112TH CONGRESS 2D SESSION

H. R. 4332

To amend section 505(j) of the Federal Food, Drug, and Cosmetic Act to extend the period for a first applicant, with respect to a generic drug, to obtain tentative approval without forfeiting the 180-day exclusivity period, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

March 29, 2012

Mr. Pallone (for himself and Mr. Guthrie) introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

- To amend section 505(j) of the Federal Food, Drug, and Cosmetic Act to extend the period for a first applicant, with respect to a generic drug, to obtain tentative approval without forfeiting the 180-day exclusivity period, and for other purposes.
 - 1 Be it enacted by the Senate and House of Representa-
 - 2 tives of the United States of America in Congress assembled,
 - 3 SECTION 1. SHORT TITLE.
 - 4 This Act may be cited as the "Generic Drug Applica-
 - 5 tion Review Fairness Act of 2012".

1	SEC. 2. EXTENSION OF PERIOD FOR FIRST APPLICANT TO
2	OBTAIN TENTATIVE APPROVAL WITHOUT
3	FORFEITING 180-DAY EXCLUSIVITY PERIOD.
4	(a) Extension of Period.—
5	(1) In General.—Subclause (IV) of section
6	505(j)(5)(D)(i) of the Federal Food, Drug, and Cos-
7	metic Act (21 U.S.C. $355(j)(5)(D)(i)$) is amended to
8	read as follows:
9	"(IV) Failure to obtain ten-
10	TATIVE APPROVAL.—The first appli-
11	cant fails to obtain tentative approva
12	of the application within 60 months
13	after the date on which—
14	"(aa) the application is filed
15	and initially contains a certifi-
16	cation described in paragraph
17	(2)(A)(vii)(IV), or
18	"(bb) the application is
19	amended to first contain such a
20	certification,
21	unless the failure is caused by ϵ
22	change in or a review of the require-
23	ments for approval of the application
24	imposed after the date on which the
25	application is so filed or amended.".
26	(2) Applicability.—

1	(A) In General.—Subject to subsection
2	(b), the amendment made by paragraph (1) ap-
3	plies—
4	(i) only with respect to an application
5	that is filed under section 505(j) of the
6	Federal Food, Drug, and Cosmetic Act (21
7	U.S.C. 355(j)) on or after the day that is
8	30 months prior to the date of the enact-
9	ment of this Act; and
10	(ii) only if no certification under para-
11	graph (2)(A)(vii)(IV) of such section
12	505(j) was made before such day with re-
13	spect to the listed drug (as such term is
14	used in such section 505(j)).
15	(B) CERTAIN APPLICATIONS.—If an appli-
16	cation was filed under section 505(j) of the
17	Federal Food, Drug, and Cosmetic Act (21
18	U.S.C. 355(j)) prior to the day specified in sub-
19	paragraph (A)(i) and, on such day, contained a
20	certification described in paragraph
21	(2)(A)(vii)(IV), the application shall be subject
22	to paragraph (5)(D)(i)(IV) of such section
23	505(j) as in effect on the day before the date
24	of the enactment of this Act.

(b)	INCREMENTAL REDUCTION OF EXTENDED PE-
RIOD.—	
	(1) Period duration.—
	(A) Effective on October 1, 2013, sub-
	clause (IV) of section $505(j)(5)(D)(i)$ of the
	Federal Food, Drug, and Cosmetic Act (21
	U.S.C. $355(j)(5)(D)(i)$, as amended by sub-
	section (a)(1), is amended by striking "60
	months" and inserting "54 months".
	(B) Effective on October 1, 2014, sub-
	clause (IV) of section $505(j)(5)(D)(i)$ of the
	Federal Food, Drug, and Cosmetic Act (21
	U.S.C. $355(j)(5)(D)(i)$, as amended by sub-
	paragraph (A), is amended by striking "54
	months" and inserting "48 months".
	(C) Effective on October 1, 2015, sub-
	clause (IV) of section $505(j)(5)(D)(i)$ of the
	Federal Food, Drug, and Cosmetic Act (21
	U.S.C. $355(j)(5)(D)(i)$, as amended by sub-
	paragraph (B), is amended by striking "48
	months" and inserting "42 months".
	(D) Effective on October 1, 2016, sub-
	clause (IV) of section $505(j)(5)(D)(i)$ of the
	Federal Food, Drug, and Cosmetic Act (21

U.S.C. 355(j)(5)(D)(i), as amended by sub-

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1	paragraph (C), is amended by striking "42
2	months" and inserting "36 months".
3	(E) Effective on October 1, 2017, sub-
4	clause (IV) of section $505(j)(5)(D)(i)$ of the
5	Federal Food, Drug, and Cosmetic Act (21
6	U.S.C. $355(j)(5)(D)(i)$, as amended by sub-
7	paragraph (D), is amended by striking "36
8	months" and inserting "30 months".
9	(2) Applicability.—
10	(A) The amendments made by subpara-
11	graphs (A), (B), (C), and (D) of paragraph (1)
12	apply only with respect to an application under
13	section 505(j) of the Federal Food, Drug, and
14	Cosmetic Act (21 U.S.C. 355(j)) that—
15	(i) is filed and initially contains a cer-
16	tification described in paragraph
17	(2)(A)(vii)(IV) during the period of one
18	fiscal year beginning on the effective date
19	of the respective amendment; or
20	(ii) is amended to initially contain
21	such a certification during such period.
22	(B) The amendment made by paragraph
23	(1)(E) applies only with respect to an applica-
24	tion under section 505(j) of the Federal Food,

1	Drug, and Cosmetic Act (21 U.S.C. 355(j))
2	that—
3	(i) is filed and initially contains a cer-
4	tification described in paragraph
5	(2)(A)(vii)(IV) on or after October 1,
6	2017; or
7	(ii) is amended to initially contain
8	such a certification on or after October 1,
9	2017.
10	(c) Conforming Amendment.—Subparagraph (G)
11	of section $505(q)(1)$ of the Federal Food, Drug, and Cos-
12	metic Act (21 U.S.C. 355(q)(1)) is amended by striking
13	"the 30-month period" and inserting "the period".
14	SEC. 3. MAINTENANCE OF OFFICE OF GENERIC DRUGS AT
15	CDER; REPORTING.
15 16	CDER; REPORTING. Section 505(j) of the Federal Food, Drug, and Cos-
16 17	Section 505(j) of the Federal Food, Drug, and Cos-
16 17	Section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) is amended by adding at the
16 17 18	Section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) is amended by adding at the end the following:
16 17 18 19	Section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) is amended by adding at the end the following: "(11) Office of Generic Drugs.—
16 17 18 19 20	Section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) is amended by adding at the end the following: "(11) Office of Generic Drugs.— "(A) Office.—The Secretary shall main-
116 117 118 119 220 221	Section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) is amended by adding at the end the following: "(11) Office of Generic Drugs.— "(A) Office.—The Secretary shall maintain the Office of Generic Drugs as a separate

1	"(B) REPORTING.—The Director of the
2	Office of Generic Drugs shall report directly to
3	the Director of the Center for Drug Evaluation
4	and Research.".

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