The Identification of Yeasts from Clinical Material

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Medical mycology is presently emphasizing sophisticated methods for isolation and identification of yeasts from clinical material. Many factors have contributed to this trend toward sophistication, including an increase in the numbers and kinds of isolated yeasts which are unfamiliar to the clinical mycologist. Currently approximately 70 species in 10 genera are recognized (Lodder, 1970). These yeasts include species of Rhodotorula (Lodder, 1970; Louria et al., 1967; and Blair et al., 1970), Kluyveromyces (Lodder, 1970), Saccharomyces (Lodder, 1970), Hansenula (Lodder, 1970), Pichia (Lodder, 1970), Debaryomyces (Lodder, 1970), Sporobolomyces (Lodder, 1970), and 20 species of Candida (Haley, 1971; Lodder, 1970; and Painter and Isenberg, 1973). Other examples include species of Cryptococcus other than neoformans (Blair et al., 1970; Haley, 1971), including a red informo-minitarius (Lodder, 1970) and an unusual species of Torulopsis (Haley, 1971; Lodder, 1970). These unfamiliar isolates may be caused by new drugs, drug combinations, or underlying diseases. Whatever the cause, more saprophytic yeasts are associated with clinical material and/or disease processes than before.

Not only is the clinical mycologist currently concerned with the isolation of more yeasts with opportunistic potential, but also with a growing trend toward dual infections, primarily yeast and bacterial. Dual infections have been seen in Wadsworth V.A. patients undergoing home dialysis.

In addition to unfamiliar species, the clinical mycology laboratory discovers occasional aberrant yeast isolates, possibly produced by certain drugs, which only the well-trained mycologist may recognize. Experiences with atypical fungi revealed a C. neoformans without a capsule, a Geotrichum sp. that formed atypical arthropores, and strains of C. albicans that failed to form typical chlamydomespores on corn meal tween agar.

All of the above-mentioned problems indicate that a clinical mycology laboratory interested in species identification must be adequately equipped to isolate and identify well-known as well as unusual yeasts. Although the physician determines the significance of the various yeasts isolated, the clinical mycologist aids in this regard by designating yeast medically important sexual stages and imperfect forms, and characteristics of the sexual stages in clinical material, are described.

Included in this report is a guide to yeast identification which relies on the Luecke plate, a modified Dalmau plate.

**Index Descriptions:** Yeast Identification, Non-Filamentous Fungi, Mycology in Medicine.

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the initial isolate. The yeast must be isolated and purified. The same emulsified portion of the yeast colony in sterile distilled water which was used for direct microscopic examination may be used here. A loopful of this emulsified specimen is streaked on Bacto Sabouraud dextrose agar and incubated at room temperature to obtain isolated colonies. A yeast suspension in water separates the individual yeast cells even before streaking, thus allowing more complete isolations. If the culture is mixed, different colony types will be seen in the area where single colonies are separated. Each different colony should undergo a second streaking on Sabouraud’s agar as a purity check (Haley, 1971).

Pure cultures are maintained on Bacto yeast malt agar (YM) or Bacto malt agar (MA), but before various methods for identification are utilized, each pure colony is subcultured to YM agar. If possible two successive transfers to YM agar are made before further identification studies are undertaken (Lodder, 1970).

**India Ink Preparation to Detect Capsules.** This preparation aids in the detection of capsules, a significant taxonomic character.

a. Place a loopful of India ink (Sanford) on a clean slide and a loopful of sterile distilled water close to the loopful of India ink.

b. Make a light emulsion of the yeast in the drop of water.

c. Mount with a cover slip such that the edge of the cover slip is placed at the outer edge of the drop of India ink and allowed to fall across both the ink and the yeast suspension, taking care to avoid air bubbles.

d. Immediately examine the area of the slide where the ink and water suspension meet. Capsules are seen easily in this area.

**Characterization of Vegetative Growth on Corn Meal Tween Agar (Luecke Plate).** The type of vegetative growth can be detected by growing the yeast isolates on corn meal tween agar, or Luecke plates. The Luecke plate is a combination of Wickerham's (1951) description of the Dalmau plate (Difco, 1972b), and our modification of Haley's technique (1971).

A corn meal tween plate (CMT) (Difco, 1972a; Walker and Huppert, 1960) is divided into four sections (Figure 1). Two sections are used for unknowns, the third and fourth for the two controls. A pure colony of yeast is touched with the tip of a sterile inoculation needle and a single streak is made on the CMT plate from near the edge of the petri dish down the center of the section of agar. The streak barely scarifies the medium (streak 1 in Figure 1). Starting from the edge of the petri dish a sterile applicator stick streaks back and forth along streak 1 (streak 2 in Figure 1). Although the span of streak 2 may be greater near the edge of the petri dish, note that it becomes narrow toward the center. A sterile square cover slip is placed over streaks 1 and 2 so that the streaks and a portion of the unstreaked medium are completely covered by the cover slip. This is an important and necessary step since observation of growth ultimately takes place at the edge of streak 2 (at the junction of the streaked and unstreaked medium under the cover slip). It is here that typical yeast morphology is seen and purity ascertained. The plate is incubated at 25°C for 48-72 hours in a moist chamber and then examined for typical blastospores and/or hyphae and/or arthrospores.

A control plate with *C. albicans* and one with a *Tricho* sporon sp. are also streaked to compare with each plate containing unknowns.

![Figure 1. Method of streaking a Luecke plate.](https://scholarworks.uni.edu/pias/vol81/iss1/9)
Growth immediately surrounding the discs indicates assimilation of the sugar by the yeast; clear areas around the discs indicate a lack of assimilation.

Positive controls: Candida guilliermondii for assimilation of all sugars, except lactose; C. pseudotropicalis for assimilation of lactose.

Negative controls: Candida krusei.

Fermentation Tests. These studies are performed according to Wickerham (1951). The medium contains 0.45% yeast extract, 0.75% peptone, and approximately 1 mg bromthymol blue sufficient to produce a green color in the medium. The yeast extract-peptone broth is dispensed in 2 ml quantities in 12 x 150 mm screw-capped tubes. Durham tubes (50 x 6 mm) are added and the medium is autoclaved 15 minutes at 15 p.s.i. Sugars are prepared individually as 6% aqueous solutions, except raffinose which is prepared as a 12% solution. Each is sterilized with a Swinnex-47 millipore filter and stored until needed. One ml quantities are added to 2 ml of yeast-peptone broth.

The fermentation broth is inoculated with 0.1 ml of inoculum A. Tubes are shaken and incubated at room temperature or at 34-37°C, and examined regularly for 10-14 days.

Durham tubes are used to detect CO₂ production resulting from fermentative action. A pH change simply indicates the assimilation of a carbon compound.

To detect false negatives due to supersaturation of the broth, all tubes giving an acid reaction are shaken slightly and the cap vented. A rapid release of gas frequently follows.

Carbohydrates most helpful in identification include the following:

Hexose sugars—glucose (GLU), galactose (GAL);
Disaccharides—sucrose (USC), maltose (MAL), lactose (LAC), melibiose (MEL*), trehalose (TRE), cellobiose;
Tri saccharides—raffinose (RAF); and
Polysaccharides—inulin (INU).

The above 10 carbohydrates are used since keys to the majority of the species of the various genera are based on the fermentation and assimilation of these sugars.

Two Candida species, C. tropicalis and C. pseudotropicalis, may be necessary to run as positive controls with unknown yeasts; Cryptococcus species may serve as negative controls. Although each species ferments only several carbohydrates, the two together will provide fermentation of all the listed carbohydrates.

Nitrate Assimilation Test. The medium to test nitrate assimilation is prepared as follows. To Bacto yeast carbon base (11.7 gm) is added 100 ml distilled water. YCB is sterilized via the Swinnex-47 millipore filter and stored in the refriger-ator until needed. The final medium, prepared by mixing 100 ml of sterile YCB with 900 ml of melted 2.25% sterile agar, is poured into petri dishes while still warm but not hot to touch after 0.1 ml of inoculum A is added to the petri dish. The mixture of warm YCB and inoculum A is swirled gently to mix, and allowed to cool.

One disc containing 1.0% KNO₃ is placed in the center of the inoculated YCB prepared above. The discs are prepared by soaking overnight in a petri dish containing approximately 20 ml of 1.0% KNO₃ and by decanting completely the excess nitrate, after which, the next day, the discs and the petri dish are autoclaved and dried. The petri dish is sealed with masking tape to avoid dehydration during storage at room temperature. The YCB plate with disc is incubated at 25°C. After 24 to 48 hours, growth around the disc where KNO₃ has diffused is a positive assimilation test.

Controls: positive control—Cryptococcus albidus; negative control—Torulopsis glabrata or Candida albicans.

Starch Production. The test for formation of starch has been employed primarily to separate the genus Torulopsis from Cryptococcus (lodder, 1970). The ability to produce starch is often associated with the utilization of inositol and the ability to hydrolyze urea.

The YNBs containing various sugar discs are used to test for starch production. After an incubation period of 14 days, one drop of Lugol's iodine solution is added to the area around the glucose disc. Strains producing starch will turn the area blue to purple to green depending upon the intensity of starch.

A positive and a negative control are necessary. Starch results may be confusing unless both controls are used. The negative control may reveal a small black granule at the edge of the glucose disc, but this does not indicate positivity. A black ring around the disc with a diffusing black-green halo indicates starch production.

Positive control—Cryptococcus sp.; negative control—Toru­lopsis sp.

Medium for Ascospore Production. Two media are used to test for ascospore production. Gorodkowa's medium (lodder, 1970) consists of 0.1% glucose, 1% peptone, 0.5% sodium chloride, and 2% agar. The medium is steamed to dissolve the agar and dispensed in 90 ml aliquots, autoclaved at 15 p.s.i. for 15 minutes, and poured into petri dishes as needed. Cultures are incubated at room temperature and observed for 4-6 weeks for ascospores.

Bacto malt agar (lodder, 1970) is inoculated concomitantly with Gorodkowa's medium for ascospore production. (Malt agar also serves as a good medium for pigment production.)

Urease Production. Urea hydrolysis is detected by inoculation of Christensen's urea agar slants (dolan, 1971) with one loopful of yeast and incubation at 30°C for 72 hours. In the presence of urease, the color of the medium changes to pink or red.

Positive control: Rhodotorula sp. or Cryptococcus sp.

Yeast Identification Guide. A guide for yeast identification is presented (Figure 2). Important steps in methodology are included. Spaces are provided for additional carbohydrates if more are needed to conclusively identify yeasts to species. When unknowns are identified, these guides may be filed alphabetically according to patient's name for future reference.

RESULTS AND DISCUSSION

Table 1 lists the genera and species of 3,474 yeasts isolated during a three-year study, the number and percent of each species isolated, and the various types of clinical material from which the species were isolated. Rare yeasts are indicated by few collections. Three sexual forms are represented, including Pichia, Kluyveromyces, and Sporobolomyces; the latter may be a Basidiomycete (lodder, 1970). In

* Use in fermentation tests only when raffinose is fermented.
Identification of Yeasts

Name of patient ... Identification No. ...
Date of specimen ... Specimen ... Ward ...
Initial isolate
1. Microscopic: Mixed ... Pure ...
2. India ink for capsules: Yes ... No ...
Pure culture
3. Luecke plate for vegetative growth characteristics.
   Blastospores only ... Pseudohyphae ...
   True hyphae ...
   Arthrospores ...
   Aberrant ...
   Control = C. albicans
   Control = Trichosporon sp.
4. Malt agar for pigmentation and ascospores.
   Pigment ... Ascospores ...
   Pigment control = Rhodotorula sp.
   Ascospore control = Saccharomyces sp.
5. Corodkowa agar for ascospores.
   Incubate at 25°C for 3-6 weeks.
   Ascospores ...
   Control = Saccharomyces sp.
6. Growth at 37°C, 45-48°C, 25°C ...
7. Starch production
   Unknown ...
   Positive control = Cryptococcus sp.
   Negative control = Torulopsis sp.
8. Production of urease (48 hrs. at 30°C).
   Unknown ...
   Positive control = Cryptococcus sp.
   Negative control = Torulopsis sp.
9. Assimilation of NO₃ (48-96 hrs. at room temperature).
   Unknown ...
   Positive control = Cryptococcus albidus var. albidus.
   Negative control = C. albicans or Torulopsis glabrata.
10. Assimilation of carbohydrates (examine at 24 and 48 hrs.,
     incubate at 30°C).

   glu* gal suc mal lac mel raf inu tre
Control 1 (positive) + - + + + + + + + + C. pseudotropicalis
Control 2 (positive) + + + + + + + + + + C. guilliermondii
Control 3 (negative) + + + + + + + + + + C. krusei
Unknown

11. Fermentation of carbohydrates (examine at 1, 2, 3, 4, 7, 10 day; incubate at 37°C).

   Control 1 (positive) + + + + + + + + + + C. tropicalis
   Control 2 (positive) + + + + + + + + + + C. pseudotropicalis
   Control 3 (negative) + + + + + + + + + + Cryptococcus sp.
   Unknown
   Identification ...
   Date ...
   * glu = glucose
   gal = galactose
   suc = sucrose
   mal = maltose
   lac = lactose
   mel = melibiose
   ino = inositol
   raf = raffinose
   inu = inulin
   tre = trehalose
   + = Acid and gas.

Table 1. Yeasts Isolated During a Three-Year Period.

<table>
<thead>
<tr>
<th>Organism</th>
<th>Number of Strains</th>
<th>Percent of Total</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>Candida albicans</td>
<td>2,136</td>
<td>62.0</td>
<td>Sputum, lung biopsy, autopsy, wounds, pleural fluid, stool, ulcer, genitals, urine, blood</td>
</tr>
<tr>
<td>Candida tropicalis</td>
<td>627</td>
<td>18.0</td>
<td>Peritoneal fluid, sputum, wounds, blood, urine, bronchial washings</td>
</tr>
<tr>
<td>Torulopsis glabrata</td>
<td>111</td>
<td>3.2</td>
<td>Sputum, blood, urine, lung biopsy, bronchial washings, catheter</td>
</tr>
<tr>
<td>Candida stellatoidea</td>
<td>94</td>
<td>2.7</td>
<td>Sputum, blood, catheter, wounds</td>
</tr>
<tr>
<td>Geotrichum candidum</td>
<td>60</td>
<td>1.7</td>
<td>Stool, trachea, pleural fluid, lung abscess, lung biopsy, urine</td>
</tr>
<tr>
<td>Candida pseudotropicalis</td>
<td>40</td>
<td>1.2</td>
<td>Sputum, trachea, urine, blood, peritoneal fluid, endocarditis, ulcer, bronchial washings</td>
</tr>
<tr>
<td>Candida lipolytica</td>
<td>33</td>
<td>0.9</td>
<td>Throat swabs, wound, foot, catheter, autopsy</td>
</tr>
<tr>
<td>Candida krusei</td>
<td>28</td>
<td>0.8</td>
<td>Sputum, skin, urine, blood, trachea</td>
</tr>
<tr>
<td>Trichosporon fermentans</td>
<td>27</td>
<td>0.7</td>
<td>Sputum, bronchial washings, catheter</td>
</tr>
<tr>
<td>Candida rugosa</td>
<td>26</td>
<td>0.7</td>
<td>Sputum, bronchial washings, catheter, urine</td>
</tr>
<tr>
<td>Candida parapsilosis</td>
<td>25</td>
<td>0.7</td>
<td>Fistula, wounds, sputum</td>
</tr>
<tr>
<td>Trichosporon pululans</td>
<td>25</td>
<td>0.7</td>
<td>Urine, sputum, bronchial washings</td>
</tr>
<tr>
<td>Trichosporon capitatum</td>
<td>24</td>
<td>0.7</td>
<td>Urine, wounds, trachea, gastric washings</td>
</tr>
<tr>
<td>Candida guilliermondii</td>
<td>22</td>
<td>0.6</td>
<td>Lung biopsy, gastric washings, sputum</td>
</tr>
<tr>
<td>Trichosporon cutaneum</td>
<td>22</td>
<td>0.6</td>
<td>Urine, sputum, abdominal fluid, throat swabs, gastric washings</td>
</tr>
<tr>
<td>Torulopsis holmii</td>
<td>21</td>
<td>0.6</td>
<td>Foot, bronchial washings, catheters, wounds</td>
</tr>
<tr>
<td>Candida brumptii</td>
<td>19</td>
<td>0.6</td>
<td>Conjunctiva, catheter, skin (burn), urine</td>
</tr>
<tr>
<td>Trichosporon inkin</td>
<td>16</td>
<td>0.5</td>
<td>Sputum, throat swabs, skin lesion</td>
</tr>
<tr>
<td>Rhodotorula rubra</td>
<td>14</td>
<td>0.4</td>
<td>Leg ulcer, pleural fluid, scrotal tumor, sputum, urine</td>
</tr>
<tr>
<td>Torulopsis candida</td>
<td>14</td>
<td>0.4</td>
<td>Foot lesion, sputum, urine</td>
</tr>
<tr>
<td>Cryptococcus neoformans</td>
<td>13</td>
<td>0.4</td>
<td>CSF, skin</td>
</tr>
<tr>
<td>Geotrichum versiforme</td>
<td>13</td>
<td>0.4</td>
<td>Stool, sputum, bronchial washings</td>
</tr>
<tr>
<td>Sporobolomyces roseus</td>
<td>7</td>
<td>0.2</td>
<td>Sputum, lung biopsy, trachea</td>
</tr>
<tr>
<td>Trichosporon variabile</td>
<td>7</td>
<td>0.2</td>
<td>Sputum, skin lesion, urin</td>
</tr>
<tr>
<td>Saccharomyces cerevisiae</td>
<td>7</td>
<td>0.2</td>
<td>Sputum, wound, bronchial washings</td>
</tr>
<tr>
<td>Torulopsis sphaerica</td>
<td>7</td>
<td>0.2</td>
<td>Sputum</td>
</tr>
<tr>
<td>Candida ater</td>
<td>7</td>
<td>0.2</td>
<td>Sputum, urine, skin</td>
</tr>
<tr>
<td>Saccharomyces telluris</td>
<td>7</td>
<td>0.2</td>
<td>Sputum, urine</td>
</tr>
<tr>
<td>Cryptococcus albidus</td>
<td>7</td>
<td>0.2</td>
<td>Lung biopsy, urine</td>
</tr>
</tbody>
</table>

Figure 2. A guide for yeast identification.

https://scholarworks.uni.edu/pias/vol81/iss1/9
addition to these genera, Lodder (1970) cites *Hansenula* and *Debaryomyces* as genera having medical importance. Their fermentation and assimilation characteristics are listed in Lodder (1970) and serve to assist in speciation of the various sexual forms. One area of difficulty in speciation of the various sexual stages may lie in the recognition of ascospores.

### Description of Sexual Stages

Taxonomically, of the perfect genera *Saccharomyces*, *Kluveromyces*, *Lodderomyces*, *Pichia*, *Hansenula*, and *Debaryomyces*, Lodder (1970) considers *Debaryomyces* the most primitive. From this genus probably arose the other five genera.

Two species of *Debaryomyces*, *D. marama* and *D. hansenii*, are the sexual stages of *Torulopsis candida*. Members of the genus *Debaryomyces* are small yeasts containing lipid globules which reproduce by multiple budding on CMT. Pseudomycelia are slight or absent. After two days in malt agar, cells are spherical to short-oval, single, in pairs or short chains. A one-month-old culture appears greyish white to yellowish.

Asci are formed by conjugation of the mother cell and the bud. The warts which develop on the resulting ascospore (produced on Gorodkowa’s agar) are difficult to see under the light microscope, but can be seen well under oil or with phase microscopy. Usually one (rarely two) ascospore is produced in the ascus. Members of this genus ferment weakly or not at all. *D. marama* produces oval ascospores, 1-4 per ascus, while *D. hansenii* contains 1-2 spherical ascospores in its ascus. Both species of *Debaryomyces* are perfect forms of *Torulopsis*.

*Saccharomyces*, a strong fermenter, reproduces asexually by budding. Blastospores are spherical, oblong, ellipsoidal, cylindrical, or elongate. True mycelia are absent but pseudohyphae may form. Asci, however, contain 1-4 spherical ascospores. Asci do not rupture at maturity. *Torulopsis botnia* is the imperfect form of *S. telluris*.

*Hansenula*, a second genus thought to have arisen from *Debaryomyces*, produces blastospores which bud on a narrow base at various places on the spherical, ellipsoidal, oblong, cylindrical, or elongated cells. At times these cells bear long tapers at one or both ends which are markedly elongated to thread-like. Primitive pseudohyphae are found on CMT under the cover slip and consist of cylindroidal cells which are highly branched and produce a few chains of cells (Blastospores). True hyphae are not produced on CMT.

Asci of *Hansenula* sp. have the shape of vegetative blastospores within which are produced 1-4 smooth walled ascospores which are hat-shaped, hemispherical, spherical, or Saturn-shaped. Ascospores are liberated upon maturity of the ascus.

*H. jadensis* is the only species of medical importance. Its ascospores are hat-shaped with a prominent brim. This species produces very few ascospores and may be the perfect stage of *C. utilis*, which is the only medically important species of *Candida* which assimilates nitrates. Assimilation and fermentation characteristics of *Hansenula* sp. can be found in Lodder (1970).

The genus *Pichia* produces blastospores of various shapes which reproduce by multilateral budding and, in most species, by pseudohyphae. True hyphae may be very limited.

Ascospores are spherical, hat- or Saturn-shaped, usually contain an oil droplet, are generally smooth, and asci contain one to four ascospores. *P. membranaceaefaciens* is the perfect form of *Candida valida* and produces ascospores which are spherical or hemispherical with or without a ledge or brim. One to four ascospores are produced per ascus. *P. guilliermondii* produces only hat-shaped ascospores.

Short descriptions of the ascosporic stages of *Hansenula* and *Pichia* have been presented. Lodder (1970) indicates that “even very primitive species of *Hansenula* and *Pichia* may produce diploid colonies; these may be distinguished from haploid colonies by differences in color, size and texture.” Asci arise from diploid colonies. Different colonies on ascospore media, after two attempts at colony purification, may be a clue that asci are present. If different colonies are revealed on ascospore media and if the yeast produces hyphae, hyphal cuttings are suggested. The hyphal tips are dug out of the medium, squashed, and streaked on malt agar for isolated colonies. The resulting colonies are observed for asci and, if found, diploid cultures are seen at the edge of the colony.

If the yeast does not produce hyphae, sectors of a colony of different textures and colors are streaked on malt agar to isolate colonies, which then can be examined for asci.

*Kluveromyces* produces blastospores which reproduce by budding. Pseudohyphae may be formed. Cells are spheroïd-al, ellipsoidal, cylindrical, or elongate. True mycelia are absent.

Asci rupture upon maturation and liberate ascospores which tend to agglutinate. Ascospores are crescentic, reniform, oblong with obtuse ends, spheroïdal, or prolate-ellipsoidal. One to numerous smooth-walled ascospores are formed within an ascus.

*Kluveromyces* is the perfect stage of *Candida madisoniensis*, *C. pseudotropicalis*, and *Torulopsis spharica*.

The blastospores of *Lodderomyces* are spheroïd-al, or ellipsoidal to cylindrical, and reproduce by budding. Pseudohyphae are abundant; they may be branched or the blastospores may be arranged in clusters or in branching clusters along pseudohyphal cells.

Mature ascospores are large, oblong, with obtuse to prolate ellipsoidal ends. They are occasionally slightly curved and not liberated from the ascus. One or rarely two ascospores are produced in each ascus. *L. elongisporus* is the only species in the genus and is considered to be the sexual phase of *Candida parapsilosis*.

*Sporobolomyces* reproduces vegetatively by simple budding and at times by the formation of pseudo- and/or true mycelia with crosswalls. Blastospores are ovoid to elongate, and on some of the vegetative cells blastospores develop. Blastospores are typically asymmetrical, kidney- or sickle-shaped, and develop in an oblique position at the tips of aerial sterigmata. Upon maturity, these spores are forcefully discharged into the air by a “droplet mechanism.” In this process, a droplet of liquid is excreted at the tip of the sterigma near the hilum of the spore, and when the volume of the drop increases, the drop is discharged, carrying the blastospore with it.
Figure 3. Scheme for the identification of yeasts from clinical material.
A red or salmon-pink color is produced by two species of medical importance on malt agar, but some species are not pigmented.

The cells forming blastospores are considered by some authors to be reduced basidios (Lodder, 1970). Although cyto-
tological data has still not clearly demonstrated a sexual stage, many authors tend to link this genus with the Basidiomycetes.

Basidiomycete sexuality has also been suggested for Candida albicans and C. stellatoidea. The chlamydospores are considered "teliospore homology" (Lodder, 1970). Telio-
or chlamydospores are markedly influenced by environmental conditions and are stimulated by nutritionally deficient media (corn meal tween). The sexual stages of C. albicans and C. stellatoidea have been classified in a separate genus, Syringospora (Lodder, 1970).

Identical physiological characteristics often serve to connect a sexual genus with an sexual one. There are, however, sexual stages which have not yet been associated with an imperfect form and imperfect forms not yet associated with sexual forms. The Identification Scheme (Figure 3) attempts to assist in the identification of either stage. For this reason a yeast may be found in two areas of the Scheme.

**Discussion of the Yeast Identification Guide** (Figure 2). The methods necessary for accuracy in yeast identification have been presented. Their importance justifies the preparation of this Guide, which is in the form of a daily work sheet on which the result of each test is reported. The mere presence of each step on the chart reminds the mycologist what test is completed and what is yet to be performed.

The negative and positive controls are important steps in the chart, for unless the mycologists are certain the systems for testing unknowns are performing adequately, the accurate identification of the unknown is not assured. Only 10 stock cultures of knowns are needed for adequate controls for all steps in the Guide. They should be maintained in stock on YM agar. The knowns used in this laboratory are indicated next to each test in the guide.

**Identification Scheme** (Figure 3). A work sheet (Guide) replete with results probably contains all the information necessary for identification of the unknown. With the use of the Identification Scheme it is possible to identify all the yeast genera isolated during our three-year study as well as those yeasts cited by Lodder (1970) and discussed earlier which have medical importance.

Utilization of the Scheme begins with the microscopic ap-
pearance of the pure unknown on the Luecke plate (corn meal tween agar) (Walker, 1960). The unknown will re-
son in one of three ways: (1) produce blastospores only; (2) produce blastospores, true and/or pseudohyphae, but not arthrospores; or (3) produce blastospores (or the appearance of blastospores), true hyphae, and arthrospores. Identity of the yeast genus, and sometimes species, follows determina-
tion of the response of the yeast on the Luecke plate.

**Production of Blastospores Only.** If an unknown reveals blastospores without true or pseudohyphae, the unknown may belong to the genus Rhodotorula, Cryptococcus, Saccharomyces, Torulopsis, or Debaryomyces. Since Kluyveromyces, Saccharomyces, and Debaryomyces are Ascomycetes, an obvious second step in identification necessitates inoculation of Corn-
dkowa agar and malt agar for production of ascospores.

(Ascosporic yeasts will be discussed later.)

Pigment production, or the lack of it, on malt agar and the presence or absence of urease will assist in identification.
Identification of Yeasts

The typical chlamydospores of *C. albicans* (Walker, 1960; Lodder, 1970). The Luecke plate therefore assures purity, and contains a "built-in" mechanism for stimulating all yeasts to reveal typical morphology, including the large number of typical chlamydospores of *C. albicans*.

Rarely do we depend on one technique or one step for the identification of any species. Additional steps are often required for the identification of atypical strains. For example, absolute dependency on corn meal tween for the identifications may be deceptive since *C. tropicalis* and *C. stellatoidea* have been known to produce chlamydospores on this medium. Furthermore, a rare strain of *C. albicans* may not produce chlamydospores (Walker and Huppert, 1959). However, morphologic characteristics of *C. stellatoidea* and *C. tropicalis* are often clues that *C. albicans* is not present on corn meal tween, notwithstanding the occasional chlamydospores which may be seen. When formed, the chlamydospores of *C. tropicalis* are scanty and not typical of those of *C. albicans*.

If a yeast fails to reveal identifiable structures on corn meal tween in 24 hours, the culture is allowed additional incubation time, and assimilation and fermentation tests are performed immediately. Additional incubation time on corn meal tween has ultimately revealed typical morphology which was shown by fermentation and assimilation tests.

A well-known procedure for *C. albicans* identification is the "germ tube test" (Blair et al., 1970). Identification is effected in a few hours by the production of "germ tubes" which are filamentous outgrowths from the blastospores or chlamydospores after exposure to mouse peritoneal fluid in *vivo* or to human or animal serum in *vitro*. This technique may be helpful for rapid identification of atypical *C. albicans* strains since atypical cultures tend to prolong the identification period. However, additional studies should be undertaken to determine whether these atypical strains would produce typical germ tubes. This technique would also be helpful in ruling out the presence of *C. albicans* in the event a *C. tropicalis* or *C. stellatoidea* produced chlamydospores on corn meal tween. However, our identification steps include assimilation and fermentation studies on all atypical *Candida* isolates and all species of *Candida* which do not produce the typical morphological characteristics of *C. albicans* on corn meal tween. *C. stellatoidea* and *C. tropicalis* would be identified using these techniques, and therefore additional methods are not necessary.

To summarize, we prefer the use of corn meal tween agar to culture purity and to stimulate typical morphologic characteristics, including those of *C. albicans*. If *Candida* species other than *albicans* are present, or if atypical chlamydospore-producing yeasts are present, fermentation and assimilation tests are performed while the yeast undergoes prolonged incubation on corn meal tween. These steps serve to identify each isolate to species while the one-step germ tube technique merely ascertains the presence of *C. albicans*. Since our procedures not only identify *C. albicans* but other yeast genera to species a test which identifies only one species is not necessary.

When typical *C. albicans* chlamydospores and morphology are not produced on corn meal tween, sugar assimilation and fermentation tests are warranted. Only one medically important species of *Candida*, *C. utilis*, assimilates nitrate (Lodder, 1970). This is the only genus of this group (*Candida, Pichia, Hansenula, Lodderomyces, Sporobolomyces, and Kluyveromyces*) which does not produce ascospores or ballisto-
spores. If the unknown assimilates nitrate but is not C. utilis, other genera in this group must be considered. Assimilation and fermentation results will help to identify the Candida species (Lodder, 1970). The genera Lodderomyces, Pichia, and Kluyveromyces, and the majority of Candida species cannot assimilate nitrates. Ascospore media results are important in identification of those genera which may or may not assimilate nitrate. Of those genera which assimilate nitrate, Hansenula produces hat- or Saturn-shaped ascospores and grows well at 45-48°C. These characteristics differentiate Hansenula, Candida, and important species of Sporobolomyces.

Hansenula jadinii has been isolated from clinical material and is the perfect stage of C. utilis. Its fermentation and assimilation characteristics are seen in Lodder (1970).

Sporobolomyces is differentiated from Candida species in its production of characteristic ballistospores. The latter have been described under the description of sexual stages. Two species of Sporobolomyces have attained importance in medical mycology. Their fermentation and assimilation characteristics are listed in Lodder (1970).

Production of Blastospores, Blastospore-Like Cells, True Hyphae and Arthrospores. The presence of these characteristics on the Luecke plate indicates that the unknown belongs to one of two genera, Geotrichum or Trichosporon. Geotrichum, although often yeast-like in culture, is not classified as a yeast. Yet its blastospore-like arthrospores often seen in primary isolates are frequently interpreted as yeast cells. Growth of Geotrichum sp. on CMT for three days or more will reveal characteristic arthrospores. Depending on the species, a Geotrichum may reveal asymmetric germ tube formation. But this characteristic is often confused with the zigzag fashion of arthropore production in Trichosporon species. Trichosporon, however, produces budding blastospores in addition to arthrospores, which serves to differentiate Trichosporon from Geotrichum, which does not form budding cells.

Those species of Trichosporon recovered from clinical specimens in this laboratory are listed in Table 1.

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