ESB 5666 - H COMM AMD
By Committee on Judiciary

ADOPTED 04/16/2013

Strike everything after the enacting clause and insert the following:

"**Sec. 1.** RCW 7.71.030 and 2012 c 165 s 1 are each amended to read as follows:

(1) If the limitation on damages under RCW 7.71.020 and P.L. 99-660 Sec. 411(1) does not apply, this section shall provide the exclusive ((~~remedy~~)) remedies in any lawsuit by a health care provider for any action taken by a professional peer review body of health care providers as defined in RCW 7.70.020((~~, that is found to be based on matters not related to the competence or professional conduct of a health care provider~~)).

(2) ((~~Actions~~)) Remedies shall be limited to appropriate injunctive relief, and damages shall be allowed only for lost earnings directly attributable to the action taken by the professional peer review body, incurred between the date of such action and the date the action is functionally reversed by the professional peer review body.

(3) Reasonable attorneys' fees and costs shall be awarded if approved by the court under RCW 7.71.035.

(4) The statute of limitations for actions under this section shall be one year from the date of the action of the professional peer review body.

Sec. 2. RCW 70.41.200 and 2007 c 273 s 22 and 2007 c 261 s 3 are each reenacted and amended to read as follows:

(1) Every hospital shall maintain a coordinated quality improvement program for the improvement of the quality of health care services rendered to patients and the identification and prevention of medical malpractice. The program shall include at least the following:

(a) The establishment of ((~~a~~)) one or more quality improvement committees with the responsibility to review the services rendered in

the hospital, both retrospectively and prospectively, in order to improve the quality of medical care of patients and to prevent medical malpractice. ((~~The~~)) Different quality improvement committees may be established as a part of a quality improvement program to review different health care services. Such committees shall oversee and coordinate the quality improvement and medical malpractice prevention program and shall ensure that information gathered pursuant to the program is used to review and to revise hospital policies and procedures;

(b) A process, including a medical staff privileges sanction procedure which must be conducted substantially in accordance with medical staff bylaws and applicable rules, regulations, or policies of the medical staff through which credentials, physical and mental capacity, professional conduct, and competence in delivering health care services are periodically reviewed as part of an evaluation of staff privileges;

(c) ((~~The~~)) A process for the periodic review of the credentials, physical and mental capacity, professional conduct, and competence in delivering health care services of all ((~~persons~~)) other health care providers who are employed or associated with the hospital;

(d) A procedure for the prompt resolution of grievances by patients or their representatives related to accidents, injuries, treatment, and other events that may result in claims of medical malpractice;

(e) The maintenance and continuous collection of information concerning the hospital's experience with negative health care outcomes and incidents injurious to patients including health care-associated infections as defined in RCW 43.70.056, patient grievances, professional liability premiums, settlements, awards, costs incurred by the hospital for patient injury prevention, and safety improvement activities;

(f) The maintenance of relevant and appropriate information gathered pursuant to (a) through (e) of this subsection concerning individual physicians within the physician's personnel or credential file maintained by the hospital;

(g) Education programs dealing with quality improvement, patient safety, medication errors, injury prevention, infection control, staff responsibility to report professional misconduct, the legal aspects of

patient care, improved communication with patients, and causes of malpractice claims for staff personnel engaged in patient care activities; and

(h) Policies to ensure compliance with the reporting requirements of this section.

(2) Any person who, in substantial good faith, provides information to further the purposes of the quality improvement and medical malpractice prevention program or who, in substantial good faith, participates on the quality improvement committee shall not be subject to an action for civil damages or other relief as a result of such activity. Any person or entity participating in a coordinated quality improvement program that, in substantial good faith, shares information or documents with one or more other programs, committees, or boards under subsection (8) of this section is not subject to an action for civil damages or other relief as a result of the activity. For the purposes of this section, sharing information is presumed to be in substantial good faith. However, the presumption may be rebutted upon a showing of clear, cogent, and convincing evidence that the information shared was knowingly false or deliberately misleading.

(3) Information and documents, including complaints and incident reports, created specifically for, and collected and maintained by, a quality improvement committee are not subject to review or disclosure, except as provided in this section, or discovery or introduction into evidence in any civil action, and no person who was in attendance at a meeting of such committee or who participated in the creation, collection, or maintenance of information or documents specifically for the committee shall be permitted or required to testify in any civil action as to the content of such proceedings or the documents and information prepared specifically for the committee. This subsection does not preclude: (a) In any civil action, the discovery of the identity of persons involved in the medical care that is the basis of the civil action whose involvement was independent of any quality improvement activity; (b) in any civil action, the testimony of any person concerning the facts which form the basis for the institution of such proceedings of which the person had personal knowledge acquired independently of such proceedings; (c) in any civil action by a health care provider regarding the restriction or revocation of that individual's clinical or staff privileges, introduction into evidence

information collected and maintained by quality improvement committees regarding such health care provider; (d) in any civil action, disclosure of the fact that staff privileges were terminated or restricted, including the specific restrictions imposed, if any and the reasons for the restrictions; or (e) in any civil action, discovery and introduction into evidence of the patient's medical records required by regulation of the department of health to be made regarding the care and treatment received.

(4) Each quality improvement committee shall, on at least a semiannual basis, report to the governing board of the hospital in which the committee is located. The report shall review the quality improvement activities conducted by the committee, and any actions taken as a result of those activities.

(5) The department of health shall adopt such rules as are deemed appropriate to effectuate the purposes of this section.

(6) The medical quality assurance commission or the board of osteopathic medicine and surgery, as appropriate, may review and audit the records of committee decisions in which a physician's privileges are terminated or restricted. Each hospital shall produce and make accessible to the commission or board the appropriate records and otherwise facilitate the review and audit. Information so gained shall not be subject to the discovery process and confidentiality shall be respected as required by subsection (3) of this section. Failure of a hospital to comply with this subsection is punishable by a civil penalty not to exceed two hundred fifty dollars.

(7) The department, the joint commission on accreditation of health care organizations, and any other accrediting organization may review and audit the records of a quality improvement committee or peer review committee in connection with their inspection and review of hospitals. Information so obtained shall not be subject to the discovery process, and confidentiality shall be respected as required by subsection (3) of this section. Each hospital shall produce and make accessible to the department the appropriate records and otherwise facilitate the review and audit.

(8) A coordinated quality improvement program may share information and documents, including complaints and incident reports, created specifically for, and collected and maintained by, a quality improvement committee or a peer review committee under RCW 4.24.250

with one or more other coordinated quality improvement programs maintained in accordance with this section or RCW 43.70.510, a coordinated quality improvement committee maintained by an ambulatory surgical facility under RCW 70.230.070, a quality assurance committee maintained in accordance with RCW 18.20.390 or 74.42.640, or a peer review committee under RCW 4.24.250, for the improvement of the quality of health care services rendered to patients and the identification and prevention of medical malpractice. The privacy protections of chapter 70.02 RCW and the federal health insurance portability and accountability act of 1996 and its implementing regulations apply to the sharing of individually identifiable patient information held by a coordinated quality improvement program. Any rules necessary to implement this section shall meet the requirements of applicable federal and state privacy laws. Information and documents disclosed by one coordinated quality improvement program to another coordinated quality improvement program or a peer review committee under RCW 4.24.250 and any information and documents created or maintained as a result of the sharing of information and documents shall not be subject to the discovery process and confidentiality shall be respected as required by subsection (3) of this section, RCW 18.20.390 (6) and (8), 74.42.640 (7) and (9), and 4.24.250.

(9) A hospital that operates a nursing home as defined in RCW 18.51.010 may conduct quality improvement activities for both the hospital and the nursing home through a quality improvement committee under this section, and such activities shall be subject to the provisions of subsections (2) through (8) of this section.

(10) Violation of this section shall not be considered negligence per se.

Sec. 3. RCW 70.41.230 and 1994 sp.s. c 9 s 744 are each amended to read as follows:

(1) Prior to granting or renewing clinical privileges or association of any physician or hiring a physician, a hospital or facility approved pursuant to this chapter shall request from the physician and the physician shall provide the following information:

(a) The name of any hospital or facility with or at which the physician had or has any association, employment, privileges, or practice during the prior five years: PROVIDED, That the hospital may

request additional information going back further than five years, and the physician shall use his or her best efforts to comply with such a request for additional information;

(b) ((~~If such association, employment, privilege, or practice was discontinued, the reasons for its discontinuation~~)) Whether the physician has ever been or is in the process of being denied, revoked, terminated, suspended, restricted, reduced, limited, sanctioned, placed on probation, monitored, or not renewed for any professional activity listed in (b)(i) through (x) of this subsection, or has ever voluntarily or involuntarily relinquished, withdrawn, or failed to proceed with an application for any professional activity listed in (b)(i) through (x) of this subsection in order to avoid an adverse action or to preclude an investigation or while under investigation relating to professional competence or conduct:

(i) License to practice any profession in any jurisdiction;

(ii) Other professional registration or certification in any jurisdiction;

(iii) Specialty or subspecialty board certification;

(iv) Membership on any hospital medical staff;

(v) Clinical privileges at any facility, including hospitals, ambulatory surgical centers, or skilled nursing facilities;

(vi) Medicare, medicaid, the food and drug administration, the national institute of health (office of human research protection), governmental, national, or international regulatory agency, or any public program;

(vii) Professional society membership or fellowship;

(viii) Participation or membership in a health maintenance organization, preferred provider organization, independent practice association, physician-hospital organization, or other entity;

(ix) Academic appointment;

(x) Authority to prescribe controlled substances (drug enforcement agency or other authority);

(c) Any pending professional medical misconduct proceedings or any pending medical malpractice actions in this state or another state, the substance of the allegations in the proceedings or actions, and any additional information concerning the proceedings or actions as the physician deems appropriate;

(d) The substance of the findings in the actions or proceedings and any additional information concerning the actions or proceedings as the physician deems appropriate;

(e) A waiver by the physician of any confidentiality provisions concerning the information required to be provided to hospitals pursuant to this subsection; and

(f) A verification by the physician that the information provided by the physician is accurate and complete.

(2) Prior to granting privileges or association to any physician or hiring a physician, a hospital or facility approved pursuant to this chapter shall request from any hospital with or at which the physician had or has privileges, was associated, or was employed, during the preceding five years, the following information concerning the physician:

(a) Any pending professional medical misconduct proceedings or any pending medical malpractice actions, in this state or another state;

(b) Any judgment or settlement of a medical malpractice action and any finding of professional misconduct in this state or another state by a licensing or disciplinary board; and

(c) Any information required to be reported by hospitals pursuant to RCW 18.71.0195.

(3) The medical quality assurance commission shall be advised within thirty days of the name of any physician denied staff privileges, association, or employment on the basis of adverse findings under subsection (1) of this section.

(4) A hospital or facility that receives a request for information from another hospital or facility pursuant to subsections (1) and (2) of this section shall provide such information concerning the physician in question to the extent such information is known to the hospital or facility receiving such a request, including the reasons for suspension, termination, or curtailment of employment or privileges at the hospital or facility. A hospital, facility, or other person providing such information in good faith is not liable in any civil action for the release of such information.

(5) Information and documents, including complaints and incident reports, created specifically for, and collected, and maintained by a quality improvement committee are not subject to discovery or introduction into evidence in any civil action, and no person who was

in attendance at a meeting of such committee or who participated in the creation, collection, or maintenance of information or documents specifically for the committee shall be permitted or required to testify in any civil action as to the content of such proceedings or the documents and information prepared specifically for the committee. This subsection does not preclude: (a) In any civil action, the discovery of the identity of persons involved in the medical care that is the basis of the civil action whose involvement was independent of any quality improvement activity; (b) in any civil action, the testimony of any person concerning the facts which form the basis for the institution of such proceedings of which the person had personal knowledge acquired independently of such proceedings; (c) in any civil action by a health care provider regarding the restriction or revocation of that individual's clinical or staff privileges, introduction into evidence information collected and maintained by quality improvement committees regarding such health care provider; (d) in any civil action, disclosure of the fact that staff privileges were terminated or restricted, including the specific restrictions imposed, if any and the reasons for the restrictions; or (e) in any civil action, discovery and introduction into evidence of the patient's medical records required by regulation of the department of health to be made regarding the care and treatment received.

(6) Hospitals shall be granted access to information held by the medical quality assurance commission and the board of osteopathic medicine and surgery pertinent to decisions of the hospital regarding credentialing and recredentialing of practitioners.

(7) Violation of this section shall not be considered negligence per se.

Sec. 4. RCW 70.230.080 and 2007 c 273 s 9 are each amended to read as follows:

(1) Every ambulatory surgical facility shall maintain a coordinated quality improvement program for the improvement of the quality of health care services rendered to patients and the identification and prevention of medical malpractice. The program shall include at least the following:

(a) The establishment of ((~~a~~)) one or more quality improvement committees with the responsibility to review the services rendered in

the ambulatory surgical facility, both retrospectively and prospectively, in order to improve the quality of medical care of patients and to prevent medical malpractice. ((~~The~~)) Different quality improvement committees may be established as a part of the quality improvement program to review different health care services. Such committees shall oversee and coordinate the quality improvement and medical malpractice prevention program and shall ensure that information gathered pursuant to the program is used to review and to revise the policies and procedures of the ambulatory surgical facility;

(b) A process, including a medical staff privileges sanction procedure which must be conducted substantially in accordance with medical staff bylaws and applicable rules, regulations, or policies of the medical staff through which credentials, physical and mental capacity, professional conduct, and competence in delivering health care services are periodically reviewed as part of an evaluation of staff privileges;

(c) The periodic review of the credentials, physical and mental capacity, and competence in delivering health care services of all persons who are employed or associated with the ambulatory surgical facility;

(d) A procedure for the prompt resolution of grievances by patients or their representatives related to accidents, injuries, treatment, and other events that may result in claims of medical malpractice;

(e) The maintenance and continuous collection of information concerning the ambulatory surgical facility's experience with negative health care outcomes and incidents injurious to patients, patient grievances, professional liability premiums, settlements, awards, costs incurred by the ambulatory surgical facility for patient injury prevention, and safety improvement activities;

(f) The maintenance of relevant and appropriate information gathered pursuant to (a) through (e) of this subsection concerning individual practitioners within the practitioner's personnel or credential file maintained by the ambulatory surgical facility;

(g) Education programs dealing with quality improvement, patient safety, medication errors, injury prevention, staff responsibility to report professional misconduct, the legal aspects of patient care, improved communication with patients, and causes of malpractice claims for staff personnel engaged in patient care activities; and

(h) Policies to ensure compliance with the reporting requirements of this section.

(2) Any person who, in substantial good faith, provides information to further the purposes of the quality improvement and medical malpractice prevention program or who, in substantial good faith, participates on the quality improvement committee is not subject to an action for civil damages or other relief as a result of such activity. Any person or entity participating in a coordinated quality improvement program that, in substantial good faith, shares information or documents with one or more other programs, committees, or boards under subsection (8) of this section is not subject to an action for civil damages or other relief as a result of the activity. For the purposes of this section, sharing information is presumed to be in substantial good faith. However, the presumption may be rebutted upon a showing of clear, cogent, and convincing evidence that the information shared was knowingly false or deliberately misleading.

(3) Information and documents, including complaints and incident reports, created specifically for, and collected and maintained by, a quality improvement committee are not subject to review or disclosure, except as provided in this section, or discovery or introduction into evidence in any civil action, and no person who was in attendance at a meeting of such committee or who participated in the creation, collection, or maintenance of information or documents specifically for the committee shall be permitted or required to testify in any civil action as to the content of such proceedings or the documents and information prepared specifically for the committee. This subsection does not preclude: (a) In any civil action, the discovery of the identity of persons involved in the medical care that is the basis of the civil action whose involvement was independent of any quality improvement activity; (b) in any civil action, the testimony of any person concerning the facts which form the basis for the institution of such proceedings of which the person had personal knowledge acquired independently of such proceedings; (c) in any civil action by a health care provider regarding the restriction or revocation of that individual's clinical or staff privileges, introduction into evidence of information collected and maintained by quality improvement committees regarding such health care provider; (d) in any civil action, disclosure of the fact that staff privileges were terminated or

restricted, including the specific restrictions imposed, if any, and the reasons for the restrictions; or (e) in any civil action, discovery and introduction into evidence of the patient's medical records required by rule of the department to be made regarding the care and treatment received.

(4) Each quality improvement committee shall, on at least a semiannual basis, report to the management of the ambulatory surgical facility, as identified in the facility's application, in which the committee is located. The report shall review the quality improvement activities conducted by the committee, and any actions taken as a result of those activities.

(5) The department shall adopt such rules as are deemed appropriate to effectuate the purposes of this section.

(6) The medical quality assurance commission, the board of osteopathic medicine and surgery, or the podiatric medical board, as appropriate, may review and audit the records of committee decisions in which a practitioner's privileges are terminated or restricted. Each ambulatory surgical facility shall produce and make accessible to the commission or board the appropriate records and otherwise facilitate the review and audit. Information so gained is not subject to the discovery process and confidentiality shall be respected as required by subsection (3) of this section. Failure of an ambulatory surgical facility to comply with this subsection is punishable by a civil penalty not to exceed two hundred fifty dollars.

(7) The department and any accrediting organization may review and audit the records of a quality improvement committee or peer review committee in connection with their inspection and review of the ambulatory surgical facility. Information so obtained is not subject to the discovery process, and confidentiality shall be respected as required by subsection (3) of this section. Each ambulatory surgical facility shall produce and make accessible to the department the appropriate records and otherwise facilitate the review and audit.

(8) A coordinated quality improvement program may share information and documents, including complaints and incident reports, created specifically for, and collected and maintained by, a quality improvement committee or a peer review committee under RCW 4.24.250 with one or more other coordinated quality improvement programs maintained in accordance with this section or RCW 43.70.510 or

70.41.200, a quality assurance committee maintained in accordance with RCW 18.20.390 or 74.42.640, or a peer review committee under RCW 4.24.250, for the improvement of the quality of health care services rendered to patients and the identification and prevention of medical malpractice. The privacy protections of chapter 70.02 RCW and the federal health insurance portability and accountability act of 1996 and its implementing regulations apply to the sharing of individually identifiable patient information held by a coordinated quality improvement program. Any rules necessary to implement this section shall meet the requirements of applicable federal and state privacy laws. Information and documents disclosed by one coordinated quality improvement program to another coordinated quality improvement program or a peer review committee under RCW 4.24.250 and any information and documents created or maintained as a result of the sharing of information and documents are not subject to the discovery process and confidentiality shall be respected as required by subsection (3) of this section, RCW 18.20.390 (6) and (8), 70.41.200(3), 74.42.640 (7) and (9), and 4.24.250.

(9) An ambulatory surgical facility that participates in a coordinated quality improvement program under RCW 43.70.510 shall be deemed to have met the requirements of this section.

(10) Violation of this section shall not be considered negligence per se.

Sec. 5. RCW 70.230.140 and 2007 c 273 s 15 are each amended to read as follows:

(1) Prior to granting or renewing clinical privileges or association of any practitioner or hiring a practitioner, an ambulatory surgical facility approved pursuant to this chapter shall request from the practitioner and the practitioner shall provide the following information:

(a) The name of any hospital, ambulatory surgical facility, or other facility with or at which the practitioner had or has any association, employment, privileges, or practice <u>during the prior five years: PROVIDED, That the ambulatory surgical facility may request additional information going back further than five years, and the physician shall use his or her best efforts to comply with such a request for additional information</u>;

(b) ((~~If such association, employment, privilege, or practice was discontinued, the reasons for its discontinuation~~)) Whether the physician has ever been or is in the process of being denied, revoked, terminated, suspended, restricted, reduced, limited, sanctioned, placed on probation, monitored, or not renewed for any professional activity listed in (b)(i) through (x) of this subsection, or has ever voluntarily or involuntarily relinquished, withdrawn, or failed to proceed with an application for any professional activity listed in (b)(i) through (x) of this subsection in order to avoid an adverse action or to preclude an investigation or while under investigation relating to professional competence or conduct:

(i) License to practice any profession in any jurisdiction;

(ii) Other professional registration or certification in any jurisdiction;

(iii) Specialty or subspecialty board certification;

(iv) Membership on any hospital medical staff;

(v) Clinical privileges at any facility, including hospitals, ambulatory surgical centers, or skilled nursing facilities;

(vi) Medicare, medicaid, the food and drug administration, the national institute of health (office of human research protection), governmental, national, or international regulatory agency, or any public program;

(vii) Professional society membership or fellowship;

(viii) Participation or membership in a health maintenance organization, preferred provider organization, independent practice association, physician-hospital organization, or other entity;

(ix) Academic appointment;

(x) Authority to prescribe controlled substances (drug enforcement agency or other authority);

(c) Any pending professional medical misconduct proceedings or any pending medical malpractice actions in this state or another state, the substance of the allegations in the proceedings or actions, and any additional information concerning the proceedings or actions as the practitioner deems appropriate;

(d) The substance of the findings in the actions or proceedings and any additional information concerning the actions or proceedings as the practitioner deems appropriate;

(e) A waiver by the practitioner of any confidentiality provisions concerning the information required to be provided to ambulatory surgical facilities pursuant to this subsection; and

(f) A verification by the practitioner that the information provided by the practitioner is accurate and complete.

(2) Prior to granting privileges or association to any practitioner or hiring a practitioner, an ambulatory surgical facility approved under this chapter shall request from any hospital or ambulatory surgical facility with or at which the practitioner had or has privileges, was associated, or was employed, during the preceding five years, the following information concerning the practitioner:

(a) Any pending professional medical misconduct proceedings or any pending medical malpractice actions, in this state or another state;

(b) Any judgment or settlement of a medical malpractice action and any finding of professional misconduct in this state or another state by a licensing or disciplinary board; and

(c) Any information required to be reported by hospitals or ambulatory surgical facilities pursuant to RCW 18.130.070.

(3) The medical quality assurance commission, board of osteopathic medicine and surgery, podiatric medical board, or dental quality assurance commission, as appropriate, shall be advised within thirty days of the name of any practitioner denied staff privileges, association, or employment on the basis of adverse findings under subsection (1) of this section.

(4) A hospital, ambulatory surgical facility, or other facility that receives a request for information from another hospital, ambulatory surgical facility, or other facility pursuant to subsections (1) and (2) of this section shall provide such information concerning the physician in question to the extent such information is known to the hospital, ambulatory surgical facility, or other facility receiving such a request, including the reasons for suspension, termination, or curtailment of employment or privileges at the hospital, ambulatory surgical facility, or facility. A hospital, ambulatory surgical facility, other facility, or other person providing such information in good faith is not liable in any civil action for the release of such information.

(5) Information and documents, including complaints and incident reports, created specifically for, and collected and maintained by, a

quality improvement committee are not subject to discovery or introduction into evidence in any civil action, and no person who was in attendance at a meeting of such committee or who participated in the creation, collection, or maintenance of information or documents specifically for the committee shall be permitted or required to testify in any civil action as to the content of such proceedings or the documents and information prepared specifically for the committee. This subsection does not preclude: (a) In any civil action, the discovery of the identity of persons involved in the medical care that is the basis of the civil action whose involvement was independent of any quality improvement activity; (b) in any civil action, the testimony of any person concerning the facts which form the basis for the institution of such proceedings of which the person had personal knowledge acquired independently of such proceedings; (c) in any civil action by a health care provider regarding the restriction or revocation of that individual's clinical or staff privileges, introduction into evidence information collected and maintained by quality improvement committees regarding such health care provider; (d) in any civil action, disclosure of the fact that staff privileges were terminated or restricted, including the specific restrictions imposed, if any, and the reasons for the restrictions; or (e) in any civil action, discovery and introduction into evidence of the patient's medical records required by rule of the department to be made regarding the care and treatment received.

(6) Ambulatory surgical facilities shall be granted access to information held by the medical quality assurance commission, board of osteopathic medicine and surgery, or podiatric medical board pertinent to decisions of the ambulatory surgical facility regarding credentialing and recredentialing of practitioners.

(7) Violation of this section shall not be considered negligence per se."

Correct the title.

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