



Technical Bulletin

Curian® Campy Rapid Fluorescent Immunoassay

TO: Medical Staff and Clients

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DATE: June 2, 2025

SUBJECT: Replacing *Campylobacter* Culture with Curian® Campy

Effective Monday, June 2, 2025, the *Campylobacter* component of stool testing will be performed using the Curian® Campy rapid fluorescent immunoassay.

As of June 2, Diagnostic Laboratory Services, Inc. (DLS) will begin offering the FDA cleared Curian® Campy rapid fluorescent immunoassay for the detection of a *Campylobacter*-specific antigen in stool specimens, which replaces *Campylobacter* culture in a routine stool culture. Orders for Stool Culture, Unit Code 622, will be redirected to a new Panel Code 8471 that includes the Stool Culture, Shiga Toxin and *Campylobacter* antigen.

Campylobacter infection is one of the most widespread infectious diseases and recovery in culture is challenging at best. Culture of *Campylobacter* requires specialized media, a microaerophilic environment, a dedicated incubator set to 42°C and multiple days for biochemical confirmation. The Curian® Campy rapid fluorescent immunoassay addresses these culture limitations by offering:

- Same day turn-around-time
- Removes the subjectivity of culture
- Increased sensitivity¹

Profile code: UC 8471 (Stool culture, Shiga Toxin and Campy Antigen)

Acceptable Specimen: Stool (Fecal swab or Cary-Blair)

Optimal Specimen: Fecal swab, 1 swab; or Cary-Blair

Ambient and Refrigerated Stability: 4 days

¹Buss JE, Cresse M, Doyle S, Buchan BW, Craft DW, Young S. *Campylobacter* culture fails to correctly detect *Campylobacter* in 30% of positive patient stool specimens compared to non-cultural methods. *Eur J Clin Microbiol Infect Dis*. 2019 Jun;38(6):1087-1093. doi: 10.1007/s10096-019-03499-x. Epub 2019 Feb 19. PMID: 30783889; PMCID: PMC6520473.

Please refer any questions to:

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Diagnostic Laboratory Services, Inc.

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