*Brochothrix*, a New Genus Tentatively Placed in the Family *Lactobacillaceae*

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**Microbacterium thermosphactum** McLean and Sulzbacher differs to such an extent from the type species of *Microbacterium, M. lacticum* Orla-Jensen, that it cannot be retained in this genus. Recent studies have shown that *M. thermosphactum* strains form a distinct taxon worthy of genus status. A new genus, *Brochothrix*, is established for the species *B. thermosphacta* (McLean and Sulzbacher) comb. nov. This genus is tentatively placed in the family *Lactobacillaceae*. *B. thermosphacta* is the type species of the genus, and ATCC 11509 is designated to be the type strain of *B. thermosphacta*.

In all of the studies, the closest associates of the *M. thermosphactum* group were the taxa representing the genera *Listeria* and *Lactobacillus*, but the relationship is not close enough to indicate that *M. thermosphactum* strains could be accommodated as a species in either of these genera.

On the basis of all of this information, we consider that the bacteria now referred to as *M. thermosphactum* should be reclassified in a new genus, for which we propose the name *Brochothrix* (brochos Gr. n. loop; thrix Gr. n. thread; M.L. fem. n. *Brochothrix* loop thread), composed of one species, *B. thermosphacta* (McLean and Sulzbacher) comb. nov. We designate strain SW26 (= ATCC 11509 = NCIB 10018), one of the original isolates of McLean and Sulzbacher (15), to be the type strain of *B. thermosphacta*.

**Description of the New Genus *Brochothrix***

The salient characters of this genus, based on the literature descriptions of *Microbacterium thermosphactum* Orla-Jensen (1, 2, 4–10, 15, 20, 22, 25) and our own observations, are as follows. Gram stains of exponential-phase cultures show rods occurring singly, in short chains, or in long, kinked, filamentous-like chains which bend and loop to give characteristic knotted masses. In stationary-phase cultures, the rods give rise to coccoid forms which, on subculture onto a suitable medium, develop into rod forms (5). The length of the rod is from 1 to 2 μm; the diameter is from 0.6 to 0.75 μm. Gram positive, but some cells lose the ability to retain the Gram stain. Nonmotile. No endospores are produced. Nonhemolytic. Optimum growth temperature, 20 to 22 °C. Growth occurs at 1 and 30 °C but not usually at 35 to 37 °C. Not heat resistant. Facultative anaerobes. Glucose metabolism is fermentative. Acid without gas.
usually produced from arabinose, cellobiose, dulcitol, glucose, lactose, maltose, mannitol, sucrose, and xylose. Methyl red and Voges-Proskauer tests are positive, H$_2$S and indole are not produced, nitrate is rarely reduced (2, 9), gelatin is not liquefied, and exogenous urea and citrate are not utilized. Catalase and cytochromes are present, but their production depends on the growth medium and temperature of incubation. Both are present in measurable amounts when the bacteria are grown on APT (Difco) medium at 20 °C, but weaker or negative reactions are obtained at the same temperature on HIA (Difco) medium. Negative results are frequently obtained if the bacteria are grown on either medium and incubated at 30 °C (5).

Cell walls contain meso-diaminopimelic acid, glutamic acid, and alanine but not arabinose or galactose (21, 22). The guanine plus cytosine content of the DNA determined by estimation of melting point is 36 mol% (4).

The type species is *B. thermosphacta* (McLean and Sulzbacher) comb. nov.

**Description of *B. thermosphacta***

This description is based on studies of the type strain SW26 (= ATCC 11509 = NCIB 10018) (2, 4, 5, 15, 21, 23) and our own observations.

Surface colonies on nutrient agar (1 to 2 days) are circular, 0.75 mm in diameter, convex with entire margin, and not pigmented; the surface is glossy. In older cultures, the edge breaks up and the center of the colony becomes raised to give a "fried-egg" appearance.

Cultures may contain colonies of both types at one time (2). Colony size is markedly increased if glucose is included in the medium. Gram stains of exponential- and stationary-phase cultures show the growth cycle described for the genus (5). Optimum growth temperature, 20 to 22 °C. Growth at 1 °C but not at 37 °C. Does not survive heating at 63 °C for 5 min. Facultatively anaerobic. Glucose metabolism is fermentative; lactic acid [mainly L(+)], with only small amounts of bi-products, is produced. Acid but no gas produced from arabinose, cellobiose, fructose, glucose, glycerol, lactose, maltose, mannitol, mannose, melezitose, rhamnose, salicin, sorbitol, sucrose, and xylose. Weak or delayed (7 days) acid production from adonitol, dulcitol, galactose, inositol, and melibiose. No acid produced from sorbose. Milk is made slightly acid, otherwise unchanged. Methyl red and Voges-Proskauer positive. H$_2$S and indole not produced, sodium hippurate not hydrolyzed, nitrate not reduced, gelatin not liquefied, casein not digested, and deoxyribonu-

clease not produced. Does not hydrolyze tweens 20, 40, 60, and 80. Exogenous urea and citrate not utilized. Enzymes of the tricarboxylic acid cycle almost totally absent (4). Cell walls contain meso-diaminopimelic acid, glutamic acid, and alanine but not arabinose or galactose (21, 22). The major phospholipids are phosphatidylglycerol, bis-phosphatidylglycerol, and phosphatidylethanolamine; the glycolipid fraction contains acetylated glucose and small amounts of a glycosyl diglyceride. The fatty-acid components are predominantly C$_{15}$ and C$_{17}$ branched-chain isomers (23). The guanine plus cytosine content of the DNA is 36 mol% (4).

**Family Relationship of the Genus Brochothrix**

The higher taxonomic affiliation of the genus *Brochothrix* could be controversial. Most numerical taxonomy studies (7, 11; Wilkinson, Ph.D. thesis) indicate a fairly close relationship to the lactic acid bacteria, and this relationship is reinforced by the results of enzyme (4, 14) and DNA base-ratio studies (4). The apparent close relationship to *Kurthia* indicated by the numerical taxonomy study of Davis and Newton (8) was almost certainly due to the choice of tests used in this study.

We consider, in the light of present knowledge (3, 18), that the presence of catalase and cytochromes in *B. thermosphacta* is not sufficient reason to exclude it from the family Lactobacillaceae. Although Schleifer (21) was of the opinion that the presence of meso-diaminopimelic acid in the cell wall indicated a closer relationship to the corynebacteria, other studies (12, 13, 26) have shown that meso-diaminopimelic acid also occurs in the cell walls of certain lactobacilli, and in a later paper Schleifer and Kandler (22) stated that *B. thermosphacta* is not closely related to the animal corynebacteria because its cell walls do not contain arabinose and galactose. However, Shaw and Stead (23) have reported that the lipid composition of *B. thermosphacta* is incompatible with that of the Lactobacillaceae.

We do not yet have sufficient knowledge to judge the weight that should be given to these latter findings in the context of the very strong evidence provided by other criteria that *B. thermosphacta* should be classified in the family Lactobacillaceae, and we suggest, therefore, that *Brochothrix* be placed in this family for the present.

**REPRINT REQUESTS**

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LITERATURE CITED