BIOMARK Laboratories-INDIA www.biomarklabs.com TECHNICAL SHEET

B1234 NYC AGAR BASE								
Formula								
Ingredients :	gms	s/lit.						
Proteose peptone	15.00							
Corn starch	1.00							
Glucose	5.00							
Sodium chloride	5.0	C						
Dipotassium hydr	ogen phosphate 4.00)						
Potassium dihydr								
Agar	20.0	00						
Final pH (at 25°C) : 7.4 <u>+</u> 0.2								
Directions :								
Suspend 25.50 grams in 320 ml distilled water. Heat to boiling to dissolve the medium completely.								
Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Avoid overheating. Cool to 45-50°C								
and add aseptical	ly 100 ml of sedimented l	norse b	lood cells and	60 ml of citrated horse plasma along with				
rehydrated contents of 1 vial of NYC Supplement (BF125) and 1 vial of Yeast Autolysate Supplement								
(BF126). Mix well and pour into sterile Petri plates.								
Principle :								
Proteose peptone	, horse plasma, haemogic	bin pro	ovide nutrients	for the growth of N. gonorrhoeae and N.				
meningitidis . Ph	meningitidis . Phosphate buffers the medium. The selective supplement added contains the antibiotics							
vancomycin, colis	stin, nystatin and trimet	noprim,	, to suppress	the accompanying flora. Vancomycin is				
innibitory for gra	am-positive bacteria. Col	istin in	inibits gramne	egative bacteria, including Pseudomonas				
species, while Proteus is inhibited by trimethoprim. The combination of trimethoprim and colistin acts								
synergistically ag	Jainst gram-negative ba		tarch neutraliz	zes the toxic metabolites produced by				
Neissena. The ye	ast autorysate suppleme	nt Iumm	is the COZ rec	quirements needed to enhance weissena				
growth of connor	hilling opposed Also pro		metabolized by	y gonococci to produce sufficient CO2 for				
growin of caphor	banding both size and put	sence (or yeast autory	sale reduces the lag phase of growth of				
	Inducting both size and hul	inder of	colonies.					
		Croom	to vollow					
		Lomog	Jean to yellow					
Appearance :		tomogeneous Free Flowing powder						
(11) Kenyarated medium		7110	7.4 + 0.2					
	utodaving/heating)	7.4 ± 0	7.4 ± 0.2					
	utoclaving/heating):	Cleant ye						
Clarity (post autoclaving/heating) : C			lear to slightly opalescent					
(III)Q.C. Test Microbiological								
Cultural chara	cteristics observed after in	i presei	nce of 5-10% (CO2 and 70% numidity with added				
Sealmented no	Drse Diood cells and citrate		e plasma along) with renydrated contents of 1 vial of NYC				
Supplement (1	SF125) and I vial of Yeast	Autory	sate Suppleme	ent (BF126), after an incubation at 35-				
	o nours.							
MICROURGANISM (ATCC)								
Haemophilus	(19418)	goo						
Neisseria gono	orrnoea (19424)	goo	od-luxuriant					
Neisseria mei	ningitiais (13090)	goo	od-luxuriant					
Streptococcus pneumoniae (6303)			od-luxuriant					
Streptococcus pyogenes(19615)			od-luxuriant					
Pseudomonas aeruginosa (27853)			ne-poor					
Proteus mirabilis(13883) none-poor								
Precautions : 1. For Laboratory Use.								
2. Follow proper, established laboratory procedures in handling and disposing of								
infectious materials.								
Limitations :	1. Since the nutritional re	nce the nutritional requirements of organisms vary, some strains may be						
	ncountered that fail to grow or grow poorly on this medium.							
Jse : It is recommended for the selective isolation of gonococci .								
Storage :	Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.							
Packing :	500 gm. bottle							

BIOMARK Laboratories-INDIA www.biomarklabs.com TECHNICAL SHEET

Product profile:	Reconstitution	nstitution Quantity on		Supplement	Sterilization
		Preparation (500g)			
B1234	51.0g/l	9.80 L	7.4 ± 0.2	NYC Supplement	121°C / 15 minutes
				(BF125) and 1 vial	
				of Yeast	
				Autolysate	
				Supplement	
				(BF126)	