§ 17.367 Republic of the Philippines to print forms.

The Secretary of National Defense of the Republic of the Philippines will, with the concurrence of the Secretary of Veterans Affairs, print all forms for applications for hospitalization, forms for physical examination reports, forms for billings for services rendered, and such other forms as may be necessary and incident to the efficient execution of the program governed by the provisions of 38 U.S.C. 1724 and 1732, and 38 CFR 17.36 through 17.40 and §§ 17.350 through 17.370. The forms will be used whenever applicable in the general operation of the program.

[33 FR 5301, Apr. 3, 1968, as amended at 61 FR 21969, May 13, 1996]

§ 17.369 Inspections.

The U.S. Department of Veterans Affairs, through authorized representatives, has the right under the agreements cited in § 17.350, to inspect the Veterans Memorial Medical Center, its premises and all appurtenances and records to determine completeness and correctness of such records, and to determine according to the provisions of the cited agreements whether standards maintained conform to the necessary requirements.

[33 FR 5301, Apr. 3, 1968, as amended at 47 FR 58251, Dec. 30, 1982]

§ 17.370 Termination of payments.

Payments may be terminated if the U.S. Department of Veterans Affairs determines the Veterans Memorial Medical Center has not replaced and upgraded as needed equipment during the period in which the agreements cited in § 17.50 are in effect or has not rehabilitated the existing physical plant and facilities to place the medical center on a sound and effective operating basis, or has not maintained the medical center in a well-equipped and effective operating condition. Payments, however, will not be stopped unless the Veterans Memorial Medical Center has been given at least 60 days advance written notice of intent to stop payments.

(Authority: 38 U.S.C. 1732)

[33 FR 5301, Apr. 3, 1968, as amended at 47 FR 58251, Dec. 30, 1982]

CONFIDENTIALITY OF HEALTHCARE QUALITY ASSURANCE REVIEW RECORDS

AUTHORITY: 38 U.S.C. 5705.

SOURCE: 59 FR 53355, Oct. 24, 1994, unless otherwise noted.

§ 17.500 General.

(a) Section 5705, title 38, United States Code was enacted to protect the integrity of the VA's medical quality assurance program by making confidential and privileged certain records and documents generated by this program and information contained therein. Disclosure of quality assurance records and documents made confidential and privileged by 38 U.S.C. 5705 and the regulations in §§ 17.500 through 17.511 may only be made in accordance with the provisions of 38 U.S.C. 5705 and those regulations.

(b) The purpose of the regulations in §§ 17.500 through 17.511 is to specify and provide for the limited disclosure of those quality assurance documents which are confidential under the provisions of 38 U.S.C. 5705.

(c) For purposes of the regulations in §§ 17.500 through 17.511, the VA's medical quality assurance program consists of systematic healthcare reviews carried out by or for VA for the purpose of improving the quality of medical care or improving the utilization of healthcare resources in VA medical facilities. These review activities may involve continuous or periodic data collection and may relate to either the structure, process, or outcome of health care provided in the VA.

(d) Nothing in the regulations in §§ 17.500 through 17.511 shall be construed as authority to withhold any record or document from a committee or subcommittee of either House of Congress or any joint committee or subcommittee of Congress, if such record or document pertains to any matter within the jurisdiction of such committee or joint committee.

(e) The regulations in §§17.500 through 17.511 do not waive the sovereign immunity of the United States, and do not waive the confidentiality provisions and disclosure restrictions of 38 U.S.C. 5705.

(Authority: 38 U.S.C. 5705)

§ 17.501 Confidential and privileged documents.

(a) Documents and parts of documents are considered confidential and privileged if they were produced by or for the VA in the process of conducting systematic healthcare reviews for the purpose of improving the quality of health care or improving the utilization of healthcare resources in VA healthcare facilities and meet the criteria in paragraphs (b) and (c) of this section. The four classes of healthcare quality assurance reviews with examples are:

(1) Monitoring and evaluation reviews conducted by a facility:

(i) Medical records reviews,

(ii) Drug usage evaluations,

(iii) Blood usage reviews,

(iv) Surgical case/invasive procedure reviews,

(v) Service and program monitoring including monitoring performed by individual services or programs, several services or programs working together, or individuals from several services or programs working together as a team,

(vi) Mortality and morbidity reviews,

(vii) Infection control review and surveillance,

(viii) Occurrence screening,

(ix) Tort claims peer reviews (except reviews performed to satisfy the requirements of a governmental body or a professional health care organization which is licensing practitioners or monitoring their professional performance),

(x) Admission and continued stay reviews,

(xi) Diagnostic studies utilization reviews,

(xii) Reports of special incidents (VA Form 10-2633 or similar forms) and follow-up documents unless developed during or as a result of a Board of Investigation;

(2) Focused reviews which address specific issues or incidents and which are designated by the reviewing office at the outset of the review as protected by 38 U.S.C. 5705 and the regulations in §§17.500 through 17.511; focused reviews may be either:

(i) Facility focused reviews;

(ii) VA Central Office or Regional focused reviews;

(3) VA Central Office or Regional general oversight reviews to assess facility compliance with VA program requirements if the reviews are designated by the reviewing office at the outset of the review as protected by 38 U.S.C. 5705 and the regulations in §§17.500 through 17.511; and

(4) Contracted external reviews of care, specifically designated in the contract or agreement as reviews protected by 38 U.S.C. 5705 and the regulations in §§17.500 through 17.511.

(b) The Under Secretary for Health, Regional Director or facility Director will describe in advance in writing those quality assurance activities included under the classes of healthcare quality assurance reviews listed in paragraph (a) of this section. Only documents and parts of documents resulting from those activities which have been so described are protected by 38 U.S.C. 5705 and the regulations in §§17.500 through 17.511. If an activity is not described in a VA Central Office or Regional policy document, this requirement may be satisfied at the facility level by description in advance of the activity and its designation as protected in the facility quality assurance plan or other policy document.

(c) Documents and parts of documents generated by activities which meet the criteria in paragraphs (a) and (b) of this section shall be confidential and privileged only if they:

(1) Identify, either implicitly or explicitly, individual practitioners, patients, or reviewers except as provided in paragraph (g)(6) of this section; or

(2) Contain discussions relating to the quality of VA medical care or utilization of VA medical resources by healthcare evaluators during the course of a review of quality assurance information or data, even if they do not identify practitioners, patients, or reviewers; or

(3) Are individual committee, service, or study team minutes, notes, reports, memoranda, or other documents either