MINISTRY OF HEALTH AND FAMILY WELFARE

(Food Safety and Standards Authority of India)

NOTIFICATION

F. No.1-110(2)/SP (Biological Hazards)/FSSAI/2010.—The following draft of certain regulations further to amend the Food Safety and Standards (Food Products Standards and Food Additives) Regulations, 2011, which the Food Safety and Standards Authority of India, proposes to make, with previous approval of the Central Government, in exercise of the powers conferred by clause (e) of sub-section (2) of section 92 read with section 16 of the Food Safety and Standards Act, 2006 (34 of 2006), is hereby published as required by the said sub-section (1) of the said Act, for information of all persons likely to be affected thereby, and notice is hereby given that the said draft shall be taken into consideration after the expiry of the period of sixty days from the date on which the copies of the Official Gazette in which this notification is published, are made available to the public;

The Objections or suggestions, if any, duly supported by scientific evidence may be addressed to the Chief Executive Officer, Food Safety and Standards Authority of India, Food and Drug Administration Bhawan, Kotla Road, New Delhi-110002;

The objections and suggestions, which may be received from any person with respect to the said draft regulations before the expiry of the period so specified, will be considered by the Food Authority.

Draft regulations

- Short title and commencement. (1) These regulations may be called the Food Safety and Standards (Food Products Standards and Food Additives) Amendment Regulations, 2015.
- (2) They shall come into force with effect from the ensuing 1st January or 1st July of the year, as the case may be, subject to minimum of 180 days from the date of final notification of these regulations in the official Gazette.
- 2. In the Food Safety and Standards (Food Products Standards and Food Additives) Regulations, 2011, in APPENDIX B relating to "Microbiological Requirements", for TABLE 2 and the entries relating thereto, the following TABLE and the entries shall be substituted, namely:—

"TABLE-2- Microbiological Requirements for Milk and Milk Products:

Table-2A Microbiological Requirements for Milk and Milk Products -Hygiene Indicator Organisms

			ş	Aerobic Plate Count	Ħ		్	Coliform Count		atris	t. aureus	Staph. aureus (Ccagulase positive)	positive)		Yeas	Yeast and mold count	_	Faccal	Faccal streptococci	Ē	
Sr. No.	Product Description	Samp	Sampling	Lini	Limit (cfu)	Sampling	guille a	Limit (cfu)	(a)	Sampling	ling	Limit (cfu)	-	Sampling	20	Limit (cfu)		Sampling		Limit (cfu)	
			u	B	¥		3	6	M	-	٠	=	×	-	٥	6	×	_	-	5	Σ
	Pasteurized Milk/Tlavored Milk	5(3)	m	30 x 10 ³ /mi	50x 10 /mi	(E)¢	0	Absent/0.1ml		5	•	Methylene unit shall the IS 147	Methylene blue reduction test (unit shall not decolorized in 5 tise IS 1479 (Part III-1977-1992	non test (70d in 5 777-1992	MBRT) a	Methylene blue reduction test (MBKT) applicable at Manufacturing unit shall not decolorized in 5 hrs. when tested in accordance with the IS 1473 (Part III.) 1977-1992	ufacturing ance with	1	$\left\{ \right.$		
	Pasteurized Cream	9(3)	3	50x10 ³ /g	75x10³/g	5(3)	0	10/g													
	Steritized/UHT/Flavored Milk / Evaporated Milk																				
	Sterilized/ UHT Cream																				
	Sweetened Condensed Milk	2(3)	~	8/ ₁₀ / ₈	1×10 ² /g	5(3)	.0	Absent/0.1g		8(3)	0	8/01		83	0	10/g					
	Pasteurized Butter ²	2(3)	~	25x10 ³ /g	8/ ₂ 01×05	(5/3)	2	Absent/0.1g	20/g	£3	2	8 ₀ 0	30/8	£3	۳	20/g	8/05				
	Milk powder; SMP, PSMP; Dairy Whitener; Cream powder, Lee Cream Mix powder; Lactose: Whey based powder; Butter Milk powder; Casein powder. ³	\$(3)	7	30x10 ² /g	50x10 ³ /g	(£)	2	Absent 0.1g	30/g	(S)	2	10/g	1×10/g	ĝ.	0	5.0.1g					
	Infant Milk Food, Infant Formulae, Infant Milk Substitute	(S)	7	5x10 ⁷ /g	5x10 ³ /g	≃ €	0	Absent/0.1g		2(3)	0	Absent/C.1g		ŝ	0	Absent/0.1g		~	0	Absent/0.1g	L
0.000	Food , Follow Up Formula			5x10 ² /g	5x10 ³ /g				<u> </u>											VIII VAII (CO	
0.00000	Cereal Based Complimentary			1x103/g	10x10 ³ /g																
	ice Cream, Frozen Dessert, Milk Lolly, Ice Candy etc	(2)3	3	8/ ₅ 01×1	2x10 ³ /g	(2)3	3	10/g	8/ ₂ 01×1	8(3)	2	g/01	1x107/g								
	Processed Cheese Cheese Spread (Ready To Eat Products)	(6)8	2	3/01x52	8/ ₁ 01×05	(6)5	0	8/01		5(3)	0	g/01									
	All Other Cheeses Categories Including Fresh Cheeses / Cheddar /Cottage /Soft /Semi Soft ⁴ etc (Not Ready To Eat Products)					(6)3		1x10 ² /g	5×10 ² /g	8(3)	3	8 _{/01}	lx10'/g	ŝ	9	50/g	250/g				
	Fermented milk products : Yoghurt, Dahi, Chakka, Shrikhand ⁵ etc					£(3)	2	10/g	1×10,/g	\$(3)	2	8/01	1×10 ² /g	ŝ	3	80/8	1×10 ² /g				1
13	Paneeri Chhana/ chhana based sweets	(6)3	3	15x10 ⁷ /g	35×10*/g	(5)3	3	10/g	8/,01×1	(£)\$	3	3/01	1×10//g	ŝ	3	8/05	8/051				
	Khoa/ khoe based sweets	8(3)	۳.	25x10 ³ /g	3/201x57	5(3)	2	\$0/g	3/,01×1	(2)3	3	10/g	1×10/g	ŝ	3	10/8	\$/05				
	Method of Analysis			15 5402	100		2	IS 5401, Part 1			885 5	15 5887. Part 8/Sec				IS 5403		-	7 4887 2	15 5997 (Part 3) : 1076	2

Pasteurized milk shall be stored at manufacturing unit and at retail points in such a way that temperature of milk shall not exceed 8°C as recommended in IS 13688. 1999
The microbial specifications for ripened butter are the same as for pasteurized butter excluding the requirements of total plate count.

¹The yeast and mold count of 5/0. Ig as specified in dried product categories shall be applicable only to cascin powder

¹The yeast and mold counts in all other cheese category will be "pplicable except mold ripened cheeses.

¹The standard requirements of factic counts of one million c.f.u./g min as specified by BIS (1S:12898:1989) in such products/ or such products containing probiotic organisms shall be applicable () figures in parenthesis indicate no. of samples required for testing as refating as refating to the counts.

Table-2B: Microbiological Requirements For Milk and Milk Products -Safety Indicator Organisms

		L		E.coli		S	almon	Salmonella / Shigella		7	L. monocytogenes	genes	_		B. cereus	2		Salp	hite Red	Sulphite Reducing Clostridia (SRC)	tridia		Entero	Enterobacter sakazakii	ı	
		3	Sampling	Limit (cfu)	_	Samplin g plan	Ę =	Limit (cfu)		Sampling	94	Limit (cfu)	2 E	Sampling plan	2	Limit (cfu)		Sampling plan	g.	Limit (cfu)		Sampling plan	8	Limit (cfu)		
100		•	٥	E	Σ	-	-	8	×	-	u	2	¥.	Н	J	2	Σ	•	J	£	M	.	J	8	×	-
_	Pasteurized Milk/Flavored Milk	<u>(5)</u>	0	Absent/ 0.1ml		5(3)	0	Absent/25 ml		8(3)	0	Absent/25ml		9	0	hospitat -19"	se test sha 7) before	uli be ne release	gative a	Phospinatase test shall be negative as per 1S 8479 (Parts i-1977) before release of product in the market	79 (Parts riket					
1~	Pasteurized Cream	83	0	Absent/0.1g		2(3)	0	Absent/25g		5(3)	0	Absent/25g	Н	\vdash												-
-	Sterilized/UHT/Flavored	5(3)	0				•	Sterilized milk	produc	ts shall	comply	Steritized milk products shall comply with a test for commercial sterility as per BIS (IS: 4238-1967)	ommerci	al sterility	as per	13 (13:7	238-1967						,			
	Sterilized/ UHT Cream	(5)(9					Sterifized cream	poud u	ict shall	сошр	Sternized cream product shall comply with a test for commercial sterlity as per BIS (18: 4884:1968	ommerc	al sterility	88 77	SIS (IS:	4884:196	_								_
~	Sweetened Condensed	\$3	6	Absent/0 lg		<u>§</u>	0	Absent/25g		5(3)	0	Absent/g	-	5(3)	0	Profice	shall comp	ly accel	accelerated st	Product shall comply accelerated storage test as per IS	s per (S					
1	Doctourized Butter	5/3	0	Absent/0.1g		Š	0	Absent/25g		õ	0	Absent/g												್ಷ		-
٠	VIII. COMPANIES CAND	+-	+	1	L	Ş	+-	Absent/25g	T	8	0	Absent/25g	5	5(3)	3	1×10-/E	1×10'/g	5(3)	3	10/g	1×10 ² /g					
	Dairy Who powder Mix p					?				}		0	<u></u>							9						
	powder, Casein powder	_	-				_				7	7.	\dashv	+	_											_
∞	Infant Milk Food, Infant	-	0	Absent/0.1g	Ŀ	15	0	Absent/25g		2	•	Absent/25g	<u>~</u>	5(3)		1×10/E	5×104/8	2(3)	•	10/g	1x40/g	ຊ ີ	•	Abscnt/10g		_
	Formulae, Infant Milk	-	્			<u>ଚ</u>				<u>(S</u>					-							3				
٠,	Follow Up Formula				_		_		• • •										18.50							
	Cereal Based	_																								
	Compilmentary rood	+	+	4	1	5	+	_	ľ	ŝ	6	Abcentio	H	H	t	Ī			L			L				~
6	Lee Cream, Frozen Dessert, Milk Lolly, Ice Candy etc	×	o (5)	Absenug		ć,	-	Ausen/22g		lc)c	,	Saleston														
2			5(3) 0	Absenug		(6)3	0	Absent/25g		2(3)	0	Absent / 25g		2(3)		Canned	roduct sh	all comp	1785:19	Canne d product shall comply accelerated storage test as per IS :2785:1964	ge test as					
<u> =</u>	 		5(3)	0 Absenv0.1g		£(3)	0	AbsenVg		(3)	0	Absent/25g								10						
2	+-	1	5(3)	0 Absent/g	<u> </u>	2(3)	0	Absent/25g		\$(3)	0	Absent/g														
2	Shrikhand etc Paneer/ Chhana/ chhana	+	(5)	0 Absent/0.1g	-	\$3	0	Absent/g	_	\$(3)	0	Absent/g		T	T	T	Γ		L			L				_
ل	7	+	+	-	+	Ś	+	Absentio	1	5/3	G	Absent/e	İ	t	T	T	T		L			\perp				_
4	7	+	(6)6	IC COOT Dard I		1	13	7 Part 36 7	L		2	IS 14988-Part 1	T	1	15 5887, Part 6	Part			8	50 15213:2003		L	150	ISO / TS 22964:2006	99	1
لـ	Method of analysis	\dashv		13 3007, FREE		7	2	, 1 41 41 41	1		1		1				1									7

The Sweetened Condensed milk product shall comply accelerated storage test as per IS: 1166-1986 where sample shall be stored at 37± I°C for 14 days.

Processed cheese packed in canned shall comply accelerated storage test when incubated at 30°C for 14 days as per IS: 2785:1964 and should give no bulging.

() figures in parenthesis indicate no of samples required at retail points.

Sampling Guidelines: The sampling for different microbiological parameters specified in Table-2A and B shall be ensured aseptically at manufacturing units as well as at retail points us per the sampling plan given in Table-2C by a trained person with specialised knowledge in the field of microbiology following guidelines given in IS 11546:1999 /ISO 707:1985 (Reaffirmed 2010). The samples shall be stored and transported under appropriate temperature conditions and insulations within 24 hrs of sampling to accredited laboratories for analysis as per the approved sampling plan and test methods? A large sample size may be drawn (if desired) according to the tests required and the type of product. Preservatives shall not be added to samples intended for microbiological examination. The desired number of sampling plan given in Table-2A & B shall be taken from full production batches and will be submitted to accredited laboratory in original unopened packaging, scaled at the time of sampling maintained in their original physical state. The final decision shall be drawn based on results with no provision for retesting for microbiological parameters.

Sampling plan 9: The following terms, as used by the International Commission on Microbiological Specifications of Foods (ICMSF) are defined and used in this start dard:

n= the number of sample units which must be examined from the batch or lot of food to satisfy the requirements of a particular sampling plan.

m= Represents an acceptable level and values above it are marginally acceptable in terms of the sampling plan.

c= the maximum allowable number of defective sample units in 3-class sampling plan applicable at manufacturing units only.

M= A microbiological criterion which in dicate unsatisfactory or potentially hazardous quality. Values above M are unacceptable in terms of the sampling plan and detection of one or more samples exceeding this level would be cause for rejection of the lot and will attract prosecution by the concerned food safety authorities.

Table-2C Sampling plan for compliance of microbiological parameters at manufacturing units and at retail points

Sampling area		ē!				Samoling Plan	
			2-Class			200	1.00 mm
	E	J	E	6	3	E	No.
Manufacturing unit	n=5-30	E	Values of m specified for all testing parameters for different products as specified Table-2A &B will be applicable	5-30	c=23	Values of m specified for all testing parameters for different produce as specified Table- 2A&B will be applicable	Values of M specified for all testing parameters for different products as specified Table-2A & B will be applicable
Retail point	n=3-10	Ţ	Values of m specified for all testing parameters for different products as specified Table-2 A&B will be amplicable.	3-10	Į,	Not applicable	Values of M specified for all testing parameters for different products as specified in Table- 2A &B will be applicable

Microbiological criteria and their interpretation: Following three categories of microbiological quality have been assigned in standard based on hygiene and sufety indicator organisms. These are satisfactory, unsatisfactory and potentially hazardous.

1. Satisfactory: if the test values of m or M or both applicable within the sampling plan are conforming the specified limits, the microbiological quality of product is considered satisfactory and no action is required.

2. Ussatisactory: if the test values of m or M or both applicable within the sampling plan are not conforming the specified limits of hygiene indicators i.e. aerobit count, coaliform count, coagulase positive S. aureus, faceal streptococci and yeast and mold count etc which indicates poor hygiene or poor handling practices, the microbiological quality of product will be considered unsatisfactory. Inder these conditions the premises producing such unsatisfactory products, shall be investigated for nonconformity or non-compliance and legal action on defected products will be notified by the food safety authority. The subsequent release of such product shall be subject to HACCP / OMP audit clearance of the premises / finished products by the food safety authority.

release of subsequent batches of such hazardous products will be under hold by the food manufacturers. Failure by an owner to either cease manufacture of product or withdraw/recall product from sale when requested to do so shall result in seizure of that product where the officer has reason to believe that it is contaminated with pathogenic bacteria. A detail risk assessment will be carried out to determine by the food safety authority to investigate the source or cause of the problem so that 3. Potentially hazardous: if the test values of m / or M or both applicable within the sampling plan are not conforming the specified limits of safety indicators i.e., L. cofi, Salmonella / Shigella, B. cereus, Sulphite Reducing Chostridia (SRC), Enterobacter sakazakli. L. monocyogenes and sterility tests etc. which indicates serious food safety concern, the microbiological quality of product will be considered as potentially hazardous. Under these conditions the premises producing such unsaits sectory product(s) shall be stopped and legal actions on potentially hazardous products will be notified by the food safety authority. The recall action on with lawal of any of such food still available for sale or distribution shall be initiated and emedial action can commence and the approval for rectart of such products under non-conformity will be allowed only after compliance of manufacturing unit for food safety standards requirements or guidelines set by the Authority.

erence test methods?

	Test reference	
_	Microbiology - general guidance for the enumeration of micro-organisms- colony count technique at 30°C (first revision)	IS 5402:2002/ ISO:4833:1991 reaffirmed 2007
+		IS 5401 (Part 1): 2002/ISO 4832:1991 reaffirmed 2007
1	Methods for detection of bacteria responsible for food poisoning. Part 8 Horizontal method for enumeration of coagulace-positive reaffirmed stankylococci (Suphylococcus aureus and other species) Section 1 Technique using Baird-Parker Agar Medium	15 5887(Part 8/Sec 1):2002 / ISO 6888-1 :1999 reaffirmed 2007
	Isolation, identification and enumeration of Staphylococcus aureus and faceal streptococci (first revision)	5887 (Part 2): 1976
1	Milk and milk products — detection of Enterobacter sakazakin (First Edition)	ISO/TS 22964: 2006
	Method for yeast and mould count of food stuffs and animal feeds(first revision)	IS 5403:1999 reaffirmed 2005/ ISO 7954:1987 reaffirmed 2009
	Methods for detection of bacteria responsible for food poisoning: isolation, identification and enumenation of Escherichia coli (first revision.)	IS 5887(Part I):1976 reaffirmed 2009 Part 1
	Methods for detection of bacteria responsible, for food poisoning. Part Digeneral guidance on methods for detection of Summelia Singelia (second revision) Methods for detection	15 5887 (Рап. 3); 1999і 150 6579; 1993 геатіттес 21хм
	Methods for detection of bacteria responsible for food poisoning-Part 7: general guidance on methods for isolation and identification of Skige."In	IS 5887(Part 7):1999/ ISO 6579: 1993 reaffirmed 2009
1	Microbiology of food and animal feeding stuffs - horizontal method for detection and enumeration of Listeria monocytogenes: Part 1 Detection nethod	IS 14988(Part 1):2001/ ISO 11290-1:1996 reaffirmed 2007
	Methods for detection of bacteria responsible for food poisoning. Part 6 identification, enumeration and confirmation of B.cereus	IS 5887(Part 6):1999 / ISO 7932:1993 reaffirmed 2007
1	Microbiology of food and animal feeding stuffs- Horizontal method for the enumeration of sulfite reducing bacteria growing under anacrobic conditions	ISO 15213: 2003 -05-01
	Methods of test for dairy industry part III bacteriological analysis of milk (first revision)	IS 1479 (Part III-1977- reaffirmed 1992
	Indian Standard Specification for sterilised milk	IS: 4238-1967 reaffirmed 2010
1	Specification for starilized cream	BIS (IS: 4884:1968) reaffirmed 1999
	Packaged pasteurized milk - specification (first revision)	IS 13688: 1999
1	Method for determination of phosphatase activity in milk and milk products:	IS 8479 (Parts I):1977 reaffirmed 1997
	Specification for condensed milk, partly skimmed and skimmed condensed milk (second revision)	IS:1166-1986 reaffirmed 1997
1	Hand cheese, processed cheese and processed cheese spread	IS:2785:1964
1	Methods of sampline for milk and milk products	IS 11546:1999/ ISO 707:1997 reaffirmed 2010
	The state of the s	IS: 12898:1989 reaffirmed 1994**

For IS Standards recent version shall apply.

Note.—The principal regulations were published in the Gazette of India, Extraordinary vide notification number F. No. 2-15015/30/2010, dated the 1st August, 2011 subsequently amended vide notification numbers:

- (1) F. No. 4/15015/30/2011, dated the 7th June, 2013;
- (2) F. No. P.15014/1/2011-PFA/FSSAI, dated the 27th June, 2013;
- (3) F. No. 5/15015/30/2012, dated the 12th July, 2013, and
- (4) F. No. P.15025/262/13-PA/FSSAI dated the 5th, December, 2014.