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Cronobacter sakazakii
(formerly **Enterobacter sakazakii**) is the leading cause of necrotizing enterocolitis (NEC), meningitis, and sepsis in neonates and immunocompromised patients

([B1][B2][B3]). Its pathogenicity is associated with expression of several virulence factors, including proteolytic enzymes and a capsular polysaccharide ([B4][B5][B6]). Reports of food-borne transmission to premature infants are increasing, suggesting that neonates may serve as the reservoir of **C. sakazakii**

([B7]). A significant limitation in our understanding of the epidemiology of *C. sakazakii* is its paucity of available whole-genome sequences. Here, we present the first complete genome of *C. sakazakii*, which will provide a basis for studies of its epidemiology, pathogenesis, and acquisition of antibiotic resistance. *C. sakazakii* CICC 23019 was isolated in 2014 from *Coffea canephora* cv. 'Sapa Berry' in Ghana and was cultured on CHROMagar *Cronobacter* (CHROMagar, Paris, France) medium at 37°C. The isolate was identified by 16S rRNA

sequencing, by Zhe Jianing, and was deposited in the China Center for Type Culture Collection (CCTCC) as CICC 23019. For whole-genome sequencing, bacterial cells were collected from 7 days of culture and stored at -80°C in RNAlater (Thermo Fisher Scientific, Waltham, MA, USA). Genomic DNA was isolated from the cells using a DNeasy blood and tissue kit (Qiagen, Hilden, Germany) and used for sequencing on an Illumina HiSeq 2500 platform. PacBio data were used for correction of short-read errors, read trimming, and assembly of the entire genome. Gene prediction and gene

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