## **NOTES**

## Black Beauty Out of Mycobacterium fortuitum Cruz

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A black-pigmented strain developed spontaneously from a typical strain of *Mycobacterium fortuitum*.

In 1958, our strain no. 1000 was received from C. H. Collins, Public Health Service, County Hall, London, as his strain no. 3 of *Mycobacterium* species, a strain described as nonpigmented (Fig. 1A) and arylsulfatase-positive. Shortly after its arrival, cultures of the strain were lyophilized. Our tests and observations (3), completed in 1959, showed strain no. 1000 to be a typical strain of *M. fortuitum* (Table 1).

After 10 years of storage at 4 C, during a routine examination of our lyophilized cultures, strain no. 1000 was revived on a slant of yeast-dextrose-agar (9). One black-pigmented colony appeared among several isolated colonies above the confluent growth on the slant. The black colony was picked, and growth of the resulting deeply pigmented strain is shown in Fig. 1B. In addition to its black growth, this strain produced some dark, soluble pigment. Cultures of the strain were strongly acid-fast, and its morphology was typical of the more rapidly growing mycobacteria. Except for less activity on trehalose, the black-pigmented strain had the same physiological characteristics as its parent strain (Table 1).

The pigmentation of cultures of *M. fortuitum* on various media, varying from straw-colored to black, has been described (1, 2, 5-8, 10). Occasionally, old cultures of *M. fortuitum* on glycerol-agar (containing soil extract) formed a deep black soluble pigment (4); the growth, however, remained whitish, and subcultures of these pigment-forming old cultures did not blacken the medium.

The black-pigmented daughter strain from our strain no. 1000 is the only *M. fortuitum* to produce black growth that we have observed. As shown in Fig. 1C, sectors of the original whitish growth appeared in some cultures of the black strain, and the dark pigment is not a stable property of the strain.

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## LITERATURE CITED

- Cerbón, J., and L. F. Bojalil. 1961. Physiological relationships of rapidly growing mycobacteria. J. Gen. Microbiol. 25.7-15
- Cerbón, J., and A. Trujillo. 1963. A comparison of methods for the classification of mycobacteria, utilization of carbon sources and deamidase tests. Amer. Rev. Resp. Dis. 88:546– 550.
- Gordon, R. E. 1966. Some strains in search of a genus— Corynebacterium, Mycobacterium, Nocardia or what? J. Gen. Microbiol. 43:329-343.
- Gordon, R. E., and J. M. Mihm. 1959. A comparison of four species of mycobacteria. J. Gen. Microbiol. 21:736-748.
- Kushner, D. S., S. McMillen, and M. Senderi. 1957. Atypical acid-fast bacilli. II. Mycobacterium fortuitum: bacteriologic characteristics and pathenogenicity for laboratory animals. Amer. Rev. Tuberc. 76:108-122.
- Moore, M., and J. B. Frerichs. 1953. An unusual acid-fast infection of the knee with subcutaneous, abscess-like lesions of the gluteal region. J. Invest. Dermatol. 20:133-169.
- Ross, A. J. 1960. Mycobacterium salmoniphilium sp. nov. from salmonoid fishes. Amer. Rev. Resp. Dis. 81:241-250.
- Tsukamura, M. 1965. Salicylate degradation test for differentiation of Mycobacterium fortuitum from other mycobacteria. J. Gen. Microbiol. 41:309-315.
- Waksman, S. A. 1950. The actinomycetes, p. 196. Chronica Botanica Co., Waltham, Mass.
- Wells, A. Q., E. Agius, and N. Smith. 1955. Mycobacterium fortuitum. Amer. Rev. Tuberc. 72:53-63.

Fig. 1. Growth of Mycobacterium fortuitum on yeast-dextrose-agar. (A) Original strain no. 1000; (B, C) strain no. 1000 black. Incubation, 3 weeks.  $\times$  2.

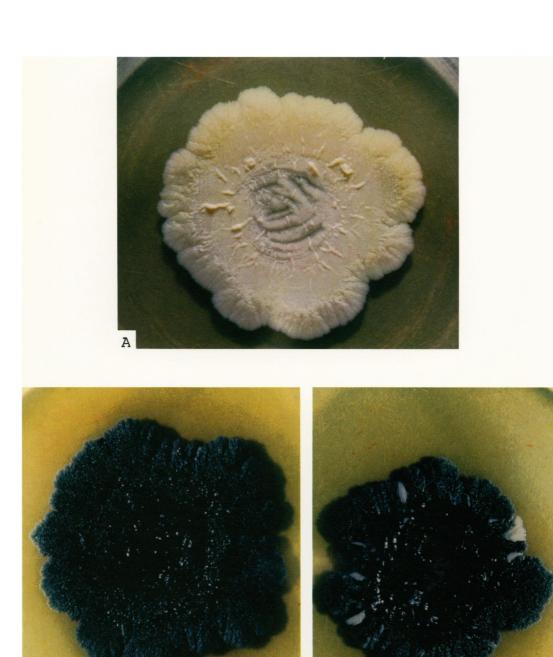


TABLE 1. Some physiological reactions of Mycobacterium fortuitum

Property	Strain no. 1000 (1959)	Strain no. 1000 black (1968)	150 Strains of M. fortuitum
Decomposition of			
Casein	_ a	_	$- (0)^{b}$
Tyrosine	- 1	_	- (0)
Urea	+	+	+ (94)
Deamination of phenyl-			/13
alanine	_	_	- (1)
Hydrolysis of			+ (94)
Starch	+ +	+ +	+ (94) + (95)
Growth at	+		T (93)
45 C			<b>–</b> (5)
40 C	+	+	$\pm (70)$
28 C	1	+	+ (100)
Survival at 60 C, 4 hr	<u> </u>	<u>'</u>	- (0)
Acid from			(0)
Arabinose	_	_	- (2)
Dulcitol.	_	_	-(0)
Erythrito!	_	_	-(12)
Galactose	_	_	<b>–</b> (8)
Glucose	+	+	+ (98)
Inositol	_	_	<b>–</b> (9)
Lactose	_	_	- (0)
Mannitol	_	_	<b>=</b> (28)
Mannose	+	+	+ (98)
$\alpha$ -m-D-Glucoside	_	_	<b>–</b> (0)
Raffinose	_	_	- (1)
Rhamnose	_	_	- (0)
Sorbitol	_	_	- (3)
Trehalose	+	Trc	+ (96)
Xylose	_	_	- (3)
Utilization of			(2)
Benzoate	_	_	- (3)
Citrate.	+	+	± (82)
Lactate	+	+	+ (98) + (97)
Mucate	+ - -	+	+ (97)  - (0)
Oxalate			-(0)
Succinate	+	+	+ (98)
Growth on dyes	1	'	1 (50)
Methyl violet	+	+	+ (99)
Pyronin	+		+ (100)
Color change of Mac-	'		, (====)
Conkey agar	+	+	+ (98)
Resistance to			
Penicillin, 10 units	+	+	+ (100)
Bacitracin, 10 units	+	+	+ (85)
Production of arylsul-			
fatase, 3 days	+	+	+ (96)
Growth in $(0.2\%)$ sali-			, /02\3
cylate broth	i —	1 -	$\pm (83)^d$

<sup>&</sup>quot; Symbols: +, 85 to 100% of strains positive; ±, 50 to 84° c of strains positive;  $\mp$ , 15 to 49% of strains positive; -, 0 to 14% of strains positive.

<sup>b</sup> Numbers in parentheses represent per cent

positive strains.

<sup>&</sup>lt;sup>c</sup> Trace.

<sup>&</sup>lt;sup>d</sup> Only 52 strains were tested.