

contact surfaces shall be smooth, easily cleaned and maintained in good repair. The pump and hose cabinet shall be fully enclosed with tight fitting doors and the inlet and outlet shall be provided with dust covers to give adequate protection from road dust. Tank manholes should be equipped with an adequate filtering system during loading and unloading. New and replacement transport tanks shall comply with 3-A Sanitary Standards for Stainless Steel Automotive Milk and Milk Products Transportation Tanks for Bulk Delivery and/or Farm Pick-up Service.

(3) *Facilities for cleaning and sanitizing.* Enclosed or covered facilities (as climatic conditions require) shall be available for washing and sanitizing of transport tanks, piping, and accessories, at central locations or at all plants that receive or ship milk or milk products in transport tanks.

(d) *Transfer of milk to transport tank.* Milk shall be transferred under sanitary conditions from farm bulk tanks through stainless steel piping or approved tubing. The sanitary piping and tubing shall be capped when not in use.

QUALITY SPECIFICATIONS FOR RAW MILK

§ 58.132 Basis for classification.

The quality classification of raw milk for manufacturing purposes from each producer shall be based on an organoleptic examination for appearance and odor, a drug residue test, and quality control tests for sediment content, bacterial estimate and somatic cell count. All milk received from producers shall not exceed the Food and Drug Administration's established limits for pesticide, herbicide and drug residues. Producers shall be promptly notified of any shipment or portion thereof of their milk that fails to meet any of these quality specifications.

[58 FR 26912, May 6, 1993]

§ 58.133 Methods for quality and wholesomeness determination.

(a) *Appearance and odor.* The appearance of acceptable raw milk shall be normal and free of excessive coarse sediment when examined visually or by an acceptable test procedure. The milk shall not show any abnormal condition

(including, but not limited to, curdled, ropy, bloody or mastitic condition), as indicated by sight or other test procedures. The odor shall be fresh and sweet. The milk shall be free from objectionable feed and other off-odors that adversely affect the finished product.

(b) *Somatic cell count.* (1) A laboratory examination to determine the level of somatic cells shall be made at least four times in each 6-month period at irregular intervals on milk received from each patron.

(2) A confirmatory test for somatic cells shall be done when a herd sample exceeds either of the following screening test results:

(i) California Mastitis Test—Weak Positive (CMT 1).

(ii) Wisconsin Mastitis Test—WMT value of 18 mm.

(3) The confirmatory test for somatic cells shall be performed by using one of the following procedures:

(i) Direct Microscopic Somatic Cell Count (Single Strip Procedure). Pyronin Y-methyl green stain shall be used for goat milk.

(ii) Electronic Somatic Cell Count.

(iii) Optical Somatic Cell Count.

(4) The results of the confirmatory test for somatic cells shall be the official result.

(5) Whenever the confirmatory somatic cell count indicates the presence of more than 1,000,000 somatic cells per ml., the following procedures shall be applied:

(i) The producer shall be notified with a warning of the excessive somatic cell count.

(ii) Whenever two of the last four consecutive somatic cell counts exceed 1,000,000 per ml., the appropriate state regulatory authority shall be notified and a written notice given to the producer. This notice shall be in effect as long as two of the last four consecutive samples exceed 1,000,000 per ml.

(6) An additional sample shall be taken after a lapse of 3 days but within 21 days of the notice required in paragraph (b)(5)(ii) of this section. If this sample also exceeds 1,000,000 per ml., subsequent milkings shall not be accepted for market until satisfactory compliance is obtained. Shipment may

be resumed and a temporary status assigned to the producer by the appropriate State regulatory agency when an additional sample of herd milk is tested and found satisfactory. The producer may be assigned a full reinstatement status when three out of four consecutive somatic cell count tests do not exceed 1,000,000 per ml. The samples shall be taken at a rate of not more than two per week on separate days within a 3-week period.

(c) *Drug residue level.* (1) USDA-approved plants shall not accept for processing any milk testing positive for drug residue. All milk received at USDA-approved plants shall be sampled and tested, prior to processing, for beta lactam drug residue. When directed by the regulatory agency, additional testing for other drug residues shall be performed. Samples shall be analyzed for beta lactams and other drug residues by methods evaluated by the Association of Official Analytical Chemists (AOAC) and accepted by the Food and Drug Administration (FDA) as effective in determining compliance with "safe levels" or established tolerances. "Safe levels" and tolerances for particular drugs are established by the FDA. Other test methods evaluated by the Virginia Polytechnic Institute and State University, or by other institutions using equivalent evaluation procedures, and determined to demonstrate accurate compliance results, may be employed on a temporary basis until they are evaluated by the AOAC and accepted or rejected by the FDA.

(2) Individual producer milk samples for beta lactam drug residue testing shall be obtained from each milk shipment as follows:

(i) *Milk in farm bulk tanks.* A sample shall be taken at each farm and shall include milk from each farm bulk tank.

(ii) *Milk in cans.* A sample shall be formed separately at the receiving plant for each can milk producer included in a delivery, and shall be representative of all milk received from the producer.

(3) Load milk samples for beta lactam drug residue testing shall be obtained from each milk shipment as follows:

(i) *Milk in bulk milk pickup tankers.* A sample shall be taken from the bulk milk pickup tanker after its arrival at the plant and prior to further commingling.

(ii) *Milk in cans.* A sample representing all of the milk received on a shipment shall be formed at the plant, using a sampling procedure that includes milk from every can on the vehicle.

(4) *Follow-up to positive-testing samples.* (i) When a load sample tests positive for drug residue, the appropriate State regulatory agency shall be notified immediately of the positive test result and of the intended disposition of the shipment of milk containing the drug residue.

(ii) Each individual producer sample represented in the positive-testing load sample shall be singly tested to determine the producer of the milk sample testing positive for drug residue. Identification of the producer responsible for producing the milk testing positive for drug residue, and details of the final disposition of the shipment of milk containing the drug residue, shall be reported immediately to the appropriate agency.

(iii) Milk shipment from the producer identified as the source of milk testing positive for drug residue shall cease immediately and may resume only after a sample from a subsequent milking does not test positive for drug residue.

[50 FR 34672, Aug. 27, 1985, as amended at 58 FR 26912, May 6, 1993]

§ 58.134 Sediment content.

(a) *Method of testing.* Methods for determining the sediment content of the milk of individual producers shall be those described in the latest edition of Standard Methods for the Examination of Dairy Products. Sediment content shall be based on comparison with applicable charts of the United States Sediment Standards for Milk and Milk Products, subpart T, § 58.2728 through 58.2732, of this part.

(b) *Sediment content classification.* Milk shall be classified for sediment content, regardless of the results of the appearance and odor examination required in § 58.133(a), as follows: