The September 19, 2007 final rule provided an analysis on the impact of small entities (72 FR 53642-3). The analysis published in the final rule remains valid. Since this interim final rule with comment merely updates the Congressional authority under which the rule operates, we have determined that this rule will not have a significant impact on small entities based on the overall effect on revenues.

Section 1102(b) of the Act requires us to prepare a regulatory impact analysis if a rule may have a significant impact on the operations of a substantial number of small rural hospitals. This analysis must conform to the provisions of section 604 of the RFA. For purposes of section 1102(b) of the Act, we define a small rural hospital as a hospital that is located outside of a Metropolitan statistical Area (superseded by Core Based Statistical Areas) and has fewer than 100 beds. This rule affects those small rural hospitals that have been cited for a deficiency based on noncompliance with required conditions of participation and for which a revisit is needed to ensure that the deficiency has been corrected. We identified in the September 19, 2007 final rule that for the effective period of that rule that less than 3 percent of all hospitals may be assessed a revisit user fee and that less than 1 percent of those hospitals would be rural hospitals (72 FR 53643). The analysis published in the final rule remains valid. Since this interim final rule with comment merely updates the Congressional authority under which the rule operates, we maintain that this rule will not have a

significant impact on small rural hospitals.

Section 202 of the Unfunded Mandates Reform Act of 1995 also requires that agencies assess anticipated costs and benefits before issuing any rule whose mandates require spending in any one year of \$100 million in 1995 dollars, updated annually for inflation. That threshold level is currently approximately \$120 million. This interim final rule with comment will have no mandated effect on State, local, or tribal governments and the impact on the private sector is estimated to be less than \$120 million and will only effect those Medicare providers or suppliers for which a revisit user fee is assessed based on the need to conduct a revisit survey to ensure deficient practices that were cited have been corrected.

Executive Order 13132 establishes certain requirements that an agency must meet when it promulgates a proposed rule (and subsequent final rule) that imposes substantial direct requirement costs on State and local governments, preempts State law, or otherwise has Federalism implications. This interim final rule with comment will not substantially affect State or local governments. This rule establishes user fees for providers and suppliers for which CMS has identified deficient practices and requires a revisit to assure that corrections have been made. Therefore, we have determined that this interim final rule with comment will not have a significant affect on the rights, roles, and responsibilities of State or local governments.

B. Impact on Providers/Suppliers

There is no change on the impact on providers and suppliers with the

publication of this interim final rule with comment. The impact remains as discussed in the final rule (72 FR 53643).

Final Fee Schedule for Onsite and Offsite Revisit Surveys

The FY 2007 fee schedule published on September 19, 2007 (72 FR 53647) in the final rule will be retained. As noted in the final rule, the published fee schedule will be utilized by CMS for the assessment of such fees until such time as a new fee schedule notice is proposed and published in final form. The calculations utilized to determine the fee as identified in the final rule will be the same (72 FR 53645–6). We will continue to assess a flat fee based on provider or supplier type and type of revisit survey conducted. Table A below identifies the final fee schedule.

TABLE A.—FINAL FEE SCHEDULE

Facility	Fee assessed per offsite revisit survey	Fee assessed per onsite revisit survey
SNF & NF Hospitals ...	\$168	\$2,072
HHA	168	2,554
Hospice	168	1,613
ASC	168	1,736
RHC	168	1,669
ESRD	168	851
		1,490

Costs for All Revisit User Fees Assessed

We anticipated that the combined costs for all providers and suppliers for all revisit surveys in FY 2007 would total approximately \$37.3 million on an annual basis, with onsite revisit surveys amounting to approximately \$34.6 million and offsite revisit surveys

totaling approximately \$2.7 million. (72 FR 53645). However, actual fees assessed in FY 2007 were much less than this amount, since CMS did not charge for revisits that occurred prior to publication of the final regulation. Since we continue to operate under this same estimate for FY 07, we provide below monthly estimates of the impact for the period of the current Continuing Resolution in Tables B and C. For the period of the current Continuing Resolution, we will use the FY 2007 fee schedule established in the final rule for the assessment of fees until a new fee schedule notice is proposed and published as final.

In Table B below, we provide the projected costs for the period of this current Continuing Resolution based on the fee schedule of the final rule. We expect the combined costs for all providers and suppliers for all onsite revisit surveys for the period of this current Continuing Resolution to total approximately \$2.9 million. We first multiplied the total number of onsite revisit surveys in one year by the expected revisit user fees assessed per revisits as finalized in Table A above, estimated by provider or supplier, to obtain the annual cost of revisit surveys. We then divided this number by 12 to obtain the monthly cost per provider or supplier of onsite revisit surveys to obtain the total costs for onsite revisit surveys for the period of the current Continuing Resolution (roughly 1 month). We then totaled all providers and suppliers to achieve the total costs for all onsite revisit surveys for the period of this current Continuing Resolution.

TABLE B.—ONSITE REVISIT SURVEYS—ESTIMATED MONTHLY COSTS

Facility	Monthly number of onsite revisit surveys	Fee assessed per onsite revisit surveys (hrs × \$112)	Monthly costs for onsite revisit surveys *
SNF & NF	1,191	\$2,072	\$2,467,061
Hospitals	48	2,554	122,379
HHA	89	1,613	143,557
Hospice	21	1,736	37,035
ASC	8	1,669	13,213
RHC	12	851	10,567
ESRD	58	1,490	86,668
Total	1,427	2,880,480

* Monthly costs may differ from the multiple of monthly revisits and fee per revisit due to rounding. The time period of this CR is roughly 1 month. Total numbers of onsite revisit surveys were rounded up based on FY 2006 actual data presented in the final rule.

We expect the combined costs for all providers and suppliers for all offsite revisit surveys to total \$229,250 for the period of the current Continuing

Resolution. In Table C below, we first estimated by provider or supplier the number of offsite revisit surveys expected for an entire fiscal year, and

multiplied this number by the expected revisit user fee of \$168 per offsite revisit survey to obtain the annual cost of surveys. We then divided this number

by 12 to obtain the monthly cost of period of the current Continuing
 offsite revisit surveys to obtain the total Resolution (roughly 1 month).
 costs for offsite revisit surveys for the

TABLE C.—OFFSITE REVISIT SURVEYS—ESTIMATED MONTHLY COSTS

Facility	Monthly number of offsite revisit surveys	Fee assessed per offsite revisit survey (\$112 × 1.5 hrs)	Monthly costs for offsite revisit surveys*
SNF & NF	1,262	\$168	\$211,932
Hospitals	23	168	3,892
HHA	43	168	7,238
Hospice	4	168	714
ASC	8	168	1,302
RHC	6	168	938
ESRD	19	168	3,234
Total	1,365	229,250

* Monthly costs may differ from the multiple of monthly revisits and fee per revisit due to rounding. The time period of this CR is roughly 1 month.

As shown in Table D below, we provide the aggregate costs expected as projected for the entire FY 2007, as well as the costs we would expect to offset for the period of the current Continuing Resolution.

TABLE D.—TOTAL COSTS COMBINED FOR ALL REVISITS SURVEYS PER FISCAL YEAR & PERIOD OF CR

	FY 2007	Period of CR*
Onsite Revisit Surveys	\$34,565,760	\$2,880,480
Offsite Revisit Surveys	2,751,000	229,250
Total Costs All Revisits	37,316,760	3,109,730

* CR period's costs are based on CR period revisit surveys rounded up to the nearest whole number as shown in Table B & C.

E. Alternatives Considered

We considered a number of alternatives to the revisit user fee program. Such alternatives were discussed in the final rule published on September 19, 2007 (72 FR 53647). We affirm the continuing validity of that analysis. The current Continuing Resolution provides CMS with the authority to continue projects or activities as was otherwise provided for in FY 2007, and as such CMS is required to publish an interim final rule with comment. This interim final rule with comment merely updates the Congressional authority under which the rule operates.

In accordance with Executive Order 12866, this rule has been reviewed by the Office of Management and Budget.

List of Subjects in 42 CFR Part 488

Administrative practice and procedure, Health facilities, Medicare, Reporting and Recording requirements.

■ For the reasons set forth in the preamble, the Centers for Medicare & Medicaid Services amends 42 CFR chapter IV, part 488 as set forth below:

PART 488—SURVEY, CERTIFICATION, AND ENFORCEMENT PROCEDURES

■ 1. The authority citation for part 488 is revised to read as follows:

Authority: Secs. 1102 and 1871 of the Social Security Act, unless otherwise noted (42 U.S.C. 1302 and 1395(hh)); Continuing Resolution Pub. L. 101–16 Division B of HR 3222.

(Catalog of Federal Domestic Assistance Program No. 93.778, Medical Assistance Program)

(Catalog of Federal Domestic Assistance Program No. 93.773, Medicare—Hospital Insurance; and Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: December 3, 2007.

Kerry Weems,

Acting Administrator, Centers for Medicare & Medicaid Services.

Approved: December 13, 2007.

Michael O. Leavitt,

Secretary.

[FR Doc. 07–6093 Filed 12–14–07; 12:13 pm]

BILLING CODE 4120–01–P

DEPARTMENT OF COMMERCE

National Oceanic and Atmospheric Administration

50 CFR Part 660

[Docket No. 060824226 6322 02]

RIN 0648–AW34

Magnuson-Stevens Act Provisions; Fisheries Off West Coast States; Pacific Coast Groundfish Fishery; Biennial Specifications and Management Measures; Inseason Adjustments

AGENCY: National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Commerce.

ACTION: Final rule; inseason adjustments to biennial groundfish management measures; request for comments.

SUMMARY: This final rule announces inseason changes to management measures in the commercial Pacific Coast groundfish fishery. These actions, which are authorized by the Pacific Coast Groundfish Fishery Management Plan (FMP), are intended to allow fisheries to access more abundant