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	neotame and their derivative compounds, in all of their trade
4	names, are poisonous and deleterious food additives due to their
5	neurotoxic 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 neotame and their derivative compounds
12	in order to protect and ensure the public health, safety and
13	welfare.
14	SECTION 2. Section 328-1, Hawaii Revised Statutes, is
15	amended by adding two new definitions to be appropriately
16	inserted and to read as follows:

"_"Aspartame" means the artificial sweetener with the

18 technical name L-aspartyl-L-phenylalanine methyl ester.



1	"Neo	tame" means the artificial sweetener with the technical
2	name N-[N	-(3,3-dimethylbutyl)-L-alpha-aspartyl]-L-phenylalanine
3	1-methyl	ester."
4	SECT	ION 3. Section 328-6, Hawaii Revised Statutes, is
5	amended t	o read as follows:
6	"§32	8-6 Prohibited acts. The following acts and the
7	causing t	hereof within the State by any person are prohibited:
8	(1)	The manufacture, sale, delivery, holding, or offering
9		for sale of any food, drug, device, or cosmetic that
10		is adulterated or misbranded;
11	(2)	The adulteration or misbranding of any food, drug,
12		device, or cosmetic;
13	(3)	The receipt in commerce of any food, drug, device, or
14		cosmetic that is adulterated or misbranded, and the
15		delivery or proffered delivery thereof for pay or
16		otherwise;
17	(4)	The sale, delivery for sale, holding for sale, or
18		offering for sale of any article in violation of
19		section 328-11, 328-12, or 328-17;
20	(5)	The dissemination of any false advertisement;
21	(6)	The refusal to permit entry or inspection, or to
22		permit the taking of a sample, as authorized by

1		sections 328-22 and 328-23 to 328-27, or to permit
2		access to or copying of any record as authorized by
3		section 328-23;
4	(7)	The giving of a guaranty or undertaking which guaranty
5		or undertaking is false, except by a person who relied
6		on a guaranty or undertaking to the same effect signed
7		by, and containing the name and address of the person
8		residing in the State from whom the person received in
9		good faith the food, drug, device, or cosmetic;
10	(8)	The removal or disposal of a detained or embargoed
11		article in violation of sections 328-25 to 328-27;
12	(9)	The alteration, mutilation, destruction, obliteration,
13		or removal of the whole or any part of the labeling
14		of, or the doing of any other act with respect to a
15		food, drug, device, or cosmetic, if the act is done
16		while the article is held for sale and results in the
17		article being adulterated or misbranded;
18	(10)	Forging, counterfeiting, simulating, or falsely
19		representing, or without proper authority using any
20		mark, stamp, tag, label, or other identification
21		device authorized or required by rules adopted under

1		this part or regulations adopted under the Federal
2		Act;
3	(11)	The use, on the labeling of any drug or in any
4		advertisement relating to the drug, of any
5		representation or suggestion that an application with
6		respect to the drug is effective under section 328-17,
7		or that the drug complies with that section;
8	(12)	The use by any person to the person's own advantage,
9		or revealing other than to the department of health or
10		to the courts when relevant in any judicial proceeding
11		under this part, any information acquired under
12		authority of section 328-11, 328-12, 328-17, or 328-
13		23, concerning any method or process which as a trade
14		secret is entitled to protection;
15	(13)	In the case of a prescription drug distributed or
16		offered for sale in this State, the failure of the
17		manufacturer, packer, or distributor thereof to
18		maintain for transmittal, or to transmit, to any
19		practitioner who makes written request for information
20		as to the drug, true and correct copies of all printed
21		matter which is required to be included in any package
22		in which that drug is distributed or sold, or such

1		othe	r printed matter as is approved under the Federal
2		Act.	Nothing in this paragraph shall be construed to
3		exem	pt any person from any labeling requirement
4		impo	sed by or under other provisions of this part;
5	(14)	(A)	Placing or causing to be placed upon any drug or
6			device or container thereof, with intent to
7			defraud, the trade name or other identifying
8			mark, or imprint of another or any likeness of
9			any of the foregoing; or
10		(B)	Selling, dispensing, disposing of, or causing to
11			be sold, dispensed, or disposed of, or concealing
12			or keeping in possession, control, or custody,
13			with intent to sell, dispense, or dispose of, any
14			drug, device, or any container thereof, with
15			knowledge that the trade name or other
16			identifying mark or imprint of another or any
17			likeness of any of the foregoing has been placed
18			thereon in a manner prohibited by subparagraph
19			(A); or
20		(C)	Making, selling, disposing of, or causing to be
21			made, sold, or disposed of, or keeping in
22			possession, control, or custody, or concealing,

1		with intent to defraud, any punch, die, plate, or
2		other thing designed to print, imprint, or
3		reproduce that trade name or other identifying
4		mark or imprint of another or any likeness of any
5		of the foregoing upon any drug, device, or
6		container thereof;
7	(15)	Except as provided in part VI and section 461-1,
8		dispensing or causing to be dispensed a different drug
9		or brand of drug in place of the drug or brand of drug
10		ordered or prescribed without express permission in
11		each case of the person ordering or prescribing;
12	(16)	The distribution in commerce of a consumer commodity
13		as defined in this part, if such commodity is
14		contained in a package, or if there is affixed to that
15		commodity a label, which does not conform to this part
16		and of rules adopted under authority of this part;
17		provided that this prohibition shall not apply to
18		persons engaged in business as wholesale or retail
19		distributors of consumer commodities except to the
20		extent that such persons:
21		(A) Are engaged in the packaging or labeling of such
22		commodities; or

1		(B) Prescribe or specify by any means the manner in
2		which such commodities are packaged or labeled;
3	(17)	The selling or dispensing in restaurants, soda
4		fountains, drive-ins, lunch wagons, or similar public
5		eating establishments of imitation milk and imitation
6		milk products in place of fresh milk and fresh milk
7		products respectively; of liquid or dry products which
8		simulate cream but do not comply with content
9		requirements for cream in place of cream; of non-dairy
10		frozen desserts which do not comply with content
11		requirements for dairy frozen desserts in place of
12		dairy frozen desserts; and of any other imitation food
13		or one made in semblance of a genuine food in place of
14		such genuine food, unless the consumer is notified by
15		either proper labeling or conspicuous posted signs or
16		conspicuous notices on menu cards and advertisements
17		informing of such substitution, to include but not
18		limited to the substitution of imitation milk in milk
19		shake and malted milk drinks;
20	(18)	Wilfully and falsely representing or using any
21		devices, substances, methods, or treatment as
22		effective in the diagnosis, cure, mitigation,

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

	INTRODUCED BY: Mele Carrell
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6	SECTION 6. This Act shall take effect upon its approval.
5	and stricken. New statutory material is underscored.
4	SECTION 5. Statutory material to be repealed is bracketed
3	containing aspartame or neotame prior to January 1, 2010.
2	delivery, holding, or offering for sale of any food product
1	SECTION 4. This section shall not apply to the sale,

JAN 2 3 2009

Report Title:

Artificial Sweetener; Aspartame; Neotame; Ban

Description:

Bans the use of the artificial sweeteners aspartame and neotame in food products.