WAC 246-335-635 Bill of rights. A hospice agency at the time of admission must provide each patient, designated family member, or legal representative with a written bill of rights affirming each patient's right to:

(1) Receive effective pain management and symptom control and quality services from the hospice agency for services identified in the plan of care;

(2) Be cared for by appropriately trained or credentialed personnel, contractors and volunteers with coordination of services;

(3) A statement advising of the right to ongoing participation in the development of the plan of care;

(4) Choose his or her attending physician;

(5) A statement advising of the right to have access to the department's listing of licensed hospice agencies and to select any licensee to provide care, subject to the individual's reimbursement mechanism or other relevant contractual obligations;

(6) A listing of the total services offered by the hospice agency and those being provided to the patient;

(7) Refuse specific services;

(8) The name of the individual within the hospice agency responsible for supervising the patient's care and the manner in which that individual may be contacted;

(9) Be treated with courtesy, respect, and privacy;

(10) Be free from verbal, mental, sexual, and physical abuse, neglect, exploitation, discrimination, and the unlawful use of restraint or seclusion;

(11) Have property treated with respect;

(12) Privacy and confidentiality of personal information and health care related records;

(13) Be informed of what the hospice agency charges for services, to what extent payment may be expected from health insurance, public programs, or other sources, and what charges the patient may be responsible for paying;

(14) A fully itemized billing statement upon request, including the date of each service and the charge. Agencies providing services through a managed care plan are not required to provide itemized billing statements;

(15) Be informed about advanced directives and POLST and the agency's scope of responsibility;

(16) Be informed of the agency's policies and procedures regarding the circumstances that may cause the agency to discharge a patient;

(17) Be informed of the agency's policies and procedures for providing back-up care when services cannot be provided as scheduled;

(18) A description of the agency's process for patients and family to submit complaints to the hospice agency about the services and care they are receiving and to have those complaints addressed without retaliation;

(19) Be informed of the department's complaint hotline number to report complaints about the licensed agency or credentialed health care professionals; and

(20) Be informed of the DSHS end harm hotline number to report suspected abuse of children or vulnerable adults.

(21) The hospice agency must ensure that the patient rights under this section are implemented and updated as appropriate.

[Statutory Authority: RCW 70.127.120 and 43.70.250. WSR 18-06-093, § 246-335-635, filed 3/6/18, effective 4/6/18.]