



AMR TECHNICAL SCORECARD

VETERINARY

Bacterial Culture, Detection,
Identification and Antimicrobial
Susceptibility Testing of Milk
Samples



Version 1.1 – August 2021





Score

Section	Sum of	Curren	nt Audit	Previous audit	
	maximum	Date:		Date:	
	points ¹	Currer	nt audit	Previou	ıs audit
		SC	ore	SCO	ore
1. Documents and Records			%		%
2. Management Reviews			%		%
3. Organization and Personnel			%		%
4. Client Management and Customer Service			%		%
5. Equipment			%		%
6. Evaluation and Audits			%		%
7. Purchasing and Inventory			%		%
8. Process Control and Internal and External Quality Assessment			%		%
9. Information Management			%		%
10. Corrective Action			%		%
11. Occurrence Management and Process Improvement			%		%
12. Facilities and Safety			%		%
Milk Module Total			%		%
Milk Module Stars ²					

 $^{^{\}mbox{\tiny 1}}$ Total number of points of all questions minus points for questions answered with NA. $^{\mbox{\tiny 2}}$ No Stars: < 55%

¹ Star: 55% - 64%

² Stars: 65% - 74%

³ Stars: 75% - 84%

⁴ Stars: 85% - 94%

⁵ Stars: ≥95%

A. General Information

Name of assessor(s)			
Title & organization of assessor			
Name of laboratory being assessed			
Date, type and scope of last assessment?	Date	Туре	Score
Internal			
External			
Did the last assessment include assessment of bacterial culture of feces?		Y/N	

B. Technical Information

M-A How many milk cultures and molecular tests were performed last year^{3,4}?

M-A How many milk o			ulture				cular ⁵		Clinical diagnosis	Active surveillanc
										е
	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Entire year	Entire year
Farms: Bulk milk										
S. aureus										
S. agalactiae										
S. uberis										
S. dysgalactiae										
C. bovis										
K. pneumoniae										
E. coli										
P. aeruginosa										
Mycoplasma spp.										
Other isolates (specify)										
Other isolates (specify)										
Other isolates (specify)										
Farms: Individual cows										
S. aureus										
S. agalactiae										
S. uberis										
S. dysgalactiae										
C. bovis										
K. pneumoniae										
E. coli										
P. aeruginosa										
Mycoplasma spp.										
Other isolates (specify)										
Other isolates (specify)										
Other isolates (specify)										
Unknown / other®										
S. aureus										
S. agalactiae										
S. uberis										
S. dysgalactiae										
C. bovis										
K. pneumoniae										

³ It is highly recommended that assessors obtain the necessary permission to review the laboratory data. However, if assessors are unable to review the laboratory data this question is NOT compulsory for completion of the assessment.

⁴ Refer to the World Organisation for Animal Health (OIE) for further information on OIE listed diseases and other diseases of importance: https://www.oie.int/en/international-standard-setting/terrestrial-manual/access-online/

⁵ Molecular tests performed on milk for the detection of bacterial milk pathogens.

⁶ If the laboratory cannot distinguish between samples originating from farms or slaughterhouses, the number of organisms isolated should be recorded as "Unknown/other".

E. coli					
P. aeruginosa					
Mycoplasma spp.					
Other isolates (specify)					
Other isolates (specify)					
Other isolates (specify)					
TOTAL NUMBER OF					
ISOLATES					
TOTAL NUMBER OF					
MILK CULTURES					
PERFORMED					
TOTAL NUMBER OF					
CONTAMINATED MILK					
CULTURES					
TOTAL NUMBER OF					
NEGATIVE MILK					
CULTURES					

Q = Quarter

М-В	Are there any significant variations (> 20%) in the number of milk culture or molecular tests performed or organisms isolated each quarter? If 'Yes', please explain

Section 1: Documents & Records

All generic requirements apply, see SLIPTA Section 1. In addition, assessors should review the following:

SLIPT			N	Υ	Р	N	Comments	Score
SLIPT A 1.5	M1.1	Does the laboratory have documentation covering the following processes? a) Production of Blood Agar, MacConkey Agar or other media for milk pathogen isolation? b) Microscopic examination and somatic cell count c) Processing of milk culture and molecular tests	N A	Y	P	N	Comments	Score 2
		d) Detection, identification and AST of milk pathogens e) Reporting of milk culture and molecular test results f) Interlaboratory comparison or proficiency testing						
		(PT) g) Laboratory safety	-					
1.5	M1.2	Are the documents complete, in-date and witnessed by all staff performing milk culture and molecular tests ⁷ ?						2
1.5	M1.3	Are the following processes documented? a) Rejection criteria for milk samples? b) How to identify potential pathogens on all primary media? (SOP should describe colony appearance of						3

 $^{^{7}}$ See ISO15189:2012 Clause 5.5.3 for minimum requirements for a technical Standard Operating Procedure (SOP).

⁸ For more information see OIE Manual of Diagnostic Tests and Vaccines for Terrestrial Animals 2019 <u>chapter 1.1.2:</u> <u>Collection, submission and storage of diagnostic specimens</u>

		potential pathogens and define how to proceed when a potential pathogen					
		is encountered)					
	c)	Semi-quantitative					
	'	culture for milk					
		samples?					
	d)	Which organisms to					
		identify based on					
		relative quantities (>					
	e)	10 ⁴ CFU / ml)? Instructions for					
		referral of milk					
		culture and					
		molecular tests not					
		performed at the					
		laboratory?					
	f)	Instructions for					
		handling samples received after					
		hours?					
	g)	Instructions for					
	0,	referral of bacterial					
		isolates for					
		identification and					
	In V	AST?					
	h)	Instructions on how to determine AST					
		conversions for					
		automated, disk					
		diffusion,					
		Etest/Gradient and					
		microdilution AST?					
	i)	Definition of rare/					
		unexpected AST results?					
	j)	Confirmatory tests					
	17	for unusual or					
		unexpected patient					
		AST results?					
	k)	Turnaround time for					
		milk culture and					
		molecular tests ⁹ ?					
Section	1: Docui	ments & Records S	ubto	tal			7

Section 2: Management Reviews

⁹ From sample collection to reporting.

Section 3: Organization & Personnel

All generic requirements apply, see SLIPTA Section 3. In addition, assessors should review the following:

followin	g:							
SLIPT			N	Υ	P	N	Comments	Score
Α			Α					
3.6	M3.1	Is there evidence that laboratory staff have been trained in the following ¹⁰ :						
		a) Microscopic examination and somatic cell count						
		b) Processing of milk samples for culture and molecular tests						
		c) Detection / identification and AST of milk pathogens						3
		d) Interpretation of milk culture and molecular test results						
		e) Reporting of milk culture and molecular test results						
		f) QC for milk culture and molecular tests						
		g) Laboratory safety						
3.7	M3.2	Is there evidence that laboratory staff are following the procedures described in the laboratory documentation?":						
		a) Microscopic examination and somatic cell count						2
		b) Processing of milk samples for culture and molecular tests						3
		c) Interpretation of milk culture and molecular test results						
		d) Identification and AST of milk						

Review training records, competency assessment forms and duty rosters. Pay attention to date of training and scope of training compared with techniques being performed.

11 Directly observe procedures being performed compared to the SOP.

	pathogens						
	e) Reporting of milk						
	culture and						
	molecular test						
	results						
Section 3: Organization & Personnel Subtotal							

Section 4: Client Management & Customer Service

All generic requirements apply, see SLIPTA Section 4. In addition, assessors should review the following:

SLIPT A	Ĭ		N A	Υ	Р	N	Comments	Score
4.1	M4.1	Is there evidence that the laboratory has provided clients with information/instructions on milk collection, storage and transportation to the laboratory?						3
4.1	M4.2	Is there evidence that the laboratory has provided clients with information/instructions on interpretation of milk microscopy, culture and molecular results and AST?						2
Section	n 4: Cl	ient Management & C	usto	mer	Serv	ice S	Subtotal	5

Section 5: Equipment

Section 6: Evaluation and Audits

Section 7: Purchasing & Inventory

All generic requirements apply, see SLIPTA Section 7. In addition, assessors should review the following:

SLIPT A	ŭ		N A	Υ	Р	N	Comments	Score
7.10	M7.1	Is all media for bacterial culture isolation, identification and AST stored correctly and in date (from date of manufacture media						2

SLIPT A		N A	Υ	Р	N	Comments	Score
	must be stored at 2-8°C) ¹² ?						
	Blood Agar						
	MacConkey agar						
	M-17 or equivalent						
	Mueller Hinton						
Section	7: Purchasing & Inventory	Sub	total				2

Section 8: Process Control

All generic requirements apply, see SLIPTA Section 8. In addition, assessors should review the following:

SLIPT	191		N	Υ	Р	N	Comments	Score
Α			Α					
SPECIA	MEN CO	LLECTION						
8.5	M8.1	If milk samples will reach the laboratory more than 2 hours post collection, are they transported to the laboratory on ice?						2
		Y CONTROL				T		
8.8	M8.2	Does the laboratory perform QC testing on all media before use ¹³ ?						
		Blood agar						
		Do QC records for blood agar plates demonstrate that they are checked for their ability to support growth of fastidious organisms such as <i>S. pneumoniae</i> ?						
		Do QC records for blood agar plates demonstrate that they are checked for their ability to show beta, alpha, and gamma hemolysis?						3
		MacConkey agar (MAC)	1		1		
		Do QC records for MAC plates demonstrate that they are checked for their ability to suppress growth of Gram -positive organisms while						

¹² According to manufacturer's requirements.
13 This includes in-house made or purchased from commercial sources.

SLIPT A			N A	Υ	Р	N	Comments	Score
		allowing the growth of Gram -negative organisms? Do QC records for MAC plates demonstrate that they are checked for their ability to allow visualization of lactose fermentation?						
		M-17 or equivalent						-
		Do QC records for M-17 or equivalent agar plates demonstrate their ability to support lactic streptococci?						
		Mueller Hinton Agar (MF	lA)					
		Do QC records demonstrate that MHA plates are checked for their ability to grow S. aureus & E. coli?						
8.8	M8.3	Does the laboratory: a) Perform sterility and performance tests for every batch of culture media using certified reference strains as controls? b) Source their						
		reference strains from an authorized supplier (e.g. ATCC)? c) Store, culture and sub-culture the reference strains in accordance with the specification from						3
8.10	M8.4	the supplier? Does the laboratory determine the cause of failed QC (root cause analysis), perform corrective actions and measure their effectiveness?						2
		LK CULTURE PROCEDURI	E					
8.7	M8.5	Are all milk samples processed within 2 hours of collection, or a						2

SLIPT			N	Υ	Р	N	Comments	Score
Α			Α					
		maximum of 4 hours after collection if						
		transported on ice?						
8.7	M8.6	Does the laboratory						
0.7	IVIO.U	perform a somatic cell						
		count on all milk						
		specimens prior to						
		inoculation on culture						2
		media in order to						_
		determine the number of						
		somatic cells in the						
		milk?						
8.7	M8.7	Does the laboratory						
		have a procedure for						
		rechecking somatic cell						
		counts to ensure						
		consistency of						2
		microscopic						
		observations/interpretati						
		ons among all personnel performing microscopy?						
8.7	M8.8	Does the laboratory		-				
0.7	IVIO.O	perform a bacterial						
		culture on all milk						
		samples (or those with						2
		somatic cell counts >10 ⁵						_
		white blood cells/mL) as						
		per their policy?						
8.7	M8.9	Is milk plated using a						2
		calibrated 1µL loop?						2
8.7	M8.10	Are the following media						
		used for primary culture						
		of milk?						
		Blood Agar or						2
		equivalent						_
		MacConkey Agar or						
		equivalent	-				_	
0.7	N/O 11	M-17 or equivalent ¹⁴ Are media used for						
8.7	M8.11	primary culture of milk						
		incubated aerobically at						2
		35-37°C for at least 18						2
		hours?						
8.7	M8.12	Does the lab use						
		appropriate criteria for						2
		determining						

¹⁴ M-17 Agar media is proposed for the cultivation, detection and enumeration of lactic streptococci in milk and dairy products. The M-17 media are superior to other comparable culture media for the cultivation of the fastidious species *Streptococcus cremoris, Streptococcus diacetilactis and Streptococcus lactis*.

SLIPT A			N A	Y	Р	N	Comments	Score
		contamination of a milk culture specimen? (polymicrobial culture/no predominant colonies >10 ⁴ CFU)						
	ERIAL ID 8							
8.7	M8.13	Does the laboratory perform identification tests (ID) for at least the following milk pathogens? S. aureus S. agalactiae S. uberis S. dysgalactiae C. bovis K. pneumoniae E. coli P. aeruginosa Mycoplasma spp						2
8.7	M8.14	Does the laboratory perform AST on at least the following milk pathogens using an approved test method? S. aureus S. agalactiae S. uberis S. dysgalactiae C. bovis K. pneumoniae E. coli P. aeruginosa Mycoplasma spp						2
8.7	M8.15	Is the following testing performed for <i>S. aureus</i> identification? ¹⁵ • Catalase • Coagulase (slide or tube) • Mannitol Salt Agar (MSA) • Dnase						2

¹

¹⁵ If the laboratory performs penicillin AST, it is recommended that *S. aureus* isolates with penicillin zones sizes or MICs in the susceptible range are tested for B-lactamase production using the zone-edge test or a nitrocefin test before being reported as penicillin susceptible.

SLIPT			N	Υ	Р	N	Comments	Score
Α			Α					
8.7	M8.16	Does S. aureus AST						
		include the following						
		antibiotics16:						2
		Cefoxitin						
		Vancomycin						
8.7	M8.17	Does the laboratory						
		detect						
		methicillin/nafcillin						2
		resistance in S. aureus						
		using oxacillin disk?						
8.7	M8.18	Is the following testing						
		performed for						
		Streptococcus sp.						
		identification?						
		Bacitracin						2
		Pyrrolidonyl						
		Arylamidase (PYR)						
		Bile solubility						
		Catalase						
		Optochin						
8.7	M8.19	Does Streptococcus sp.						
		AST include the						
		following antibiotics:						
		Oxacillin						2
		Co-trimoxazole						
		Ceftriaxone or						
		cefotaxime						
8.7	M8.20	Is the following testing						
		performed for						
		Corynebacterium sp.						
		identification?						2
		Catalase						
		Nitrate reduction						
		Lipid dependency						
8.7	M8.21	Does Corynebacterium						
		sp. AST include the						
		following antibiotics:						2
		Penicillin						
		Erythromycin						
		Tetracycline						

¹⁶ If oxacillin and cefoxitin results are discrepant for *S. aureus* (one is susceptible and one is resistant), the laboratory should repeat the testing. Note: oxacillin testing should always be tested by MIC (not disc diffusion). If the results remain discrepant, oxacillin should be reported as resistant.

SLIPT			N	Υ	Р	N	Comments	Score
Α			Α					
8.7	M8.22	Is the following testing performed to identify Gram negative bacilli? Oxidase Indole Methyl Red Voges Proskauer Citrate Triple Sugar Iron or Kligler Iron Urease Motility						2
8.7	M8.23	Is Gram negative bacteria AST done as per current CLSI Vet/ VetCAST guidelines for diagnostic testing and CLSI/EUCAST guidelines for surveillance testing? ¹⁷						2
8.7	M8.24	Does the laboratory test for <i>Mycoplasma spp.</i> using molecular methods?						2
8.7	M8.25	Does the laboratory use Combination Disk Test or another equivalent method for Extended Spectrum Beta- Lactamase (ESBL) screening ^{18,19} ?						2
8.7	M8.26	Does the laboratory use Combination Disk Test or another equivalent method for carbapenemase screening?						2
		ORY COMPARISON, PT A	ND E	XTER	NAL (QUAL	ITY ASSURANCE (EQA))
8.14	M8.27	Is the laboratory enrolled in an interlaboratory comparison and/or PT program for milk culture and/or molecular tests for organism identification and AST?						2

 $^{^{17}}$ See user guide section 3.4.5 for links to CLSI, EUCAST, CLSI veterinary and VetCAST guidelines. 18 J Clin Microbiol. 2013 Sep; 51(9): 2986–2990.

¹⁹https://www.eucast.org/fileadmin/src/media/PDFs/EUCAST_files/Resistance_mechanisms/EUCAST_detection_of_resista nce_mechanisms_170711.pdf

SLIPT A			N A	Υ	Р	N	Comments	Score
8.14	M8.28	Did the laboratory pass the last 3 rounds of interlaboratory comparison or PT program testing?						2
8.14	M8.29	Does the laboratory receive onsite supervision visits as part of the EQA program for milk culture and/or molecular tests?						2
Sectio	n 8: Pro	ocess Control Subtota	al					60

Section 9: Information Management

All generic requirements apply, see SLIPTA Section 9. In addition, assessors should review the following:

9.3 M9.1 Does the final report for milk culture list the organisms for which the specimen was and was not cultured20? 9.3 M9.2 Does the laboratory report alert organisms which include at least the following:21 • ESBL producing organisms • Methicillin-resistant S. aureus (MRSA) • Colistin Resistant E.								y,	tollowing
9.3 M9.1 Does the final report for milk culture list the organisms for which the specimen was and was not cultured ²⁰ ? 9.3 M9.2 Does the laboratory report alert organisms which include at least the following: ²¹ • ESBL producing organisms • Methicillin-resistant S. aureus (MRSA) • Colistin Resistant E.	Score	Comments	N	Р	Y				
milk culture list the organisms for which the specimen was and was not cultured ²⁰ ? 9.3 M9.2 Does the laboratory report alert organisms which include at least the following: ²¹ • ESBL producing organisms • Methicillin-resistant S. aureus (MRSA) • Colistin Resistant E.						Α			
report alert organisms which include at least the following: ²¹ • ESBL producing organisms • Methicillin-resistant S. aureus (MRSA) • Colistin Resistant E.	2						milk culture list the organisms for which the specimen was and was	M9.1	9.3
Vancomycin resistant <i>S. aureus</i>	2						report alert organisms which include at least the following: ²¹ • ESBL producing organisms • Methicillin-resistant S. aureus (MRSA) • Colistin Resistant E. coli • Vancomycin	M9.2	9.3
Section 9: Information Management Subtotal	4			al	btot	nt Su	: Information Manageme	n 9: In	Sectio

Section 10: Identification of Non-conformities, Corrective and Preventive Actions

²⁰ The laboratory should inform the veterinarian on the report what organisms were excluded during the culture process. This may be either by choice of media or incubation conditions (e.g. anaerobic organisms). Assessors should review a number of laboratory reports to determine how results are reported. Procedures should be consistent with the laboratory's SOPs.

Alert organisms are organisms with significant public health threat and/or organisms that are notifiable.

Section 11: Occurrence/Incident Management & Process Improvement

All generic requirements apply, see SLIPTA Section 11. In addition, assessors should review the following

followin	g:							
SLIPT			N	Υ	Р	N	Comments	Score
Α			Α					
11.4 & 11.5	M11.1	Are the following performance indicators collected ²² ?						
		Number of milk culture and molecular tests performed (disaggregated by type) Farms: Bulk						
		Farms: Bulk milk Farms: Individual cows				-		
		o Unknown/ referred ²³			-			
		Number and percentage of samples for bacterial milk culture or molecular tests rejected (disaggregated by reason e.g. leaked, insufficient volume) (target <1%)						3
		Number and percentage of milk cultures with somatic cell counts > 10 ⁵ cells/ml						
		Number of milk culture and molecular tests where pathogens were identified/isolated (disaggregated by type)						
		o S. aureus						_
		S. agalactiaeS. uberis		_				-

 $^{^{22}}$ It may not be possible for laboratories to determine the origin of the milk samples if this is not collected on the

laboratory requisition form.

23 If the laboratory can't distinguish the origin of the milk samples, the number of organisms isolated should be recorded as "Unknown/referred".

SLIPT A		N A	Υ	Р	N	Comments	Score
	 S. dysgalactiae C. bovis K. pneumoniae E. coli P. aeruginosa Mycoplasma spp Number and percentage of contaminated milk culture tests Milk culture and molecular test TAT²⁴ 						
Section 11: Subtotal	Occurrence/Incident M	ana	geme	ent 8	c Pro	cess Improvement	3

Section 12: Facilities and Biosafety

²⁴ From sample collection to reporting.





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