COUNSEL

- 1.1 Senator moves to amend S.F. No. 73 as follows:
- 1.2 Page 10, line 1, delete "(a)"
- 1.3 Page 10, delete line 15
- 1.4 Page 11, line 23, delete everything after "cancer" and insert a semicolon
- 1.5 Page 11, delete lines 24 to 26
- 1.6 Page 12, line 12, delete everything after "<u>illness</u>" and insert "<u>; or</u>"
- 1.7 Page 12, delete lines 13 to 16
- 1.8 Page 18, line 19, delete everything after "one" and insert "who is a patient or caregiver
- 1.9 of a patient in the medical cannabis program;"
- 1.10 Page 18, delete line 20
- 1.11 Page 91, line 7, delete "<u>valid for one year</u>" and insert "<u>permanent</u>"
- 1.12 Page 91, delete lines 8 and 9
- 1.13 Page 96, line 28, after the semicolon, insert "<u>or</u>"
- 1.14 Page 96, line 29, delete "<u>or</u>"
- 1.15 Page 96, delete line 30
- 1.16 Page 97, line 18, after "products" insert "on behalf of the patient" and delete the first
- 1.17 comma and insert "; that a patient store the patient's supply of medical cannabis flower or
- 1.18 medicinal cannabinoid products in a locked container accessible only to the patient, the
- 1.19 patient's designated caregiver, or the patient's parent, legal guardian, or spouse;" and delete
- 1.20 "<u>, and</u>" and insert "<u>; and</u>"

1.21 Page 97, line 20, after the period, insert "<u>Nothing in this subdivision requires facilities</u> 1.22 and providers listed in this subdivision to adopt such restrictions."

- 1.23 Page 97, after line 20, insert:
- 1.24 "(b) No facility or provider listed in this subdivision may unreasonably limit a patient's
- 1.25 access to or use of medical cannabis flower or medical cannabiniod products to the extent
- 1.26 that such use is authorized under sections 342.42 to 342.56. No facility or provider listed
- 1.27 in this subdivision may prohibit a patient access to or use of medical cannabis flower or
- 1.28 medical cannabinoid products due solely to the fact that cannabis is a Schedule I drug
- 1.29 pursuant to the federal Uniform Controlled Substances Act. If a federal regulatory agency,
- 1.30 the United States Department of Justice, or the federal Centers for Medicare and Medicaid

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2.1	Services takes one of the following actions, a facility or provider may suspend compliance
2.2	with this paragraph until the regulatory agency, the United States Department of Justice, or
2.3	the federal Centers for Medicare and Medicaid Services notifies the facility or provider that
2.4	it may resume permitting the use of medical cannabis flower or medical cannabinoid products
2.5	within the facility or in the provider's service setting:
2.6	(1) a federal regulatory agency or the United States Department of Justice initiates
2.7	enforcement action against a facility or provider related to the facility's compliance with
2.8	the medical cannabis program; or
2.9	(2) a federal regulatory agency, the United States Department of Justice, or the federal
2.10	Centers for Medicare and Medicaid Services issues a rule or otherwise provides notification
2.11	to the facility or provider that expressly prohibits the use of medical cannabis in health care
2.12	facilities or otherwise prohibits compliance with the medical cannabis program."
2.13	Page 97, line 21, delete "(b)" and insert "(c)"
2.14	Page 97, line 27, delete everything after the period
2.15	Page 97, delete lines 28 to 31
2.16	Page 97, before line 32, insert:
2.17	"Subd. 3. Child care facilities. A proprietor of a family or group family day care program
2.18	must disclose to parents or guardians of children cared for on the premises of the family or
2.19	group family day care program, if the proprietor permits the smoking or use of medical
2.20	cannabis on the premises, outside of its hours of operation. Disclosure must include posting

2.21 <u>on the premises a conspicuous written notice and orally informing parents or guardians.</u>"