

beneficiary. The Act changes the existing provisions that require a nonavailability statement (NAS) for inpatient delivery but do not require an NAS for outpatient prenatal, outpatient delivery and post-partum care. The change will provide for continuity of care for maternity patients. Beneficiaries will need one NAS for the entire episode of maternity care that shall remain valid until 42 days following termination of the pregnancy.

Regulatory Procedure

Executive order 12866 requires certain regulatory assessments for any significant regulatory action, defined as one which would result in an annual effect on the economy of \$100 million or more, or have other substantial impacts. The Regulatory Flexibility Act (RFA) requires that each Federal agency prepare, and make available for public comment, a regulatory flexibility analysis when the agency issues a regulation which would have a significant impact on a substantial number of small entities. This Interim Final Rule is not a significant regulatory action under E.O. 12866, nor would it have a significant impact on small entities. The changes set forth in the interim final rule are minor revision to the existing regulation.

The interim final rule will not impose additional information collection requirements on the public under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501-3511). This rule is being issued as an interim final rule, with comment period, as an exception to our standard practice of soliciting public comments prior to issuance. The Assistant Secretary of Defense (Health Affairs) has determined that following the standard practice in this case would be impracticable, unnecessary, and contrary to the public interest. This determination is based on several factors. First, this change directly implements a statutory amendment enacted by Congress expressively for this purpose. Second, this rule implements the statutory policy without embellishment. All public comments are invited.

List of Subject in 32 CFR Part 199

Claims, Handicapped, Health insurance, Military personnel.

PART 199—[AMENDED]

Accordingly, 32 CFR 199 is amended as follows:

1. The authority citation for Part 199 continues to read as follows:

Authority: 5 U.S.C. 301 and 10 U.S.C. Chapter 55.

2. Section 199.4(a) is amended by revising paragraphs (a)(9) and (a)(9)(i)(B).

§ 199.4 Basic program benefits.

(a) * * *

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(9) *Nonavailability Statements within a 40-mile catchment area.* In some geographic locations, it is necessary for CHAMPUS beneficiaries not enrolled in TRICARE Prime to determine whether the required medical care can be provided through a Uniformed Services facility. If the required care cannot be provided, the hospital commander, or designee, will issue a Nonavailability Statement (DD form 1251). Except for emergencies, a Nonavailability Statement should be issued before medical care is obtained from a civilian source. Failure to secure such a statement may waive the beneficiary's rights to benefits under CHAMPUS.

(i) * * *

(A) * * *

(B) For CHAMPUS beneficiaries who are not enrolled in TRICARE Prime, an NAS is required for services in connection with non-emergency inpatient hospital care and outpatient and inpatient maternity care if such services are available at a facility of the Uniformed Services located within a 40-mile radius of the residence of the beneficiary, except that an NAS is not required for services otherwise available at a facility of the Uniformed Services located within a 40-mile radius of the beneficiary's residence when another insurance plan or program provides the beneficiary primary coverage for the services. For maternity care, an NAS is required for services related to outpatient prenatal, outpatient or inpatient delivery, and outpatient post-partum care subsequent to the visit that confirms the pregnancy. The requirement for an NAS does not apply to beneficiaries enrolled in TRICARE Prime, even when those beneficiaries use the point-of-service option under § 199.17(n)(3).

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Dated: December 16, 1999.

L.M. Bynum,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

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DEPARTMENT OF DEFENSE

Department of the Air Force

32 CFR Part 806b

[Air Force Instruction 37-132]

Air Force Privacy Act Program

AGENCY: Department of the Air Force, DOD

ACTION: Final rule.

SUMMARY: The Department of the Air Force is adopting the exemption rule published on October 18, 1999, at 64 FR 56181 as final. No comments were received during the sixty day comment period.

EFFECTIVE DATE: December 17, 1999.

FOR FURTHER INFORMATION CONTACT: Mrs. Anne Rollins at (703) 588-6187.

SUPPLEMENTARY INFORMATION:

Executive Order 12866, 'Regulatory Planning and Review'

It has been determined that this Privacy Act rule is not a significant regulatory action. The rule does not:

(1) Have an annual effect to the economy of \$100 million or more; or adversely affect in a material way the economy; a section of the economy; productivity; competition; jobs; the environment; public health or safety; or state, local, or tribal governments or communities;

(2) Create a serious inconsistency or otherwise interfere with an action taken or planned by another Agency;

(3) Materially alter the budgetary impact of entitlements, grants, user fees, or loan programs or the rights and obligations of recipients thereof;

(4) Raise novel legal or policy issues arising out of legal mandates, the President's priorities, or the principles set forth in this Executive Order.

Public Law 96-354, 'Regulatory Flexibility Act' (5 U.S.C. 601)

It has been certified that this Privacy Act rule is not subject to the Regulatory Flexibility Act (5 U.S.C. 601) because it would not, if promulgated, have a significant economic impact on a substantial number of small entities.

Public Law 96-511, 'Paperwork Reduction Act' (44 U.S.C. Chapter 35)

It has been certified that this Privacy Act rule does not impose any reporting or record keeping requirements under the Paperwork Reduction Act of 1995.

List of subjects in 32 CFR part 806b

Privacy.

Accordingly, 32 CFR part 806b is revised to read as follows:

PART 806b—AIR FORCE PRIVACY ACT PROGRAM

1. The authority citation for 32 CFR Part 806b continues to read as follows:

Authority: Pub. L. 93-579, 88 Stat 1896 (5 U.S.C. 552a).

2. Appendix C to Part 806b is amended by adding paragraph (b)(21) as follows:

* * * * *

b. Specific exemptions.* * *

(21) *System identifier and name:*

F036 AF DP G, Military Equal Opportunity and Treatment.

(i) *Exemption:* Investigatory material compiled for law enforcement purposes may be exempt pursuant to 5 U.S.C. 552a(k)(2). However, if an individual is denied any right, privilege, or benefit for which he would otherwise be entitled by Federal law or for which he would otherwise be eligible, as a result of the maintenance of the information, the individual will be provided access to the information except to the extent that disclosure would reveal the identity of a confidential source. Portions of this system of records may be exempt pursuant to 5 U.S.C. 552a(d), (e)(4)(H), and (f).

(iii) *Authority:* 5 U.S.C. 552a(k)(2)

(iv) *Reasons:* (1) From subsection (d) because access to the records contained in this system would inform the subject of an investigation of the existence of that investigation, provide the subject of the investigation with information that might enable him to avoid detection, and would present a serious impediment to law enforcement. In addition, granting individuals access to information collected while an Equal Opportunity and Treatment clarification/investigation is in progress conflicts with the just, thorough, and timely completion of the complaint, and could possibly enable individuals to interfere, obstruct, or mislead those clarifying/investigating the complaint.

(2) From subsection (e)(4)(H) because this system of records is exempt from individual access pursuant to subsection (k) of the Privacy Act of 1974.

(3) From subsection (f) because this system of records has been exempted from the access provisions of subsection (d).

(4) Consistent with the legislative purpose of the Privacy Act of 1974, the Department of the Air Force will grant access to nonexempt material in the records being maintained. Disclosure will be governed by the Department of the Air Force's Privacy Instruction, but will be limited to the extent that the identity of confidential sources will not be compromised; subjects of an

investigation of an actual or potential violation will not be alerted to the investigation; the physical safety of witnesses, informants and law enforcement personnel will not be endangered, the privacy of third parties will not be violated; and that the disclosure would not otherwise impede effective law enforcement. Whenever possible, information of the above nature will be deleted from the requested documents and the balance made available. The controlling principle behind this limited access is to allow disclosures except those indicated above. The decisions to release information from this system will be made on a case-by-case basis.

Dated: December 16, 1999.

L.M. Bynum,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

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ENVIRONMENTAL PROTECTION AGENCY**40 CFR Parts 52 and 70**

[Region VII Tracking No. MO 083-1083a; FRL-6510-9]

Approval and Promulgation of Implementation Plans; State of Missouri

AGENCY: Environmental Protection Agency (EPA).

ACTION: Direct final rule.

SUMMARY: The Environmental Protection Agency (EPA) is taking final action to approve certain portions of the State Implementation Plan (SIP) revisions submitted by the state of Missouri and as revisions to the part 70 (operating permits) program. These revisions established emission and service fees for 1997 and 1998 and clarify language regarding reporting requirements, emission calculations and verification.

DATES: This direct final rule is effective on February 22, 2000 without further notice, unless EPA receives adverse comment by January 24, 2000. If adverse comment is received, EPA will publish a timely withdrawal of the direct final rule in the **Federal Register** and inform the public that the rule will not take effect.

ADDRESSES: All comments should be addressed to: Kim Johnson, Environmental Protection Agency, Air Planning and Development Branch, 901 North 5th Street, Kansas City, Kansas 66101.

Copies of the state submittal(s) are available at the following addresses for inspection during normal business hours: Environmental Protection Agency, Air Planning and Development Branch, 901 North 5th Street, Kansas City, Kansas 66101; and the Environmental Protection Agency, Air and Radiation Docket and Information Center, Air Docket (6102), 401 M Street, SW, Washington, DC 20460.

FOR FURTHER INFORMATION CONTACT: Kim Johnson, Environmental Protection Agency, Air Planning and Development Branch, 901 North 5th Street, Kansas City, Kansas 66101, (913) 551-7975.

SUPPLEMENTARY INFORMATION:**Background***What is a SIP?*

Section 110 of the Clean Air Act (CAA) requires states to develop air pollution regulations and control strategies to ensure that state air quality meets the national ambient air quality standards established by EPA. These ambient standards are established under section 109 of the CAA, and they currently address six criteria pollutants. These pollutants are: carbon monoxide, nitrogen dioxide, ozone, lead, particulate matter (PM), and sulfur dioxide.

Each state must submit these regulations and control strategies to EPA for approval and incorporation into the Federally enforceable SIP.

The CAA requires each state to have a Federally approved SIP which protects air quality primarily by addressing air pollution at its point of origin. These SIPs can be extensive, containing state regulations or other enforceable documents and supporting information such as emission inventories, monitoring networks, and modeling demonstrations.

What is the Federal Approval Process for a SIP?

In order for state regulations to be incorporated into the Federally enforceable SIP, states must formally adopt the regulations and control strategies consistent with state and Federal requirements. This process generally includes a public notice, public hearing, public comment period, and a formal adoption by a state-authorized rulemaking body.

Once a state rule, regulation, or control strategy is adopted, the state submits it to EPA for inclusion into the SIP. EPA must provide public notice and seek additional public comment regarding the proposed Federal action on the state submission. If adverse comments are received, they must be