



## SINGLE AND PARALLEL DYE-BASED REAL-TIME PCR DETECTION OF FOODBORNE PATHOGENS *SALMONELLA ENTERICA* AND *CAMPYLOBACTER JEJUNI*

Nur Areena Chin<sup>1</sup>, Arifah Arina Syairah Janudin<sup>1</sup>, Noor Faizah Mohd-Naim<sup>2</sup>, Pooja Shivanand<sup>3</sup>, Nur Thaqifah Salihah<sup>4</sup>, Minhaz Uddin Ahmed<sup>1</sup>✉

<sup>1</sup>Universiti Brunei Darussalam, Faculty of Science, Biosensors and Nanobiotechnology Laboratory, Integrated Science Building, Jalan Tungku Link, Gadong, BE1410, Brunei Darussalam

<sup>2</sup>Universiti Brunei Darussalam, PAPRSB Institute of Health Sciences, Jalan Tungku Link, Gadong, BE1410, Brunei Darussalam

<sup>3</sup>Universiti Brunei Darussalam, Faculty of Science, Environmental and Life Sciences Programme, Jalan Tungku Link, Gadong, BE1410, Brunei Darussalam

<sup>4</sup>Universiti Islam Sultan Sharif Ali, Jalan Pasar Baharu, Gadong, BE1310, Brunei Darussalam

✉ [minhaz.ahmed@ubd.edu.bn](mailto:minhaz.ahmed@ubd.edu.bn)

<https://doi.org/10.34302/crpjfst/2023.15.4.4>

---

### Article history:

**Received:** 9 September 2022

**Accepted:** 10 October 2023

### Keywords:

*Food safety;*

*Foodborne bacteria;*

*Real-time PCR;*

*Dye-based;*

*Artificial contamination.*

### ABSTRACT

*Salmonella enterica* and *Campylobacter jejuni* are some of the common foodborne pathogens causing gastrointestinal illnesses worldwide. The development of sensitive and specific detection methods is essential to ensure food safety. Dye-based real-time PCR assay using SYBR<sup>TM</sup> GreenER<sup>TM</sup> dye was developed for the detection of *Salmonella enterica* and *Campylobacter jejuni*. Designed primer sets specifically targeting the genes *ompF* and *omp50* in *Salmonella* and *Campylobacter*, respectively, were utilised in the study. The assay was able to detect *Salmonella* and *Campylobacter* at as low as 50 fg/μl and 10 fg/μl, respectively. Specificity analysis performed using 16 different bacterial strains to check for cross-reactivity with the respective bacteria found the assay to be specific to *Salmonella* and *Campylobacter*. The assay successfully detected *Salmonella enterica* in inoculated food at as low as 5 fg/reaction for some food samples. Meanwhile, the detection limit for *Campylobacter jejuni* in all inoculated food samples was 2000 fg/reaction. The coefficient variations (CV%) of the assays for both pathogens indicated that the assays were highly reproducible. Therefore, the developed real-time PCR assays for both *Salmonella enterica* and *Campylobacter jejuni* detections were specific and sensitive and can be used for rapid screening to detect these foodborne pathogens.

---