

ENAC	.S:
	58-85-101, Utah Code Annotated 1953
	58-85-102, Utah Code Annotated 1953
	58-85-103 , Utah Code Annotated 1953
	58-85-104, Utah Code Annotated 1953
Be it en	acted by the Legislature of the state of Utah:
	Section 1. Section 58-67-501 is amended to read:
	58-67-501. Unlawful conduct.
	(1) "Unlawful conduct" includes, in addition to the definition in Section 58-1-501:
	(a) buying, selling, or fraudulently obtaining, any medical diploma, license, certificate,
or regis	tration;
	(b) aiding or abetting the buying, selling, or fraudulently obtaining of any medical
diploma	a, license, certificate, or registration;
	(c) substantially interfering with a licensee's lawful and competent practice of medicine
in accor	rdance with this chapter by:
	(i) any person or entity that manages, owns, operates, or conducts a business having a
direct o	r indirect financial interest in the licensee's professional practice; or
	(ii) anyone other than another physician licensed under this title, who is engaged in
direct c	linical care or consultation with the licensee in accordance with the standards and ethics
of the p	rofession of medicine; or
	(d) entering into a contract that limits a licensee's ability to advise the licensee's
patients	fully about treatment options or other issues that affect the health care of the licensee's
patients	
	(2) "Unlawful conduct" does not include:
	(a) establishing, administering, or enforcing the provisions of a policy of accident and
health i	nsurance by an insurer doing business in this state in accordance with Title 31A,
Insuran	ce Code;
	(b) adopting, implementing, or enforcing utilization management standards related to
paymen	t for a licensee's services, provided that:
	(i) utilization management standards adopted, implemented, and enforced by the payer

have been approved by a physician or by a committee that contains one or more physicians; and

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- (ii) the utilization management standards does not preclude a licensee from exercising independent professional judgment on behalf of the licensee's patients in a manner that is independent of payment considerations;
- (c) developing and implementing clinical practice standards that are intended to reduce morbidity and mortality or developing and implementing other medical or surgical practice standards related to the standardization of effective health care practices, provided that:
- (i) the practice standards and recommendations have been approved by a physician or by a committee that contains one or more physicians; and
- (ii) the practice standards do not preclude a licensee from exercising independent professional judgment on behalf of the licensee's patients in a manner that is independent of payment considerations;
 - (d) requesting or recommending that a patient obtain a second opinion from a licensee;
- (e) conducting peer review, quality evaluation, quality improvement, risk management, or similar activities designed to identify and address practice deficiencies with health care providers, health care facilities, or the delivery of health care;
- (f) providing employment supervision or adopting employment requirements that do not interfere with the licensee's ability to exercise independent professional judgment on behalf of the licensee's patients, provided that employment requirements that may not be considered to interfere with an employed licensee's exercise of independent professional judgment include:
- (i) an employment requirement that restricts the licensee's access to patients with whom the licensee's employer does not have a contractual relationship, either directly or through contracts with one or more third-party payers; or
- (ii) providing compensation incentives that are not related to the treatment of any particular patient;
- (g) providing benefit coverage information, giving advice, or expressing opinions to a patient or to a family member of a patient to assist the patient or family member in making a decision about health care that has been recommended by a licensee; [or]
 - (h) in compliance with Section 58-85-103:
 - (i) obtaining an investigational drug or investigational device;
 - (ii) administering the investigational drug to an eligible patient; or

90	(iii) treating an eligible patient with the investigational drug or investigational device;
91	<u>or</u>
92	[(h)] (i) any otherwise lawful conduct that does not substantially interfere with the
93	licensee's ability to exercise independent professional judgment on behalf of the licensee's
94	patients and that does not constitute the practice of medicine as defined in this chapter.
95	Section 2. Section 58-67-502 is amended to read:
96	58-67-502. Unprofessional conduct.
97	(1) "Unprofessional conduct" includes, in addition to the definition in Section
98	58-1-501:
99	[(1)] (a) using or employing the services of any individual to assist a licensee in any
100	manner not in accordance with the generally recognized practices, standards, or ethics of the
101	profession, state law, or division rule;
102	[(2)] (b) making a material misrepresentation regarding the qualifications for licensure
103	under Section 58-67-302.7; or
104	[(3)] <u>(c)</u> violating the dispensing requirements of Section 58-17b-309 or Chapter 17b,
105	Part 8, Dispensing Medical Practitioner and Dispensing Medical Practitioner Clinic Pharmacy,
106	if applicable.
107	(2) "Unprofessional conduct" does not include, in compliance with Section 58-85-103:
108	(a) obtaining an investigational drug or investigational device;
109	(b) administering the investigational drug to an eligible patient; or
110	(c) treating an eligible patient with the investigational drug or investigational device.
111	Section 3. Section 58-68-501 is amended to read:
112	58-68-501. Unlawful conduct.
113	(1) "Unlawful conduct" includes, in addition to the definition in Section 58-1-501:
114	(a) buying, selling, or fraudulently obtaining any osteopathic medical diploma, license,
115	certificate, or registration; and
116	(b) aiding or abetting the buying, selling, or fraudulently obtaining of any osteopathic
117	medical diploma, license, certificate, or registration;
118	(c) substantially interfering with a licensee's lawful and competent practice of medicine
119	in accordance with this chapter by:
120	(i) any person or entity that manages, owns, operates, or conducts a business having a

direct or indirect financial interest in the licensee's professional practice; or

- (ii) anyone other than another physician licensed under this title, who is engaged in direct clinical care or consultation with the licensee in accordance with the standards and ethics of the profession of medicine; or
- (d) entering into a contract that limits a licensee's ability to advise the licensee's patients fully about treatment options or other issues that affect the health care of the licensee's patients.
 - (2) "Unlawful conduct" does not include:
- (a) establishing, administering, or enforcing the provisions of a policy of accident and health insurance by an insurer doing business in this state in accordance with Title 31A, Insurance Code;
- (b) adopting, implementing, or enforcing utilization management standards related to payment for a licensee's services, provided that:
- (i) utilization management standards adopted, implemented, and enforced by the payer have been approved by a physician or by a committee that contains one or more physicians; and
- (ii) the utilization management standards does not preclude a licensee from exercising independent professional judgment on behalf of the licensee's patients in a manner that is independent of payment considerations;
- (c) developing and implementing clinical practice standards that are intended to reduce morbidity and mortality or developing and implementing other medical or surgical practice standards related to the standardization of effective health care practices, provided that:
- (i) the practice standards and recommendations have been approved by a physician or by a committee that contains one or more physicians; and
- (ii) the practice standards do not preclude a licensee from exercising independent professional judgment on behalf of the licensee's patients in a manner that is independent of payment considerations;
 - (d) requesting or recommending that a patient obtain a second opinion from a licensee;
- (e) conducting peer review, quality evaluation, quality improvement, risk management, or similar activities designed to identify and address practice deficiencies with health care providers, health care facilities, or the delivery of health care;
 - (f) providing employment supervision or adopting employment requirements that do

152	not interfere with the licensee's ability to exercise independent professional judgment on behalf
153	of the licensee's patients, provided that employment requirements that may not be considered to
154	interfere with an employed licensee's exercise of independent professional judgment include:
155	(i) an employment requirement that restricts the licensee's access to patients with
156	whom the licensee's employer does not have a contractual relationship, either directly or
157	through contracts with one or more third-party payers; or
158	(ii) providing compensation incentives that are not related to the treatment of any
159	particular patient;
160	(g) providing benefit coverage information, giving advice, or expressing opinions to a
161	patient or to a family member of a patient to assist the patient or family member in making a
162	decision about health care that has been recommended by a licensee; [or]
163	(h) in compliance with Section 58-85-103:
164	(i) obtaining an investigational drug or investigational device;
165	(ii) administering the investigational drug to an eligible patient; or
166	(iii) treating an eligible patient with the investigational drug or investigational device;
167	<u>or</u>
168	[(h)] (i) any otherwise lawful conduct that does not substantially interfere with the
169	licensee's ability to exercise independent professional judgment on behalf of the licensee's
170	patients and that does not constitute the practice of medicine as defined in this chapter.
171	Section 4. Section 58-68-502 is amended to read:
172	58-68-502. Unprofessional conduct.
173	(1) "Unprofessional conduct" includes, in addition to the definition in Section
174	58-1-501:
175	[(1)] (a) using or employing the services of any individual to assist a licensee in any
176	manner not in accordance with the generally recognized practices, standards, or ethics of the
177	profession, state law, or division rule; or
178	[(2)] (b) violating the dispensing requirements of Section 58-17b-309 or Chapter 17b,
179	Part 8, Dispensing Medical Practitioner and Dispensing Medical Practitioner Clinic Pharmacy,
180	if applicable.
181	(2) "Unprofessional conduct" does not include, in compliance with Section 58-85-103:
182	(a) obtaining an investigational drug or investigational device;

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183	(b) administering the investigational drug to an eligible patient; or
184	(c) treating an eligible patient with the investigational drug or investigational device.
185	Section 5. Section 58-85-101 is enacted to read:
186	CHAPTER 85. UTAH RIGHT TO TRY ACT
187	<u>58-85-101.</u> Title.
188	This chapter is known as the "Utah Right to Try Act."
189	Section 6. Section 58-85-102 is enacted to read:
190	<u>58-85-102.</u> Definitions.
191	As used in this chapter:
192	(1) "Eligible patient" means an individual who has been diagnosed with a terminal
193	illness by a physician.
194	(2) "Physician" means an individual who is licensed under:
195	(a) Title 58, Chapter 67, Utah Medical Practice Act; or
196	(b) Title 58, Chapter 68, Utah Osteopathic Medical Practice Act.
197	(3) "Insurer" means the same as that term is defined in Section 31A-1-301.
198	(4) "Investigational device" means a device that:
199	(a) meets the definition of "investigational device" in 21 C.F.R. Sec. 812.3; and
200	(b) has successfully completed the United States Food and Drug Administration Phase
201	1 testing for an investigational device described in 21 C.F.R. Part 812.
202	(5) "Investigational drug" means a drug that:
203	(a) meets the definition of "investigational new drug" in 21 C.F.R. Sec. 312.3; and
204	(b) has successfully completed the United States Food and Drug Administration Phase
205	1 testing for an investigational new drug described in 21 C.F.R. Part 312.
206	(6) "Terminal illness" means a condition of a patient that:
207	(a) is serious or life-threatening;
208	(b) as determined by a physician, is likely to pose a greater risk to the patient than the
209	risk posed to the patient by treatment with an investigational drug or investigational device;
210	<u>and</u>
211	(c) presents the patient with no treatment option that is satisfactory or comparable to
212	treatment with an investigational drug or device.
213	Section 7. Section 58-85-103 is enacted to read:

214	58-85-103. Right to request investigational drug or device.
215	(1) An eligible patient may obtain an investigational drug through an agreement with
216	the investigational drug's manufacturer and the eligible patient's physician that provides:
217	(a) for the transfer of the investigational drug from the manufacturer to the physician;
218	and
219	(b) that the physician will administer the investigational drug to the patient.
220	(2) An eligible patient may obtain an investigational device through an agreement with
221	the investigational device's manufacturer and the eligible patient's physician that provides:
222	(a) for the transfer of the investigational device from the manufacturer to the physician
223	and
224	(b) that the physician will use the investigational device to treat the patient.
225	Section 8. Section 58-85-104 is enacted to read:
226	58-85-104. Insurance coverage No right of action.
227	This chapter does not:
228	(1) require an insurer to cover the cost of:
229	(a) administering an investigational drug under this chapter; or
230	(b) treating a patient with an investigational device under this chapter;
231	(2) prohibit an insurer from covering the cost of:
232	(a) administering an investigational drug under this chapter; or
233	(b) treating a patient with an investigational device under this chapter;
234	(3) require a manufacturer of an investigational drug or investigational device to agree
235	to make an investigational drug or investigational device available to an eligible patient or an
236	eligible patient's physician;
237	(4) require a physician to agree to:
238	(a) administer an investigational drug to an eligible patient under this chapter; or
239	(b) treat an eligible patient with an investigational device under this chapter; or
240	(5) create a private right of action for any harm done to an eligible patient:
241	(a) resulting from the eligible patient's use of an investigational drug or investigational
242	device, against:
243	(i) a manufacturer of an investigational drug or investigational device under this
244	chapter;

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245	(ii) a physician who administers an investigational drug or treats an eligible patient
246	with an investigational device under this chapter; or
247	(iii) a hospital where a physician administers an investigational drug to an eligible
248	patient or treats an eligible patient with an investigational device under this chapter;
249	(b) against a physician or hospital, for the physician's or hospital's refusal to:
250	(i) administer an investigational drug to an eligible patient under this chapter; or
251	(ii) treat an eligible patient with an investigational device under this chapter; or
252	(c) against a manufacturer, for the manufacturer's refusal to provide an eligible patient
253	with an investigational drug or an investigational device under this chapter.

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Office of Legislative Research and General Counsel