CHARM® II BETA-LACTAM ASSAYS

APPENDIX N BULK MILK TANKER SCREENING TEST FORM

Competitive (Raw Commingled Cow Milk and Pasteurized White Milks) IMS #9-C2
Sequential (Raw Commingled Cow and Goat Milk) IMS #9-C3
Quantitative (Raw Commingled Cow Milk) IMS #9-C4
Cloxacillin (Raw Commingled Cow Milk) IMS #9-C9

[Unless otherwise stated all tolerances are ±5%]

GENERAL REQUIREMENTS

1.	See	ee Appendix N General Requirements (App. N GR) items 1-8 & 15					
			SAMPLES				
2.	See	App	p. N GR item 9				
			APPARATUS & REAGENTS				
3.	Equ	ipm	ent				
	a.	Ana	alyzer heater for 13 x 100 mm tubes				
		1.	85±2°C for Competitive Assay				
		2.	65±2°C for Sequential Assay				
		3.	55±2°C for Quantitative Assay				
		4.	35±2°C for Cloxacillin Assay				
		5.	Temperature checked by electronic display, or by placing accuracy checked temperature measuring device in tube containing liquid (bulb submersed) in heating unit; maintain records				
		6.	Or, use 6 inch partial immersion thermometer placed directly into small thermometer well in middle of heating unit; maintain records				
		7.	Temperature measuring device for each incubator (App. N item 3)				
	b.	Mix	ker, Maxi-mixer II or equivalent				
	C.	Cer	ntrifuge, Whisperfuge [®] or Heraeus [®] (3400 rpm) or equivalent				
	d.	Scii	intillation counter. Charm II or equivalent				

	e.	Scintillation fluid dispenser, set to dispense 3 mL						
		1.	·	months with Class A graduated cylinder and				
	f.	Cot	ton swabs					
	g.	Bor	osilicate test tubes, 13	x 100 mm				
	h.	Plas	stic stoppers for tubes					
	i.	Pipe	ettors - Fixed Volume	or Electronic (see App. N GR item 7)				
		1.	300 µL and appropria	ate tips				
		2.	5.0 mL and appropria	ate tips				
	j.	Tim	er					
١.	Rea	igent	s					
	a.	Scintillation fluid – Optifluor or equivalent supplied by manufacturer of test kits						
	b.	Competitive, Sequential or Quantitative Assay						
		Reagent blister packages: microbial binder (green) tablet, tracer reagent (yellow) tablet						
			Lot #:	Exp. Date:				
		2.	0.008 IU/mL Penicilli	n G standard				
			Lot #:	Exp. Date:				
		3.	Zero control standard	d				
			Lot #:	Exp. Date:				
	C.	Clo	xacillin Assay					
		1.	Reagent blister pack tracer reagent (blue)	ages: microbial/antibody binder (white) tablet, tablet				
			Lot #:	Exp. Date:				
		2.	10 ppb Cloxacillin sta	andard				
			Lot #:	Exp. Date:				

		3.	Zero control standard	
			Lot #: Exp. Date:	
5.	Rea	igeni	t stability	
	a.	All t	tablet reagents stored at -15°C or below	
	b.		sitive Control – Lyophilized 0.008 IU/mL penicillin G or 10 ppb oxacillin standard for Cloxacillin assay	
		1.	Reconstitute with 100 mL (measured) Negative Control (allow to sit 15 min prior to use or aliquotting)	
			Lab Prep. Date: Lab Exp. Date:	
		2.	For Quantitative Only: Dilute reconstituted 0.008 IU/mL Penicillin G standard 1:4 with Zero Control Standard	
		3.	Use within 48 hours when stored at 0.0-4.5°C	
		4.	Or, aliquot within 24 hours and freeze at -15°C or colder in a non frost-free freezer or in an insulated foam container in a frost-free freezer; use within 2 months	
			Lab Prep. Date: Lab Exp. Date:	
			a. Thaw and use within 24 hours. Store at 0.0-4.5°C	
	C.		gative Control – Lyophilized Zero Control Standard (ZCS) or ernatively, raw milk qualified to test similar to ZCS	
			Lab Prep. Date: Lab Exp. Date:	
		1.	Reconstitute ZCS according to manufacture instructions. (Allow to sit 15 min prior to use or aliquotting)	
			 To qualify raw milk, test sample 3 times and average results. Average must be within ± 10% of ZCS 	
			Lab Prep. Date: Lab Exp. Date:	
		2.	Use within 72 hours when stored at 0.0-4.5°C	
		3.	Or, aliquot within 24 hours and freeze at -15°C or colder in a non frost-free freezer or in an insulated foam container in a frost-free freezer; use within 2 months	
			Lab Prep. Date: Lab Exp. Date:	
			a. Thaw and use within 24 hours. Store at 0.0-4.5°C	

	d.	Scir	ntillation fluid expires s	ix (6)	months after opening	
		Date	e Opened:	L	ab Exp. Date:	
					TECHNIQUE	
6.	nev lact	v lot (of reagents. Steps 6, creening methods ar	7, an	ol Average to be determined for each d 8 are for the various Charm betas operator choice which method is	
	a.	Con	npetitive Assay Contro	l Poir	nt (CP) and Negative Control Average	
		1.	Run six 0.008 IU/mL Pen G	2.	Run three Negative Controls	
			Penicillin G		Negative Control	
	+	2. 3. 4. 5. 6. Av.		2. 3.		
	b.	Seq	uential Assay Control	Point	(CP) and Negative Control Average	
		1.	Run six 0.008 IU/mL Pen G	2.	Run three Negative Controls	
			Penicillin G		Negative Control	
	+	1. 2. 3. 4. 5. 6. Av. -25% CP		2. 3.		

C.	Quantitative Assay Control Point (CP) and Negative Control Average				
	1.	Run six Negative Controls	2.		
		Negative Control		Penicillin G	
_	3. 4. 5. 6.		2. 3.		
d.	Clox	kacillin Assay Control	Point	(CP) and Zero Control Average	
	1.	Run six 10 ppb Cloxacillin	2.	Run three Negative Controls	
		Cloxacillin		Negative Control	
+	1. 2. 3. 4. 5. 6. Av. 15% CP		2.		
Acc	epta	bility of Control Poin	t Det	erminations	
a.		ny of the 6 control poin that determination	nt dete	erminations deviate from the average,	
	1.	For Competitive Ass	ay ca	nnot deviate by more than ±15%	
	2.	For Sequential Assa	y can	not deviate by more than ±25%	
	3.	For Quantitative Ass	ay ca	nnot deviate by more than ±15%	
	4.	For Cloxacillin Assay	/ canr	not deviate by more than ±15%	
b.		e re-determined value average and proceed		thin the allowed deviation recalculate testing	

7.

	C.	If the value is not within allowed deviation, run another set of six (6) standards			
	d.	A co	ommon control point for multiple analysts may be used		
		1.	Control point determination performed by one analyst only		
		2.	Control point determination rotated and inclusive of all certified/approved analysts		
		3.	If daily performance check fails and is not resolved by using fresh controls, technique should be reviewed for consistency and corrective action taken as necessary		
8.	Dail	ly Pe	rformance and Operation Check (also see App. N GR item 10)		
	a.		e negative control tests ±20% (±15% for Quantitative Assay) ablished for each new kit lot		
	b.	The	positive control tests less than or equal to the control point		
	C.	If th	ese conditions are not met re-determine control point(s)		
		1.	Conditions met, proceed with testing		
		2.	Conditions not met, discontinue testing and seek technical assistance		
9.	Bet	a-lac	tam (all except Cloxacillin) Test Procedures		
	a.	Lab	el test tubes, one for each test sample		
	b.	Add	1 1 green tablet to each tube		
	C.	Add	I 300 μL water to each tube		
	d.	and	akup tablets in tubes by mixing tubes 10 times on mixer in a rise fall motion in 10 sec, if necessary continue mixing, green tablets st be completely suspended before proceeding		
	e.	vort	milk sample(s)/control(s) 25 times in 7 sec with a 1 ft movement or ex for 10 sec at maximum setting, use within 3 min (samples must n appropriate container to allow the use of vortexing)		
	f.	Add	1 5.0 mL of mixed sample/control to corresponding tube		
		1.	Using pipettor (item 3.i.2) with new tip for each sample/control, draw up 5 mL avoiding foam or bubbles		
		2.	Remove tip from liquid		
		3.	Expel test portion into appropriate tube		

g.	Con	npetit	tive Assay			
	1.		following steps must be completed within 40 sec (all sample es being assayed)			
		a.	Add yellow tablet to each tube			
		b.	Vortex tubes 10 times in a rise and fall motion in 10 sec (yellow tablets do not breakup)			
	2.	Incu	ubate tubes for 3 min at 85±2°C			
	3.		move tubes and centrifuge for 3 min; optionally for 5 min me time used to determine control point)			
	4.	Skip	o to item 11			
h.	Seq	uenti	ial Assay			
	1.	Vort	tex tubes 10 times in a rise and fall motion in 10 sec			
	2.	Incu	ubate tubes for 2 min at 65±2°C			
	3.		e following steps must be completed within 40 sec (all sample es being assayed)			
		a.	Add yellow tablet to each tube			
		b.	Vortex tubes as in item 9.h.1 above			
	4.	Incu	ubate tubes for 2 min at 65±2°C			
	5.		move tubes and centrifuge for 3 min; optionally for 5 min me time used to determine control point)			
	6.	Skip	o to item 11			
i.	Qua	ıntitat	tive Assay			
	1.	Vort	tex tubes 10 times in a rise and fall motion in 10 sec			
	2.	Incubate tubes for 7 min at 55±2°C				
	3.		following steps must be completed within 40 sec (all sample es being assayed)			
		a.	Add yellow tablet to each tube			
		b.	Vortex tubes as in item 1 above			
	1	Inci	thate tubes for 2 min at 55±2°C			

		5.	Remove tubes and centrifuge for 3 min; optionally for 5 min (same time used to determine control point)	
		6.	Skip to item 11	
10.	Clo	xacill	lin Test Procedure	
	a.	Con	npetitive Assay	
		1.	Mix milk sample(s)/control(s) 25 times in 7 sec with a 1 ft movement or vortex for 10 sec at maximum setting, use within 3 min (samples must be in appropriate containers to allow the use of vortexing)	
		2.	Fill identified test tubes ¾ full with milk samples, avoiding foam and bubbles, and centrifuge for 5 min	
		3.	Cool tubes to 0.0-4.5°C	
		4.	Label empty test tubes, one for each test sample	
		5.	Add 1 white tablet to each new empty tube	
		6	Add 300 µL water to each tube	
		7.	Breakup tablets in tubes by vortexing tubes 10 times on mixer in a rise and fall motion in 10 sec, if necessary continue vortexing, white tablets must be completely suspended before proceeding	
		8.	Draw up 5.0 mL of centrifuged sample/control from below the fat layer	
			a. Use new tip for each sample/control	
			b. Remove tip from liquid	
			c. Expel test portion into appropriate tube	
		9.	The following steps must be completed within 40 sec (all sample tubes being assayed)	
			a. Add blue tablet to each tube	
			b. Vortex tubes 10 times in a rise and fall motion in 10 sec (blue tablets do not breakup)	
		10.	Incubate tubes for 3 min at 35±2°C	
		11	Remove tubes and centrifuge for 5 min	

11.		er Centrifugation Step in Beta-Lactam (9.g.3, 9.h.5, and 9.i.5) and xacillin (10.a.11) Test Procedures					
	a.	Immediately pour off milk					
	b.	While still draining tubes, remove fat ring with 2 or more cotton swabs, continue until dry, do not touch pellet (do not go much below the fat ring)					
	C.	Add 300 µL of water to tubes and break up pellets using vortex mixer					
	d.	Pellets must be completely suspended before proceeding to next step					
	e. Add 3 mL of scintillation fluid to each tube, cap and vortex or shake until uniformly mixed						
	f.	Count tubes on scintillation counter for 1 min using [14C] channel					
	g.	Record counts as counts per minute (CPM)					
12.	Inte	rpretation					
	a.	If the beta-lactam assay (not applicable to Cloxacillin Assay) result in the analyzer is at least 50 points greater than the control point, then the sample result is Negative (NF)					
	b.	If Cloxacillin assay result is greater than the control then the sample is Negative (NF)					
	C.	If the beta-lactam assay result in the analyzer is less than or equal to the control point then the sample is Presumptive Positive					
	d.	If the beta-lactam assay (not applicable to Cloxacillin Assay) result in the analyzer is less than 50 points greater than the control point, then the sample must be re-counted					
		If on re-count the result is greater than the control point, then the sample is Negative (NF)					
		If on re-count the result is equal to or less than the control point, then the sample is Presumptive Positive					
13.	Con and Ass or C	fication of Initial Positive Samples (see App. N GR item 11); Ifirmation of Presumptive Positive Samples (see App. N GR item 12); Producer Traceback (see App. N GR item 13). For Quantitative ay: PROMPTLY retest the SAME sample using the Sequential Assay Competitive Assay, and when these beta-lactam assays give Not nd [NF] the Cloxacillin Assay is required					
11	Don	orting (see Ann. N.G.P. item 14)					

15.	Handling of Exempt Quantities of Radioactive Materials						
	a. No mouth pipetting						
	 No smoking, eating or use of cosmetics while reagents are being handled 						
	C.	Nuclear Regulatory Commission (NRC) licensed facilities must meet requirements as they relate to the use of gloves, other protective measures, and handling of wastes					
	d. Wash hands thoroughly after handling reagents						
	e.	Wipe up spills immediately and thoroughly					
	f. Properly dispose of all contaminated waste						