# 2016 -- S 3013 SUBSTITUTE A

======= LC005936/SUB A =======

# STATE OF RHODE ISLAND

# IN GENERAL ASSEMBLY

## JANUARY SESSION, A.D. 2016

# $A\ N\quad A\ C\ T$

## RELATING TO HEALTH AND SAFETY - UTILIZATION REVIEW

Introduced By: Senator Roger Picard

Date Introduced: May 19, 2016

Referred To: Senate Health & Human Services

It is enacted by the General Assembly as follows:

1	SECTION 1. Chapter 23-17.12 of the General Laws entitled "Health Care Services -
2	Utilization Review Act" is hereby repealed in its entirety.
3	CHAPTER 23-17.12
4	Health Care Services – Utilization Review Act
5	23-17.12-1. Purpose of chapter The purpose of the chapter is to:
6	(1) Promote the delivery of quality health care in a cost effective manner;
7	(2) Foster greater coordination between health care providers, patients, payors and
8	utilization review entities;
9	(3) Protect patients, businesses, and providers by ensuring that review agents are
10	qualified to perform utilization review activities and to make informed decisions on the
11	appropriateness of medical care; and
12	(4) Ensure that review agents maintain the confidentiality of medical records in
13	accordance with applicable state and federal laws.
14	23-17.12-2. Definitions As used in this chapter, the following terms are defined as
15	<del>follows:</del>
16	(1) "Adverse determination" means a utilization review decision by a review agent not to
17	authorize a health care service. A decision by a review agent to authorize a health care service in
18	an alternative setting, a modified extension of stay, or an alternative treatment shall not constitute
19	an adverse determination if the review agent and provider are in agreement regarding the

1 decision. Adverse determinations include decisions not to authorize formulary and nonformulary 2 medication. (2) "Appeal" means a subsequent review of an adverse determination upon request by a 3 4 patient or provider to reconsider all or part of the original decision. 5 (3) "Authorization" means the review agent's utilization review, performed according to subsection 23-17.12-2(20), concluded that the allocation of health care services of a provider. 6 given or proposed to be given to a patient was approved or authorized. 7 8 (4) "Benefit determination" means a decision of the enrollee's entitlement to payment for 9 covered health care services as defined in an agreement with the payor or its delegate. 10 (5) "Certificate" means a certificate of registration granted by the director to a review 11 <del>agent.</del> 12 (6) "Complaint" means a written expression of dissatisfaction by a patient, or provider. The appeal of an adverse determination is not considered a complaint. 13 14 (7) "Concurrent assessment" means an assessment of the medical necessity and/or appropriateness of health care services conducted during a patient's hospital stay or course of 15 16 treatment. If the medical problem is ongoing, this assessment may include the review of services 17 after they have been rendered and billed. This review does not mean the elective requests for 18 clarification of coverage or claims review or a provider's internal quality assurance program 19 except if it is associated with a health care financing mechanism. 20 (8) "Department" means the department of health. 21 (9) "Director" means the director of the department of health. 22 (10) "Emergent health care services" has the same meaning as that meaning contained in the rules and regulations promulgated pursuant to chapter 12.3 of title 42 as may be amended 23 from time to time and includes those resources provided in the event of the sudden onset of a 24 25 medical, mental health, or substance abuse or other health care condition manifesting itself by 26 acute symptoms of a severity (e.g. severe pain) where the absence of immediate medical attention 27 could reasonably be expected to result in placing the patient's health in serious jeopardy, serious 28 impairment to bodily or mental functions, or serious dysfunction of any body organ or part. 29 (11) "Patient" means an enrollee or participant in all hospital or medical plans seeking 30 health care services and treatment from a provider. 31 (12) "Payor" means a health insurer, self-insured plan, nonprofit health service plan, 32 health insurance service organization, preferred provider organization, health maintenance 33 organization or other entity authorized to offer health insurance policies or contracts or pay for 34 the delivery of health care services or treatment in this state.

1 (13) "Practitioner" means any person licensed to provide or otherwise lawfully providing 2 health care services, including, but not limited to, a physician, dentist, nurse, optometrist, 3 podiatrist, physical therapist, clinical social worker, or psychologist. 4 (14) "Prospective assessment" means an assessment of the medical necessity and/or 5 appropriateness of health care services prior to services being rendered. (15) "Provider" means any health care facility, as defined in § 23-17-2 including any 6 mental health and/or substance abuse treatment facility, physician, or other licensed practitioners 7 8 identified to the review agent as having primary responsibility for the care, treatment, and 9 services rendered to a patient. 10 (16) "Retrospective assessment" means an assessment of the medical necessity and/or 11 appropriateness of health care services that have been rendered. This shall not include reviews 12 conducted when the review agency has been obtaining ongoing information. 13 (17) "Review agent" means a person or entity or insurer performing utilization review 14 that is either employed by, affiliated with, under contract with, or acting on behalf of: 15 (i) A business entity doing business in this state; 16 (ii) A party that provides or administers health care benefits to citizens of this state, 17 including a health insurer, self insured plan, non profit health service plan, health insurance 18 service organization, preferred provider organization or health maintenance organization 19 authorized to offer health insurance policies or contracts or pay for the delivery of health care 20 services or treatment in this state; or 21 (iii) A provider. 22 (18) "Same or similar specialty" means a practitioner who has the appropriate training 23 and experience that is the same or similar as the attending provider in addition to experience in 24 treating the same problems to include any potential complications as those under review. 25 (19) "Urgent health care services" has the same meaning as that meaning contained in the rules and regulations promulgated pursuant to chapter 12.3 of title 42 as may be amended 26 27 from time to time and includes those resources necessary to treat a symptomatic medical, mental 28 health, or substance abuse or other health care condition requiring treatment within a twenty four 29 (24) hour period of the onset of such a condition in order that the patient's health status not 30 decline as a consequence. This does not include those conditions considered to be emergent 31 health care services as defined in subdivision (10). 32 (20) "Utilization review" means the prospective, concurrent, or retrospective assessment 33 of the necessity and/or appropriateness of the allocation of health care services of a provider, 34 given or proposed to be given to a patient. Utilization review does not include:

1	-(i) Elective requests for the clarification of coverage; or
2	-(ii) Benefit determination; or
3	(iii) Claims review that does not include the assessment of the medical necessity and
4	appropriateness; or
5	(iv) A provider's internal quality assurance program except if it is associated with a
6	health care financing mechanism; or
7	(v) The therapeutic interchange of drugs or devices by a pharmacy operating as part of a
8	licensed inpatient health care facility; or
9	(vi) The assessment by a pharmacist licensed pursuant to the provisions of chapter 19 of
10	title 5 and practicing in a pharmacy operating as part of a licensed inpatient health care facility in
11	the interpretation, evaluation and implementation of medical orders, including assessments and/or
12	comparisons involving formularies and medical orders.
13	(21) "Utilization review plan" means a description of the standards governing utilization
14	review activities performed by a private review agent.
15	(22) "Health care services" means and includes an admission, diagnostic procedure,
16	therapeutic procedure, treatment, extension of stay, the ordering and/or filling of formulary or
17	nonformulary medications, and any other services, activities, or supplies that are covered by the
18	patient's benefit plan.
19	(23) "Therapeutic interchange" means the interchange or substitution of a drug with a
20	dissimilar chemical structure within the same therapeutic or pharmacological class that can be
21	expected to have similar outcomes and similar adverse reaction profiles when given in equivalent
22	doses, in accordance with protocols approved by the president of the medical staff or medical
23	director and the director of pharmacy.
24	23-17.12-3. General certificate requirements (a) A review agent shall not conduct
25	utilization review in the state unless the department has granted the review agent a certificate.
26	(b) Individuals shall not be required to hold separate certification under this chapter
27	when acting as either an employee of, an affiliate of, a contractor for, or otherwise acting on
28	behalf of a certified review agent.
29	(c) The department shall issue a certificate to an applicant that has met the minimum
30	standards established by this chapter, and regulations promulgated in accordance with it,
31	including the payment of any fees as required, and other applicable regulations of the department.
32	(d) A certificate issued under this chapter is not transferable, and the transfer of fifty
33	percent (50%) or more of the ownership of a review agent shall be deemed a transfer.

1 adopt regulations necessary to implement the provisions of this chapter.

1	adopt regulations necessary to implement the provisions of this chapter.
2	(f) The director of health is authorized to establish any fees for initial application,
3	renewal applications, and any other administrative actions deemed necessary by the director to
4	implement this chapter.
5	(g) The total cost of certification under this title shall be borne by the certified entities
6	and shall be one hundred and fifty percent (150%) of the total salaries paid to the certifying
7	personnel of the department engaged in those certifications less any salary reimbursements and
8	shall be paid to the director to and for the use of the department. That assessment shall be in
9	addition to any taxes and fees otherwise payable to the state.
10	(h) The application and other fees required under this chapter shall be sufficient to pay
11	for the administrative costs of the certificate program and any other reasonable costs associated
12	with carrying out the provisions of this chapter.
13	(i) A certificate expires on the second anniversary of its effective date unless the
14	certificate is renewed for a two (2) year term as provided in this chapter.
15	(j) Any systemic changes in the review agents operations relative to certification
16	information on file shall be submitted to the department for approval within thirty (30) days prior
17	to implementation.
18	23-17.12-4. Application process (a) An applicant requesting certification or
19	recertification shall:
20	(1) Submit an application provided by the director; and
21	(2) Pay the application fee established by the director through regulation and § 23-17.12-
22	<del>3(f).</del>
23	(b) The application shall:
24	(1) Be on a form and accompanied by supporting documentation that the director
25	requires; and
26	(2) Be signed and verified by the applicant.
27	(c) Before the certificate expires, a certificate may be renewed for an additional two (2)
28	<del>years.</del>
29	(d) If a completed application for recertification is being processed by the department, a
30	certificate may be continued until a renewal determination is made.
31	(e) In conjunction with the application, the review agent shall submit information that
32	the director requires including:
33	(1) A request that the state agency regard specific portions of the standards and criteria
34	or the entire document to constitute "trade secrets" within the meaning of that term in § 38-2-

1 <del>2(4)(i)(B);</del>

2	(2) The policies and procedures to ensure that all applicable state and federal laws to
3	protect the confidentiality of individual medical records are followed;
4	(3) A copy of the materials used to inform enrollees of the requirements under the health
5	benefit plan for seeking utilization review or pre-certification and their rights under this chapter,
6	including information on appealing adverse determinations;
7	(4) A copy of the materials designed to inform applicable patients and providers of the
8	requirements of the utilization review plan;
9	(5) A list of the third party payors and business entities for which the review agent is
10	performing utilization review in this state and a brief description of the services it is providing for
11	each client; and
12	(6) Evidence of liability insurance or of assets sufficient to cover potential liability.
13	(f) The information provided must demonstrate that the review agent will comply with
14	the regulations adopted by the director under this chapter.
15	23-17.12-5. General application requirements An application for certification or
16	recertification shall be accompanied by documentation to evidence the following:
17	(1) The requirement that the review agent provide patients and providers with a summary
18	of its utilization review plan including a summary of the standards, procedures and methods to be
19	used in evaluating proposed or delivered health care services;
20	(2) The circumstances, if any, under which utilization review may be delegated to any
21	other utilization review program and evidence that the delegated agency is a certified utilization
22	review agency delegated to perform utilization review pursuant to all of the requirements of this
23	<del>chapter;</del>
24	(3) A complaint resolution process consistent with subsection 23-17.12-2(6) and
25	acceptable to the department, whereby patients, their physicians, or other health care providers
26	may seek resolution of complaints and other matters of which the review agent has received
27	written notice;
28	(4) The type and qualifications of personnel (employed or under contract) authorized to
29	perform utilization review, including a requirement that only a practitioner with the same license
30	status as the ordering practitioner, or a licensed physician or dentist, is permitted to make a
31	prospective or concurrent adverse determination;
32	(5) The requirement that a representative of the review agent is reasonably accessible to
33	patients, patient's family and providers at least five (5) days a week during normal business in
34	Rhode Island and during the hours of the agency's review operations;

1 (6) The policies and procedures to ensure that all applicable state and federal laws to 2 protect the confidentiality of individual medical records are followed; 3 (7) The policies and procedures regarding the notification and conduct of patient 4 interviews by the review agent; 5 (8) The requirement that no employee of, or other individual rendering an adverse determination for, a review agent may receive any financial incentives based upon the number of 6 7 denials of certification made by that employee or individual; 8 (9) The requirement that the utilization review agent shall not impede the provision of 9 health care services for treatment and/or hospitalization or other use of a provider's services or 10 facilities for any patient; 11 (10) Evidence that the review agent has not entered into a compensation agreement or 12 contract with its employees or agents whereby the compensation of its employees or its agents is 13 based upon a reduction of services or the charges for those services, the reduction of length of 14 stay, or utilization of alternative treatment settings; provided, nothing in this chapter shall prohibit 15 agreements and similar arrangements; and 16 (11) An adverse determination and internal appeals process consistent with § 23-17.12-9 and acceptable to the department, whereby patients, their physicians, or other health care 17 18 providers may seek prompt reconsideration or appeal of adverse determinations by the review 19 agent. 20 23-17.12-6. Denial, suspension, or revocation of certificate. -- (a) The department may 21 deny a certificate upon review of the application if, upon review of the application, it finds that the applicant proposing to conduct utilization review does not meet the standards required by this 22 23 chapter or by any regulations promulgated pursuant to this chapter. 24 (b) The department may revoke a certificate and/or impose reasonable monetary penalties not to exceed five thousand dollars (\$5,000) per violation in any case in which: 25 26 (1) The review agent fails to comply substantially with the requirements of this chapter 27 or of regulations adopted pursuant to this chapter; 28 (2) The review agent fails to comply with the criteria used by it in its application for a 29 certificate; or 30 (3) The review agent refuses to permit examination by the director to determine 31 compliance with the requirements of this chapter and regulations promulgated pursuant to the 32 authority granted to the director in this chapter; provided, however, that the examination shall be 33 subject to the confidentiality and "need to know" provisions of subdivisions 23-17.12-9(c)(4) and 34 (5). These determinations may involve consideration of any written grievances filed with the

1 department against the review agent by patients or providers.

2 (c) Any applicant or certificate holder aggrieved by an order or a decision of the 3 department made under this chapter without a hearing may, within thirty (30) days after notice of 4 the order or decision, make a written request to the department for a hearing on the order or 5 decision pursuant to § 42-35-15. (d) The procedure governing hearings authorized by this section shall be in accordance 6 with §§ 42-35-9 -- 42-35-13 as stipulated in § 42-35-14(a). A full and complete record shall be 7 8 kept of all proceedings, and all testimony shall be recorded but need not be transcribed unless the 9 decision is appealed pursuant to § 42-35-15. A copy or copies of the transcript may be obtained 10 by any interested party upon payment of the cost of preparing the copy or copies. Witnesses may 11 be subpoenaed by either party. 12 23-17.12-7. Judicial review. -- Any person who has exhausted all administrative 13 remedies available to him or her within the department, and who is aggrieved by a final decision 14 of the department under § 23-17.12 6, is entitled to judicial review pursuant to §§ 42-35-15 and 42-35-16. 15 16 23-17.12-8. Waiver of requirements. -- (a) Except for utilization review agencies 17 performing utilization review activities to determine the necessity and/or appropriateness of 18 substance abuse and mental health care, treatment or services, the department shall waive all the 19 requirements of this chapter, with the exception of those contained in §§ 23-17.12-9, (a)(1) (3), 20 (5), (6), (8), (b)(1)-(6), and (c)(2)-(6), 23-17.12-12, and 23-17.12-14, for a review agent that has 21 received, maintains and provides evidence to the department of accreditation from the utilization 22 review accreditation commission (URAC) or other organization approved by the director. The 23 waiver shall be applicable only to those services that are included under the accreditation by the utilization review accreditation commission or other approved organization. 24 25 (b) The department shall waive the requirements of this chapter only when a direct 26 conflict exists with those activities of a review agent that are conducted pursuant to contracts with 27 the state or the federal government or those activities under other state or federal jurisdictions. 28 (c) The limitation in subsection 23-17.12-8(b) notwithstanding, the department may 29 waive or exempt all or part of the requirements of this chapter by mutual written agreement with 30 a state department or agency when such waiver or exemption is determined to be necessary and 31 appropriate to the administration of a health care related program. The department shall 32 promulgate such regulations as deemed appropriate to implement this provision. 33 23-17.12-8.1. Variance of statutory requirements. [Repealed effective July 1, 2015.] -34 - (a) The department is authorized to issue a statutory variance from one or more of the specific

1	requirements of this chapter to a review agent where it determines that such variance is necessary
2	to permit the review agent to evaluate and address practitioner billing and practice patterns when
3	the review agent believes in good faith that such patterns evidence the existence of fraud or
4	abuse. Any variance issued by the department pursuant to this section shall be limited in
5	application to those services billed directly by the practitioner. Prior to issuing a statutory
6	variance the department shall provide notice and a public hearing to ensure necessary patient and
7	health care provider protections in the process. Statutory variances shall be issued for a period not
8	to exceed one year and may be subject to such terms and conditions deemed necessary by the
9	department.
10	(b) On or before January 15th of each year, the department shall issue a report to the
11	general assembly summarizing any review agent activity as a result of a waiver granted under the
12	provisions of this section.
13	23-17.12-9. Review agency requirement for adverse determination and internal
14	appeals (a) The adverse determination and appeals process of the review agent shall conform
15	to the following:
16	(1) Notification of a prospective adverse determination by the review agent shall be
17	mailed or otherwise communicated to the provider of record and to the patient or other
18	appropriate individual as follows:
19	(i) Within fifteen (15) business days of receipt of all the information necessary to
20	complete a review of non-urgent and/or non-emergent services;
21	-(ii) Within seventy two (72) hours of receipt of all the information necessary to complete
22	a review of urgent and/or emergent services; and
23	-(iii) Prior to the expected date of service.
24	(2) Notification of a concurrent adverse determination shall be mailed or otherwise
25	communicated to the patient and to the provider of record period as follows:
26	(i) To the provider(s) prior to the end of the current certified period; and
27	(ii) To the patient within one business day of making the adverse determination.
28	(3) Notification of a retrospective adverse determination shall be mailed or otherwise
29	communicated to the patient and to the provider of record within thirty (30) business days of
30	receipt of a request for payment with all supporting documentation for the covered benefit being
31	reviewed.
32	(4) A utilization review agency shall not retrospectively deny authorization for health
33	care services provided to a covered person when an authorization has been obtained for that
34	service from the review agent unless the approval was based upon inaccurate information

1 material to the review or the health care services were not provided consistent with the provider's 2 submitted plan of care and/or any restrictions included in the prior approval granted by the review 3 agent. 4 (5) Any notice of an adverse determination shall include: 5 (i) The principal reasons for the adverse determination, to include explicit documentation of the criteria not met and/or the clinical rationale utilized by the agency's clinical reviewer in 6 making the adverse determination. The criteria shall be in accordance with the agency criteria 7 8 noted in subsection 23-17.12-9(d) and shall be made available within the first level appeal 9 timeframe if requested unless otherwise provided as part of the adverse determination notification 10 process; 11 (ii) The procedures to initiate an appeal of the adverse determination, including the name 12 and telephone number of the person to contract with regard to an appeal; 13 (iii) The necessary contact information to complete the two way direct communication 14 defined in subdivision 23-17.12-9(a)(7); and (iv) The information noted in subdivision 23-27.12-9(a)(5)(i)(ii)(iii) for all verbal 15 16 notifications followed by written notification to the patient and provider(s). 17 (6) All initial retrospective adverse determinations of a health care service that had been 18 ordered by a physician, dentist or other practitioner shall be made, documented and signed 19 consistent with the regulatory requirements which shall be developed by the department with the 20 input of review agents, providers and other affected parties. 21 (7) A level one appeal decision of an adverse determination shall not be made until an 22 appropriately qualified and licensed review physician, dentist or other practitioner has spoken to, or otherwise provided for, an equivalent two-way direct communication with the patient's 23 attending physician, dentist, other practitioner, other designated or qualified professional or 24 25 provider responsible for treatment of the patient concerning the medical care, with the exception of the following: 26 27 (i) When the attending provider is not reasonably available; 28 (ii) When the attending provider chooses not to speak with agency staff; 29 (iii) When the attending provider has negotiated an agreement with the review agent for 30 alternative care; and/or 31 (iv) When the attending provider requests a peer to peer communication prior to the 32 adverse determination, the review agency shall then comply with subdivision 23 - 17.12 - 9(c)(1) in 33 responding to such a request. Such requests shall be on the case specific basis unless otherwise 34 arranged for in advance by the provider.

1 (8) All initial, prospective and concurrent adverse determinations of a health care service 2 that had been ordered by a physician, dentist or other practitioner shall be made, documented and signed by a licensed practitioner with the same licensure status as the ordering practitioner or a 3 4 licensed physician or dentist. This does not prohibit appropriately qualified review agency staff 5 from engaging in discussions with the attending provider, the attending provider's designee or appropriate health care facility and office personnel regarding alternative service and treatment 6 options. Such a discussion shall not constitute an adverse determination provided though that any 7 8 change to the provider's original order and/or any decision for an alternative level of care must be made and/or appropriately consented to by the attending provider or the provider's designee 9 10 responsible for treating the patient. 11 (9) The requirement that, upon written request made by or on behalf of a patient, any 12 adverse determination and/or appeal shall include the written evaluation and findings of the reviewing physician, dentist or other practitioner. The review agent is required to accept a verbal 13

request made by or on behalf of a patient for any information where a provider or patient can
 demonstrate that a timely response is urgent.

(b) The review agent shall conform to the following for the appeal of an adverse
 determination:

18 (1) The review agent shall maintain and make available a written description of the 19 appeal procedure by which either the patient or the provider of record may seek review of 20 determinations not to authorize a health care service. The process established by each review 21 agent may include a reasonable period within which an appeal must be filed to be considered and 22 that period shall not be less than sixty (60) days.

(2) The review agent shall notify, in writing, the patient and provider of record of its
 decision on the appeal as soon as practical, but in no case later than fifteen (15) or twenty-one
 (21) business days if verbal notice is given within fifteen (15) business days after receiving the
 required documentation on the appeal.

27 (3) The review agent shall also provide for an expedited appeals process for emergency
 28 or life threatening situations. Each review agent shall complete the adjudication of expedited
 29 appeals within two (2) business days of the date the appeal is filed and all information necessary

30 to complete the appeal is received by the review agent.

31 (4) All first level appeals of determinations not to authorize a health care service that had
32 been ordered by a physician, dentist, or other practitioner shall be made, documented, and signed
33 by a licensed practitioner with the same licensure status as the ordering practitioner or a licensed
34 physician or a licensed dentist.

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1	(5) All second level appeal decisions shall be made, signed, and documented by a
2	licensed practitioner in the same or a similar general specialty as typically manages the medical
3	condition, procedure, or treatment under discussion.
4	(6) The review agent shall maintain records of written appeals and their resolution, and
5	shall provide reports as requested by the department.
6	(c) The review agency must conform to the following requirements when making its
7	adverse determination and appeal decisions:
8	(1) The review agent must assure that the licensed practitioner or licensed physician is
9	reasonably available to review the case as required under subdivision 23-17.12-9(a)(7) and shall
10	conform to the following:
11	(i) Each agency peer reviewer shall have access to and review all necessary information
12	as requested by the agency and/or submitted by the provider(s) and/or patients;
13	(ii) Each agency shall provide accurate peer review contact information to the provider at
14	the time of service, if requested, and/or prior to such service, if requested. This contact
15	information must provide a mechanism for direct communication with the agency's peer
16	reviewer;
17	(iii) Agency peer reviewers shall respond to the provider's request for a two way direct
18	communication defined in subdivision 23-17.12-9(a)(7)(iv) as follows:
19	(A) For a prospective review of non-urgent and non-emergent health care services, a
20	response within one business day of the request for a peer discussion;
21	(B) For concurrent and prospective reviews of urgent and emergent health care services,
22	a response within a reasonable period of time of the request for a peer discussion; and
23	(C) For retrospective reviews, prior to the first level appeal decision.
24	(iv) The review agency will have met the requirements of a two-way direct
25	communication, when requested and/or as required prior to the first level of appeal, when it has
26	made two (2) reasonable attempts to contact the attending provider directly.
27	(v) Repeated violations of this section shall be deemed to be substantial violations
28	pursuant to § 23-17.12-14 and shall be cause for the imposition of penalties under that section.
29	(2) No reviewer at any level under this section shall be compensated or paid a bonus or
30	incentive based on making or upholding an adverse determination.
31	(3) No reviewer under this section who has been involved in prior reviews of the case
32	under appeal or who has participated in the direct care of the patient may participate as the sole
33	reviewer in reviewing a case under appeal; provided, however, that when new information has
34	been made available at the first level of appeal, then the review may be conducted by the same

1 reviewer who made the initial adverse determination.

2 (4) A review agent is only entitled to review information or data relevant to the utilization review process. A review agent may not disclose or publish individual medical records 3 4 or any confidential medical information obtained in the performance of utilization review 5 activities. A review agent shall be considered a third party health insurer for the purposes of § 5-37.3-6(b)(6) of this state and shall be required to maintain the security procedures mandated in § 6 5-37.3-4(c). 7 8 (5) Notwithstanding any other provision of law, the review agent, the department, and all 9 other parties privy to information which is the subject of this chapter shall comply with all state and federal confidentiality laws, including, but not limited to, chapter 37.3 of title 5 10 11 (Confidentiality of Health Care Communications and Information Act) and specifically § 5-37.3-12 4(c), which requires limitation on the distribution of information which is the subject of this 13 chapter on a "need to know" basis, and § 40.1-5-26. 14 (6) The department may, in response to a complaint that is provided in written form to 15 the review agent, review an appeal regarding any adverse determination, and may request 16 information of the review agent, provider or patient regarding the status, outcome or rationale 17 regarding the decision. 18 (d) The requirement that each review agent shall utilize and provide upon request, by 19 Rhode Island licensed hospitals and the Rhode Island Medical Society, in either electronic or 20 paper format, written medically acceptable screening criteria and review procedures which are 21 established and periodically evaluated and updated with appropriate consultation with Rhode 22 Island licensed physicians, hospitals, including practicing physicians, and other health care providers in the same specialty as would typically treat the services subject to the criteria as 23 follows: 24 25 (1) Utilization review agents shall consult with no fewer than five (5) Rhode Island 26 licensed physicians or other health care providers. Further, in instances where the screening 27 criteria and review procedures are applicable to inpatients and/or outpatients of hospitals, the 28 medical director of each licensed hospital in Rhode Island shall also be consulted. Utilization 29 review agents who utilize screening criteria and review procedures provided by another entity 30 may satisfy the requirements of this section if the utilization review agent demonstrates to the 31 satisfaction of the director that the entity furnishing the screening criteria and review procedures 32 has complied with the requirements of this section. 33 (2) Utilization review agents seeking initial certification shall conduct the consultation

34 for all screening and review criteria to be utilized. Utilization review agents who have been

certified for one year or longer shall be required to conduct the consultation on a periodic basis
 for the utilization review agent's highest volume services subject to utilization review during the
 prior year; services subject to the highest volume of adverse determinations during the prior year;
 and for any additional services identified by the director.

5 (3) Utilization review agents shall not include in the consultations as required under
6 paragraph (1) of this subdivision, any physicians or other health services providers who have
7 financial relationships with the utilization review agent other than financial relationships for
8 provisions of direct patient care to utilization review agent enrollees and reasonable compensation
9 for consultation as required by paragraph (1) of this subdivision.

(4) All documentation regarding required consultations, including comments and/or
 recommendations provided by the health care providers involved in the review of the screening
 criteria, as well as the utilization review agent's action plan or comments on any
 recommendations, shall be in writing and shall be furnished to the department on request. The
 documentation shall also be provided on request to any licensed health care provider at a nominal
 cost that is sufficient to cover the utilization review agent's reasonable costs of copying and
 mailing.

17 (5) Utilization review agents may utilize non-Rhode Island licensed physicians or other
18 health care providers to provide the consultation as required under paragraph (1) of this
19 subdivision, when the utilization review agent can demonstrate to the satisfaction of the director
20 that the related services are not currently provided in Rhode Island or that another substantial
21 reason requires such approach.

(6) Utilization review agents whose annualized data reported to the department
 demonstrate that the utilization review agent will review fewer than five hundred (500) such
 requests for authorization may request a variance from the requirements of this section.

25 23-17.12-10. External appeal requirements. -- (a) In cases where the second level of appeal to reverse an adverse determination is unsuccessful, the review agent shall provide for an 26 27 external appeal by an unrelated and objective appeal agency, selected by the director. The director 28 shall promulgate rules and regulations including, but not limited to, criteria for designation, 29 operation, policy, oversight, and termination of designation as an external appeal agency. The 30 external appeal agency shall not be required to be certified under this chapter for activities 31 conducted pursuant to its designation. 32 (b) The external appeal shall have the following characteristics:

33 (1) The external appeal review and decision shall be based on the medical necessity for
 34 the health care or service and the appropriateness of service delivery for which authorization has

1 been denied.

2 (2) Neutral physicians, dentists, or other practitioners in the same or similar general specialty as typically manages the health care service shall be utilized to make the external appeal 3 4 decisions. (3) Neutral physicians, dentists, or other practitioners shall be selected from lists: 5 (i) Mutually agreed upon by the provider associations, insurers, and the purchasers of 6 health services; and 7 8 (ii) Used during a twelve (12) month period as the source of names for neutral physician, 9 dentist, or other practitioner reviewers. 10 (4) The neutral physician, dentist, or other practitioner may confer either directly with 11 the review agent and provider, or with physicians or dentists appointed to represent them. 12 (5) Payment for the appeal fee charged by the neutral physician, dentist, or other 13 practitioner shall be shared equally between the two (2) parties to the appeal; provided, however, 14 that if the decision of the utilization review agent is overturned, the appealing party shall be reimbursed by the utilization review agent for their share of the appeal fee paid under this 15 16 subsection. 17 (6) The decision of the external appeal agency shall be binding; however, any person who is aggrieved by a final decision of the external appeal agency is entitled to judicial review in 18 19 a court of competent jurisdiction. 20 23-17.12-11. Repealed ... --23-17.12-12. Reporting requirements. -- (a) The department shall establish reporting 21 22 requirements to determine if the utilization review programs are in compliance with the 23 provisions of this chapter and applicable regulations. (b) By November 14, 2014, the department shall report to the general assembly 24 25 regarding hospital admission practices and procedures and the effects of such practices and 26 procedures on the care and wellbeing of patients who present behavioral healthcare conditions on an emergency basis. The report shall be developed with the cooperation of the department of 27 28 behavioral healthcare, developmental disabilities, and hospitals and of the department of children, youth, and families, and shall recommend changes to state law and regulation to address any 29 30 necessary and appropriate revisions to the department's regulations related to utilization review 31 based on the Federal Mental Health Parity and Addiction Equity Act of 2008 (MHPAEA) and the 32 Patient Protection and Affordable Care Act, Pub. L. 111-148, and the state's regulatory 33 interpretation of parity in insurance coverage of behavioral healthcare. These recommended or 34 adopted revisions to the department's regulations shall include, but not be limited to:

1	(1) Adverse determination and internal appeals, with particular regard to the time
2	necessary to complete a review of urgent and/or emergent services for patients with behavioral
3	health needs;
4	-(2) External appeal requirements;
5	(3) The process for investigating whether insurers and agents are complying with the
6	provisions of chapter 17.12 of title 23 in light of parity in insurance coverage for behavioral
7	healthcare, with particular regard to emergency admissions; and
8	(4) Enforcement of the provisions of chapter 17.12 of title 23 in light of insurance parity
9	for behavioral healthcare.
10	23-17.12-13. Lists The director shall periodically provide a list of private review
11	agents issued certificates and the renewal date for those certificates to all licensed health care
12	facilities and any other individual or organization requesting the list.
13	23-17.12-14. Penalties A person who substantially violates any provision of this
14	chapter or any regulation adopted under this chapter or who submits any false information in an
15	application required by this chapter is guilty of a misdemeanor and on conviction is subject to a
16	penalty not exceeding five thousand dollars (\$5,000).
17	23-17.12-15. Annual report The director shall issue an annual report to the governor
18	and the general assembly concerning the conduct of utilization review in the state. The report
19	shall include a description of utilization programs and the services they provide, an analysis of
20	complaints filed against private review agents by patients or providers and an evaluation of the
21	impact of utilization review programs on patient access to care.
22	23-17.12-16. Fees The proceeds of any fees, monetary penalties, and fines collected
23	pursuant to the provisions of this chapter shall be deposited as general revenues.
24	-
25	23-17.12-17. Severability If any provision of this chapter or the application of any
26	provision to any person or circumstance shall be held invalid, that invalidity shall not affect the
27	provisions or application of this chapter which can be given effect without the invalid provision
28	or application, and to this end the provisions of this chapter are declared to be severable.
29	SECTION 2. Chapter 23-17.13 of the General Laws entitled "Health Care Accessibility
30	and Quality Assurance Act" is hereby repealed in its entirety.
31	CHAPTER 23-17.13
32	Health Care Accessibility and Quality Assurance Act
33	23-17.13-1. Purpose The legislature declares that:
34	(1) It is in the best interest of the public that those individuals and care entities involved

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1	with the delivery of plan coverage in our state meet the standards of this chapter to insure
2	accessibility and quality for the state's patients;
3	(2) Nothing in the legislation is intended to prohibit a health care entity or contractor
4	from forming limited networks of providers; and
5	(3) It is a vital state function to establish these standards for the conduct of health plans
6	by a health care entity in Rhode Island.
7	23-17.13-2. Definitions As used in this chapter:
8	(1) "Adverse decision" means any decision by a review agent not to certify an admission,
9	service, procedure, or extension of stay. A decision by a reviewing agent to certify an admission,
10	service, or procedure in an alternative treatment setting, or to certify a modified extension of stay,
11	shall not constitute an adverse decision if the reviewing agent and the requesting provider are in
12	agreement regarding the decision.
13	-(2) "Contractor" means a person/entity that:
14	(i) Establishes, operates or maintains a network of participating providers;
15	(ii) Contracts with an insurance company, a hospital or medical or dental service plan, an
16	employer, whether under written or self insured, an employee organization, or any other entity
17	providing coverage for health care services to administer a plan; and/or
18	(iii) Conducts or arranges for utilization review activities pursuant to chapter 17.12 of
19	this title.
20	(3) "Direct service ratio" means the amount of premium dollars expended by the plan for
21	covered services provided to enrollees on a plan's fiscal year basis.
22	(4) "Director" means the director of the department of health.
23	(5) "Emergency services" has the same meaning as the meaning contained in the rules
24	and regulations promulgated pursuant to chapter 12.3 of title 42, as may be amended from time to
25	time, and includes the sudden onset of a medical or mental condition that the absence of
26	immediate medical attention could reasonably be expected to result in placing the patient's health
27	in serious jeopardy, serious impairment to bodily or mental functions, or serious dysfunction of
28	any bodily organ or part.
29	(6) "Health care entity" means a licensed insurance company, hospital, or dental or
30	medical service plan or health maintenance organization, or a contractor as described in
31	subdivision (2), that operates a health plan.
32	(7) "Health care services" includes, but is not limited to, medical, mental health,
33	substance abuse, and dental services.
24	

34 (8) "Health plan" means a plan operated by a health care entity as described in

1 subdivision (6) that provides for the delivery of care services to persons enrolled in the plan 2 through: (i) Arrangements with selected providers to furnish health care services; and/or 3 4 (ii) Financial incentives for persons enrolled in the plan to use the participating providers 5 and procedures provided for by the plan. 6 (9) "Provider" means a physician, hospital, pharmacy, laboratory, dentist, or other state licensed or other state recognized provider of health care services or supplies, and whose services 7 8 are recognized pursuant to 213(d) of the Internal Revenue Code, 26 U.S.C. § 213(d), that has 9 entered into an agreement with a health care entity as described in subdivision (6) or contractor as described in subdivision (2) to provide these services or supplies to a patient enrolled in a plan. 10 11 (10) "Provider incentive plan" means any compensation arrangement between a health 12 care entity or plan and a provider or provider group that may directly or indirectly have the effect 13 of reducing or limiting services provided with respect to an individual enrolled in a plan. 14 (11) "Qualified health plan" means a plan that the director of the department of health 15 certified, upon application by the program, as meeting the requirements of this chapter. 16 (12) "Qualified utilization review program" means utilization review program that meets 17 the requirements of chapter 17.12 of this title. 18 (13) "Most favored rate clause" means a provision in a provider contract whereby the 19 rates or fees to be paid by a health plan are fixed, established or adjusted to be equal to or lower 20 than the rates or fees paid to the provider by any other health plan or third party payor. 21 23-17.13-3. Certification of health plans. -- (a) Certification process. 22 (1) Certification. 23 (i) The director shall establish a process for certification of health plans meeting the 24 requirements of certification in subsection (b). 25 (ii) The director shall act upon the health plan's completed application for certification 26 within ninety (90) days of receipt of such application for certification. 27 (2) Review and recertification. To ensure compliance with subsection (b), the director 28 shall establish procedures for the periodic review and recertification of qualified health plans not 29 less than every five (5) years; provided, however, that the director may review the certification of 30 a qualified health plan at any time if there exists evidence that a qualified health plan may be in 31 violation of subsection (b). 32 (3) Cost of certification. The total cost of obtaining and maintaining certification under 33 this title and compliance with the requirements of the applicable rules and regulations are borne 34 by the entities so certified and shall be one hundred and fifty percent (150%) of the total salaries

1	paid to the certifying personnel of the department engaged in those certifications less any salary
2	reimbursements and shall be paid to the director to and for the use of the department. That
3	assessment shall be in addition to any taxes and fees otherwise payable to the state.
4	(4) Standard definitions To help ensure a patient's ability to make informed decisions
5	regarding their health care, the director shall promulgate regulation(s) to provide for standardized
6	definitions (unless defined in existing statute) of the following terms in this subdivision,
7	provided, however, that no definition shall be construed to require a health care entity to add any
8	benefit, to increase the scope of any benefit, or to increase any benefit under any contract:
9	-(i) Allowable charge;
10	-(ii) Capitation;
11	-(iii) Co-payments;
12	-(iv) Co-insurance;
13	-(v) Credentialing;
14	-(vi) Formulary;
15	-(vii) Grace period;
16	-(viii) Indemnity insurance;
17	-(ix) In-patient care;
18	-(x) Maximum lifetime cap;
19	-(xi) Medical necessity;
20	-(xii) Out-of-network;
21	-(xiii) Out-patient;
22	-(xiv) Pre-existing conditions;
23	-(xv) Point of service;
24	-(xvi) Risk sharing;
25	-(xvii) Second opinion;
26	(xviii) Provider network;
27	-(xix) Urgent care.
28	(b) Requirements for certification The director shall establish standards and procedures
29	for the certification of qualified health plans that conduct business in this state and who have
30	demonstrated the ability to ensure that health care services will be provided in a manner to assure
31	availability and accessibility, adequate personnel and facilities, and continuity of service, and has
32	demonstrated arrangements for ongoing quality assurance programs regarding care processes and
33	outcomes; other standards shall consist of, but are not limited to, the following:
34	(1) Prospective and current enrollees in health plans must be provided information as to

34 (1) Prospective and current enrollees in health plans must be provided information as to

1 the terms and conditions of the plan consistent with the rules and regulations promulgated under 2 chapter 12.3 of title 42 so that they can make informed decisions about accepting and utilizing the health care services of the health plan. This must be standardized so that customers can compare 3 4 the attributes of the plans, and all information required by this paragraph shall be updated at intervals determined by the director. Of those items required under this section, the director shall 5 also determine which items shall be routinely distributed to prospective and current enrollees as 6 listed in this subsection and which items may be made available upon request. The items to be 7 8 disclosed are:

9 (i) Coverage provisions, benefits, and any restriction or limitations on health care
 10 services, including but not limited to, any exclusions as follows: by category of service, and if
 11 applicable, by specific service, by technology, procedure, medication, provider or treatment
 12 modality, diagnosis and condition, the latter three (3) of which shall be listed by name.

13 (ii) Experimental treatment modalities that are subject to change with the advent of new 14 technology may be listed solely by the broad category "Experimental Treatments". The 15 information provided to consumers shall include the plan's telephone number and address where 16 enrollees may call or write for more information or to register a complaint regarding the plan or 17 coverage provision.

18 (2) Written statement of the enrollee's right to seek a second opinion, and reimbursement
 19 if applicable.

20 (3) Written disclosure regarding the appeals process described in § 23 17.12 1 et seq.
21 and in the rules and regulations for the utilization review of care services, promulgated by the
22 department of health, the telephone numbers and addresses for the plan's office which handles
23 complaints as well as for the office which handles the appeals process under § 23 17.12 1 et seq.
24 and the rules and regulations for the utilization of health.

(4) Written statement of prospective and current enrollees' right to confidentiality of all
 health care record and information in the possession and/or control of the plan, its employees, its
 agents and parties with whom a contractual agreement exists to provide utilization review or who
 in any way have access to care information. A summary statement of the measures taken by the
 plan to ensure confidentiality of an individual's health care records shall be disclosed.

30 (5) Written disclosure of the enrollee's right to be free from discrimination by the health

31 plan and the right to refuse treatment without jeopardizing future treatment.

32 (6) Written disclosure of a plan's policy to direct enrollees to particular providers. Any
 33 limitations on reimbursement should the enrollee refuse the referral must be disclosed.

33 limitations on reimbursement should the enrollee refuse the referral must be disclosed.

34 (7) A summary of prior authorization or other review requirements including

preauthorization review, concurrent review, post service review, post payment review and any
 procedure that may lead the patient to be denied coverage for or not be provided a particular
 service.

4 (8) Any health plan that operates a provider incentive plan shall not enter into any
5 compensation agreement with any provider of covered services or pharmaceutical manufacturer
6 pursuant to which specific payment is made directly or indirectly to the provider as an
7 inducement or incentive to reduce or limit services, to reduce the length of stay or the use of
8 alternative treatment settings or the use of a particular medication with respect to an individual
9 patient, provided however, that capitation agreements and similar risk sharing arrangements are
10 not prohibited.

(9) Health plans must disclose to prospective and current enrollees the existence of
 financial arrangements for capitated or other risk sharing arrangements that exist with providers
 in a manner described in paragraphs (i), (ii), and (iii):

(i) "This health plan utilizes capitated arrangements, with its participating providers, or
 contains other similar risk sharing arrangements;

(ii) This health plan may include a capitated reimbursement arrangement or other similar
 risk sharing arrangement, and other financial arrangements with your provider;

18 (iii) This health plan is not capitated and does not contain other risk sharing

19 arrangements."

20 (10) Written disclosure of criteria for accessing emergency health care services as well
 21 as a statement of the plan's policies regarding payment for examinations to determine if

emergency health care services are necessary, the emergency care itself, and the necessary services following emergency treatment or stabilization. The health plan must respond to the request of the treating provider for post-stabilization treatment by approving or denying it as soon

25 as possible.

(11) Explanation of how health plan limitations impact enrollees, including information
 on enrollee financial responsibility for payment for co-insurance, co-payment, or other non covered, out-of-pocket, or out-of-plan services. This shall include information on deductibles and

- 29 benefits limitations including, but not limited to, annual limits and maximum lifetime benefits.
- 30 (12) The terms under which the health plan may be renewed by the plan enrollee,

31 including any reservation by the plan of any right to increase premiums.

32 (13) Summary of criteria used to authorize treatment.

- 33 (14) A schedule of revenues and expenses, including direct service ratios and other
- 34 statistical information which meets the requirements set forth below on a form prescribed by the

1	director.
2	-(15) Plan costs of health care services, including but not limited to all of the following:
3	-(i) Physician services;
4	-(ii) Hospital services, including both inpatients and outpatient services;
5	-(iii) Other professional services;
6	(iv) Pharmacy services, excluding pharmaceutical products dispensed in a physician's
7	office;
8	(v) Health education;
9	(vi) Substance abuse services and mental health services.
10	(16) Plan complaint, adverse decision, and prior authorization statistics. This statistical
11	data shall be updated annually:
12	(i) The ratio of the number of complaints received to the total number of covered
13	persons, reported by category, listed in paragraphs (b)(15)(i) (vi);
14	(ii) The ratio of the number of adverse decisions issued to the number of complaints
15	received, reported by category;
16	(iii) The ratio of the number of prior authorizations denied to the number of prior
17	authorizations requested, reported by category;
18	(iv) The ratio of the number of successful enrollee appeals to the total number of appeals
19	filed.
20	(17) Plans must demonstrate that:
21	(i) They have reasonable access to providers, so that all covered health care services will
22	be provided. This requirement cannot be waived and must be met in all areas where the health
23	<del>plan has enrollees;</del>
24	(ii) Urgent health care services, if covered, shall be available within a time frame that
25	meets standards set by the director.
26	(18) A comprehensive list of participating providers listed by office location, specialty if
27	applicable, and other information as determined by the director, updated annually.
28	(19) Plans must provide to the director, at intervals determined by the director, enrollee
29	satisfaction measures. The director is authorized to specify reasonable requirements for these
30	measures consistent with industry standards to assure an acceptable degree of statistical validity
31	and comparability of satisfaction measures over time and among plans. The director shall publish
32	periodic reports for the public providing information on health plan enrollee satisfaction.
33	(c) Issuance of certification.
34	(1) Upon receipt of an application for certification the director shall polify and afford

34 (1) Upon receipt of an application for certification, the director shall notify and afford

1 the public an opportunity to comment upon the application.

2	(2) A health care plan will meet the requirements of certification, subsection (b) by
3	providing information required in subsection (b) to any state or federal agency in conformance
4	with any other applicable state or federal law, or in conformity with standards adopted by an
5	accrediting organization provided that the director determines that the information is substantially
6	similar to the previously mentioned requirements and is presented in a format that provides a
7	meaningful comparison between health plans.
8	(3) All health plans shall be required to establish a mechanism, under which providers,
9	including local providers participating in the plan, provide input into the plan's health care policy,
10	including technology, medications and procedures, utilization review criteria and procedures,
11	quality and credentialing criteria, and medical management procedures.
12	(4) All health plans shall be required to establish a mechanism under which local
13	individual subscribers to the plan provide input into the plan's procedures and processes regarding
14	the delivery of health care services.
15	(5) A health plan shall not refuse to contract with or compensate for covered services an
16	otherwise eligible provider or non-participating provider solely because that provider has in good
17	faith communicated with one or more of his or her patients regarding the provisions, terms or
18	requirements of the insurer's products as they relate to the needs of that provider's patients.
19	(6) (i) All health plans shall be required to publicly notify providers within the health
20	plans' geographic service area of the opportunity to apply for credentials. This notification
21	process shall be required only when the plan contemplates adding additional providers and may
22	be specific as to geographic area and provider specialty. Any provider not selected by the health
23	plan may be placed on a waiting list.
24	(ii) This credentialing process shall begin upon acceptance of an application from a
25	provider to the plan for inclusion.
26	(iii) Each application shall be reviewed by the plan's credentialing body.
27	(iv) All health plans shall develop and maintain credentialing criteria to be utilized in
28	adding providers from the plans' network. Credentialing criteria shall be based on input from
29	providers credentialed in the plan and these standards shall be available to applicants. When
30	economic considerations are part of the decisions, the criteria must be available to applicants.
31	Any economic profiling must factor the specialty utilization and practice patterns and general
32	information comparing the applicant to his or her peers in the same specialty will be made
33	available. Any economic profiling of providers must be adjusted to recognize case mix, severity
34	of illness, age of patients and other features of a provider's practice that may account for higher

1	than or lower than expected costs. Profiles must be made available to those so profiled.
2	(7) A health plan shall not exclude a provider of covered services from participation in
3	its provider network based solely on:
4	(i) The provider's degree or license as applicable under state law; or
5	(ii) The provider of covered services lack of affiliation with, or admitting privileges at a
6	hospital, if that lack of affiliation is due solely to the provider's type of license.
7	(8) Health plans shall not discriminate against providers solely because the provider
8	treats a substantial number of patients who require expensive or uncompensated medical care.
9	(9) The applicant shall be provided with all reasons used if the application is denied.
10	(10) Plans shall not be allowed to include clauses in physician or other provider contracts
11	that allow for the plan to terminate the contract "without cause"; provided, however, cause shall
12	include lack of need due to economic considerations.
13	(11) (i) There shall be due process for non-institutional providers for all adverse
14	decisions resulting in a change of privileges of a credentialed non institutional provider. The
15	details of the health plan's due process shall be included in the plan's provider contracts.
16	(ii) A health plan is deemed to have met the adequate notice and hearing requirement of
17	this section with respect to a non-institutional provider if the following conditions are met (or are
18	waived voluntarily by the non-institutional provider):
19	(A) The provider shall be notified of the proposed actions and the reasons for the
20	
20	proposed action.
20	proposed action. (B) The provider shall be given the opportunity to contest the proposed action.
21	(B) The provider shall be given the opportunity to contest the proposed action.
21 22	(B) The provider shall be given the opportunity to contest the proposed action. (C) The health plan has developed an internal appeals process that has reasonable time
21 22 23	(B) The provider shall be given the opportunity to contest the proposed action. (C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.
21 22 23 24	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not</li> </ul>
<ul> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> </ul>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time</li> <li>limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not</li> <li>provided by the provider or provider group, the plan must require that a provider or group has met</li> </ul>
<ul> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> </ul>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time</li> <li>limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not</li> <li>provided by the provider or provider group, the plan must require that a provider or group has met</li> <li>all appropriate standards of the department of business regulation.</li> </ul>
<ul> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> </ul>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not provided by the provider or provider group, the plan must require that a provider or group has met all appropriate standards of the department of business regulation.</li> <li>(13) A health plan shall not include a most favored rate clause in a provider contract.</li> </ul>
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<ol> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> <li>28</li> <li>29</li> </ol>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not provided by the provider or provider group, the plan must require that a provider or group has met all appropriate standards of the department of business regulation.</li> <li>(13) A health plan shall not include a most favored rate clause in a provider contract.</li> <li>23-17.13-4. Penalties and enforcement. (a) The director of the department of health may, in lieu of the suspension or revocation of a license, levy an administrative penalty in an</li> </ul>
<ul> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> <li>28</li> <li>29</li> <li>30</li> </ul>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not provided by the provider or provider group, the plan must require that a provider or group has met all appropriate standards of the department of business regulation.</li> <li>(13) A health plan shall not include a most favored rate clause in a provider contract.</li> <li><u>23-17.13-4. Penalties and enforcement</u> (a) The director of the department of health may, in lieu of the suspension or revocation of a license, levy an administrative penalty in an amount not less than five hundred dollars (\$500) nor more than fifty thousand dollars (\$50,000),</li> </ul>
<ul> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> <li>28</li> <li>29</li> <li>30</li> <li>31</li> </ul>	<ul> <li>(B) The provider shall be given the opportunity to contest the proposed action.</li> <li>(C) The health plan has developed an internal appeals process that has reasonable time limits for the resolution of an internal appeal.</li> <li>(12) If the plan places a provider or provider group at financial risk for services not provided by the provider or provider group, the plan must require that a provider or group has met all appropriate standards of the department of business regulation.</li> <li>(13) A health plan shall not include a most favored rate clause in a provider contract.</li> <li>23-17.13 4. Penalties and enforcement. (a) The director of the department of health may, in lieu of the suspension or revocation of a license, levy an administrative penalty in an amount not less than five hundred dollars (\$500) nor more than fifty thousand dollars (\$50,000), if reasonable notice, in writing, is given of the intent to levy the penalty and the particular health</li> </ul>

1 <del>public.</del>

1	public.
2	(b) Any person who knowingly and willfully violates this chapter shall be guilty of a
3	misdemeanor and may be punished by a fine not to exceed five hundred dollars (\$500) or by
4	imprisonment for a period not exceeding one year, or both.
5	(c) (1) If the director of health shall for any reason have cause to believe that any
6	violation of this chapter has occurred or is threatened, the director of health may give notice to the
7	particular health organization and to their representatives, or other persons who appear to be
8	involved in the suspected violation, to arrange a conference with the alleged violators or their
9	authorized representatives for the purpose of attempting to ascertain the facts relating to the
10	suspected violation, and, in the event it appears that any violation has occurred or is threatened, to
11	arrive at an adequate and effective means of correcting or preventing the violation;
12	(2) Proceedings under this subsection shall be governed by chapter 35 of title 42.
13	(d) (1) The director of health may issue an order directing a particular health
14	organization or a representative of that health organization to cease and desist from engaging in
15	any act or practice in violation of the provisions of this chapter;
16	(2) Within thirty (30) days after service of the order to cease and desist, the respondent
17	may request a hearing on the question of whether acts or practices in violation of this chapter
18	have occurred. Those hearings shall be conducted pursuant to §§ 42-35-9 through 42-35-13, and
19	judicial review shall be available as provided by §§ 42-35-15 and 42-35-16.
20	(e) In the case of any violation of the provisions of this chapter, if the director of health
21	elects not to issue a cease and desist order, or in the event of noncompliance with a cease and
22	desist order issued pursuant to subsection (d), the director of health may institute a proceeding to
23	obtain injunctive relief, or seeking other appropriate relief, in the superior court for the county of
24	Providence.
25	23-17.13-5. Severability If any section, clause, or provision of this chapter shall be
26	held either unconstitutional or ineffective in whole or in part to the extent that it is not
27	unconstitutional or ineffective, it shall be valid and effective and no other section, clause or
28	provision shall on account thereof be termed invalid or ineffective.
29	23-17.13-6. Contracts with providers for dental services (a) No contract between a
30	dental plan of a health care entity and a dentist for the provision of services to patients may
31	require that a dentist provide services to its subscribers at a fee set by the health care entity unless
32	said services are covered services under the applicable subscriber agreement. "Covered services,"
33	as used herein, means services reimbursable under the applicable subscriber agreement, subject to
34	such contractual limitations on subscriber benefits as may apply, including, for example,

1 deductibles, waiting period or frequency limitations.

2	(b) For the purposes of this section "dental plan" shall include any policy of insurance
3	which is issued by a health care entity which provides for coverage of dental services not in
4	connection with a medical plan.
5	23-17.13-7. Contracts with providers and optometric services (a) No contract
6	between an eye care provider and a company offering accident and sickness insurance as defined
7	in chapter 18 of title 27; a nonprofit medical service corporation as defined in chapter 20 of title
8	27; or a health maintenance organization as defined in chapter 41 of title 27; or a vision plan, may
9	require that an eye care provider provide services or materials to its subscribers at a fee set by the
10	insurer or vision plan unless the insurer or vision plan compensates the eye care provider for the
11	provision of such services or materials to the patient. Reimbursement paid by the insurer or vision
12	plan for covered services and materials shall not provide nominal reimbursement in order to claim
13	that services and materials are covered services.
14	(b) (1) "Services" means services and materials for which reimbursement from the vision
15	plan is provided for by an enrollee's plan contract, or for which a reimbursement would be
16	available but for the application of the enrollee's contractual limitations of deductibles,
17	copayments, or coinsurance.
18	(2) "Materials" means and includes, but is not limited to, lenses, devices containing
19	lenses, prisms, lens treatments and coatings, contact lenses, orthoptics, vision training, and
20	prosthetic devices to correct, relieve, or treat defects or abnormal conditions of the human eye or
21	its adnexa.
22	(3) "Eye care provider" means an optometrist, optician, or ophthalmologist.
23	SECTION 3. Chapter 23-17.18 of the General Laws entitled "Health Plan Modification
24	Act" is hereby repealed in its entirety.
25	CHAPTER 23-17.18
26	Health Plan Modification Act
27	23-17.18-1. Modification of health plans (a) A health plan may materially modify the
28	terms of a participating agreement it maintains with a physician only if the plan disseminates in
29	writing by mail to the physician the contents of the proposed modification and an explanation, in
30	nontechnical terms, of the modification's impact.
31	(b) The health plan shall provide the physician an opportunity to amend or terminate the
32	physician contract with the health plan within sixty (60) days of receipt of the notice of
33	modification. Any termination of a physician contract made pursuant to this section shall be
34	effective fifteen (15) calendar days from the mailing of the notice of termination in writing by

1	mail to the health plan. The termination shall not affect the method of payment or reduce the
2	amount of reimbursement to the physician by the health plan for any patient in active treatment
3	for an acute medical condition at the time the patient's physician terminates his, her, or its
4	physician contract with the health plan until the active treatment is concluded or, if earlier, one
5	year after the termination; and, with respect to the patient, during the active treatment period the
6	physician shall be subject to all the terms and conditions of the terminated physician contract,
7	including but not limited to, all reimbursement provisions which limit the patient's liability.
8	(c) Nothing in this section shall apply to accident only, specified disease, hospital
9	indemnity, Medicare supplement, long-term care, disability income, or other limited benefit
10	health insurance policies.
11	SECTION 4. Title 27 of the General Laws entitled "INSURANCE" is hereby amended
12	by adding thereto the following chapter:
13	CHAPTER 18.8
14	HEALTH CARE ACCESSIBILITY AND QUALITY ASSURANCE ACT
15	27-18.8-1. Purpose The legislature declares that:
16	(1) It is in the best interest of the public that those individuals and care entities involved
17	with the delivery of plan coverage in our state meet the standards of this chapter to insure
18	accessibility and quality for the state's patients;
19	(2) Nothing in the legislation is intended to prohibit a health care entity or contractor
20	from forming limited networks of providers; and
21	(3) It is a vital state function to establish these standards for the conduct of health plans
22	by a health care entity in Rhode Island.
23	27-18.8-2. Definitions As used in this chapter:
24	(1) "Adverse determination" means a utilization review decision by a review agent not to
25	authorize a health care service. A decision by a review agent to authorize a health care service in
26	an alternative setting, a modified extension of stay, or an alternative treatment shall not constitute
27	an adverse determination if the review agent and provider are in agreement regarding the
28	decision. Adverse determinations include decisions not to authorize formulary and non-formulary
29	medication.
30	(2) "Benefit determination" means a decision of the enrollee's entitlement to payment for
31	covered health care services as defined in an agreement with the payor or its delegate.
32	(3) "Contractor" means a person/entity that:
33	(i) Establishes, operates or maintains a network of participating providers;
34	(ii) Contracts with an insurance company, a hospital or medical or dental service plan, an

1 employer, whether under written or self insured, an employee organization, or any other entity 2 providing coverage for health care services to administer a plan; and/or 3 (iii) Conducts or arranges for utilization review activities pursuant to this chapter. 4 (4) "Commissioner" means the commissioner of the Office of the Health Insurance 5 Commissioner. (5) "Emergency services" has the same meaning as the meaning contained in the rules 6 7 and regulations promulgated pursuant to chapter 12.3 of title 42, as may be amended from time to 8 time, and includes the sudden onset of a medical or mental condition that the absence of 9 immediate medical attention could reasonably be expected to result in placing the patient's health 10 in serious jeopardy, serious impairment to bodily or mental functions, or serious dysfunction of 11 any bodily organ or part. 12 (6) "Health care entity" means a licensed insurance company, hospital, or dental or 13 medical service plan or health maintenance organization, or a contractor as described in 14 subsection (3) of this section, that operates a health plan. (7) "Health care services" includes, but is not limited to, medical, mental health, 15 16 substance abuse, and dental services. 17 (8) "Health plan" means a plan operated by a health care entity as described in subsection (6) of this section that provides for the delivery of care services to persons enrolled in the plan 18 19 through: 20 (i) Arrangements with selected providers to furnish health care services; and/or 21 (ii) Financial incentives for persons enrolled in the plan to use the participating providers 22 and procedures provided for by the plan. 23 (9) "Provider" means a physician, hospital, pharmacy, laboratory, dentist, or other state 24 licensed or other state recognized provider of health care services or supplies, and whose services are recognized pursuant to 213(d) of the Internal Revenue Code, 26 U.S.C. § 213(d), that has 25 26 entered into an agreement with a health care entity as described in subsection (6) of this section or 27 contractor as described in subsection (2) of this section to provide these services or supplies to a 28 patient enrolled in a plan. 29 (10) "Most favored rate clause" means a provision in a provider contract whereby the 30 rates or fees to be paid by a health plan are fixed, established or adjusted to be equal to or lower 31 than the rates or fees paid to the provider by any other health plan or third party payor. 32 27-18.8-3. Certification of health plans. -- (a) Certification process. 33 (1) Certification. 34 (i) The commissioner, in consultation with the director of the department of health, shall

- establish a process for certification of health plans meeting the requirements of certification in
   subsection (b) of this section.
- 3 (ii) The commissioner shall act upon the health plan's completed application for
  4 certification within ninety (90) days of receipt of such application for certification.
- (2) Review and recertification. To ensure compliance with subsection (b) of this section,
  the commissioner shall establish procedures for the periodic review and recertification of health
  plans not less than every five (5) years; provided, however, that the commissioner may review the
  certification of a health plan at any time if there exists evidence that a health plan may be in
- 9 <u>violation of subsection (b) of this section.</u>
- (3) Cost of certification. The total cost of obtaining and maintaining certification under
  this title and compliance with the requirements of the applicable rules and regulations are borne
  by the entities so certified and shall be one hundred and fifty percent (150%) of the total salaries
  paid to the certifying personnel of the office engaged in those certifications less any salary
  reimbursements and shall be paid to the commissioner to and for the use of the office. That
  assessment shall be in addition to any taxes and fees otherwise payable to the state.
  (b) Requirements for certification. The commissioner, in consultation with the director of
- 17 the department of health, shall establish standards and procedures for the certification of health
- 18 plans that conduct business in this state and who have demonstrated the ability to ensure that
- 19 health care services will be provided in a manner to assure availability and accessibility, adequate
- 20 personnel and facilities, and continuity of service, and has demonstrated arrangements for
- 21 ongoing quality assurance programs regarding care processes and outcomes; other standards shall
- 22 <u>consist of, but are not limited to, the following:</u>
- 23 (1) Plans must demonstrate that:
- 24 (i) They have reasonable access to providers, so that all covered health care services will
- 25 <u>be provided. This requirement cannot be waived and must be met in all areas where the health</u>
- 26 <u>plan has enrollees;</u>
- 27 (ii) Covered health care services shall be available within a time frame that meets
  28 standards set by the commissioner.
- 29 (2) A comprehensive list of participating providers listed by office location, specialty if
- 30 <u>applicable, and other information as determined by the commissioner, updated annually.</u>
- 31 (c) Issuance of certification.
- 32 (1) A health care plan shall meet all or some of the requirements of certification, by
- 33 providing the required certification information to any state or federal agency in conformance
- 34 with any other applicable state or federal law, or in conformity with standards adopted by an

1 accrediting organization provided that the commissioner determines that the information is 2 substantially similar to the previously mentioned requirements. 3 (2) All health plans shall be required to establish a mechanism, under which providers, 4 including local providers participating in the plan, provide input into the plan's health care policy, 5 including technology, medications and procedures, utilization review criteria and procedures, guality and credentialing criteria, and medical management procedures. 6 7 (2) All health plans shall be required to establish a mechanism under which local 8 individual subscribers to the plan provide input into the plan's procedures and processes regarding 9 the delivery of health care services. 10 (3) A health plan shall not refuse to contract with or compensate for covered services an 11 otherwise eligible provider or non-participating provider solely because that provider has in good 12 faith communicated with one or more of their patients regarding the provisions, terms or 13 requirements of the insurer's products as they relate to the needs of that provider's patients. 14 (4) The health plan provider contracting and credentialing process shall include the 15 following: 16 (i) This credentialing process shall begin upon acceptance of an application from a 17 provider to the plan for inclusion. 18 (ii) Each application shall be reviewed by the plan's credentialing body. 19 (iii) All health plans shall develop and maintain credentialing criteria to be utilized in 20 adding providers from the plans' network. Credentialing criteria shall be based on input from 21 providers credentialed in the plan and these standards shall be available to applicants. When 22 economic considerations are part of the decisions, the criteria must be available to applicants. 23 Any economic profiling must factor the specialty utilization and practice patterns and general 24 information comparing the applicant to their peers in the same specialty will be made available. 25 Any economic profiling of providers must be adjusted to recognize case mix, severity of illness, 26 age of patients and other features of a provider's practice that may account for higher than or 27 lower than expected costs. Profiles must be made available to those so profiled. 28 (5) A health plan shall not exclude a provider of covered services from participation in its 29 provider network based solely on: 30 (i) The provider's degree or license as applicable under state law; or 31 (ii) The provider of covered services lack of affiliation with, or admitting privileges at a 32 hospital, if that lack of affiliation is due solely to the provider's type of license. 33 (6) Health plans shall not discriminate against providers solely because the provider treats 34 a substantial number of patients who require expensive or uncompensated medical care.

- 1 (7) The applicant shall be provided with all reasons used if the application is denied. 2 (8) Plans shall not be allowed to include clauses in physician or other provider contracts that allow for the plan to terminate the contract "without cause"; provided, however, cause shall 3 4 include lack of need due to economic considerations. 5 (9)(i) There shall be due process for non-institutional providers for all adverse decisions resulting in a change of privileges of a credentialed non-institutional provider. The details of the 6 7 health plan's due process shall be included in the plan's provider contracts. 8 (ii) A health plan is deemed to have met the adequate notice and hearing requirement of 9 this section with respect to a non-institutional provider if the following conditions are met (or are 10 waived voluntarily by the non-institutional provider): 11 (A) The provider shall be notified of the proposed actions and the reasons for the 12 proposed action. 13 (B) The provider shall be given the opportunity to contest the proposed action. 14 (C) The health plan has developed an internal appeals process that has reasonable time 15 limits for the resolution of an internal appeal. 16 (10) A health plan shall not include a most favored rate clause in a provider contract. 17 (11) A health plan may materially modify the terms of a participating agreement it maintains with a physician only if the plan disseminates in writing by mail to the physician the 18 19 contents of the proposed modification and an explanation, in nontechnical terms, of the 20 modification's impact. 21 (12) The health plan shall provide the physician an opportunity to amend or terminate the 22 physician contract with the health plan within sixty (60) days of receipt of the notice of 23 modification. Any termination of a physician contract made pursuant to this section shall be 24 effective fifteen (15) calendar days from the mailing of the notice of termination in writing by 25 mail to the health plan. The termination shall not affect the method of payment or reduce the 26 amount of reimbursement to the physician by the health plan for any patient in active treatment 27 for an acute medical condition at the time the patient's physician terminates their physician 28 contract with the health plan until the active treatment is concluded or, if earlier, one year after 29 the termination; and, with respect to the patient, during the active treatment period the physician 30 shall be subject to all the terms and conditions of the terminated physician contract, including but 31 not limited to, all reimbursement provisions which limit the patient's liability. 32 27-18.8-4. Contracts with providers for dental services. -- (a) No contract between a dental plan of a health care entity and a dentist for the provision of services to patients may 33
- 34 require that a dentist provide services to its subscribers at a fee set by the health care entity unless

1 said services are covered services under the applicable subscriber agreement. "Covered services," 2 as used herein, means services reimbursable under the applicable subscriber agreement, subject to 3 such contractual limitations on subscriber benefits as may apply, including, for example, 4 deductibles, waiting period or frequency limitations. (b) For the purposes of this section "dental plan" shall include any policy of insurance 5 which is issued by a health care entity which provides for coverage of dental services not in 6 7 connection with a medical plan. 8 27-18.8-5. Contracts with providers and optometric services. -- (a) No contract 9 between an eye care provider and a company offering accident and sickness insurance as defined 10 in chapter 18 of title 27; a nonprofit medical service corporation as defined in chapter 20 of title 11 27; or a health maintenance organization as defined in chapter 41 of title 27; or a vision plan, may 12 require that an eye care provider provide services or materials to its subscribers at a fee set by the 13 insurer or vision plan unless the insurer or vision plan compensates the eye care provider for the 14 provision of such services or materials to the patient. Reimbursement paid by the insurer or vision 15 plan for covered services and materials shall not provide nominal reimbursement in order to claim 16 that services and materials are covered services. 17 (b)(1) "Services" means services and materials for which reimbursement from the vision 18 plan is provided for by an enrollee's plan contract, or for which a reimbursement would be 19 available but for the application of the enrollee's contractual limitations of deductibles, 20 copayments, or coinsurance. 21 (2) "Materials" means and includes, but is not limited to, lenses, devices containing 22 lenses, prisms, lens treatments and coatings, contact lenses, orthoptics, vision training, and prosthetic devices to correct, relieve, or treat defects or abnormal conditions of the human eye or 23 24 its adnexa. 25 (3) "Eye care provider" means an optometrist, optician, or ophthalmologist. 26 27-18.8-6. Penalties and enforcement. -- (a) The commissioner may, in lieu of the 27 suspension or revocation of a license, levy an administrative penalty in an amount not less than 28 five hundred dollars (\$500) nor more than fifty thousand dollars (\$50,000), if reasonable notice, 29 in writing, is given of the intent to levy the penalty and the particular health organization has a 30 reasonable time in which to remedy the defect in its operations which gave rise to the penalty 31 citation. The commissioner may augment this penalty by an amount equal to the sum that the 32 commissioner calculates to be the damages suffered by enrollees or other members of the public. 33 (b) Any person who knowingly and willfully violates this chapter shall be guilty of a 34 misdemeanor and may be punished by a fine not to exceed five hundred dollars (\$500) or by 1 imprisonment for a period not exceeding one year, or both.

2	(c)(1) If the commissioner shall for any reason have cause to believe that any violation of
3	this chapter has occurred or is threatened, the commissioner may give notice to the particular
4	health organization and to their representatives, or other persons who appear to be involved in the
5	suspected violation, to arrange a conference with the alleged violators or their authorized
6	representatives for the purpose of attempting to ascertain the facts relating to the suspected
7	violation, and, in the event it appears that any violation has occurred or is threatened, to arrive at
8	an adequate and effective means of correcting or preventing the violation;
9	(2) Proceedings under this subsection shall be governed by chapter 35 of title 42.
10	(d)(1) The commissioner may issue an order directing a particular health organization or
11	a representative of that health organization to cease and desist from engaging in any act or
12	practice in violation of the provisions of this chapter;
13	(2) Within thirty (30) days after service of the order to cease and desist, the respondent
14	may request a hearing on the question of whether acts or practices in violation of this chapter
15	have occurred. Those hearings shall be conducted pursuant to §§ 42-35-9 through 42-35-13, and
16	judicial review shall be available as provided by §§ 42-35-15 and 42-35-16.
17	(e) In the case of any violation of the provisions of this chapter, if the commissioner
18	elects not to issue a cease and desist order, or in the event of noncompliance with a cease and
19	desist order issued pursuant to subsection (d) of this section, the commissioner may institute a
20	proceeding to obtain injunctive relief, or seeking other appropriate relief, in the superior court for
21	the county of Providence.
22	27-18.8-7. Severability If any section, clause, or provision of this chapter shall be
23	held either unconstitutional or ineffective in whole or in part to the extent that it is not
24	unconstitutional or ineffective, it shall be valid and effective and no other section, clause or
25	provision shall on account thereof be termed invalid or ineffective.
26	SECTION 5. Title 27 of the General Laws entitled "INSURANCE" is hereby amended
27	by adding thereto the following chapter:
28	CHAPTER 18.9
29	HEALTH CARE SERVICES UTILIZATION REVIEW ACT
30	27-18.9-1. Purpose of chapter The purpose of the chapter is to:
31	(1) Promote the delivery of quality health care in a cost effective manner;
32	(2) Foster greater coordination between health care providers, patients, payors and
33	utilization review entities:
34	(3) Protect patients, businesses, and providers by ensuring that review agents are

1 qualified to perform utilization review activities and to make informed decisions on the 2 appropriateness of medical care; and 3 (4) Ensure that review agents maintain the confidentiality of medical records in 4 accordance with applicable state and federal laws. 5 (5) Provide for consultation by the department of health to the office of the health insurance commissioner in furtherance of the purposes of this chapter. 6 7 27-18.9-2. Definitions. -- As used in this chapter, the following terms are defined as 8 follows: 9 (1) "Adverse determination" means a utilization review decision by a review agent not to 10 authorize a health care service. A decision by a review agent to authorize a health care service in 11 an alternative setting, a modified extension of stay, or an alternative treatment shall not constitute 12 an adverse determination if the review agent and provider are in agreement regarding the 13 decision. Adverse determinations include decisions not to authorize formulary and non-formulary 14 medication. 15 (2) "Appeal" means a subsequent review of an adverse determination upon request by a 16 patient or provider to reconsider all or part of the original decision. 17 (3) "Authorization" means the review agent's utilization review, performed according to § 18 27-18.9-2(20), concluded that the allocation of health care services of a provider, given or 19 proposed to be given to a patient was approved or authorized. 20 (4) "Benefit determination" means a decision of the enrollee's entitlement to payment for 21 covered health care services as defined in an agreement with the payor or its delegate. 22 (5) "Certificate" means a certificate of registration granted by the commissioner to a 23 review agent. 24 (6) "Complaint" means a written expression of dissatisfaction by a patient, or provider. 25 The appeal of an adverse determination is not considered a complaint. (7) "Concurrent assessment" means an assessment of the medical necessity and/or 26 27 appropriateness of health care services conducted during a patient's hospital stay or course of 28 treatment. If the medical problem is ongoing, this assessment may include the review of services 29 after they have been rendered and billed. 30 (8) "Office" means the office of the health insurance commissioner. 31 (9) "Commissioner" means the health insurance commissioner. 32 (10) "Emergent health care services" has the same meaning as that meaning contained in 33 the rules and regulations promulgated pursuant to chapter 12.3 of title 42 as may be amended from time to time and includes those resources provided in the event of the sudden onset of a 34

1 medical, mental health, or substance abuse or other health care condition manifesting itself by 2 acute symptoms of a severity (e.g. severe pain) where the absence of immediate medical attention 3 could reasonably be expected to result in placing the patient's health in serious jeopardy, serious 4 impairment to bodily or mental functions, or serious dysfunction of any body organ or part. 5 (11) "Patient" means an enrollee or participant in all hospital or medical plans seeking health care services and treatment from a provider. 6 7 (12) "Payor" means a health insurer, self-insured plan, nonprofit health service plan, 8 health insurance service organization, preferred provider organization, health maintenance 9 organization or other entity authorized to offer health insurance policies or contracts or pay for 10 the delivery of health care services or treatment in this state. 11 (13) "Practitioner" means any person licensed to provide or otherwise lawfully providing 12 health care services, including, but not limited to, a physician, dentist, nurse, optometrist, 13 podiatrist, physical therapist, clinical social worker, or psychologist. 14 (14) "Prospective assessment" means an assessment of the medical necessity and/or 15 appropriateness of health care services prior to services being rendered. 16 (15) "Provider" means any health care facility, as defined in § 23-17-2 including any 17 mental health and/or substance abuse treatment facility, physician, or other licensed practitioners 18 identified to the review agent as having primary responsibility for the care, treatment, and 19 services rendered to a patient. 20 (16) "Retrospective assessment" means an assessment of the medical necessity and/or 21 appropriateness of health care services that have been rendered. This shall not include reviews 22 conducted when the review agency has been obtaining ongoing information. 23 (17) "Review agent" means a person or entity or insurer performing utilization review 24 that is either employed by, affiliated with, under contract with, or acting on behalf of: 25 (i) A business entity doing business in this state; 26 (ii) A party that provides or administers health care benefits to citizens of this state, 27 including a health insurer, self-insured plan, nonprofit health service plan, health insurance 28 service organization, preferred provider organization or health maintenance organization 29 authorized to offer health insurance policies or contracts or pay for the delivery of health care 30 services or treatment in this state; or 31 (iii) A provider not involved in the care of the patient. 32 (18) "Same or similar specialty" means a practitioner who has the appropriate training 33 and experience that is the same or similar as the attending provider in addition to experience in 34 treating the same problems to include any potential complications as those under review.

1	(19) "Urgent health care services" has the same meaning as that meaning contained in the
2	rules and regulations promulgated pursuant to chapter 12.3 of title 42 as may be amended from
3	time to time and includes those resources necessary to treat a symptomatic medical, mental
4	health, or substance abuse or other health care condition requiring treatment within a twenty-four
5	(24) hour period of the onset of such a condition in order that the patient's health status not
6	decline as a consequence. This does not include those conditions considered to be emergent
7	health care services as defined in subsection (10) of this section.
8	(20) "Utilization review" means the prospective, concurrent, or retrospective assessment
9	of the necessity and/or appropriateness of the allocation of health care services of a provider,
10	given or proposed to be given to a patient. Utilization review does not include:
11	(i) Elective requests for the clarification of coverage; or
12	(ii) Benefit determination; or
13	(iii) Claims review that does not include the assessment of the medical necessity and
14	appropriateness; or
15	(iv) A provider's internal quality assurance program except if it is associated with a health
16	care financing mechanism; or
17	(v) The therapeutic interchange of drugs or devices by a pharmacy operating as part of a
18	licensed inpatient health care facility; or
19	(vi) The assessment by a pharmacist licensed pursuant to the provisions of chapter 19 of
20	title 5 and practicing in a pharmacy operating as part of a licensed inpatient health care facility in
21	the interpretation, evaluation and implementation of medical orders, including assessments and/or
22	comparisons involving formularies and medical orders.
23	(21) "Utilization review plan" means a description of the standards governing utilization
24	review activities performed by a private review agent.
25	(22) "Health care services" means and includes an admission, diagnostic procedure,
26	therapeutic procedure, treatment, extension of stay, the ordering and/or filling of formulary or
27	non-formulary medications, and any other services, activities, or supplies that are covered by the
28	patient's benefit plan.
29	(23) "Therapeutic interchange" means the interchange or substitution of a drug with a
30	dissimilar chemical structure within the same therapeutic or pharmacological class that can be
31	expected to have similar outcomes and similar adverse reaction profiles when given in equivalent
32	doses, in accordance with protocols approved by the president of the medical staff or medical
33	director and the director of pharmacy.
34	27-18.9-3. General requirements (a) A review agent shall not conduct utilization

1 review in the state unless the office has granted the review agent a certificate. 2 (b) Individuals shall not be required to hold separate certification under this chapter when 3 acting as either an employee of, an affiliate of, a contractor for, or otherwise acting on behalf of a 4 certified review agent. 5 (c) The office shall issue a certificate to an applicant that has met the minimum standards established by this chapter, and regulations promulgated in accordance with it, including the 6 7 payment of any fees as required, and other applicable regulations of the office. 8 (d) A certificate issued under this chapter is not transferable, and the transfer of fifty 9 percent (50%) or more of the ownership of a review agent shall be deemed a transfer. 10 (e) After consultation with the payors and providers of health care, the office shall adopt 11 regulations necessary to implement the provisions of this chapter. 12 (f) The commissioner is authorized to establish any fees for initial application, renewal 13 applications, and any other administrative actions deemed necessary by the commissioner to 14 implement this chapter. 15 (g) The total cost of certification under this title shall be borne by the certified entities 16 and shall be one hundred fifty percent (150%) of the total salaries paid to the certifying personnel 17 of the office engaged in those certifications less any salary reimbursements and shall be paid to the commissioner to and for the use of the office. That assessment shall be in addition to any 18 19 taxes and fees otherwise payable to the state, and shall be paid to the commissioner to and for the 20 use of the office. 21 (h) The application and other fees required under this chapter shall be sufficient to pay 22 for the administrative costs of the certificate program and any other reasonable costs associated 23 with carrying out the provisions of this chapter. 24 (i) A certificate expires on the third anniversary of its effective date unless the certificate 25 is renewed for a three (3) year term as provided in this chapter. 26 (j) Any systemic changes in the review agents operations relative to certification 27 information on file shall be submitted to the office for approval within thirty (30) days prior to 28 implementation. For purposes of this chapter, systemic changes are further defined in regulation. 29 27-18.9-4. General application requirements. -- An application for certification or

- 30 recertification shall be accompanied by documentation to evidence the following:
- 31 (1) The requirement that the review agent provide patients and providers with a summary
- 32 of its utilization review plan including a summary of the standards, procedures and methods to be
- 33 <u>used in evaluating proposed or delivered health care services;</u>
- 34 (2) The circumstances, if any, under which utilization review may be delegated to any

1 other utilization review program and evidence that the delegated agency is a certified utilization

2 review agency delegated to perform utilization review pursuant to all of the requirements of this

3 <u>chapter</u>;

- 4 (3) A complaint resolution process consistent with § 27-18.9-2(6) and acceptable to the
  5 office, whereby patients, their physicians, or other health care providers may seek resolution of
  6 complaints and other matters of which the review agent has received written notice;
- 7 (4) The type and qualifications of personnel (employed or under contract) authoriz
- 7 (4) The type and qualifications of personnel (employed or under contract) authorized to
- 8 perform utilization review, including a requirement that only a practitioner with the same license
- 9 status as the ordering practitioner, or a licensed physician or dentist, is permitted to make a
- 10 prospective or concurrent adverse determination;
- 11 (5) The requirement that a representative of the review agent is reasonably accessible to
- 12 patients, patient's family and providers at least five (5) days a week during normal business in
- 13 <u>Rhode Island and during the hours of the agency's review operations;</u>
- (6) The policies and procedures to ensure that all applicable state and federal laws to
   protect the confidentiality of individual medical records are followed;
- 16 (7) The policies and procedures regarding the notification and conduct of patient
   17 interviews by the review agent;
- 18 (8) The requirement that no employee of, or other individual rendering an adverse
   19 determination for, a review agent may receive any financial incentives based upon the number of
- 20 denials of certification made by that employee or individual;
- (9) The requirement that the utilization review agent shall not impede the provision of
   health care services for treatment and/or hospitalization or other use of a provider's services or
- 23 <u>facilities for any patient;</u>
- (10) Evidence that the review agent has not entered into a compensation agreement or
   contract with its employees or agents whereby the compensation of its employees or its agents is
   based upon a reduction of services or the charges for those services, the reduction of length of
- 27 stay, or utilization of alternative treatment settings; provided, nothing in this chapter shall prohibit
- 28 agreements and similar arrangements; and
- 29 (11) An adverse determination and internal appeals process consistent with § 27-18.9-8
- 30 and acceptable to the office, whereby patients, their physicians, or other health care providers
- 31 may seek prompt reconsideration or appeal of adverse determinations by the review agent.
- 32 <u>27-18.9-5. Denial, suspension, or revocation of certificate. -- (a) The office may deny a</u>
- 33 certificate upon review of the application if, upon review of the application, it finds that the
- 34 <u>applicant proposing to conduct utilization review does not meet the standards required by this</u>

- 1 <u>chapter or by any regulations promulgated pursuant to this chapter.</u>
- 2 (b) The office may revoke a certificate and/or impose reasonable monetary penalties not
- 3 to exceed five thousand dollars (\$5,000) per violation in any case in which:
- 4 (1) The review agent fails to comply substantially with the requirements of this chapter or
  5 of regulations adopted pursuant to this chapter;
- 6 (2) The review agent fails to comply with the criteria used by it in its application for a
  7 certificate; or
- 8 (3) The review agent refuses to permit examination by the commissioner to determine 9 compliance with the requirements of this chapter and regulations promulgated pursuant to the 10 authority granted to the commissioner in this chapter; provided, however, that the examination 11 shall be subject to the confidentiality and "need to know" provisions of §§ 27-18.9-8 (c)(4) and 12 (c)(5). These determinations may involve consideration of any written grievances filed with the 13 office against the review agent by patients or providers.
- 14 (c) Any applicant or certificate holder aggrieved by an order or a decision of the office
- 15 made under this chapter without a hearing may, within thirty (30) days after notice of the order or
- 16 decision, make a written request to the office for a hearing on the order or decision pursuant to §
- 17 <u>42-35-15.</u>
- (d) The procedure governing hearings authorized by this section shall be in accordance
  with §§ 42-35-9 through 42-35-13 as stipulated in § 42-35-14(a). A full and complete record shall
  be kept of all proceedings, and all testimony shall be recorded but need not be transcribed unless
  the decision is appealed pursuant to § 42-35-15. A copy or copies of the transcript may be
- 22 obtained by any interested party upon payment of the cost of preparing the copy or copies.
- 23 Witnesses may be subpoenaed by either party.
- 24 <u>27-18.9-6. Judicial review. --</u> Any person who has exhausted all administrative remedies
   25 available to him or her within the office, and who is aggrieved by a final decision of the office
- 26 <u>under § 27-18.9-5, is entitled to judicial review pursuant to §§ 42-35-15 and 42-35-16.</u>
- 27 <u>27-18.9-7. Waiver of requirements. -- (a) The commissioner may waive all or part of</u>
- 28 the requirements of this chapter if the agent maintains and provides evidence of accreditation by
- 29 <u>an organization that has been approved by the commissioner and in accordance with regulation.</u>
- 30 (b) The office shall waive the requirements of this chapter only when a conflict exists
- 31 with those activities of a review agent that are conducted pursuant to contracts with the state or
- 32 the federal government or those activities under other state or federal jurisdictions.
- 33 (c) The office shall waive de minimus activity, in accordance with the regulations
- 34 <u>adopted by the commissioner.</u>

1	27-18.9-7.1. Variance of statutory requirements Statutory variances shall be issued
2	for a period not to exceed one year and may be subject to such terms and conditions deemed
3	necessary as determined by the commissioner. Prior to issuing a statutory variance the office shall
4	provide notice and public hearing to ensure necessary patient and health care provider protections
5	in the process.
6	27-18.9-8. Review agency requirement for adverse determination and internal
7	appeals (a) The adverse determination and appeals process of the review agent shall conform
8	to the following:
9	(1) Notification of a prospective adverse determination by the review agent shall be
10	mailed or otherwise communicated to the provider of record and to the patient or other
11	appropriate individual as follows:
12	(i) Within fifteen (15) calendar days of receipt of all the information necessary to
13	complete a review of non-urgent and/or non-emergent services;
14	(ii) Within seventy-two (72) hours of receipt of all the information necessary to complete
15	a review of urgent and/or emergent services; and
16	(iii) Prior to the expected date of service.
17	(2) Notification of a concurrent adverse determination shall be mailed or otherwise
18	communicated to the patient and to the provider of record period as follows:
19	(i) To the provider(s) prior to the end of the current certified period; and
20	(ii) To the patient within one business day of making the adverse determination.
21	(3) Notification of a retrospective adverse determination shall be mailed or otherwise
22	communicated to the patient and to the provider of record within thirty (30) calendar days of
23	receipt of a request for payment with all supporting documentation for the covered benefit being
24	reviewed.
25	(4) A utilization review agency shall not retrospectively deny authorization for health
26	care services provided to a covered person when an authorization has been obtained for that
27	service from the review agent unless the approval was based upon inaccurate information
28	material to the review or the health care services were not provided consistent with the provider's
29	submitted plan of care and/or any restrictions included in the prior approval granted by the review
30	agent.
31	(5) Any notice of an adverse determination shall include:
32	(i) The principal reasons for the adverse determination, to include explicit documentation
33	of the criteria not met and/or the clinical rationale utilized by the agency's clinical reviewer in
34	making the adverse determination. The criteria shall be in accordance with the agency criteria

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- 1 noted in § 27-18.9-8(d) and shall be made available within the first level appeal timeframe if
- 2 requested unless otherwise provided as part of the adverse determination notification process;
- 3 (ii) The procedures to initiate an appeal of the adverse determination, including the name
- 4 and telephone number of the person to contract with regard to an appeal;
- 5 (iii) The necessary contact information to complete the two-way direct communication
- 6 <u>defined in § 27-18.9-8(a)(7); and</u>
- 7 (iv) The information noted in § 27-18.9-8(a)(5)(i)(ii)(iii) for all verbal notifications
  8 followed by written notification to the patient and provider(s).
- 9 (6) All initial retrospective adverse determinations of a health care service that had been
- 10 ordered by a physician, dentist or other practitioner shall be made, documented and signed
- 11 consistent with the regulatory requirements which shall be developed by the office with the input
- 12 of review agents, providers and other affected parties.
- 13 (7) A level one appeal decision of an adverse determination shall not be made until an
- 14 <u>appropriately qualified and licensed review physician, dentist or other practitioner has spoken to,</u>
- 15 or otherwise provided for, an equivalent two-way direct communication with the patient's
- 16 attending physician, dentist, other practitioner, other designated or qualified professional or
- 17 provider responsible for treatment of the patient concerning the medical care, with the exception
- 18 <u>of the following:</u>
- 19 (i) When the attending provider is not reasonably available;
- 20 (ii) When the attending provider chooses not to speak with agency staff;
- 21 (iii) When the attending provider has negotiated an agreement with the review agent for
- 22 <u>alternative care; and/or</u>
- (iv) When the attending provider requests a peer to peer communication prior to the
   adverse determination, the review agency shall then comply with § 27-18.9-8(c)(1) in responding
   to such a request. Such requests shall be on the case specific basis unless otherwise arranged for
   in advance by the provider.
- 27 (8) All initial, prospective and concurrent adverse determinations of a health care service
  28 that had been ordered by a physician, dentist or other practitioner shall be made, documented and
  29 signed by a licensed practitioner with the same licensure status as the ordering practitioner or a
  30 licensed physician or dentist. This does not prohibit appropriately qualified review agency staff
  31 from engaging in discussions with the attending provider, the attending provider's designee or
  32 appropriate health care facility and office personnel regarding alternative service and treatment
  33 options. Such a discussion shall not constitute an adverse determination provided though that any
- 34 change to the provider's original order and/or any decision for an alternative level of care must be

1 made and/or appropriately consented to by the attending provider or the provider's designee 2 responsible for treating the patient. 3 (9) The requirement that, upon written request made by or on behalf of a patient, any 4 adverse determination and/or appeal shall include the written evaluation and findings of the 5 reviewing physician, dentist or other practitioner. The review agent is required to accept a verbal request made by or on behalf of a patient for any information where a provider or patient can 6 7 demonstrate that a timely response is urgent. 8 (b) The review agent shall conform to the following for the appeal of an adverse 9 determination: 10 (1) The review agent shall maintain and make available a written description of the 11 appeal procedure by which either the patient or the provider of record may seek review of 12 determinations not to authorize a health care service. The process established by each review 13 agent may include a reasonable period within which an appeal must be filed to be considered and 14 that period shall not be less than one hundred eighty (180) days of receipt of the adverse 15 determination. 16 (2) The review agent shall notify, in writing, the patient and provider of record of its 17 decision on the appeal as soon as practical, but in no case later than fifteen (15) or twenty-one 18 (21) business days if verbal notice is given within fifteen (15) business days after receiving the 19 required documentation on the appeal. 20 (3) The review agent shall also provide for an expedited appeals process for emergency 21 or life threatening situations. Each review agent shall complete the adjudication of expedited appeals within two (2) business days or seventy-two (72) hours, whichever occurs sooner, of the 22 23 date the appeal is filed and all information necessary to complete the appeal is received by the 24 review agent. 25 (4) All first level appeals of determinations not to authorize a health care service that had 26 been ordered by a physician, dentist, or other practitioner shall be made, documented, and signed 27 by a licensed practitioner with the same licensure status as the ordering practitioner or a licensed 28 physician or a licensed dentist. 29 (5) All second level appeal decisions shall be made, signed, and documented by a 30 licensed practitioner in the same or a similar general specialty as typically manages the medical 31 condition, procedure, or treatment under discussion. 32 (6) The review agent shall maintain records of written appeals and their resolution, and 33 shall provide reports as requested by the office. 34 (c) The review agency must conform to the following requirements when making its

- 1 <u>adverse determination and appeal decisions:</u>
- 2 (1) The review agent must assure that the licensed practitioner or licensed physician is reasonably available to review the case as required under § 27-18.9-8(a)(7) and shall conform to 3 4 the following: 5 (i) Each agency peer reviewer shall have access to and review all necessary information as requested by the agency and/or submitted by the provider(s) and/or patients; 6 7 (ii) Each agency shall provide accurate peer review contact information to the provider at 8 the time of service, if requested, and/or prior to such service, if requested. This contact 9 information must provide a mechanism for direct communication with the agency's peer 10 reviewer; 11 (iii) Agency peer reviewers shall respond to the provider's request for a two-way direct 12 communication defined in § 27-18.9-8(a)(7)(iv) as follows: 13 (A) For a prospective review of non-urgent and non-emergent health care services, a 14 response within one business day of the request for a peer discussion; 15 (B) For concurrent and prospective reviews of urgent and emergent health care services, a 16 response within a reasonable period of time of the request for a peer discussion; and 17 (C) For retrospective reviews, prior to the first level appeal decision. 18 (iv) The review agency will have met the requirements of a two-way direct 19 communication, when requested and/or as required prior to the first level of appeal, when it has 20 made two (2) reasonable attempts to contact the attending provider directly. 21 (v) Repeated violations of this section shall be deemed to be substantial violations 22 pursuant to § 27-18.9-5(b) and shall be cause for the imposition of penalties under that section. (2) No reviewer at any level under this section shall be compensated or paid a bonus or 23 24 incentive based on making or upholding an adverse determination. 25 (3) No reviewer under this section who has been involved in prior reviews of the case 26 under appeal or who has participated in the direct care of the patient may participate as the sole 27 reviewer in reviewing a case under appeal; provided, however, that when new information has 28 been made available at the first level of appeal, then the review may be conducted by the same 29 reviewer who made the initial adverse determination. 30 (4) A review agent is only entitled to review information or data relevant to the utilization 31 review process. A review agent may not disclose or publish individual medical records or any 32 confidential medical information obtained in the performance of utilization review activities. A 33 review agent shall be considered a third party health insurer for the purposes of § 5-37.3-6(b)(6) of this state and shall be required to maintain the security procedures mandated in § 5-37.3-4(c). 34

1	(5) Notwithstanding any other provision of law, the review agent, the office, and all other
2	parties privy to information which is the subject of this chapter shall comply with all state and
3	federal confidentiality laws, including, but not limited to, chapter 37.3 of title 5 (Confidentiality
4	of Health Care Communications and Information Act) and specifically § 5-37.3-4(c), which
5	requires limitation on the distribution of information which is the subject of this chapter on a
6	"need to know" basis, and § 40.1-5-26.
7	(6) The office may, in response to a complaint that is provided in written form to the
8	review agent, review an appeal regarding any adverse determination, and may request
9	information of the review agent, provider or patient regarding the status, outcome or rationale
10	regarding the decision.
11	(d) The review agents clinical criteria used in making its utilization review decisions shall
12	comply with the following:
13	(i) The requirement that each review agent shall provide its clinical criteria as required
14	<u>by law;</u>
15	(ii) Written clinical screening criteria and review procedures are established according to
16	nationally accepted standards and protocols that are periodically evaluated and updated; and
17	(iii) Establish a process to incorporate and consider local variations to national standard
18	identified in § 27-18.9-8(d)(ii) above to include input from local participating providers.
18 19	identified in § 27-18.9-8(d)(ii) above to include input from local participating providers. (1) The screening criteria and review procedures must comply with the requirements set
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<ol> <li>19</li> <li>20</li> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> <li>28</li> <li>29</li> </ol>	<ul> <li>(1) The screening criteria and review procedures must comply with the requirements set forth in § 27-18.9-8 (d) and must meet the satisfaction of the commissioner.</li> <li>27-18.9-9. External appeal requirments (a) In cases where the second level of appeal to reverse an adverse determination is unsuccessful, the review agent shall provide for an external appeal by an unrelated and objective appeal agency, selected by the commissioner. The commissioner shall promulgate rules and regulations including, but not limited to, criteria for designation, operation, policy, oversight, and termination of designation as an external appeal agency. The external appeal agency shall not be required to be certified under this chapter for activities conducted pursuant to its designation.</li> <li>(b) The external appeal shall have the following characteristics:         <ul> <li>(1) The external appeal review and decision shall be based on the medical necessity for</li> </ul> </li> </ul>
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- (3) The neutral physician, dentist, or other practitioner may confer either directly with the
   review agent and provider, or with physicians or dentists appointed to represent them.
- 2 review agent and provider, or with physicians of dentists appointed to represent them.
- 3 (4) Payment for the appeal fee must not exceed twenty-five dollars (\$25.00). It must be
- 4 refunded to the claimant if the adverse benefit determination (or final internal adverse benefit
  5 determination) is reversed through external review. The fee must be waived if payment of the fee
- 6 would impose an undue financial hardship. In addition, the annual limit on the filing fees for any
- 7 <u>claimant within a single plan year (in the individual market, policy year) must not exceed</u>
- 8 seventy-five dollars (\$75.00). Notwithstanding the aforementioned, this subsection shall not
- 9 apply to excepted benefits as defined in 42 U.S.C. 300 gg-91(c).
- 10 (5) The decision of the external appeal agency shall be binding; however, any person who
- 11 is aggrieved by a final decision of the external appeal agency is entitled to judicial review in a
- 12 <u>court of competent jurisdiction.</u>
- 13 **<u>27-18.9-10. Reporting requirements. --</u>** The office, in consultation with the department
- 14 of health, shall establish reporting requirements to determine if the utilization review programs
- 15 are in compliance with the provisions of this chapter and applicable regulations.
- <u>27-18.9-11. Lists. --</u> The commissioner shall periodically provide a list of private review
   agents issued certificates and the renewal date for those certificates to all licensed health care
- 18 <u>facilities and any other individual or organization requesting the list.</u>
- 27-18.9-12. Penalties. -- A person who substantially violates any provision of this
   chapter or any regulation adopted under this chapter or who submits any false information in an
   application required by this chapter is guilty of a misdemeanor and on conviction is subject to a
- 22 penalty not exceeding five thousand dollars (\$5,000).
- 23 <u>27-18.9-13. Fees. --</u> The proceeds of any monetary penalties and fines collected pursuant
- 24 to the provisions of this chapter shall be deposited as general revenues.
- 25 <u>27-18.9-14. Severability. --</u> If any provision of this chapter or the application of any
- 26 provision to any person or circumstance shall be held invalid, that invalidity shall not affect the
- 27 provisions or application of this chapter which can be given effect without the invalid provision
- 28 or application, and to this end the provisions of this chapter are declared to be severable.
- 29

SECTION 6. This act shall take effect on January 1, 2017.

LC005936/SUB A

#### **EXPLANATION**

#### BY THE LEGISLATIVE COUNCIL

#### OF

# AN ACT

# RELATING TO HEALTH AND SAFETY - UTILIZATION REVIEW

\*\*\*

1 This act would transfer Utilization Review (UR) from the department of health 2 (HEALTH) to the office of the health insurance commissioner (OHIC). In addition, changes 3 would be made to the "Health Care Accessibility and Quality Assurance Act" and the "Health 4 Plan Modification Act" to comply with the "Affordable Care Act" to reflect such transfer. 5 This act would take effect on January 1, 2017.

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