LRB9204278DJpc

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AN ACT in relation to public health.

Be it enacted by the People of the State of Illinois,represented in the General Assembly:

Section 5. The Illinois Food, Drug and Cosmetic Act is
amended by changing Sections 21 and 22.4 and adding Sections
21.3 and 21.4 as follows:

- 7 (410 ILCS 620/21) (from Ch. 56 1/2, par. 521)
- 8 Sec. 21. <u>Enforcement of Act.</u>

9 (a) The authority to promulgate regulations for the 10 efficient enforcement of this Act is vested in the Director. 11 The Director is authorized to make the regulations 12 promulgated under this Act conform, in so far as practicable, 13 with those promulgated under the Federal Act.

(b) Hearings authorized or required by this Act shall be
conducted by the Director or an officer, agent or employee
designated by him.

(c) All pesticide chemical regulations and supplements thereto or revisions thereof adopted under authority of the Federal Food, Drug and Cosmetic Act are the pesticide chemical regulations in this State, except insofar as modified or rejected by regulations for finished foods promulgated by the Director.

(d) All food additive regulations and supplements thereto or revisions thereof adopted under authority of the Federal Food, Drug and Cosmetic Act are the food additive regulations in this State, except insofar as modified or rejected by regulations promulgated by the Director.

(e) All color additive regulations and supplements
thereto or revisions thereof adopted under authority of the
Federal Food, Drug and Cosmetic Act are the color additive
regulations in this State, except insofar as modified or

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rejected by regulations promulgated by the Director.

(f) All special dietary use regulations and supplements thereto or revisions thereof adopted under authority of the Federal Food, Drug and Cosmetic Act are the special dietary use regulations in this State, except insofar as modified or rejected by regulations promulgated by the Director.

7 (g) All bottled water and vended water device regulations and supplements thereto or revisions thereof 8 9 adopted under the authority of the Federal Food, Drug and Cosmetic Act are the bottled water and vended water device 10 11 regulations in this State except insofar as modified or 12 rejected by regulations promulgated by the Director.

(h) All infant formula regulations and supplements thereto or revisions thereof adopted under the authority of the Federal Food, Drug and Cosmetic Act are the infant formula regulations in this State except insofar as modified or rejected by regulations promulgated by the Director.

All drug, food, device and cosmetic 18 (i) Good Manufacturing Practices Regulations and supplements thereto 19 or revisions thereof adopted under the authority of Federal 20 21 Food, Drug and Cosmetic Act are the food, drug, device and 22 cosmetic Good Manufacturing Practices Regulations in this 23 State, except insofar as modified or rejected by regulations promulgated by the Director. 24

(j) A federal regulation automatically adopted pursuant to this Act takes effect in this State on the date it becomes effective as a Federal regulation. No publication or hearing is required. The Director shall promulgate all other proposed regulations in compliance with the requirements of The Illinois Administrative Procedure Act.

31 (k) On and after July 1, 2002, all federal regulations 32 concerning food and supplements, or revisions to the 33 regulations adopted under the authority of the Federal Food, 34 Drug and Cosmetic Act, including 21 C.F.R. 70 through 699,

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1 are the regulations concerning food in this State, except as 2 modified or rejected by rules adopted by the Director. 3 (1) On and after July 1, 2002, the Department may 4 establish and assess civil monetary penalties against a license holder for violations of this Act or regulations 5 adopted under this Act. Before assessing a penalty, the 6 7 Department must give the alleged violator an opportunity to 8 request an administrative hearing on the matter of the alleged violation. In no circumstances may any penalty 9 10 exceed \$1,000 per day for each day the license holder remains 11 in violation. All penalties collected under this Act shall 12 be deposited into the Food and Drug Safety Fund. The Attorney General may bring a civil action in the circuit 13 court to enforce the collection of a penalty imposed under 14 15 this subsection. 16 (Source: P.A. 84-891.) 17 (410 ILCS 620/21.3 new) Sec. 21.3. Food manufacturers, processors, packers, and 18 warehouses; permits and inspections. 19 (a) It is unlawful for any person to establish, 20 21 maintain, conduct, or operate a food manufacturer, processor, or warehouse within this State without first 22 packer, 23 obtaining a permit from the Department. Permits shall expire on September 30 of each year, 24 25 unless revoked by the Department for noncompliance with the 26 rules adopted under this Act or discontinuation of the 27 facility's operation. Applications for original permits or renewals shall be 28 made on forms furnished by the Department and shall include 29 30 at least the following: (1) The name and address of the applicant, or the 31 names and addresses of the partners if the applicant is 32 33 a partnership, or the names and addresses of the officers

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1 if the applicant is a corporation, or the names and addresses of all persons having a financial interest in 2 the applicant if the applicant is a group of individuals, 3 4 association, or trust; and 5 (2) The name of the food manufacturer, processor, packer or warehouse, location, mailing address, and 6 7 telephone number. 8 An initial, non-refundable fee of \$100 must be submitted 9 to the Department with the application for a license. Each annual renewal application must be submitted with a 10 non-refundable \$100 fee, except that each facility with a 11 12 license that has been expired for more than 90 days must pay a non-refundable \$150 fee for renewal of the license. All 13 fees collected shall be deposited into the Food and Drug 14 15 Safety Fund. Those food manufacturers, processors, packers, or 16 17 warehouses permitted by the Department under Section 21.2 of this Act or under the Grade A Pasteurized Milk and Milk 18 Products Act or by the Illinois Department of Agriculture 19 under the Meat and Poultry Inspection Act or the Illinois Egg 20 21 and Egg Products Act are exempt from the permit fees in this 22 Section, but not from other provisions of this Act. 23 The Department is authorized, upon request and the non-refundable payment of \$10 per certificate, to issue 24 certificates of free sale, health certificates, or an 25 equivalent, to Illinois food, dairy, drug, cosmetic, or 26 medical device manufacturers, processors, packers, or 27 warehousers. All fees collected shall be deposited into the 28 29 Food and Drug Safety Fund. The Department shall issue an Illinois shellfish 30 31 certificate, upon request, to shellfish firms in compliance with the Interstate Shellfish Sanitation Conference. 32 33 (b) Every Illinois food manufacturer, processor, packer, or warehouse shall be inspected by the Department at a 34

1 frequency that is necessary to ensure compliance with this 2 <u>Act.</u> 3 (c) This Section applies on and after July 1, 2002. (410 ILCS 620/21.4 new) 4 5 Sec. 21.4. Units of local government; home rule. (a) The regulation, permitting, and inspection of food 6 7 manufacturers, processors, packers, warehouses, manufactured dairy farms, manufactured dairy plants, bulk milk 8 hauler-samplers, bulk milk pickup tanks, certified 9 10 pasteurizer sealers, and manufactured dairy products, but not 11 including food service establishments and retail food stores, 12 are exclusive powers and functions of the State. A home rule unit may not regulate these entities. This Section is a 13 14 denial and limitation of home rule powers and functions under 15 subsection (h) of Section 6 of Article VII of the Illinois 16 Constitution. 17 (b) This Section applies on and after July 1, 2002.

18 (410 ILCS 620/22.4) (from Ch. 56 1/2, par. 522.4)

19 Sec. 22.4. Food and Drug Safety Fund. There is created in 20 the State Treasury a special fund to be known as the Food and Drug Safety Fund. All subscription, fine, and permit fees, 21 certificate fees, and other moneys collected by the 22 23 Department of Public Health under this Act shall be deposited into the Fund. Subject to appropriation by the General 24 Assembly, moneys deposited into this Fund shall be made 25 available to the Department of Public Health to administer 26 27 the--Drug--Product--Selection-Program-or-for-other Department 28 activities related to food safety, drug safety, milk safety, or drug product selection. All interest that accrues on the 29 30 moneys in the Fund shall be deposited into the Fund.

31 (Source: P.A. 89-526, eff. 7-19-96.)