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for Reserves (including commissioned officers of the Reserve Corps of the Public Health Service) by the Secretary concerned under section 206, title 37 U.S.C., or any other provision of law;

(2) Special additional duties authorized for Reserves (including commissioned officers of the Reserve Corps of the Public Health Service) by an authority designated by the Secretary concerned and performed by them on a voluntary basis in connection with the prescribed training or maintenance activities of the units to which they are assigned.

(3) Duty (other than full-time duty) for members of the National Guard or Air National Guard of any State under the provisions of law stated in paragraph (c)(3) of this section.

(4) Inactive duty for training does not include work or study performed in connection with correspondence courses, or attendance at an educational institution in an inactive status, or duty performed as a temporary member of the Coast Guard Reserve.

[34 FR 9339, June 13, 1969, as amended at 45 FR 6934, Jan. 31, 1980; 45 FR 43169, June 26, 1980; 48 FR 56580, Dec. 22, 1983; 61 FR 21965, May 13, 1996]

§ 17.31 Duty periods defined.

Full-time duty as a member of the Women's Army Auxiliary Corps, Women's Reserve of the Navy and Marine Corps and Women's Reserve of the Coast Guard.

[34 FR 9339, June 13, 1969, as amended at 61 FR 21965, May 13, 1996]

EDITORIAL NOTE: At 61 FR 21965, May 13, 1996, § 17.31(b)(5) was redesignated as § 17.31.

PROTECTION OF PATIENT RIGHTS

§ 17.32 Informed consent and advance care planning.

(a) Definitions:

Advance Directive. Specific written statements made by a patient who has decision-making capacity regarding future health care decisions in any of the following:

(i) VA Living Will. A written statement made by a patient on an authorized VA form which sets forth the patient's wishes regarding the patient's health care treatment preferences in-

cluding the withholding and withdrawal of life-sustaining treatment.

(ii) VA Durable Power of Attorney for Health Care. A written instruction on a VA form which designates the patient's choice of health care agent.

(iii) State-Authorized Advance Directive. A non-VA living will, durable power of attorney for health care, or other advance health care planning document, the validity of which is determined pursuant to applicable State law. For the purposes of this paragraph and paragraph (h) of this section, "applicable State law" means the law of the State where the advance directive was signed, the State where the patient resided when the advance directive was signed, the State where the patient now resides, or the State where the patient is receiving treatment. VA will resolve any conflict between those State laws regarding the validity of the advance directive by following the law of the State that gives effect to the expressed wishes in the advance directive.

Close friend. Any person eighteen years or older who has shown care and concern for the patient's welfare, who is familiar with the patient's activities, health, religious beliefs and values, and who has presented a signed written statement for the record that describes that person's relationship to and familiarity with the patient.

Decision-making capacity. The ability to understand and appreciate the nature and consequences of health care treatment decisions.

Health care agent. An individual named by the patient in a Durable Power of Attorney for Health Care.

Legal guardian. A person appointed by a court of appropriate jurisdiction to make decisions for an individual who has been judicially determined to be incompetent.

Practitioner. Any physician, dentist, or health care professional who has been granted specific clinical privileges to perform the treatment or procedure. For the purpose of obtaining informed consent for medical treatment, the term practitioner includes medical and dental residents and other appropriately trained health care professionals designated by VA regardless of

whether they have been granted clinical privileges.

Signature consent. The patient's or surrogate's signature on a VA-authorized consent form.

Special guardian. A person appointed by a court of appropriate jurisdiction for the specific purpose of making health care decisions.

Surrogate. An individual, organization or other body authorized under this section to give informed consent on behalf of a patient who lacks decision-making capacity.

(b) *Policy.* Except as otherwise provided in this section, all patient care furnished under title 38 U.S.C. shall be carried out only with the full and informed consent of the patient or, in appropriate cases, a representative thereof. In order to give informed consent, the patient must have decision-making capacity and be able to communicate decisions concerning health care. If the patient lacks decision-making capacity or has been declared incompetent, consent must be obtained from the patient's surrogate. Practitioners may provide necessary medical care in emergency situations without the patient's or surrogate's express consent when immediate medical care is necessary to preserve life or prevent serious impairment of the health of the patient or others and the patient is unable to consent and the practitioner determines that the patient has no surrogate or that waiting to obtain consent from the patient's surrogate would increase the hazard to the life or health of the patient or others. In such circumstances consent is implied.

(c) *General requirements for informed consent.* Informed consent is the freely given consent that follows a careful explanation by the practitioner to the patient or the patient's surrogate of the proposed diagnostic or therapeutic procedure or course of treatment. The practitioner, who has primary responsibility for the patient or who will perform the particular procedure or provide the treatment, must explain in language understandable to the patient or surrogate the nature of a proposed procedure or treatment; the expected benefits; reasonably foreseeable associated risks, complications or side effects; reasonable and available alter-

natives; and anticipated results if nothing is done. The patient or surrogate must be given the opportunity to ask questions, to indicate comprehension of the information provided, and to grant permission freely without coercion. The practitioner must advise the patient or surrogate if the proposed treatment is novel or unorthodox. The patient or surrogate may withhold or revoke his or her consent at any time.

(d) *Documentation of informed consent.* (1) The informed consent process must be appropriately documented in the health record. In addition, signature consent is required for all diagnostic and therapeutic treatments or procedures that:

- (i) Require the use of sedation;
- (ii) Require anesthesia or narcotic analgesia;
- (iii) Are considered to produce significant discomfort to the patient;
- (iv) Have a significant risk of complication or morbidity;
- (v) Require injections of any substance into a joint space or body cavity; or
- (vi) Involve testing for Human Immunodeficiency Virus (HIV).

(2) A patient or surrogate will sign with an "X" when the patient or surrogate has a debilitating illness or disability, *i.e.*, significant physical impairment and/or difficulty in executing a signature due to an underlying health condition(s), or is unable to read and write. When the patient's or surrogate's signature is indicated by an "X," two adults must witness the act of signing. By signing, the witnesses are attesting only to the fact that they saw the patient or surrogate and the practitioner sign the form. The signed form must be filed in the patient's health record. A properly executed VA-authorized consent form is valid for a period of 60 calendar days. If, however, the treatment plan involves multiple treatments or procedures, it will not be necessary to repeat the informed consent discussion and documentation so long as the course of treatment proceeds as planned, even if treatment extends beyond the 60-day period. If there is a change in the patient's condition that might alter the diagnostic or therapeutic decision, the consent is automatically rescinded.

(3) If it is impractical to consult with the surrogate in person, informed consent may be obtained by mail, facsimile, or telephone. A facsimile copy of a signed consent form is adequate to proceed with treatment. However, the surrogate must agree to submit a signed consent form to the practitioner. If consent is obtained by telephone, the conversation must be audiotaped or witnessed by a second VA employee. The name of the person giving consent and his or her authority to act as surrogate must be adequately identified for the record.

(e) *Surrogate consent.* If the practitioner who has primary responsibility for the patient determines that the patient lacks decision-making capacity and is unlikely to regain it within a reasonable period of time, informed consent must be obtained from the patient's surrogate. Patients who are incapable of giving consent as a matter of law, i.e., persons judicially determined to be incompetent and minors not otherwise able to provide informed consent, will be deemed to lack decision-making capacity for the purposes of this section. If the patient is considered a minor in the state where the VA facility is located and cannot consent to medical treatment, consent must be obtained from the patient's parent or legal guardian. The surrogate generally assumes the same rights and responsibilities as the patient in the informed consent process. The surrogate's decision must be based on his or her knowledge of what the patient would have wanted, i.e., substituted judgment. If the patient's wishes are unknown, the decision must be based on the patient's best interest. The following persons are authorized to consent on behalf of patients who lack decision-making capacity in the following order of priority:

- (1) Health care agent;
- (2) Legal guardian or special guardian;
- (3) Next-of-kin: a close relative of the patient eighteen years of age or older, in the following priority: spouse, child, parent, sibling, grandparent, or grandchild; or
- (4) Close friend.

(f) *Consent for patients without surrogates.* (1) If none of the surrogates listed in paragraph (e) of this section are

available, the practitioner may request Regional Counsel assistance to obtain a special guardian for health care or follow the procedures outlined in this paragraph (f).

(2) Facilities may use the following process to make treatment decisions for patients who lack decision-making capacity and have no surrogate. For treatments or procedures that involve minimal risk, the practitioner must verify that no authorized surrogate can be located. The practitioner must attempt to explain the nature and purpose of the proposed treatment to the patient and enter this information in the health record. For procedures that require signature consent, the practitioner must certify that the patient has no surrogate. The attending physician and the Chief of Service (or his or her designee) must indicate their approval of the treatment decision in writing. Any decision to withhold or withdraw life-sustaining treatment for such patients must be reviewed by a multi-disciplinary committee appointed by the facility Director. The committee functions as the patient's advocate and may not include members of the treatment team. The committee must submit its findings and recommendations in a written report to the Chief of Staff who must note his or her approval of the report in writing. After reviewing the record, the facility Director may concur with the decision to withhold or withdraw life support or request further review by Regional Counsel.

(g) *Special consent situations.* In addition to the other requirements of this section, additional protections are required in the following situations.

(1) No patient will undergo any unusual or extremely hazardous treatment or procedure, e.g., that which might result in irreversible brain damage or sterilization, except as provided in this paragraph (g). Before treatment is initiated, the patient or surrogate must be given adequate opportunity to consult with independent specialists, legal counsel or other interested parties of his or her choosing. The patient's or surrogate's signature on a VA authorized consent form must be witnessed by someone who is not affiliated with the VA health care facility,

e.g., spouse, legal guardian, or patient advocate. If a surrogate makes the treatment decision, a multi-disciplinary committee, appointed by the facility Director, must review that decision to ensure it is consistent with the patient's wishes or in his or her best interest. The committee functions as the patient's advocate and may not include members of the treatment team. The committee must submit its findings and recommendations in a written report to the facility Director. The Director may authorize treatment consistent with the surrogate's decision or request that a special guardian for health care be appointed to make the treatment decision.

(2) Administration of psychotropic medication to an involuntarily committed patient against his or her will must meet the following requirements. The patient or surrogate must be allowed to consult with independent specialists, legal counsel or other interested parties concerning the treatment with psychotropic medication. Any recommendation to administer or continue medication against the patient's or surrogate's will must be reviewed by a multi-disciplinary committee appointed by the facility Director for this purpose. This committee must include a psychiatrist or a physician who has psychopharmacology privileges. The facility Director must concur with the committee's recommendation to administer psychotropic medications contrary to the patient's or surrogate's wishes. Continued therapy with psychotropic medication must be reviewed every 30 days. The patient (or a representative on the patient's behalf) may appeal the treatment decision to a court of appropriate jurisdiction.

(3) If a proposed course of treatment or procedure involves approved medical research in whole or in part, the patient or representative shall be advised of this. Informed consent shall be obtained specifically for the administration or performance of that aspect of the treatment or procedure that involves research. Such consent shall be in addition to that obtained for the administration or performance of the nonresearch aspect of the treatment or procedure and must meet the requirements for informed consent set forth in

38 CFR Part 16, *Protection of Human Subjects*.

(4) Testing for Human Immunodeficiency Virus (HIV) must be voluntary and must be conducted only with the prior informed and (written) signature consent of the patient or surrogate. Patients who consent to testing for HIV must sign VA form 10-012, "Consent for HIV Antibody Testing." This form must be filed in the patient's health record. Testing must be accompanied by pre-test and post-test counseling.

(h) *Advance health care planning*. Subject to the provisions of paragraphs (h)(1) through (h)(4) of this section, VA will follow the wishes of a patient expressed in an Advance Directive when the attending physician determines and documents in the patient's health record that the patient lacks decision-making capacity and is not expected to regain it. An advance directive that is valid in one or more States under applicable State law, as defined in paragraph (a) of this section, will be recognized throughout the VA health care system.

(1) *Witnesses*. A VA Advance Directive: Living Will and Durable Power of Attorney for Health Care must be signed by the patient in the presence of two witnesses. Neither witness may to the witness' knowledge be named in the patient's will, appointed as health care agent in the advance directive, or financially responsible for the patient's care. VA employees of the Chaplain Service, Psychology Service, Social Work Service, or nonclinical employees (*e.g.*, Medical Administration Service, Voluntary Service, or Environmental Management Service) may serve as witnesses. Other individuals employed by the VA facility in which the patient is being treated may not sign as witnesses to the advance directive. Witnesses are attesting only to the fact that they saw the patient sign the form.

(2) *Instructions in critical situations*. VA will follow the unambiguous verbal or non-verbal instructions regarding future health care decisions of a patient who has decision-making capacity when the patient is admitted to

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care when critically ill and loss of capacity may be imminent and the patient is not physically able to sign an advance directive form, or the appropriate form is not readily available. The patient's instructions must have been expressed to at least two members of the health care team. The substance of the patient's instructions must be recorded in a progress note in the patient's health record and must be co-signed by at least two members of the health care team who were present and can attest to the wishes expressed by the patient. These instructions will be given effect only if the patient loses decision-making capacity during the presenting situation.

(3) *Revocation.* A patient who has decision-making capacity may revoke an advance directive or instructions in a critical situation at any time by using any means expressing the intent to revoke.

(4) *VA policy and disputes.* Neither the treatment team nor surrogate may override a patient's clear instructions in an Advance Directive or in instructions in critical situations, except that those portions of an Advance Directive or instructions given in a critical situation that are not consistent with VA policy will not be given effect.

(Authority: 38 U.S.C. 7331-7334)

(The information collection requirements in this section have been approved by the Office of Management and Budget under control number 2900-0583)

[62 FR 53961, Oct. 17, 1997, as amended at 70 FR 71774, Nov. 30, 2005; 71 FR 68740, Nov. 28, 2006; 72 FR 10366, Mar. 8, 2007]

§ 17.33 Patients' rights.

(a) *General.* (1) Patients have a right to be treated with dignity in a humane environment that affords them both reasonable protection from harm and appropriate privacy with regard to their personal needs.

(2) Patients have a right to receive, to the extent of eligibility therefor under the law, prompt and appropriate treatment for any physical or emotional disability.

(3) Patients have the right to the least restrictive conditions necessary to achieve treatment purposes.

(4) No patient in the Department of Veterans Affairs medical care system,

except as otherwise provided by the applicable State law, shall be denied legal rights solely by virtue of being voluntarily admitted or involuntarily committed. Such legal rights include, but are not limited to, the following:

(i) The right to hold and to dispose of property except as may be limited in accordance with paragraph (c)(2) of this section;

(ii) The right to execute legal instruments (e.g., will);

(iii) The right to enter into contractual relationships;

(iv) The right to register and vote;

(v) The right to marry and to obtain a separation, divorce, or annulment;

(vi) The right to hold a professional, occupational, or vehicle operator's license.

(b) *Residents and inpatients.* Subject to paragraphs (c) and (d) of this section, patients admitted on a residential or inpatient care basis to the Department of Veterans Affairs medical care system have the following rights:

(1) *Visitations and communications.* Each patient has the right to communicate freely and privately with persons outside the facility, including government officials, attorneys, and clergymen. To facilitate these communications each patient shall be provided the opportunity to meet with visitors during regularly scheduled visiting hours, convenient and reasonable access to public telephones for making and receiving phone calls, and the opportunity to send and receive unopened mail.

(i) Communications with attorneys, law enforcement agencies, or government officials and representatives of recognized service organizations when the latter are acting as agents for the patient in a matter concerning Department of Veterans Affairs benefits, shall not be reviewed.

(ii) A patient may refuse visitors.

(iii) If a patient's right to receive unopened mail is restricted pursuant to paragraph (c) of this section, the patient shall be required to open the sealed mail while in the presence of an appropriate person for the sole purpose of ascertaining whether the mail contains contraband material, i.e., implements which pose significant risk of bodily harm to the patient or others or

any drugs or medication. Any such material will be held for the patient or disposed of in accordance with instructions concerning patients' mail published by the Veterans Health Administration, Department of Veterans Affairs, and/or the local health care facility.

(iv) Each patient shall be afforded the opportunity to purchase, at the patient's expense, letter writing material including stamps. In the event a patient needs assistance in purchasing writing material, or in writing, reading or sending mail, the medical facility will attempt, at the patient's request, to provide such assistance by means of volunteers, sufficient to mail at least one (1) letter each week.

(v) All information gained by staff personnel of a medical facility during the course of assisting a patient in writing, reading, or sending mail is to be kept strictly confidential except for any disclosure required by law.

(2) *Clothing.* Each patient has the right to wear his or her own clothing.

(3) *Personal Possessions.* Each patient has the right to keep and use his or her own personal possessions consistent with available space, governing fire safety regulations, restrictions on noise, and restrictions on possession of contraband material, drugs and medications.

(4) *Money.* Each patient has the right to keep and spend his or her own money and to have access to funds in his or her account in accordance with instructions concerning personal funds of patients published by the Veterans Health Administration.

(5) *Social Interaction.* Each patient has the right to social interaction with others.

(6) *Exercise.* Each patient has the right to regular physical exercise and to be outdoors at regular and frequent intervals. Facilities and equipment for such exercise shall be provided.

(7) *Worship.* The opportunity for religious worship shall be made available to each patient who desires such opportunity. No patient will be coerced into engaging in any religious activities against his or her desires.

(c) *Restrictions.* (1) A right set forth in paragraph (b) of this section may be restricted within the patient's treatment

plan by written order signed by the appropriate health care professional if—

(i) It is determined pursuant to paragraph (c)(2) of this section that a valid and sufficient reason exists for a restriction, and

(ii) The order imposing the restriction and a progress note detailing the indications therefor are both entered into the patient's permanent medical record.

(2) For the purpose of paragraph (c) of this section, a valid and sufficient reason exists when, after consideration of pertinent facts, including the patient's history, current condition and prognosis, a health care professional reasonably believes that the full exercise of the specific right would—

(i) Adversely affect the patient's physical or mental health,

(ii) Under prevailing community standards, likely stigmatize the patient's reputation to a degree that would adversely affect the patient's return to independent living,

(iii) Significantly infringe upon the rights of or jeopardize the health or safety of others, or

(iv) Have a significant adverse impact on the operation of the medical facility, to such an extent that the patient's exercise of the specific right should be restricted. In determining whether a patient's specific right should be restricted, the health care professional concerned must determine that the likelihood and seriousness of the consequences that are expected to result from the full exercise of the right are so compelling as to warrant the restriction. The Chief of Service or Chief of Staff, as designated by local policy, should concur with the decision to impose such restriction. In this connection, it should be noted that there is no intention to imply that each of the reasons specified in paragraphs (c)(2)(i) through (iv) of this section are logically relevant to each of the rights set forth in paragraph (b)(1) of this section.

(3) If it has been determined under paragraph (c)(2) of this section that a valid and sufficient reason exists for restricting any of the patient's rights set forth in paragraph (b) of this section, the least restrictive method for protecting the interest or interests

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specified in paragraphs (c)(2)(i) through (iv) of this section that are involved shall be employed.

(4) The patient must be promptly notified of any restriction imposed under paragraph (c) of this section and the reasons therefor.

(5) All restricting orders under paragraph (c) of this section must be reviewed at least once every 30 days by the practitioner and must be concurred in by the Chief of Service or Chief of Staff.

(d) *Restraint and seclusion of patients.*

(1) Each patient has the right to be free from physical restraint or seclusion except in situations in which there is a substantial risk of imminent harm by the patient to himself, herself, or others and less restrictive means of preventing such harm have been determined to be inappropriate or insufficient. Patients will be physically restrained or placed in seclusion only on the written order of an appropriate licensed health care professional. The reason for any restraint order will be clearly documented in the progress notes of the patient's medical record. The written order may be entered on the basis of telephonic authority, but in such an event, an appropriate licensed health care professional must examine the patient and sign a written order within an appropriate timeframe that is in compliance with current community and/or accreditation standards. In emergency situations, where inability to contact an appropriate licensed health care professional prior to restraint is likely to result in immediate harm to the patient or others, the patient may be temporarily restrained by a member of the staff until appropriate authorization can be received from an appropriate licensed health care professional. Use of restraints or seclusion may continue for a period of time that does not exceed current community and/or accreditation standards, within which time an appropriate licensed health care professional shall again be consulted to determine if continuance of such restraint or seclusion is required. Restraint or seclusion may not be used as a punishment, for the convenience of staff, or as a substitute for treatment programs.

(2) While in restraint or seclusion, the patient must be seen within appropriate timeframes in compliance with current community and/or accreditation standards:

(i) By an appropriate health care professional who will monitor and chart the patient's physical and mental condition; and

(ii) By other ward personnel as frequently as is reasonable under existing circumstances.

(3) Each patient in restraint or seclusion shall have bathroom privileges according to his or her needs.

(4) Each patient in restraint or seclusion shall have the opportunity to bathe at least every twenty-four (24) hours.

(5) Each patient in restraint or seclusion shall be provided nutrition and fluid appropriately.

(e) *Medication.* Patients have a right to be free from unnecessary or excessive medication. Except in an emergency, medication will be administered only on a written order of an appropriate health care professional in that patient's medical record. The written order may be entered on the basis of telephonic authority received from an appropriate health care professional, but in such event, the written order must be countersigned by an appropriate health care professional within 24 hours of the ordering of the medication. An appropriate health care professional will be responsible for all medication given or administered to a patient. A review by an appropriate health care professional of the drug regimen of each inpatient shall take place at least every thirty (30) days. It is recognized that administration of certain medications will be reviewed more frequently. Medication shall not be used as punishment, for the convenience of the staff, or in quantities which interfere with the patient's treatment program.

(f) *Confidentiality.* Information gained by staff from the patient or the patient's medical record will be kept confidential and will not be disclosed except in accordance with applicable law.

(g) *Patient grievances.* Each patient has the right to present grievances with respect to perceived infringement of the rights described in this section

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or concerning any other matter on behalf of himself, herself or others, to staff members at the facility in which the patient is receiving care, other Department of Veterans Affairs officials, government officials, members of Congress or any other person without fear or reprisal.

(h) *Notice of patient's rights.* Upon the admission of any patient, the patient or his/her representative shall be informed of the rights described in this section, shall be given a copy of a statement of those rights and shall be informed of the fact that the statement of rights is posted at each nursing station. All staff members assigned to work with patients will be given a copy of the statement of rights and these rights will be discussed with them by their immediate supervisor.

(i) *Other rights.* The rights described in this section are in addition to and not in derogation of any statutory, constitutional or other legal rights.

(Authority: 38 U.S.C. 501, 1721)

[47 FR 55486, Dec. 10, 1982. Redesignated at 61 FR 21965, May 13, 1996, as amended at 70 FR 67094, Nov. 4, 2005]

TENTATIVE ELIGIBILITY DETERMINATIONS

§ 17.34 Tentative eligibility determinations.

Subject to the provisions of §§ 17.36 through 17.38, when an application for hospital care or other medical services, except outpatient dental care, has been filed which requires an adjudication as to service connection or a determination as to any other eligibility prerequisite which cannot immediately be established, the service (including transportation) may be authorized without further delay if it is determined that eligibility for care probably will be established. Tentative eligibility determinations under this section, however, will only be made if:

(a) *In emergencies.* The applicant needs hospital care or other medical services in emergency circumstances, or

(b) *For persons recently discharged from service.* The application was filed within 6 months after date of honorable dis-

charge from a period of not less than 6 months of active duty.

[35 FR 6586, Apr. 24, 1970. Redesignated at 61 FR 21965, May 13, 1996, as amended at 64 FR 54212, Oct. 6, 1999]

HOSPITAL OR NURSING HOME CARE AND MEDICAL SERVICES IN FOREIGN COUNTRIES

§ 17.35 Hospital care and medical services in foreign countries.

The Secretary may furnish hospital care and medical services to any veteran sojourning or residing outside the United States, without regard to the veteran's citizenship:

(a) If necessary for treatment of a service-connected disability, or any disability associated with and held to be aggravating a service-connected disability;

(b) If the care is furnished to a veteran participating in a rehabilitation program under 38 U.S.C. chapter 31 who requires care for the reasons enumerated in 38 CFR 17.48(j)(2).

(Authority: 38 U.S.C. 1724)

[55 FR 11370, Mar. 28, 1990. Redesignated at 61 FR 21965, May 13, 1996]

ENROLLMENT PROVISIONS AND MEDICAL BENEFITS PACKAGE

§ 17.36 Enrollment—provision of hospital and outpatient care to veterans.

(a) *Enrollment requirement for veterans.*
(1) Except as otherwise provided in § 17.37, a veteran must be enrolled in the VA healthcare system as a condition for receiving the 'medical benefits package' set forth in § 17.38.

NOTE TO PARAGRAPH (a)(1): A veteran may apply to be enrolled at any time. (See § 17.36(d)(1).)

(2) Except as provided in paragraph (a)(3) of this section, a veteran enrolled under this section and who, if required by law to do so, has agreed to make any applicable copayment is eligible for VA hospital and outpatient care as provided in the "medical benefits package" set forth in § 17.38.

NOTE TO PARAGRAPH (a)(2): A veteran's enrollment status will be recognized throughout the United States.