

C&S Medium, 30 mL Vials (Modified Cary Blair)

Revision Date: October 26, 2015

INTENDED USE

C&S Medium provides a method for collecting and preserving fecal specimens for the culture of intestinal enteric bacteria. Because the medium can maintain the bacteria for 96 hours, immediate transportation and processing of the specimen is not necessary.

SUMMARY and EXPLANATION

The diagnosis of intestinal disease caused by bacterial infection is confirmed by the isolation and identification of the pathogenic agent from a fresh stool specimen. The ability to recover and identify these bacteria depends on the immediate collection, transportation, and culture of the specimen. A transport medium should be used if a delay of more than two hours is anticipated. The best recovery of *Shigella* species is obtained by inoculating the media directly at the bedside. Buffered glycerol saline may provide the best results if this is not possible. C&S transport medium is the method of choice for maintaining the viability of intestinal pathogens such as *Campylobacter* and *Vibrio* species. Buffered glycerol saline is not recommended for these bacteria. C&S Medium will maintain the pathogenic bacteria for up to 96 hours while limiting the overgrowth of normal flora.

CONTENTS

C&S Medium is a non-nutritive, buffered, isotonic solution with a pH indicator added. The phenol red indicator will turn yellow when the solution is acidic and, the conditions are not optimal for the recovery of the intended organisms. Each vial contains 15 ml of solution and a built-in sample collection spoon. The kit is available with or without a multilingual instruction sheet and resealable bag.

COLLECTION, STORAGE and TRANSPORT

1. Collection of fecal specimens for the recovery of intestinal bacteria should always be performed prior to the use of antacids, barium, bismuth, antidiarrheal medication, antibiotics, antimalarials, or oily laxatives.
2. A minimum of three specimens, obtained during the acute stage of diarrheal disease, is recommended to ensure the recovery of enteric pathogens.
3. Fecal specimens should be collected in a clean, dry, wide-mouthed container. A bedpan is ideal. However, a waxed half-pint container with a tight fitting lid; or a clean dry milk carton with the top two-thirds removed is acceptable.
4. Using the spoon built into the cap, small samples should be added to the vial. Pay particular attention to areas that appear bloody or watery. Add sufficient sample to raise the liquid level to the red fill line on the label.
5. Use the spoon to mix the sample. Recap the vial, making sure that the lid is securely fastened. Firmly shake the vial until the contents appear homogeneous.
6. Fill out the patient information on the side of the vial. Return the vial to the physician or laboratory.
7. Proper specimen collection from the patient is critical for successful isolation and identification of enteric organisms. If immediate transportation to a laboratory is not possible refrigerate at 2-8°C for up to 96 hours.

LABORATORY EXAMINATION

Caution: Because of the variety of bacteria that may be encountered, the use of disposable gloves is recommended when examining the contents of the vial. Subculture the bacteria in the following manner:

- A. Line a shallow tray with a paper towel and wet the towel with an appropriate disinfectant.

- B. Gently mix the C&S vial to resuspend the stool sample.
- C. Tap the bottom of the vial on the counter to remove any liquid remaining from the cap.
- D. Place vial upright on the wet towel and remove the cap.
- E. Inoculate the appropriate media using the appropriate receptacle.

The type of media used for identification of organisms transported in C&S Medium is arbitrary and based on the requirements of the laboratory. Refer to one or more of the references in the bibliography for a more complete discussion of the various regimens available. In general, fecal specimens should be inoculated to several media including an enrichment broth and several selective and/or differential plated media.

PRECAUTIONS

For in vitro diagnostic use:

1. The collected specimen is potentially infectious. Always practice good laboratory hygiene when handling.
2. Disinfect or sterilize the vial prior to disposal according to regulations.
3. If the solution in the vial is beyond its expiration date, appears yellow or cloudy prior to use, it should be discarded.
4. If the collected specimen exceeds the buffering capacity of the solution (indicated by a yellow color) conditions are no longer optimal for recovery of enteric bacteria.
5. Do not refrigerate the vials prior to, collection.
6. C&S solution is a mild irritant. In case of contact, flush thoroughly with water. If irritation persists contact a physician immediately.
7. C&S Medium is for use by trained and qualified personnel.
8. For single use only.
9. For prescription use only.

MATERIALS NOT PROVIDED

Materials for cultivation, isolation, identification, and other microbiological procedures of bacteria from clinical specimens are not provided. Fisher swab (23-400-122) used for pre-market test is not included. Refer to referenced laboratory standards for the cultivation, isolation, and identification of bacteria from clinical specimens

STABILITY & STORAGE

C&S Medium is stable for at least eighteen months from the date of manufacture when stored at room temperature. The user should conduct a visual inspection prior to use. If the solution has turned yellow, it should not be used.

CAS NUMBERS

Agar	9002-18-0
Reagent Water	7732-18-5
Sodium Thioglycolate	367-51-1
Calcium Chloride	10043-52-4
Na ₂ HPO ₄	7558-79-4
Sodium Chloride	7647-14-5
Phenol Red	143-74-8

VIAL INTEGRITY

Before use, inspect each vial for leakage, cracks, or other defects. Use only intact, undamaged vials for specimen collection.

QUALITY CONTROL

C&S Medium is provided as a non-sterile product. Each lot is tested for pH and bioburden to assure that the pH and level of viable organisms are within specifications.

All bacterial test isolates and testing procedures were established using criteria outlined in the Clinical and Laboratory Standards Institute's M40-A2 document, where applicable.

LIMITATIONS

1. C&S Medium is intended for the transport of enteric pathogens listed in the Performance Characteristics section and has not been validated for other types of organisms.
2. Product performance has been tested out to 96 hours at 2-8°C and 20-25°C.
3. Extreme temperature should be avoided during transportation of C&S Medium
4. C&S Medium is recommended for the collection and transport of enteric bacteriological samples only. Anaerobic bacteria, viruses, Chlamydia, Mycoplasma, and Ureaplasma require a transport medium formulated specifically for use with these organisms.
5. Reliable specimen collection and transport depends on many factors, including collection and handling techniques, specimen condition, volume, and timing. Best results are achieved when specimens are processed shortly after the time of collection.

CAUTION

Federal law restricts this device to sale by or on the order of a medical, clinical or hospital professional.

BIBLIOGRAPHY

1. Edwards, P.R. and W.H. Ewing. 1972. Identification of Enterbacteriaceae, 3rd ed. Burgess Publishing Co.: Minneapolis. pp. 28, 337-338.
2. Ewing, W.H. 1971. Transport methods for enterobacteriaceae and allied bacteria. Public Health Lab 29: 8-23.
3. Finegold, S.M. and E.J. Baron. 1986. Diagnostic Microbiology, 7th ed. C.V. Mosby Co.: St. Louis. pp. 260-278.
4. Lenette, E.H. et. Al. Manual of Clinical Microbiology, 3rd ed. ASM.
5. Sonnenwirth, A.C. 1970, "Collection and culture of specimens and guidelines for bacterial identification," in Gradwohl's Clinical Laboratory Methods and Diagnosis, 7th ed. C.V. Mosby Co.: St Louis. pp. 1149-1152.
6. Data on file, Medical Chemical Corp., Torrance, California.
7. M40-A2 Quality Control of Microbiological Transport Systems; Approved Standard-Second Edition.

PERFORMANCE CHARACTERISTICS

C&S Medium was tested for its ability to maintain pathogenic enteric organisms. Organisms in a human fecal matrix and pure culture were tested using the CLSI M40-A2 method. Clinically negative human fecal matrix was added to C&S vials that were then seeded with suspensions of representative enteric organisms, all other enteric organisms were taken from pure culture for testing. Vials were held at 2-8°C and 20-25°C and serial sampled at 0, 72, 96 and 120 hours for representative organisms and tested at 0 and 96 hours for all other enteric organisms. Recovery of viable organisms was tested with Swab Elution and Roll-Plate method. A polyester spun swab (Fisher 23-400-122) was used in the roll plate method only. Representative enteric organisms were cultured on selective media to assure accurate recovery from fecal matrix. All other enteric organisms were cultured on non-selective media for recovery.

Representative Enteric Organisms evaluated:

Salmonella enterica	ATCC 10708
Vibrio parahaemolyticus	ATCC 17802
Escherichia coli	O157 ATCC 43894

Other Enteric Organisms:

Escherichia coli	ATCC 8739	Vibrio parahaemolyticus	ATCC 17802
Staphylococcus aureus	ATCC 6538	Clostridium difficile	ATCC 9689
Pseudomonas aeruginosa	ATCC 9027	Campylobacter jejuni	ATCC 33201
Salmonella enterica	ATCC 10708	Enterococcus faecalis	ATCC 29212
Bacillus subtilis	ATCC 6633	Shigella dysenteriae	ATCC 9361

Acceptance criteria for recovery of bacteria as recommended in the CLSI document M40-A2 were followed. For the Roll-Plate Method, for the viability to be considered acceptable, there shall be ≥ 5 CFU following the specified holding time from the specific dilution that yielded zero-time plate counts closest to 300 CFU. For viability in the Swab Elution Method to be considered acceptable there shall be no more than a 3 log₁₀ ($1 \times 10^3 \pm 10\%$) decline in CFU between the zero-time CFU count and the CFU count for the predetermined endpoint.

The results of the Roll-Plate Method study and the Swab Elution Method studies are presented in Tables 1, 2, and 3 respectively. The results demonstrate the ability of C&S Medium to sustain the viability and recovery of test bacteria within acceptance criteria for at least 96 hours refrigerated at (2-8°C) and room temperature (20-25°C).

Viability performance studies also included an assessment of bacterial overgrowth at the refrigerated temperature only. Overgrowth assessment as defined in CLSI M40-A2 guideline is greater than 1 log₁₀ increase in CFU between zero-time and the holding time point at 2-8°C. There was no increase in bacterial count when the samples were stored at 2-8°C for 96 hours and analyzed by the Roll-Plate Method (Table 1.) and the Swab Elution Method (Table 2. and 3.). No overgrowth limit is defined by CLSI M40-A2 at room temperature (20-25°C) because most commercial transport media cannot control for it. C&S is no different; overgrowth at room temperature (20-25°C) was recorded as Too Numerous To Count (TNTC).

The performance criterion was no more than a 2 log increase or decrease in viable enteric organisms at 2-8°C. C&S Medium maintained counts of seeded organisms, with and without fecal matrix, within the specified limits at 2-8°C, whether evaluated by Swab Elution or Roll-Plate method.

PERFORMANCE CHARACTERISTICS (continued)

Table 1. Representative Enteric organism recovery results for C&S Medium using Roll-Plate Method.

Organism	Hold Temperature	Average CFU's Recovered: Time 0 hrs	Average CFU's Recovered: Time 72 hrs	Average CFU's Recovered: Time 96 hrs	Average CFU's Recovered: Time 120 hrs	T=120 hrs Log reduction (-) Log increase (+)
<i>Salmonella enterica</i>	2-8°C	190	192	162	97	-0.29
	20-25°C	64	**	**	**	N/A
<i>Vibrio parahaemolyticus</i>	2-8°C	220	160	63	68	-0.51
	20-25°C	130	**	**	**	N/A
<i>Escherichia. coli</i>	2-8°C	110	48	83	71	-0.19
	20-25°C	70	**	**	**	N/A

0.5 McFarland microorganism suspension diluted with fecal matrix and C&S Medium to 2.0×10^4 unless noted
 * diluted 2.0×10^5
 ** Too numerous to count

Table 2. Representative Enteric organism recovery results for C&S Medium using Swab Elution Method.

Organism	Hold Temperature	Average CFU's Recovered: Time 0 hrs	Average CFU's Recovered: Time 72 hrs	Average CFU's Recovered: Time 96 hrs	Average CFU's Recovered: Time 120 hrs	T=120 hrs Log reduction (-) Log increase (+)
<i>Salmonella enterica</i>	2-8°C	2.7×10^2	2.9×10^2	3.0×10^2	2.0×10^2	-0.13
	20-25°C	3.2×10^2	**	**	**	N/A
<i>Vibrio parahaemolyticus</i>	2-8°C	2.2×10^2	1.6×10^2	6.3×10^2	6.8×10^2	+0.49
	20-25°C	2.5×10^2	**	**	**	N/A
<i>Escherichia. coli</i>	2-8°C	1.6×10^2	8.9×10^1	1.3×10^2	9.4×10^1	-0.23
	20-25°C	1.0×10^2	6.8×10^2	5.4×10^2	7.6×10^2	+0.88

0.5 McFarland microorganism suspension diluted with fecal matrix and C&S Medium at 1:2000
 ** Too numerous to count

PERFORMANCE CHARACTERISTICS (continued)

Table 3. All Other Enteric Organism recovery results for C&S Medium using Swab Elution Method.

Organism	Hold Temperature	Average CFU's Recovered: Time 0 hrs	Average CFU's Recovered: Time 96 hrs	T=96 hrs Log reduction (-) Log increase (+)
<i>Escherichia coli</i>	2-8°C	1.6 X 10 ⁷	5.1 X 10 ⁷	0.51
	20-25°C	2.1 X 10 ⁷	2.5 X 10 ⁶	1.09
<i>Staphylococcus aureus</i>	2-8°C	1.6 X 10 ⁷	1.4 X 10 ⁷	-0.04
	20-25°C	1.5 X 10 ⁷	2.4 X 10 ⁷	0.20
<i>Pseudomonas aeruginosa</i>	2-8°C	8.1 X 10 ⁶	6.2 X 10 ⁶	-0.12
	20-25°C	9.2 X 10 ⁶	2.0 X 10 ⁶	1.34
<i>Salmonella enterica</i>	2-8°C	6.1 X 10 ⁷	5.8 X 10 ⁷	-0.02
	20-25°C	5.6 X 10 ⁷	2.1 X 10 ⁶	0.57
<i>Bacillus subtilis</i>	2-8°C	3.6 X 10 ⁶	4.0 X 10 ⁶	0.04
	20-25°C	4.6 X 10 ⁶	1.6 X 10 ⁷	0.54
<i>Vibrio parahaemolyticus</i>	2-8°C	9.8 X 10 ⁶	8.9 X 10 ⁶	-0.04
	20-25°C	9.8 X 10 ⁶	8.8 X 10 ⁶	-0.05
<i>Clostridium difficile</i>	2-8°C	1.2 X 10 ⁷	1.0 X 10 ⁷	-0.06
	20-25°C	1.1 X 10 ⁷	9.5 X 10 ⁶	-0.06
<i>Campylobacter jejuni</i>	2-8°C	5.8 X 10 ⁷	5.0 X 10 ⁷	-0.07
	20-25°C	4.0 X 10 ⁷	3.5 X 10 ⁷	-0.06
<i>Enterococcus faecalis</i>	2-8°C	2.3 X 10 ⁷	2.1 X 10 ⁷	-0.04
	20-25°C	2.2 X 10 ⁷	1.8 X 10 ⁷	-0.10
<i>Shigella dysenteriae</i>	2-8°C	2.2 X 10 ⁷	1.4 X 10 ⁷	-0.19
	20-25°C	2.4 X 10 ⁷	5.1 X 10 ⁶	-0.68

0.5 McFarland microorganism suspension diluted with C&S Medium at 1:15

Symbols Glossary:



Authorized Representative in the European Community



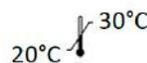
CE Marking of Conformity



Catalog Number



In Vitro Diagnostic Medical Device



Temperature Limitation



Consult Instructions For Use



Prescription Use



Manufacturer