ANAEROBIC BLOOD AGAR (CDC) w/ and w/o ADDITIVES

INTENDED USE

Remel Anaerobic Blood Agar (CDC) w/ and w/o Additives are solid media recommended for use in qualitative procedures for primary isolation and cultivation of anaerobic organisms, including fastidious strains.

SUMMARY AND EXPLANATION

This medium was formulated by V.R. Dowell and T.M. Hawkins at the Centers for Disease Control and Prevention in Atlanta, Georgia. Anaerobic Blood Agar (CDC) supports good growth and typical pigmentation of fastidious and slow-growing anaerobes, as well as other anaerobes of significant clinical importance.

PRINCIPLE

Anaerobic Blood Agar (CDC) base contains peptones which supply nitrogenous substances and amino acids necessary for the growth of anaerobic bacteria. Yeast extract provides B-complex vitamins and serves as a growth enhancer. Hemin, vitamin K, and sheep blood stimulate the growth of anaerobes. Sodium chloride is a source of essential electrolytes and maintains osmotic equilibrium. Kanamycin and vancomycin are antibiotics which inhibit gram-positive organisms and facultative anaerobic bacteria and select for gram-negative bacilli. Laked blood is used in anaerobic media to enhance pigmentation of anaerobic organisms. Neomycin aids in the isolation of anaerobic organisms, such as Bacteroides and Clostridium species, and inhibits gram-negative Enterobacteriaceae. Phenylethyl alcohol (PEA) reduces the growth of facultative gram-negative anaerobes while allowing the growth of obligate anaerobic bacteria. Paromomycin and vancomycin facilitate the recovery of fastidious, obligately anaerobic nonsporeforming gram-negative bacilli from mixed populations by inhibiting facultative and aerobic gram-negative rods. A

REAGENTS (CLASSICAL FORMULAE)*

Casein Peptone	g Vit	amin K	10.0 m	ng
Sodium Chloride5.0	g He	emin	. 5.0 m	ng
Soy Peptone5.0	g Sh	eep Blood	5	%
Yeast Extract5.0	g Ag	ar	20.0	g
L-Cystine	g De	emineralized Water10	0.00 r	ml
pH 7.5 ± 0.2 @ 25°C				

The following combinations of optional ingredients are available per liter of media:

1.	Kanamycin 100.0 mg	4.	Phenylethyl Alcohol2.5 g
	Vancomycin7.5 mg		
		5.	Paromomycin100.0 mg
2.	Laked Sheep Blood5 %		Vancomycin7.5 mg
	Kanamycin 100.0 mg		
	Vancomycin	6.	Laked Sheep Blood5 %
			Paromomycin100.0 mg
3.	Neomycin 100.0 mg		Vancomycin7.5 mg

^{*}Adjusted as required to meet performance standards.

PROCEDURE

- 1. Prior to use, reduce the plates for a minimum of 24 hours by placing them in an anaerobic environment at room temperature.
- Inoculate specimens for anaerobic culture on both selective and non-selective media as soon as possible after receipt in the laboratory; streak plates for isolation.
- 3. Incubate anaerobically at 33-37°C for 48-72 hours.
- 4. Confirm anaerobic growth by subculture to an aerobic blood agar plate.

QUALITY CONTROL

All lot numbers of Anaerobic Blood Agar (CDC) w/ and w/o Additives have been tested using the following quality control organisms and have been found to be acceptable. This quality control testing meets or exceeds CLSI standards.⁵ Testing of control organisms should be performed in accordance with established laboratory quality control procedures. If aberrant quality control results are noted, patient results should not be reported.

CONTROL INCUBATION **RESULTS** Anaerobic Blood Agar (CDC) Bacteroides fragilis ATCC® 25285 Anaerobic, up to 48 h @ 33-37°C Growth Clostridium perfringens ATCC® 13124 Anaerobic, up to 48 h @ 33-37°C Growth, beta hemolysis Fusobacterium nucleatum ATCC® 25586 Anaerobic, up to 48 h @ 33-37°C Growth Anaerobic, up to 48 h @ 33-37°C Peptostreptococcus anaerobius ATCC® 27337 Growth Prevotella melaninogenica ATCC® 25845 Anaerobic, up to 48 h @ 33-37°C Growth Escherichia coli ATCC® 25922 Ambient, 18-24 h @ 33-37°C Growth Staphylococcus aureus ATCC® 25923 Ambient, 18-24 h @ 33-37°C Growth

CONTROL	INCUBATION	RESULTS
Anaerobic KV Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
Prevotella melaninogenica ATCC® 25845	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC® 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Staphylococcus aureus ATCC® 25923	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Anaerobic LKV Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
Prevotella melaninogenica ATCC® 25845	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC® 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Staphylococcus aureus ATCC® 25923	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Anaerobic PEA Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
*Finegoldia magna ATCC® 29328	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC® 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Staphylococcus aureus ATCC® 25923	Ambient, 18-24 h @ 33-37°C	Growth
Anaerobic Neomycin Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
Clostridium perfringens ATCC® 13124	Anaerobic, up to 48 h @ 33-37°C	Growth, beta hemolysis
Prevotella melaninogenica ATCC® 25845	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC [®] 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Anaerobic PV Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
Prevotella melaninogenica ATCC® 25845	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC® 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Staphylococcus aureus ATCC® 25923	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Anaerobic LPV Blood Agar (CDC)		
Bacteroides fragilis ATCC® 25285	Anaerobic, up to 48 h @ 33-37°C	Growth
Prevotella melaninogenica ATCC® 25845	Anaerobic, up to 48 h @ 33-37°C	Growth
Escherichia coli ATCC® 25922	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
Staphylococcus aureus ATCC® 25923	Ambient, 18-24 h @ 33-37°C	Inhibition (partial to complete)
*Previously called Peptostreptococcus magnus		

INCURATION

RESULTS

BIBLIOGRAPHY

CONTROL

- Dowell, V.R., Jr. and T.M. Hawkins. 1974. Laboratory Methods in Anaerobic Bacteriology, CDC Laboratory Manual. U.S. Dept. of H.H.S. and CDC, Atlanta, GA.
- 2. Finegold, S.M., A.B. Miller, and D.J. Posnick. 1965. Ehrnährungsforschung. 10:517-528.
- 3. Dowell, V.R., Jr., C.O. Hill, and W.A. Altemeier. 1964. J. Bacteriol. 88:1811-1813.
- Dowell, V.R., Jr., G.L. Lombard, F.S. Thompson, and A.Y. Armfield. 1981. Media for Isolation, Characterization, and Identification of Obligately Anaerobic Bacteria. U.S. Dept. of H.H.S. and CDC, Atlanta, GA.
- Clinical and Laboratory Standards Institute (CLSI). 2004. Quality Control for Commercially Prepared Microbiological Culture Media; Approved Standard, 3rd ed. M22-A3. CLSI, Wayne, PA.

Refer to the front of Remel *Technical Manual of Microbiological Media* for **General Information** regarding precautions, product storage and deterioration, specimen collection, storage and transportation, materials required, quality control, and limitations.

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Printed in U.S.A.

