28	(1) As used in this section:
29	(a) "Health care provider" means the same as that term is defined in Section
30	<u>78B-3-403.</u>
31	(b) "Human cells, tissues, or cellular or tissue-based products" has the same meaning
32	as in 21 C.F.R. Sec. 1271.3 as it exists on May 1, 2024.
33	(c) (i) "Stem cell therapy" means a treatment involving the use of afterbirth placental
34	perinatal stem cells or human cells, tissues, or cellular or tissue-based products.
35	(ii) "Stem cell therapy" does not include treatment or research using human cells or
36	tissues that were derived from a fetus or embryo after an abortion.
37	(2) A health care provider whose scope of practice includes the use of stem cell therapy
38	\$→ [and] ←\$ may perform a stem cell therapy that is not approved by the United States Food and
38a	<u>Drug</u>
39	Administration, if the health care provider provides the patient with the following written
40	notice before performing the therapy:
41	"THIS NOTICE MUST BE PROVIDED TO YOU UNDER UTAH LAW. This health
42	care practitioner performs one or more stem cell therapies that have not yet been approved by
43	the United States Food and Drug Administration. You are encouraged to consult with your
44	primary care provider before undergoing a stem cell therapy."
45	(3) (a) The written notice described in Subsection (2) shall be:
46	(i) on paper that is at least eight and one-half inches by eleven inches; and
47	(ii) written in no less than forty point type.
48	(b) The health care provider shall prominently display the written notice in the entrance
49	and in an area visible to patients in the health care provider's office.
50	(4) (a) A health care provider who is required to provide written notice under
51	Subsection (2) shall obtain a signed consent form before performing the therapy.
52	(b) The consent form shall:
53	(i) be signed by the patient, or, if the patient is legally not competent, the patient's
54	representative; and
55	(ii) state, in language the patient could reasonably be expected to understand:
56	(A) the nature and character of the proposed treatment, including the treatment's United
57	States Food and Drug Administration approval status;
58	(B) the anticipated results of the proposed treatment;