(Original Signature of Member)	

118TH CONGRESS 2D SESSION

H.R.

To clarify that States do not have authority to establish or continue in effect any requirement with respect to the sale, distribution, possession, or use of less harmful alternatives to traditional tobacco products to protect public health, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

Mr.	Bergman introduced	the	following	bill;	which	was	referred	to	the
	Committee on _								

A BILL

To clarify that States do not have authority to establish or continue in effect any requirement with respect to the sale, distribution, possession, or use of less harmful alternatives to traditional tobacco products to protect public health, and for other purposes.

- 1 Be it enacted by the Senate and House of Representa-
- 2 tives of the United States of America in Congress assembled,
- 3 SECTION 1. SHORT TITLE.
- 4 This Act may be cited as the "Public Options for Un-
- 5 restricted Consumer Harm-reduction Act of 2024" or the
- 6 "POUCH Act of 2024".

1	SEC. 2. PREEMPTION OF STATE LAWS ON TOBACCO PROD-
2	UCTS FOR WHICH A MARKETING ORDER IS IN
3	EFFECT.
4	(a) In General.—Paragraph (2) of section 916(a)
5	of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
6	387p(a)) is amended to read as follows:
7	"(2) Preemption of Certain State and
8	LOCAL REQUIREMENTS.—
9	"(A) In General.—
10	"(i) Preemption.—No State or polit-
11	ical subdivision of a State may establish or
12	continue in effect with respect to a tobacco
13	product any requirement which is different
14	from, or in addition to, any requirement
15	under the provisions of this chapter relat-
16	ing to tobacco product standards, pre-
17	market review, adulteration, misbranding,
18	labeling, registration, good manufacturing
19	standards, or modified risk tobacco prod-
20	ucts.
21	"(ii) Exception.—Clause (i) does not
22	apply to requirements relating to the sale
23	of, distribution of, possession of, informa-
24	tion reporting to the State, exposure to,
25	access to, the advertising and promotion
26	of, or use of, tobacco products to or by in-

1	dividuals of any age, or relating to fire
2	safety standards for tobacco products. In-
3	formation disclosed to a State under clause
4	(i) that is exempt from disclosure under
5	section 552(b)(4) of title 5, United States
6	Code, shall be treated as a trade secret
7	and confidential information by the State.
8	"(B) Tobacco products for which a
9	MARKETING ORDER IS IN EFFECT.—No State
10	or political subdivision of a State may establish
11	or continue in effect, with respect to tobacco
12	products for which there is a marketing order
13	issued under section 910(c)(1)(A)(i) in effect,
14	any prohibition or restriction on the sale of, dis-
15	tribution of, possession of, exposure to, access
16	to, advertising and promotion of, or use of such
17	tobacco products that is different from, or in
18	addition to, any prohibition or restriction on
19	such tobacco products under the provisions of
20	this chapter.".
21	(b) Conforming Changes.—Section 916(a)(1) of
22	the Federal Food, Drug, and Cosmetic Act (21 U.S.C.
23	387p(a)(2)(A)) is amended by striking "the sale, distribu-
24	tion, possession, exposure to, access to, advertising and
25	promotion of, or use of tobacco products by individuals"

and inserting "the sale of, distribution of, possession of,
exposure to, access to, advertising and promotion of, or
use of tobacco products to or by individuals".
SEC. 3. REPORT ON PENDING TOBACCO PRODUCT APPLI-
CATIONS.
Not later than the date that is 90 days after the date
of enactment of this Act, the Commissioner of Food and
Drugs shall submit to Congress a report containing a list
specifying—
(1) each pending new tobacco product applica-
tion under section 910 of the Federal Food, Drug,
and Cosmetic Act (21 U.S.C. 387j);
(2) each pending application for a marketing
authorization order for a modified risk tobacco prod-
uct under section 911 of such Act (21 U.S.C. 387k);
and
and (3) the status of each pending application re-