

SECOND REGULAR SESSION

# SENATE BILL NO. 982

99TH GENERAL ASSEMBLY

INTRODUCED BY SENATOR WIELAND.

Read 1st time February 1, 2018, and ordered printed.

ADRIANE D. CROUSE, Secretary.

6265S.011

## AN ACT

To repeal section 376.1350, RSMo, and to enact in lieu thereof one new section relating to the protection of persons with emergency medical conditions.

*Be it enacted by the General Assembly of the State of Missouri, as follows:*

Section A. Section 376.1350, RSMo, is repealed and one new section  
2 enacted in lieu thereof, to be known as section 376.1350, to read as follows:

376.1350. For purposes of sections 376.1350 to 376.1390, the following  
2 terms mean:

3 (1) "Adverse determination", a determination by a health carrier or its  
4 designee utilization review organization that an admission, availability of care,  
5 continued stay or other health care service has been reviewed and, based upon  
6 the information provided, does not meet the health carrier's requirements for  
7 medical necessity, appropriateness, health care setting, level of care or  
8 effectiveness, and the payment for the requested service is therefore denied,  
9 reduced or terminated;

10 (2) "Ambulatory review", utilization review of health care services  
11 performed or provided in an outpatient setting;

12 (3) "Case management", a coordinated set of activities conducted for  
13 individual patient management of serious, complicated, protracted or other health  
14 conditions;

15 (4) "Certification", a determination by a health carrier or its designee  
16 utilization review organization that an admission, availability of care, continued  
17 stay or other health care service has been reviewed and, based on the information  
18 provided, satisfies the health carrier's requirements for medical necessity,  
19 appropriateness, health care setting, level of care and effectiveness;

20 (5) "Clinical peer", a physician or other health care professional who holds

21 a nonrestricted license in a state of the United States and in the same or similar  
22 specialty as typically manages the medical condition, procedure or treatment  
23 under review;

24 (6) "Clinical review criteria", the written screening procedures, decision  
25 abstracts, clinical protocols and practice guidelines used by the health carrier to  
26 determine the necessity and appropriateness of health care services;

27 (7) "Concurrent review", utilization review conducted during a patient's  
28 hospital stay or course of treatment;

29 (8) "Covered benefit" or "benefit", a health care service that an enrollee  
30 is entitled under the terms of a health benefit plan;

31 (9) "Director", the director of the department of insurance, financial  
32 institutions and professional registration;

33 (10) "Discharge planning", the formal process for determining, prior to  
34 discharge from a facility, the coordination and management of the care that a  
35 patient receives following discharge from a facility;

36 (11) "Drug", any substance prescribed by a licensed health care provider  
37 acting within the scope of the provider's license and that is intended for use in  
38 the diagnosis, mitigation, treatment or prevention of disease. The term includes  
39 only those substances that are approved by the FDA for at least one indication;

40 (12) "Emergency medical condition", the sudden and, at the time,  
41 unexpected onset of a health condition that manifests itself by symptoms of  
42 sufficient severity, **regardless of the final diagnosis that is given**, that  
43 would lead a prudent lay person, possessing an average knowledge of medicine  
44 and health, to believe that immediate medical care is required, which may  
45 include, but shall not be limited to:

46 (a) Placing the person's health in significant jeopardy;

47 (b) Serious impairment to a bodily function;

48 (c) Serious dysfunction of any bodily organ or part;

49 (d) Inadequately controlled pain; or

50 (e) With respect to a pregnant woman who is having contractions:

51 a. That there is inadequate time to effect a safe transfer to another  
52 hospital before delivery; or

53 b. That transfer to another hospital may pose a threat to the health or  
54 safety of the woman or unborn child;

55 (13) "Emergency service", a health care item or service furnished or  
56 required to evaluate and treat an emergency medical condition, which may

57 include, but shall not be limited to, health care services that are provided in a  
58 licensed hospital's emergency facility by an appropriate provider;

59 (14) "Enrollee", a policyholder, subscriber, covered person or other  
60 individual participating in a health benefit plan;

61 (15) "FDA", the federal Food and Drug Administration;

62 (16) "Facility", an institution providing health care services or a health  
63 care setting, including but not limited to hospitals and other licensed inpatient  
64 centers, ambulatory surgical or treatment centers, skilled nursing centers,  
65 residential treatment centers, diagnostic, laboratory and imaging centers, and  
66 rehabilitation and other therapeutic health settings;

67 (17) "Grievance", a written complaint submitted by or on behalf of an  
68 enrollee regarding the:

69 (a) Availability, delivery or quality of health care services, including a  
70 complaint regarding an adverse determination made pursuant to utilization  
71 review;

72 (b) Claims payment, handling or reimbursement for health care services;  
73 or

74 (c) Matters pertaining to the contractual relationship between an enrollee  
75 and a health carrier;

76 (18) "Health benefit plan", a policy, contract, certificate or agreement  
77 entered into, offered or issued by a health carrier to provide, deliver, arrange for,  
78 pay for, or reimburse any of the costs of health care services; except that, health  
79 benefit plan shall not include any coverage pursuant to liability insurance policy,  
80 workers' compensation insurance policy, or medical payments insurance issued  
81 as a supplement to a liability policy;

82 (19) "Health care professional", a physician or other health care  
83 practitioner licensed, accredited or certified by the state of Missouri to perform  
84 specified health services consistent with state law;

85 (20) "Health care provider" or "provider", a health care professional or a  
86 facility;

87 (21) "Health care service", a service for the diagnosis, prevention,  
88 treatment, cure or relief of a health condition, illness, injury or disease;

89 (22) "Health carrier", an entity subject to the insurance laws and  
90 regulations of this state that contracts or offers to contract to provide, deliver,  
91 arrange for, pay for or reimburse any of the costs of health care services,  
92 including a sickness and accident insurance company, a health maintenance

93 organization, a nonprofit hospital and health service corporation, or any other  
94 entity providing a plan of health insurance, health benefits or health services;  
95 except that such plan shall not include any coverage pursuant to a liability  
96 insurance policy, workers' compensation insurance policy, or medical payments  
97 insurance issued as a supplement to a liability policy;

98 (23) "Health indemnity plan", a health benefit plan that is not a managed  
99 care plan;

100 (24) "Managed care plan", a health benefit plan that either requires an  
101 enrollee to use, or creates incentives, including financial incentives, for an  
102 enrollee to use, health care providers managed, owned, under contract with or  
103 employed by the health carrier;

104 (25) "Participating provider", a provider who, under a contract with the  
105 health carrier or with its contractor or subcontractor, has agreed to provide  
106 health care services to enrollees with an expectation of receiving payment, other  
107 than coinsurance, co-payments or deductibles, directly or indirectly from the  
108 health carrier;

109 (26) "Peer-reviewed medical literature", a published scientific study in a  
110 journal or other publication in which original manuscripts have been published  
111 only after having been critically reviewed for scientific accuracy, validity and  
112 reliability by unbiased independent experts, and that has been determined by the  
113 International Committee of Medical Journal Editors to have met the uniform  
114 requirements for manuscripts submitted to biomedical journals or is published in  
115 a journal specified by the United States Department of Health and Human  
116 Services pursuant to Section 1861(t)(2)(B) of the Social Security Act, as amended,  
117 as acceptable peer-reviewed medical literature. Peer-reviewed medical literature  
118 shall not include publications or supplements to publications that are sponsored  
119 to a significant extent by a pharmaceutical manufacturing company or health  
120 carrier;

121 (27) "Person", an individual, a corporation, a partnership, an association,  
122 a joint venture, a joint stock company, a trust, an unincorporated organization,  
123 any similar entity or any combination of the foregoing;

124 (28) "Prospective review", utilization review conducted prior to an  
125 admission or a course of treatment;

126 (29) "Retrospective review", utilization review of medical necessity that  
127 is conducted after services have been provided to a patient, but does not include  
128 the review of a claim that is limited to an evaluation of reimbursement levels,

129 veracity of documentation, accuracy of coding or adjudication for payment;

130 (30) "Second opinion", an opportunity or requirement to obtain a clinical  
131 evaluation by a provider other than the one originally making a recommendation  
132 for a proposed health service to assess the clinical necessity and appropriateness  
133 of the initial proposed health service;

134 (31) "Stabilize", with respect to an emergency medical condition, that no  
135 material deterioration of the condition is likely to result or occur before an  
136 individual may be transferred;

137 (32) "Standard reference compendia":

138 (a) The American Hospital Formulary Service-Drug Information; or

139 (b) The United States Pharmacopoeia-Drug Information;

140 (33) "Utilization review", a set of formal techniques designed to monitor  
141 the use of, or evaluate the clinical necessity, appropriateness, efficacy, or  
142 efficiency of, health care services, procedures, or settings. Techniques may  
143 include ambulatory review, prospective review, second opinion, certification,  
144 concurrent review, case management, discharge planning or retrospective  
145 review. Utilization review shall not include elective requests for clarification of  
146 coverage;

147 (34) "Utilization review organization", a utilization review agent as  
148 defined in section 374.500.

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