Substances	Limitations
* * *	* * * *
2,2′-(2,5-Thiophenediyl)-bis(5-*tert*-butylbenzoxazole) (CAS Reg. No. 7128–64–5).	For use as an optical brightener: 1. In all polymers at levels not to exceed 0.015 percent by weight of the polymer. The finished articles are to contact food only under conditions of use A through H described in Table 2 of § 176.170(c) of this chapter. 2. In all polymers at levels not to exceed 0.05 percent by weight of the polymer. The finished articles shall contact foods only of the types identified in Table 1 of § 176.170(c) of this chapter, under Categories I, II, IV–B, VI–A, VI–B, VI–C, VII–B, and VIII under conditions of use A through H described in Table 2 of § 176.170(c) of this chapter. 3. In adhesives complying with § 175.105 of this chapter and in pressure-sensitive adhesives complying with § 175.125 of this chapter.
* * *	* * * *

Dated: January 5, 1998.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–1539 Filed 1–22–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 820

[Docket No. 90N–0172]

RIN 0910–AA09

Quality System Design Control; Open Public Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Announcement of public meeting.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following meeting: Quality System Design Control open public meeting. The topic to be discussed is the midcourse review of the new design control requirements. This action is being taken in accordance with the current good manufacturing (CGMP) final rule that appeared in the **Federal Register** of October 7, 1996.

DATES: The meeting will be held on, February 2, 1998, from 8:30 a.m. to 4:30 p.m. Written requests for oral presentations by January 28, 1998.

ADDRESSES: The meeting will be held at the National Institutes of Health (NIH), Natcher Auditorium, 45 Center Dr., Bldg. 45, Bethesda, MD. Contact for any changes: (1) Via Internet at http://www.fda.gov/cdrh/gmp, or (2) telephone toll-free at 1–800–638–2041.

FOR FURTHER INFORMATION CONTACT:

For information regarding registration: Mary Ann Fitzgerald, or

For information regarding the meeting or requests for oral presentations: Kimberly A. Trautman, Center for Devices and Radiological Health (HFZ–341), Food and Drug Administration, 2094 Gaither Rd., Rockville, MD 20850, 301–594–4648, FAX 301–594–4672.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of October 7, 1996 (61 FR 52602), FDA stated that it would hold an open public meeting in early 1998 to discuss, and to further explore any concerns industry might be having in implementing the new design control requirements. Specifically, the results of the first several months of design control inspections will be reviewed and any adjustments to the designated inspectional strategy of guidances will be addressed. Also, FDA will evaluate the information gathered at this point and determine if design control requirements, as written in the final rule, are appropriate to obtain the goals expressed in the preamble. Particular attention will be paid to clarity of information obtained, the appropriateness of the information collected with respect to the design control requirements, the manner in which the investigators are writing their observations, and any requirements that seem to be giving manufacturers a problem or where there might be misunderstandings as to what the regulation requires. It is important to note that only the requirements and issues surrounding design controls codified will be addressed.

Fax written requests for oral presentations, (including name, title, firm name, address, telephone, and fax number), and an outline of your presentation to the contact person listed above by January 28, 1998. No telephone requests will be accepted. You will be notified by facsimile whether or not the speaker's list is full. If you cannot be reached by facsimile, please note that in your request.

If you need special accommodations due to a disability, please contact Georgette Smith, NIH Conference Center, 301–496–9966, at least 7 days in advance.

Dated: January 20, 1998.

Joseph A. Levitt,

Deputy Director for Regulations Policy, Center for Devices and Radiological Health.

[FR Doc. 98–1822 Filed 1-21-98; 3:22 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF DEFENSE

Office of the Secretary

32 CFR Part 104

[DoD Instruction 1205.12]

RIN 0790–AG52

Civilian Employment and Reemployment Rights of Applicants for, and Service Members and Former Service Members of the Uniformed Services

AGENCY: Department of Defense.

ACTION: Final rule.

SUMMARY: This part identifies DoD guidelines for implementing policy, assigns responsibilities, and prescribes procedures for informing Service members of their reemployment protections. It updates, codifies, and strengthens the civilian employment rights and benefits of Service members and individuals who apply for uniformed service, and specifies the