

✓ 78 C+B
OW 4-25

M. RITA

HOUSE BILL 2613
87th GENERAL ASSEMBLY
State of Illinois

1991 and 1992

Introduced April 5, 1991, by Representative Giorgi

SYNOPSIS

(Ch. 110, par. 13-213)

Amends the Code of Civil Procedure. Provides that a cause of action based on a latent disease caused by exposure to harmful material does not accrue, for purposes of limitation periods, until the person knows or should know of the disease and its cause.

LRB8705894REnc

A BILL FOR

TIILBC 008660

HB2613

LRB8705894REmc

1	AN ACT to amend the Code of Civil Procedure by changing	64
2	Section 13-213.	
3	Be It enacted by the People of the State of Illinois,	68
4	represented in the General Assembly:	
5	Section 1. The Code of Civil Procedure is amended by	71
6	changing Section 13-213 as follows:	
7	(Ch. 110, par. 13-213)	74
8	Sec. 13-213. Product liability.	76
9	(a) As used in this Section, the term:	78
10	(1) "alteration, modification or change" or	80
11	"altered, modified, or changed" means an alteration,	81
12	modification or change that was made in the original	82
13	makeup characteristics, function or design of a product	
14	or in the original recommendations, instructions and	83
15	warnings given with respect to a product including the	84
16	failure properly to maintain and care for a product.	85
17	(2) "product" means any tangible object or goods	87
18	distributed in commerce, including any service provided	88
19	in connection with the product. Where the term "product	89
20	unit" is used, it refers to a single item or unit of a	90
21	product.	
22	(3) "product liability action" means any action.	92
23	based on the doctrine of strict liability in tort brought	93
24	against the seller of a product on account of personal	94
25	injury, (including illness, disease, disability and	95
26	death) or property, economic or other damage allegedly	96
27	caused by or resulting from the manufacture,	
28	construction, preparation, assembly, installation,	97
29	testing, makeup, characteristics, functions, design,	98
30	formula, plan, recommendation, specification,	
31	prescription, advertising, sale, marketing, packaging,	99
32	labeling, repair, maintenance or disposal of, or warning	100

THILBC 008661

HB2613

-2-

LRB8705894REnc

1 or instruction regarding any product. This definition 101
2 excludes actions brought by State or federal regulatory
3 agencies pursuant to statute. 102
4 (4) "seller" means one who, in the course of a 104
5 business conducted for the purpose, sells, distributes, 105
6 leases, assembles, installs, produces, manufactures, 106
7 fabricates, prepares, constructs, packages, labels, 107
8 markets, repairs, maintains, or otherwise is involved in
9 placing a product in the stream of commerce. 108
10 (b) Subject to the provisions of subsections (c) and (d) 110
11 no product liability action based on the doctrine of strict 111
12 liability in tort shall be commenced except within the 112
13 applicable limitations period and, in any event, within 12 113
14 years from the date of first sale, lease or delivery of 114
15 possession by a seller or 10 years from the date of first
16 sale, lease or delivery of possession to its initial user, 115
17 consumer, or other non-seller, whichever period expires 116
18 earlier, of any product unit that is claimed to have injured 117
19 or damaged the plaintiff, unless the defendant expressly has 118
20 warranted or promised the product for a longer period and the 119
21 action is brought within that period.
22 (c) No product liability action based on the doctrine of 121
23 strict liability in tort to recover for injury or damage 122
24 claimed to have resulted from an alteration, modification or 123
25 change of the product unit subsequent to the date of first 124
26 sale, lease or delivery of possession of the product unit to 125
27 its initial user, consumer or other non-seller shall be 126
28 limited or barred by subsection (b) hereof if:
29 (1) the action is brought against a seller making, 128
30 authorizing, or furnishing materials for the 129
31 accomplishment of such alteration, modification or change 130
32 (or against a seller furnishing specifications or 131
33 instructions for the accomplishment of such alteration,
34 modification or change when the injury is claimed to have 132
35 resulted from failure to provide adequate specifications 133

THILBC 008662

HB2613

-3-

LRB8705894REnc

1	or instructions), and	133
2	(2) the action commenced within the applicable	135
3	limitation period and, in any event, within 10 years from	136
4	the date such alteration, modification or change was	137
5	made, unless defendant expressly has warranted or	138
6	promised the product for a longer period and the action	
7	is brought within that period, and	139
8	(3) when the injury or damage is claimed to have	141
9	resulted from an alteration, modification or change of a	142
10	product unit, there is proof that such alteration,	143
11	modification or change had the effect of introducing into	144
12	the use of the product unit, by reason of defective	145
13	materials or workmanship, a hazard not existing prior to	
14	such alteration, modification or change.	146
15	(d) Notwithstanding the provisions of subsection (b) and	148
16	paragraph (2) of subsection (c) if the injury complained of	149
17	occurs within any of the periods provided by subsection (b)	150
18	and paragraph (2) of subsection (c), the plaintiff may bring	151
19	an action within 2 years after the date on which the claimant	152
20	knew, or through the use of reasonable diligence should have	
21	known, of the existence of the personal injury, death or	153
22	property damage, but in no event shall such action be brought	154
23	more than 8 years after the date on which such personal	155
24	injury, death or property damage occurred. In any such case,	156
25	if the person entitled to bring the action was, at the time	
26	the personal injury, death or property damage occurred, under	157
27	the age of 18 years, or under a legal disability, then the	158
28	period of limitations does not begin to run until the person	159
29	attains the age of 18 years, or the disability is removed.	160
30	(e) Replacement of a component part of a product unit	162
31	with a substitute part having the same formula or design as	163
32	the original part shall not be deemed a sale, lease or	164
33	delivery of possession or an alteration, modification or	165
34	change for the purpose of permitting commencement of a	166
35	product liability action based on the doctrine of strict	167

TIILBC 008663

HB2613

-4-

LRB8705894REMc

1 liability in tort to recover for injury or damage claimed to 167
2 have resulted from the formula or design of such product unit 168
3 or of the substitute part when such action would otherwise be 169
4 barred according to the provisions of subsection (b) of this 170
5 section.

6 (f) Nothing in this section shall be construed to create 172
7 a cause of action or to affect the right of any person to 173
8 seek and obtain indemnity or contribution. 174

9 (g) In a product liability claim against the product 176
10 seller, no limitation shall apply to the time to discover a 177
11 disease that is latent and caused by exposure to a harmful 178
12 material, in which event the action shall be deemed to have 179
13 accrued when the disease and the disease's cause have been 180
14 made known to the person or at the point the person should
15 have been aware of the disease and the disease's cause. 181

16 The term "harmful material" means any chemical substance 183
17 commonly known as asbestos, dioxins, or polychlorinated 184
18 biphenyls, whether alone or as part of any product, or any 185
19 substance that is determined to present an unreasonable risk 186
20 of injury to health or the environment by the United States 187
21 Environment Protection Agency under the federal Toxic
22 Substances Control Act, 15 U.S.C. Sec. 2601 et seq., or the 188
23 State of Illinois, and because of such risk is regulated by 189
24 the State or the Environmental Protection Agency.

25 Upon the effective date of this amendatory Act, the 191
26 provisions of this subsection shall revive the causes of 192
27 action for latent diseases caused by exposure to a harmful 193
28 material for:

29 (1) any person whose cause of action had accrued, 195
30 as defined in this subsection on or after March 3, 1987; 196
31 or

32 (2) any person who had an action pending in any 198
33 court on March 3, 1989, and because of the judicial 199
34 interpretation of a limitation period contained in this 200
35 Code, and amendments thereto, as applied to latent

TIILBC 008664

HB2613

-5-

LRB8705894REmc

1	<u>disease caused by exposure to a harmful material the:</u>	201
2	<u>(A) action was dismissed;</u>	203
3	<u>(B) dismissal of the action was affirmed; or</u>	205
4	<u>(C) action was subject to dismissal.</u>	207
5	<u>The intent of this subsection is to revive causes of</u>	209
6	<u>action for latent diseases caused by exposure to a harmful</u>	210
7	<u>material which were barred by interpretation of this Code and</u>	211
8	<u>amendments thereto in effect before this enactment.</u>	
9	<u>(h) {g}</u> The provisions of this Section 13-213 of this	213
10	Act apply to any cause of action accruing on or after January	214
11	1, 1979, involving any product which was in or entered the	215
12	stream of commerce prior to, on, or after January 1, 1979.	
13	(Source: P.A. 85-907; 86-1329.)	218

THILBC 008665