BW347	UREA AGAR BASE (CHRISTENSEN)		
Formula			
Ingredients :	**gms/lit.**		
Peptic digest of animal tissue	1.50		
Dextrose	1.00		
Sodium chloride	5.00		
Monopotassium phosphate	2.00		
Phenol red	0.012		
Agar	15.00		
Final pH (at 25°C) : 6.8 ± 0.2			
Directions :			
Suspend 24.51 grams in 950 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 50°C and aseptically add 50 ml of sterile 40% Urea Solution (BF048) and mix well. Dispense into sterile tubes and allow to set in the slanting position. Do not overheat or reheat the medium as urea decomposes very easily.			
Principle :			
Peptic digest of animal tissue provides carbon and nitrogen required for good growth of a wide variety of organisms. Dextrose is included as an energy source. Sodium Chloride maintains the osmotic balance of the medium. Urea provides a source of nitrogen for those organisms producing urease. This is indicated by a colour change of the pH indicator, Phenol red, from yellow (pH 6.8) to red to pink – red (pH 8.1). Agar is the solidifying agent.			
QC Tests – (I)Dehydrated Medium			
Colour :	Light pink		
Appearance :	Homogeneous Free Flowing powder		
(II)Rehydrated medium			
pH (post autoclaving/heating) :	6.8 ± 0.2		
Colour (post autoclaving/heating) :	Yellowish orange		
Clarity (post autoclaving/heating) :	Clear		
(III)Q.C. Test Microbiological			
Cultural characteristics observed after 18 – 24 hrs. at 35-37°C.			
MICROORGANISM (ATCC)	GROWTH	UREASE	
Enterobacter aerogenes (13048)	Luxuriant	Negative reaction, no change	
Escherichia coli (25922)	Luxuriant	Negative reaction, no change	
Proteus vulgaris (13315)	Luxuriant	Positive reaction, cerise colour	
Salmonella typhimurium (14028)	Luxuriant	Negative reaction, no change	
Proteus mirabilis (12453)	Luxuriant	Positive reaction, cerise colour	
Klebsiella pneumoniae (13883)	Luxuriant	Positive reaction, cerise colour	
Precautions :	1. For Laboratory Use.		
	2. Follow proper, established laboratory procedures in handling and disposing of infectious materials.		

Refer disclaimer Overleaf

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Limitations :	1. Since the nutritional requirements of organisms vary, some strains may be encountered that fail to grow or grow poorly on this medium.				
	2. The alkaline reaction produced in this medium after prolonged incubation may not be caused by urease activity. False positive reactions may occur due to the utilization of peptones (especially in slant agar by Pseudomonas aeruginosa. For example) or other proteins which raise the pH due to protein hydrolysis and the relese of excessive amino acid residues. To eliminate possible protein hydrolysis, perform a control test with the same test medium without urea.				
	3. Do not heat or reheat the medium because urea decomposes very easily.				
	4. Urea Agar detects rapid urease activity of only the urease – positive Proteus species. For results to be valid for the detection of Proteus, the results must be read within the first 2 to 6 hours after incubation. Urease – positive Klebsiella, in contrast, hydrolyze urea much more slowly, showing only slight penetration of the alkaline reaction into the butt of the medium in 6 hours and requiring 3 to 5 days to change the reaction of the entire butt.				
Use :	For the detection of urease production, particularly by Proteus vulgaris , Micrococci and paracolon organisms.				
Storage :	Dehydrated medium- below 30°C Prepared medium– Between 2 to 8°C.				
Packing :	500 gm. Bottle				
Product profile:	Reconstitution	Quantity on Preparation (500g)	pH (25°C)	Supplement	Sterilization
BW347	24.51 g/l	20.399L	6.8 $\pm$ 0.2	40% Urea Solution (BF048)	121^0C / 15 minutes