

Program	Statute	Section and age distinction	Regulation
The Children, Youth, and Families At-Risk Sustainable Community Projects (CYFAR SCP).	7 U.S.C. 341, <i>et seq.</i> ; 7 U.S.C. 343(d).	Section 3(d) of the Smith-Lever Act authorizes the Department to administer the CYFAR SCP. Per Program notices, CYFAR SCP supports community educational programs for at-risk children, youth, and families which are based on locally identified needs, soundly grounded in research, and which lead to the accomplishment of one of four CYFAR National Outcomes; and (2) [t]o integrate CYFAR programming into ongoing Extension programs for children, youth, and families—insuring that at-risk, low income children, youth, and families continue to be part of Extension and/or 4–H programs and have access to resources and educational opportunities.	7 CFR part 3015, 7 CFR part 3019, 7 CFR part 3430.

**Risk Management Agency**

Federal Crop Insurance Program.	7 U.S.C. 1501 .....	Per the Crop Insurance Handbook, which provides the official FCIC approved underwriting standards for policies administered by Approved Insurance Providers under the Common Crop Insurance Policy Basic Provisions, 7 CFR part 457 including the Catastrophic Risk Protection Endorsement, 7 CFR part 402, and the Actual Production History Regulation 7 CFR part 400 Subpart G for the 2014 and succeeding crop years, to be eligible for crop insurance the applicant must be of “legal majority.” Legal majority is defined as “where the individual has reached 18 years old or was conferred legal majority by a court. (1) For individuals less than 18 years of age or where legal majority has not been conferred by a court, to be eligible for crop insurance: (a) A minor must provide evidence an insurable share exists; and (b) a court-appointed guardian or parent must co-sign the application. (2) When a court-appointed guardian or parent cosigns the application: (a) An acknowledgement guaranteeing payment of the annual premium must be included with the application; and (b) a written statement describing the farming operation and the insurable share must be provided. (3) For CAT coverage only, a minor who is competent to enter into a binding contract, may insure a crop at CAT level without a co-signer; however, if not competent to enter into a binding contract, a court-appointed guardian or parent must sign the application.”	7 CFR parts 400, 402, 457.
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Dated: November 17, 2014.

**Thomas J. Vilsack,**  
*Secretary.*

[FR Doc. 2014–28452 Filed 12–9–14; 8:45 am]

**BILLING CODE P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Part 101**

[Docket No. FDA–2000–N–0011 (Formerly Docket No. 2000N–1596)]

**Uniform Compliance Date for Food Labeling Regulations**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule.

**SUMMARY:** The Food and Drug Administration (FDA or we) is establishing January 1, 2018, as the uniform compliance date for food labeling regulations that are issued between January 1, 2015, and December 31, 2016. We periodically announce uniform compliance dates for new food labeling requirements to minimize the

economic impact of label changes. On November 28, 2012, we established January 1, 2016, as the uniform compliance date for food labeling regulations issued between January 1, 2013, and December 31, 2014.

**DATES:** This rule is effective December 10, 2014. Submit electronic or written comments by February 9, 2015.

**ADDRESSES:** You may submit comments by any of the following methods:

**Electronic Submissions**

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments.

**Written Submissions**

Submit written submissions in the following ways:

- *Mail/Hand delivery/Courier* (for paper submissions): Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

*Instructions:* All submissions received must include the Docket No. (FDA–2000–N–0011) for this rulemaking. All comments received may be posted without change to <http://www.regulations.gov>, including any personal information provided. For additional information on submitting comments, see the “Comments” heading of the **SUPPLEMENTARY INFORMATION** section of this document.

*Docket:* For access to the docket to read background documents or comments received, go to <http://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

**FOR FURTHER INFORMATION CONTACT:** Michael Ellison, Center for Food Safety and Applied Nutrition (HFS–24), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 240–402–2093.

**SUPPLEMENTARY INFORMATION:**

**I. Background**

We periodically issue regulations requiring changes in the labeling of food. If the effective dates of these labeling changes were not coordinated, the cumulative economic impact on the food industry of having to respond

separately to each change would be substantial. Therefore, we periodically have announced uniform compliance dates for new food labeling requirements (see, e.g., the **Federal Register** of October 19, 1984 (49 FR 41019); December 24, 1996 (61 FR 67710); December 27, 1996 (61 FR 68145); December 23, 1998 (63 FR 71015); November 20, 2000 (65 FR 69666); December 31, 2002 (67 FR 79851); December 21, 2006 (71 FR 76599); December 8, 2008 (73 FR 74349); December 15, 2010 (75 FR 78155); and November 28, 2012 (77 FR 70885)). Use of a uniform compliance date provides for an orderly and economical industry adjustment to new labeling requirements by allowing sufficient lead time to plan for the use of existing label inventories and the development of new labeling materials.

We have determined under 21 CFR 25.30(k) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This final rule contains no collections of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

We have examined the impacts of the final rule under Executive Order 12866, Executive Order 13563, the Regulatory Flexibility Act (5 U.S.C. 601–612), and the Unfunded Mandates Reform Act of 1995 (Public Law 104–4). Executive Orders 12866 and 13563 direct Agencies to assess all costs and benefits of available regulatory alternatives and, when regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety, and other advantages; distributive impacts; and equity). We believe that this final rule is not a significant regulatory action under Executive Order 12866.

The establishment of a uniform compliance date does not in itself lead to costs or benefits. We will assess the costs and benefits of the uniform compliance date in the regulatory impact analyses of the labeling rules that take effect at that date.

The Regulatory Flexibility Act requires Agencies to analyze regulatory options that would minimize any significant economic impact of a rule on small entities. Because the final rule does not impose compliance costs on small entities, we certify that the final rule will not have a significant

economic impact on a substantial number of small entities.

Section 202(a) of the Unfunded Mandates Reform Act of 1995 requires that Agencies prepare a written statement, which includes an assessment of anticipated costs and benefits, before proposing “any rule that includes any Federal mandate that may result in the expenditure by State, local, and tribal governments, in the aggregate, or by the private sector, of \$100,000,000 or more (adjusted annually for inflation) in any one year.” The current threshold after adjustment for inflation is \$141 million, using the most current (2013) Implicit Price Deflator for the Gross Domestic Product. We do not expect this final rule to result in any 1-year expenditure that would meet or exceed this amount.

We have analyzed this final rule in accordance with the principles set forth in Executive Order 13132. We have determined that the rule does not contain policies that have substantial direct effects on the States, on the relationship between the National Government and the States, or on the distribution of power and responsibilities among the various levels of government. Accordingly, we have concluded that the rule does not contain policies that have federalism implications as defined in the Executive Order and, consequently, a federalism summary impact statement is not required.

This action is not intended to change existing requirements for compliance dates contained in final rules published before January 1, 2015. Therefore, all final rules published by FDA in the **Federal Register** before January 1, 2015, will still go into effect on the date stated in the respective final rule. We generally encourage industry to comply with new labeling regulations as quickly as feasible, however. Thus, when industry members voluntarily change their labels, it is appropriate that they incorporate any new requirements that have been published as final regulations up to that time.

In rulemaking that began with publication of a proposed rule on April 15, 1996 (61 FR 16422), and ended with a final rule on December 24, 1996, we provided notice and an opportunity for comment on the practice of establishing uniform compliance dates by issuance of a final rule announcing the date. Receiving no comments objecting to this practice, FDA finds any further advance notice and opportunity for comment or delayed effective date unnecessary for establishment of the uniform compliance date. We have previously invited comment on the practice of

establishing uniform compliance dates by issuing a final rule, and interested parties will have an opportunity to comment on the compliance date for each individual food labeling regulation as part of the rulemaking process for that regulation. Nonetheless, under 21 CFR 10.40(e)(1), we are providing an opportunity for comment on whether the uniform compliance date established by this final rule should be modified or revoked.

The new uniform compliance date will apply only to final FDA food labeling regulations that require changes in the labeling of food products and that publish after January 1, 2015, and before December 31, 2016. Those regulations will specifically identify January 1, 2018, as their compliance date. All food products subject to the January 1, 2018, compliance date must comply with the appropriate regulations when initially introduced into interstate commerce on or after January 1, 2018. If any food labeling regulation involves special circumstances that justify a compliance date other than January 1, 2018, we will determine for that regulation an appropriate compliance date, which will be specified when the final regulation is published.

## II. Comments

Interested persons may submit either electronic comments regarding this document to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: December 4, 2014.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2014–28829 Filed 12–9–14; 8:45 am]

**BILLING CODE 4164–01–P**

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## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 52

[EPA–R05–OAR–2014–0123; FRL–9920–13–Region 5]

### Approval and Promulgation of Air Quality Implementation Plans; Illinois; Withdrawal of Direct Final Rule

**AGENCY:** Environmental Protection Agency (EPA).