

Starch Agar (i23214)

For the cultivation and differentiation of a variety of microorganisms based on amylase production.

Industry: Food / Clinical

Principles & Uses

Starch Agar, initially developed for the cultivation of Neisseria, serves as a medium for detecting starchhydrolyzing microorganisms in both food and clinical samples. The addition of Gram lodine to a 48-hour culture on Starch Agar reveals starch hydrolysis by forming a colorless zone around colonies. A blue or purple zone indicates the absence of starch hydrolysis. The size of the clear zone correlates with the starchhydrolyzing activity of the studied strain. Beef extract within the medium contributes nitrogen, vitamins, carbon, and amino acids. Starch, reacting with Gram lodine, produces a blue color, and organisms with amylase activity cause a clearing around the isolate while leaving the rest of the medium blue. While initially developed for Bacillus cereus identification, Starch Agar finds application in broader microbial studies.

Composition (gr/L)

Beef Extract 3 g, Soluble Starch 10 g, Agar 12 g. Final pH at 25° C 7.5 ± 0.2

Preparation from dehydrated Powder

Suspend 25 g of the powder in 1 L of distilled water. Mix thoroughly. Autoclave at 121°C for 15 minutes.

Quality Control

Dehydrated Appearance: Light Beige, free-flowing, homogeneous.

Prepared Appearance: Light amber, slightly opalescent.

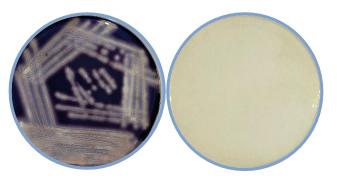
Reaction of 2.5% Solution at 25°C: pH 7.5 ± 0.2

Cultural Response

Cultural response was observed after 40 - 48 hours of incubation at $35 \pm 2^{\circ}$ C.

Organism (ATCC*)	Recovery	Starch Hydrolysis
Bacillus subtilis (6633)	Good	+
Escherichia coli (25922)	Good	-
Staphylococcus aureus (25923)	Good	-

*ATCC is a registered trade mark of the American Type Culture Collection.



Bacillus subtilis (left). Prepared Culture Media (right).

Storage

Keep the container at 15-30 °C. Store prepared medium at 2-8 °C.