JAN 18 2008

### A BILL FOR AN ACT

RELATING TO FOOD.

#### BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF HAWAII:

- 1 SECTION 1. The legislature finds it is imperative for the
- 2 public health, safety and welfare to declare that aspartame and
- 3 its derivative compounds, in all of their trade names, are
- 4 poisonous and deleterious food additives due to their neurotoxic
- 5 and carcinogenic metabolites.
- 6 The legislature finds that federal authorities have not
- 7 intended to or expressed an intention to occupy and preempt
- 8 areas of concern regarding the prohibition of toxic, neurotoxic,
- 9 carcinogenic, poisonous or deleterious food additives, and
- 10 therefore the legislature may prohibit the sale of products
- 11 containing aspartame and its derivative compounds in order to
- 12 protect and ensure the public health, safety and welfare.
- 13 SECTION 2. Section 328-1, Hawaii Revised Statutes, is
- 14 amended by adding a new definition to be appropriately inserted
- 15 and to read as follows:
- 16 ""Aspartame" means the artificial sweetener with the
- 17 technical name L-aspartyl-L-phenylalanine methyl ester."

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1	SECT	'ION 3. Section 328-6, Hawaii Revised Statutes, is
2	amended t	o read as follows:
3	"§32	8-6 Prohibited acts. The following acts and the
4	causing t	hereof within the State by any person are prohibited:
5	(1)	The manufacture, sale, delivery, holding, or offering
6		for sale of any food, drug, device, or cosmetic that
7		is adulterated or misbranded;
8	(2)	The adulteration or misbranding of any food, drug,
9		device, or cosmetic;
10	(3)	The receipt in commerce of any food, drug, device, or
11		cosmetic that is adulterated or misbranded, and the
12		delivery or proffered delivery thereof for pay or
13		otherwise;
14	(4)	The sale, delivery for sale, holding for sale, or
15		offering for sale of any article in violation of
16		section 328-11, 328-12, or 328-17;
17	(5)	The dissemination of any false advertisement;
18	(6)	The refusal to permit entry or inspection, or to
19		permit the taking of a sample, as authorized by
20		sections 328-22 and 328-23 to 328-27, or to permit
21		access to or copying of any record as authorized by
22		section 328-23;

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1	(7)	The giving of a guaranty or undertaking which guaranty
2		or undertaking is false, except by a person who relied
3		on a guaranty or undertaking to the same effect signed
4		by, and containing the name and address of the person
5		residing in the State from whom the person received in
6		good faith the food, drug, device, or cosmetic;
7	(8)	The removal or disposal of a detained or embargoed
8		article in violation of sections 328-25 to 328-27;
9	(9)	The alteration, mutilation, destruction, obliteration,
10		or removal of the whole or any part of the labeling
11		of, or the doing of any other act with respect to a
12		food, drug, device, or cosmetic, if the act is done
13		while the article is held for sale and results in the
14		article being adulterated or misbranded;
15	(10)	Forging, counterfeiting, simulating, or falsely
16	,	representing, or without proper authority using any
17		mark, stamp, tag, label, or other identification
18		device authorized or required by rules adopted under
19		this part or regulations adopted under the Federal
20		Act;
21	(11)	The use, on the labeling of any drug or in any
22		advertisement relating to the drug, of any

I		representation or suggestion that an application with
2		respect to the drug is effective under section 328-17,
3		or that the drug complies with that section;
4	(12)	The use by any person to the person's own advantage,
5		or revealing other than to the department of health or
6		to the courts when relevant in any judicial proceeding
7		under this part, any information acquired under
8		authority of section 328-11, 328-12, 328-17, or 328-
9		23, concerning any method or process which as a trade
10		secret is entitled to protection;
11	(13)	In the case of a prescription drug distributed or
12		offered for sale in this State, the failure of the
13		manufacturer, packer, or distributor thereof to
14		maintain for transmittal, or to transmit, to any
15		practitioner who makes written request for information
16		as to the drug, true and correct copies of all printed
17		matter which is required to be included in any package
18		in which that drug is distributed or sold, or such
19		other printed matter as is approved under the Federal
20		Act. Nothing in this paragraph shall be construed to
21		exempt any person from any labeling requirement
22		imposed by or under other provisions of this part;

1	(14) (A)	Placing or causing to be placed upon any drug or
2		device or container thereof, with intent to
3		defraud, the trade name or other identifying
4		mark, or imprint of another or any likeness of
5		any of the foregoing; or
6	(B)	Selling, dispensing, disposing of, or causing to
7		be sold, dispensed, or disposed of, or concealing
8		or keeping in possession, control, or custody,
9		with intent to sell, dispense, or dispose of, any
10		drug, device, or any container thereof, with
11		knowledge that the trade name or other
12		identifying mark or imprint of another or any
13		likeness of any of the foregoing has been placed
14		thereon in a manner prohibited by subparagraph
15		(A); or
16	(C)	Making, selling, disposing of, or causing to be
17		made, sold, or disposed of, or keeping in

possession, control, or custody, or concealing,

reproduce that trade name or other identifying

mark or imprint of another or any likeness of any

other thing designed to print, imprint, or

with intent to defraud, any punch, die, plate, or

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1		of the foregoing upon any drug, device, or
2		container thereof;
3	(15)	Except as provided in part VI and section 461-1,
4		dispensing or causing to be dispensed a different drug
5		or brand of drug in place of the drug or brand of drug
6		ordered or prescribed without express permission in
7		each case of the person ordering or prescribing;
8	(16)	The distribution in commerce of a consumer commodity
9		as defined in this part, if such commodity is
10		contained in a package, or if there is affixed to that
11		commodity a label, which does not conform to this part
12		and of rules adopted under authority of this part;
13		provided that this prohibition shall not apply to
14		persons engaged in business as wholesale or retail
15		distributors of consumer commodities except to the
16		extent that such persons:
17		(A) Are engaged in the packaging or labeling of such
18		commodities; or
19		(B) Prescribe or specify by any means the manner in
20		which such commodities are packaged or labeled;
21	(17)	The selling or dispensing in restaurants, soda
22		fountains, drive-ins, lunch wagons, or similar public

	eating establishments of imitation milk and imitation
	milk products in place of fresh milk and fresh milk
	products respectively; of liquid or dry products which
	simulate cream but do not comply with content
	requirements for cream in place of cream; of non-dairy
	frozen desserts which do not comply with content
	requirements for dairy frozen desserts in place of
	dairy frozen desserts; and of any other imitation food
	or one made in semblance of a genuine food in place of
	such genuine food, unless the consumer is notified by
	either proper labeling or conspicuous posted signs or
	conspicuous notices on menu cards and advertisements
	informing of such substitution, to include but not
	limited to the substitution of imitation milk in milk
	shake and malted milk drinks;
(18)	Wilfully and falsely representing or using any
	devices, substances, methods, or treatment as
	effective in the diagnosis, cure, mitigation,
	treatment, or alleviation of cancer. This paragraph
	shall not apply to any person who depends exclusively
	upon prayer for healing in accordance with teachings
	of a hone fide religious sect denomination or

1		organization, nor to a person who practices such
2		teachings;
3	(19)	The selling or offering for sale at any food facility
4		which serves or sells over the counter directly to the
5		consumer an unlabeled or unpackaged food that is a
6		confectionery which contains alcohol in excess of one-
7		half of one per cent by weight unless the consumer is
8		notified of that fact by either proper labeling or
9		conspicuous posted signs or conspicuous notices on
10		menu cards and advertisements;
11	(20)	The sale to a person below the age of twenty-one years
12		of any food which is a confectionery which contains
13		alcohol in excess of one-half of one per cent by
14		weight[-];
15	(21)	After December 31, 2008, the manufacture, sale or
16		delivery or holding or offering for sale of any food
17		containing any amount of aspartame and its derivative
18		compounds in any of their trade names."
19	SECT	ION 4. This section shall not apply to the sale,
20	delivery,	holding, or offering for sale of any food product
21	containing	g aspartame prior to January 1, 2009.

1 SECTION 5. Statutory material to be repealed is bracketed

2 and stricken. New statutory material is underscored.

3 SECTION 6. This Act shall take effect upon its approval.

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INTRODUCED BY:

### Report Title:

Artificial Sweetener; Aspartame; Ban; Food

### Description:

Bans the use of the artificial sweetener aspartame in food products.