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		L.D. 1618
	2	DATE: 6/9/5 (Filing No. H-675)
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	6	Reproduced and distributed under the direction of the Clerk of the House.
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		STATE OF MAINE
1	10	HOUSE OF REPRESENTATIVES 122ND LEGISLATURE
נ	12	FIRST SPECIAL SESSION
	• •	
Ŀ	14	HOUSE AMENDMENT "A" to COMMITTEE AMENDMENT "A" to H.P.
1	16	1141, L.D. 1618, Bill, "An Act Regarding Advertising by Drug Manufacturers"
1	18	Manulacculers
	-	Amend the amendment by inserting after the title the
2	20	following:
. 2	22	'Amend the bill by striking out the title and substituting the following:
2	24	
2	26	'An Act Regarding Advertising by Drug Manufacturers and Disclosure of Clinical Trials''
2	28	Further amend the amendment in the first paragraph after the
-		title in the first line (page 1, line 22 in amendment) by
3	30	striking out the following: "Amend" and inserting in its place
		the following: 'Further amend'
3	32	
	2.4	Further amend the amendment in section 1 in that part
3	34	designated " $$2700-A."$ in subsection 3 by striking out all of the first indented paragraph (page 2, lines 27 to 36 in amendment)
3	36	and inserting in its place the following:
3	38	3. <u>Disclosure of clinical trials of prescription drugs.</u>
		Beginning October 15, 2005, a manufacturer or labeler of
4	40	prescription drugs that is required to report marketing costs for prescription drugs pursuant to section 2698-A shall post, with
4	42	regard to those prescription drugs, on the publicly accessible
		Internet website of the federal National Institutes of Health or
4	44	its successor agency or another publicly accessible website the
		following information concerning any clinical trial that the
4	46	manufacturer conducted or sponsored on or after October 15, 2002;'

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## HOUSE AMENDMENT

HOUSE AMENDMENT "A" to COMMITTEE AMENDMENT "A" to H.P. 1141, L.D. 1618

## SUMMARY

4 amendment removes the provision that requires a This manufacturer of prescription drugs to disclose certain information regarding clinical trials of prescription drugs 6 before regulated advertisements for those prescription drugs may 8 be presented and replaces it with a requirement that a manufacturer or labeler of prescription drugs that is required to report marketing costs for prescription drugs pursuant to the 10 Maine Revised Statutes, Title 22, section 2698-A disclose certain information regarding clinical trials of prescription drugs. 12

14 16 18 SPONSORED BY: 20 (Representative LER

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