

SPECIAL PUBLICATIONS FROM THE SETO
MARINE BIOLOGICAL LABORATORY

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OF
THE JAPANESE ANTARCTIC RESEARCH EXPEDITION

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ON SOME YEASTS FROM
THE ANTARCTIC REGION

BY

MASAMI SONEDA

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SIRAHAMA, WAKAYAMA-KEN
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AS far as ascertained, the report of the yeasts from the Antarctic region could not be found. However, the present author isolated five species of yeasts from some soily materials of the Antarctic region which were collected by Dr. H. FUKUSHIMA, Dr. R. HAGA and Dr. T. MATSUDA, all were the members of the 3rd and 4th Japanese Antarctic Research Expedition. The results of the isolation are shown in Table 1. The isolation was followed to the ordinary agar-plating method; soil-suspension was poured on the malt-agar plates and kept at 20°C.

Table I. Yeasts from the Antarctic materials.

Yeast isolated	Place (date)	Number of strain
<i>Cryptococcus albidus</i>	West Ongul Isl. (4th)	1
<i>Cryptococcus laurentii</i>	Showa Base (3rd)	1
" "	Showa Base, the Continent and West Ongul Isl. (4th)	4
<i>Rhodotorula mucilaginosa</i>	the Continent and West Ongul Isl. (4th)	5
<i>Torulopsis famata</i>	Showa Base (3rd)	1
<i>Trichosporon cutaneum</i>	Showa Base (3rd)	1

Among the above five species, *Cryptococcus laurentii* and *Rhodotorula mucilaginosa* were dominant members.

The present author has some doubt that all members of yeasts, here treated, existed in the Antarctic soil when collected and they are also present there even now. But it should be speculated that above two dominant species may present in soil of the Antarctic region, and they can grow under lower temperature condition as far as tested.

On relationships between yeasts and low-temperature, HANSEN (1881, 1903) demonstrated that yeasts are often very resistant to the cold; LOCHHEAD and FARRELL (1930) reported that some yeasts could remain viable in the soil of Canadian winter; STILLE (1950) examined that yeast cells are not distributed at -195°C under the same operation; LUND (1954) examined that yeast survived freezing in grape juice several months under the same condition.

Cryptococcus albidus (SAITO) SKINNER, in HENRICI's Molds, yeasts and Actinomycetes, 2nd ed. p. 288. (1947)

syn. *Torula albida* SAITO.

Morphological properties: After 3 days in malt extract at 20°C, cells are

usually spherical $(3.0\sim 8.0) \times (3.0\sim 8.0)\mu$, single or in pairs (Fig. 1), a sediment and thin ring are formed, the cells are surrounded by capsule; after one month, a ring and sediment are present.

After one month on malt agar at 20°C , the colony is yellowish white to white, somewhat bullated, soft, mucous and slightly shiny.

Physiological properties: Oxidative; Assimilate glucose, galactose, maltose, saccharose, lactose and ethanol; potassium nitrate is well assimilated; not split arbutin; starch like compounds is formed in capsule.

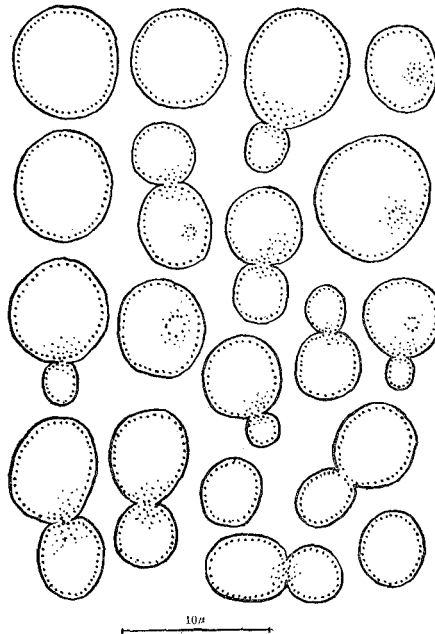


Fig. 1. *Cryptococcus albidus* (Antarctic strain)
3 days in malt extract.

This strain is different from the original description of SAITO (1922) on assimilation of potassium nitrate, but LODDER (1934) examined the above SAITO's strain and found that it assimilated potassium nitrate.

***Cryptococcus laurentii* (KUFFERATH) SKINNER**, in HENRICI's Molds, yeasts and Actinomycetes, 2nd ed. p. 288 (1947)

syn. *Torula laurentii* KUFFERATH.

Morphological properties: After 3 days in malt extract at 20°C , growth is slow, cells are almost ovoid or seldom ameoboid, $(3.0\sim 7.2) \times (3.2\sim 8.6)\mu$, single or in pairs (Fig. 2), a sediment is formed, after one month at 20°C , thin ring and slimy sediment are produced. The cells are surrounded by capsule.

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After one month on malt agar at 20°C, the color is whitish yellow to light orange, soft, slimy, smooth and somewhat glistening; margin smooth.

Slide culture on potato agar at 20°C, usually no pseudomycelium, seldom few large polymorphous cells are observed.

Physiological properties: Oxidative; glucose, galactose, maltose and saccharose are assimilated, lactose and ethanol are not assimilated; potassium nitrate is not assimilated; arbutin is not splitted; starch-like compound is formed.

This strain characteristically can grows at lower temperature; maximum temperature is 20°C, grows at 5°C after 7 days. This strain has sometimes developed ameboidal cells in malt extract, which resemble those of *Cryptococcus*

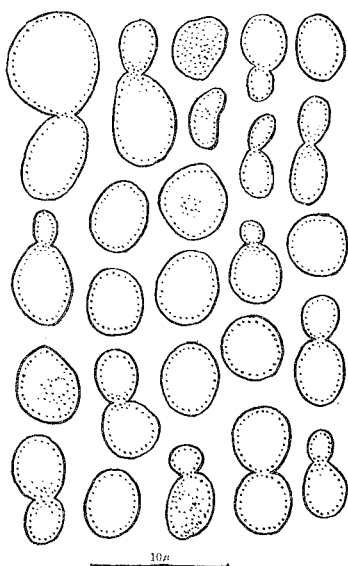


Fig. 2. *Cryptococcus laurentii* (Antarctic strain)
3 days in malt extract.

laurentii var. *flavescens*; shape and size of the cells are ranked between that of *Cryptococcus laurentii* and *Cryptococcus laurentii* var. *magnus*. However, the present author gave the name, *Cryptococcus laurentii*, for this strain, because this two features are not so important for yeast classification.

Rhodotorula mucilaginosa (JORG.) HARRISON, in Trans. Roy. Soc., V, 22:187 (1928); KOBAYASHI, Y., K. TUBAKI and M. SONEDA, in Bull. Nat. Sci. Mus. (Tokyo) 33:47 (1955); SONEDA, M., in Nagaoa 6:14 (1959).

syn. *Torula mucilaginos*a SAITO.

Morphological properties: After 3 days at 25°C, cells are almost ovoid (2.8~4.2) × (4.0~7.2), single or in pairs, (Fig. 3) a sediment and a ring are for-

med, after one month at 20°C the ring becomes thicker and pinkish color.

After one month at 20°C on malt agar, the streak is soft, smooth and somewhat mucous; the color of streak is red to whitish red; margin smooth.

Physiological properties: Oxidative; glucose, galactose, maltose, saccharose and ethanol are assimilated, lactose is not assimilated; potassium nitrate is not assimilated: arbutin is not splitted.

The characteristics of this strain resemble the original description of JÖRGENSEN (1909) and also the standard description of LODDER and KREGER-VAN RIJ (1952), but optimum temperature of the present strain for the growth is rather low; it is 20°C as far as tested.

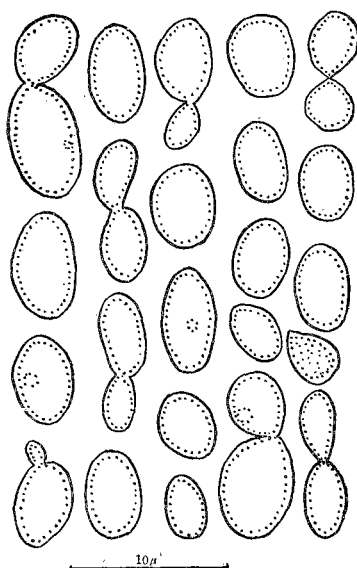


Fig. 3. *Rhodotorula mucilaginosa* (Antarctic strain) 3 days in malt extract.

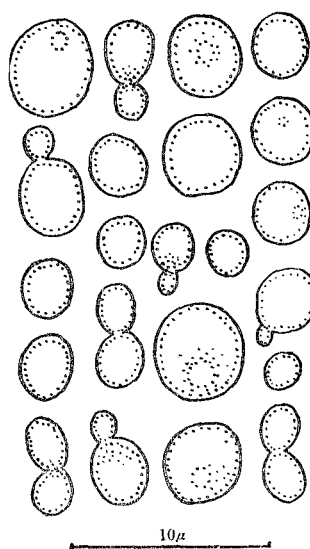


Fig. 4. *Torulopsis famata* (Antarctic strain) 3 days in malt extract.

***Torulopsis famata* (HARRISON) LODDER et K-VAN RIJ**, *The Yeasts*, p. 417 (1952); SONEDA, M., in *Nagaoa* 6:13 (1959).

syn. *Mycotorula famata* HARRISON.

Morphological properties: After 3 days in malt extract at 25°C, cells are spherical or ovoid, $(2.0\sim 4.8) \times (2.2\sim 5.2)\mu$, single or in pairs (Fig. 4), after one month at 20°C, sediment and a ring are produced.

After one month on malt agar at 20°C, the colony is whitish or yellowish white, almost smooth, only with some radial stripes, soft and not much glistening; margin smooth.

Slide culture on potato agar, pseudomycelial cells are not observed.

Physiological properties: Oxidative; glucose, galactose, maltose and saccharose

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are assimilated, lactose is not assimilated; potassium nitrate is not assimilated.

This strain resembles the original description of HARRISON (1928) on morphological and physiological properties.

Trichosporon cutaneum (DE BEURM, GOUGEROT et VAUCHER) OTA, in *Ann. parasitol, humaine et Copée* 4:1 (1926); SONEDA, M., in *Nagaoa* 6:15 (1959).
syn. *Oidium cutaneum* DE BEURM, GOUGEROT et VAUCHER.

Morphological properties: After 3 days in malt extract at 20°C, cells are spherical, ovoid and cylindrical, $(2.4\sim 7.0)\times(4.2\sim 10.0)\mu$, but may be longer, single or in pairs (Fig. 5), occasionally pseudomycelial cells and arthrospores appear, after one month at 20°C, the culture consists of thick, tough, wrinkled pellicle and membranous sediment.

After one month on malt agar at 20°C, the steak is dry, dull, not hairy, raised, whitish yellow to brownish white and wrinkled.

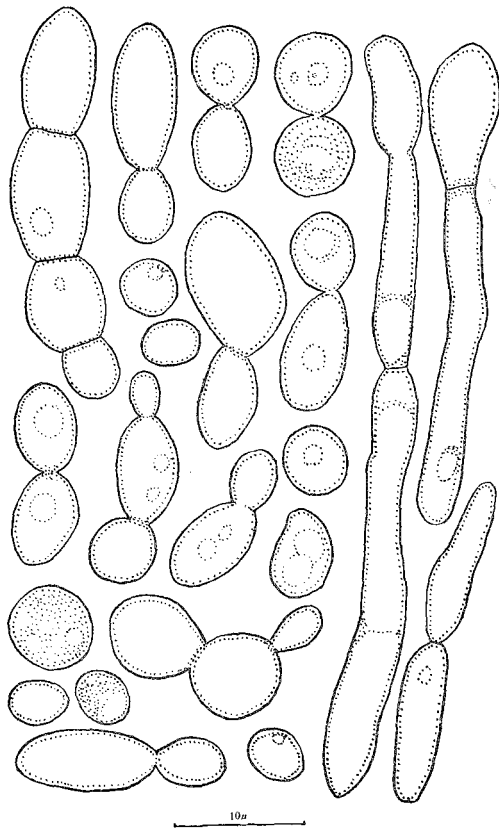


Fig. 5. *Trichosporon cutaneum* (Antarctic strain)
3 days in malt extract.

Slide culture on potato agar, the pseudomycelium, arthrospores and blastospores are developed abundantly. Chains of spherical or cylindrical cells occur on pseudomycelium, and it is splitted up into variable length arthrospores that may give raised to a zig-zag formation (Fig. 6). Sometimes large polymorphous cells are formed at intercalary cells.

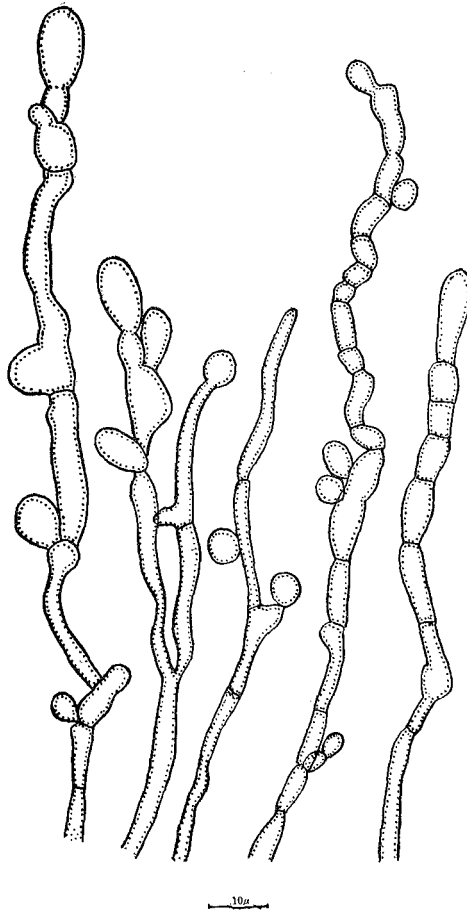


Fig. 6. *Trichosporon cutaneum* (Antarctic strain)
Slide culture on potato agar after 7 days.

Physiological properties: Oxidative; assimilate glucose, galactose, maltose, saccharose and lactose; potassium nitrate is not assimilated; ethanol is not assimilated; under appropriate conditions "Starch-like compounds" is formed.

This strain is not so different from the standard description of LODDER et KREGER-VAN RIJ, but growth on malt agar is more or less reduced, and optimum temperature for the growth is 20°C.

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RESULTS AND DISCUSSION

It seems to be difficult to decide whether these yeasts, originally isolated from the above Antarctic materials, may present in the Antarctic or not. But the high rate of the isolation and the growth under lower temperature may suggest the important problems for discussing the ecology of these Antarctic yeasts. Table II shows the rate of the comparative growth between the Antarctic strains and the strains which are presenting in our collection.

Table II. Effect of temperature for growth.
(After 7 days on malt agar)

Species	Temp.	5°C	12°C	20°C	25°C	30°C
<i>Cr. laurentii</i> (Antarctic strain)		+	+	++	—	—
	" (NI. 7353)	—	—	+++	++	—
<i>Rh. mucilaginoso</i> (Antarctic strain)		+	+	+++	+	—
	" (NI. 7203)	—	+	+++	+++	+++
<i>Trich. cutaneum</i> (Antarctic strain)		—	++	+++	++	—
	" (NI. 7461)	—	—	++	+++	+++
<i>Cr. albidus</i> (Antarctic strain)		—	+	++	++	—
	" (NI. 7349)	—	+	++	++	—
<i>T. famata</i> (Antarctic strain)		—	+	++	+++	+
	" (NI. 7577)	—	+	++	+++	+

The result indicates that these Antarctic yeasts may be divided into the following three groups from view-point of their minimum, optimum and maximum temperature.

- (1) Optimum temperature of growth is 20°C; growth well at even 5°C.
- (2) Optimum temperature of growth is 20°C; can not grow at 5°C.
- (3) Optimum temperature is about 25°C; can not grow at 5°C; not so different from type strain.

Cryptococcus laurentii and *Rhodotorula mucilaginoso* may be included in (1); minimum, optimum and maximum temperature of these strains are lower as compared with other strains of type cultures.

Trichosporon cutaneum is in (2); it grows under low-temperative condition as compared with other the two species, *Cryptococcus albidus* and *Torulopsis famata*.

Many thanks are due to Dr. Y. KOBAYASI and Dr. K. TUBAKI for their instructions and advices, and also to Dr. H. FUKUSHIMA, Yokohama Municipal University, for giving the materials.

SUMMARY

- (1) Five species of yeasts were isolated from soily materials of the Antarctic region, collected by members of the 3rd and 4th Japanese Antarctic Research Expedition.
- (2) *Cryptococcus laurentii* and *Rhodotorula mucilaginosa* seem to be dominant species among them and can grow at lower temperature as far as tested.

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Postscript ; Since this manuscript went to press, M. E. DI MENNA has published a paper on the Antarctic yeasts ("Yeasts from Antarctic", in Jour. Gen. Microbiol. **23**: 295-300, 1960) in which she described eight species of yeasts, *Candida*, *Cryptococcus*, *Rhodotorula* and *Sporobolomyces*.

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