Food and Drug Administration, HHS

§ 1.235

How and when do you cancel your facility’s registration information?

(a) Notification of registration cancellation. A facility cancelling its registration must do so within 60 calendar days of the reason for cancellation (e.g., facility ceases operations, ceases providing food for consumption in the United States, or the facility is sold to a new owner).

(b) Cancellation requirements. The cancellation of a facility’s registration must include the following information:

(1) The facility’s registration number;

(2) Whether the facility is domestic or foreign;

(3) The facility name and address;

(4) The name, address, and e-mail address (if available) of the individual submitting the cancellation; and

(5) A statement certifying that the information submitted is true and accurate, and that the person submitting the cancellation is authorized by the facility to cancel its registration.

(c) Electronic cancellation. (1) To cancel your registration electronically, you must cancel at http://www.fda.gov/furls.

(2) Once you complete your electronic cancellation, FDA will automatically provide you with an electronic confirmation of your cancellation.

(3) Your registration will be considered cancelled once FDA transmits your cancellation confirmation.

(d) Cancellation by mail or fax. If, for example, you do not have reasonable access to the Internet through any of
the methods described in §1.231(a), you may cancel your facility’s registration by mail or fax.

(1) You must cancel your registration using Form 3537a. You may obtain a copy of this form by writing to the U.S. Food and Drug Administration (HFS–681), 5600 Fishers Lane, Rockville, MD 20857, or by requesting the form by phone at 1–877–FDA–3882 (1–877–332–3882).

(2) When you receive the form, you must completely and legibly fill out the form and either mail it to the address in paragraph (d)(1) of this section or fax it to 301–436–2804 or 1–800–573–0846.

(3) If the information on the form is incomplete or illegible when FDA receives it, FDA will return the form to you for revision, provided that your mailing address or fax number is legible and valid. When returning a cancellation form for revision, FDA will use the means by which the cancellation was received by the agency (i.e., by mail or fax).

(4) FDA will enter complete and legible mailed and faxed cancellations into its registration system, along with CD-ROM cancellations, as soon as practicable, in the order FDA receives them.

(5) If the information on the form is incomplete or illegible when FDA receives it, FDA will return the form to you for revision, provided that your mailing address or fax number is legible and valid. When returning a cancellation form for revision, FDA will use the means by which the cancellation was received by the agency (i.e., by mail or fax).

(6) You must mail the CD-ROM to U.S. Food and Drug Administration (HFS–681), 5600 Fishers Lane, Rockville, MD 20857.

(7) If FDA receives a CD-ROM that does not comply with these specifications, it will return the CD-ROM to the registrant unprocessed.

(8) FDA will enter CD-ROM submissions that meet the specifications into its registration system, along with complete and legible mailed and faxed submissions, as soon as practicable, in the order FDA receives them.

(9) For each facility on the CD-ROM, FDA will mail to the preferred mailing address a copy of the cancellation(s) as entered and confirmation of the cancellation.

(10) If any information you previously submitted was incorrect at the time of submission, you must immediately resubmit your cancellation.

(11) Your registration will be considered cancelled once FDA enters your facility’s cancellation data into the registration system and the system generates a confirmation.

1.240 What other registration requirements apply?

In addition to the requirements of this subpart, you must comply with the registration regulations found in part

Additional Provisions

§ 1.240 What other registration requirements apply?

In addition to the requirements of this subpart, you must comply with the registration regulations found in part