Mycotic "Pseudotumors" of the Breast

Report of Four Cases

Karlhanns Salfelder, MD, Jan Schwarz, MD

Four patients are reported with mycotic pseudotumors of the breast. Two of the granulomas were due to involvement by histoplasmosis, one by blastomycosis, and one by cryptococcosis. In the three patients who were treated surgically, tumor was suspected preoperatively in two and an infected cyst in one. The fourth granuloma, that due to cryptococcosis, was unsuspected clinically and discovered at autopsy. Surgical excision was effective therapy in those so treated. A critical review of the literature revealed only one other previously published case (cryptococcic mastitis) of deep mycosis of the breast.

The breast has rarely been reported as a site of fungal infection and, of the cases on record, only one is acceptable as deep mycosis. However, since from one single laboratory four cases of mycotic mastitis were reported in a nine-year-period, the true incidence must be higher than recorded, and this differential diagnosis must be considered in breast "tumors" that presently are looked for and detected more actively with the help of roentgenographic methods.

REPORT OF CASES

CASE 1.-A cryptococcosis was found in both breasts of a young woman with a ten-month history of disseminated lupus erythematosus with nephritis and renal failure. She was treated with cortisone and dialyzed on several occasions. On autopsy, disseminated cryptococcosis was found in lungs, brain, meninges, liver, spleen, ovaries, and breast. Numerous cryptococci were seen in the breast tissue (Fig 1). A mild meningitis was observed across the cerebellum. As is frequently the case in cryptococcosis, no inflammatory response was seen in the other tissue (Table).

CASE 2.-A unilateral blastomycosis of the breast clinically appearing as an abscess was diagnosed preoperatively as (infected?) para-areolar cyst. After excision, drainage, and topical antifungal treatment, complete healing took place. Fungus cells with single buds morphologically consistent with Blastomyces dermatitidis were seen, as well as a characteristic tissue reaction (Fig 2). After eight years, the patient is well and there are no signs of disease in the breast or elsewhere in the body (Table).

CASE 3.-A unilateral mastitis was found in a young woman with a large caseous nodule in the breast. Preoperatively, the diagnosis was breast tumor. Yeast cells of Histoplasma capsulatum were seen in the histologic sections (Fig 3); the titer in the complement fixation test was 1:32. The patient is well and the breast normal 1" years after the resection (Table).

CASE 4.-The findings were clinically interpreted as breast tumor. The mass in the lateral mammary region was excised and found to be lymph nodes of the anterior axillary area, with granulomas. In the numerous, partly necrotic epithelioid cell granulomas, only single fungus cells could be seen (Fig 4); on culture, H capsulatum grew and the results of serologic study were abnormal. The patient is alive and apparently well (telephone report); a clinical re-examination could not be performed (Table).

COMMENT

Cryptococcic mastitis has been described only once before, also at autopsy. The diagnosis was made on re-examination of an old museum specimen. The fungus cells in our patient, situated in the stroma of the breast without any inflammatory reaction, could easily be overlooked. Lack of inflammatory response is known as one form of "reaction" in cryptococcosis. The bilateral character of breast involvement points to a hematogenous pathway. The two reports on blastomycotic mastitis in the literature require only brief comments; neither case justifies its inclusion as blastomycotic. Jung, after discussing a case of disseminated "blastomycosis," described a "sporotrichoma" of the breast. The first case has no breast involvement; the second is not blastomycotic.

Kühlmeier & Kreitner4 identified from a granulomatous breast lesion a yeast (Torulopsis inconspicua). This organism is not an accepted obligatory pathogen and has no relation to blastomycosis (in America identified by the etiologic agent B dermatitidis).

Beside these two reports, no mention of involvement of the breast in blastomycosis was found in the literature.

In our patient with blastomycosis, we assume that a pri-
<table>
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<tr>
<th>Patient No./Age, yr</th>
<th>Duration, Clinical Findings, &quot;Preoperative Diagnosis&quot;</th>
<th>Gross Appearance &amp; Histologic Findings</th>
<th>Other Findings</th>
<th>Therapy &amp; Follow-Up</th>
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<tbody>
<tr>
<td>1/20</td>
<td>10 mo</td>
<td>Not recognizable grossly; microscopically: pale yeast cells of different sizes with capsule; positive stain with mucicarmine;</td>
<td>Disseminated lupus erythematous with generalized cryptococcosis</td>
<td>Steroids; death</td>
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<td>2/30</td>
<td>2 mo; para-areolar swelling fluctuation &amp; deep induration; &quot;infected cyst&quot;</td>
<td>Suppurated mass of 4 cm diameter; purulent &amp; granulomatous inflammation with destruction of breast tissue; epithelioid cell granulomas &amp; multinucleated giant cells; yeast cells 10µ to 15µ with occasional single, broad-based buds, intensely stained with Grocott &amp; Gridley procedures; consistent with B dermatitidis</td>
<td>Clinically well; normal chest roentgenograms</td>
<td>Surgical excision; amphotericin B ointment; well after 8 yr</td>
</tr>
<tr>
<td>3/21</td>
<td>Slow-growing mass; &quot;breast tumor&quot;</td>
<td>Firm nodule 3 cm in diameter, para-areolar; partly caseated granulomatous mass replacing breast tissue; epithelioid cell granulomas, multinucleated giant cells, lymphocytes; Grocott- positive ovoid yeast cells 2µ to 4µ; occasional buds</td>
<td>Complement fixation test (CFT) (histoplasmosis) 1:32; no evidence of systemic or pulmonary</td>
<td>Surgical excision; well 2 yr later</td>
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<tr>
<td>4/55</td>
<td>Mass noted 6 mo ago, slowly increasing; &quot;breast tumor? axillary lymphadenopathy?&quot;</td>
<td>Nodular mass with individual &quot;tumors&quot; up to 5 cm from anterior axillary line, clinically &amp; on xerogram considered breast mass; excision of axillary fat 13:13:5 cm with large brown lymph nodes; massive epithelioid cell granulomas in huge lymph nodes with discrete necrosis; rare organisms on Grocott stain, 2u, to 4u,</td>
<td>Culture positive for H capsulatum; CFT (histoplasmosis) 1:4; no evidence systemic histoplasmosis</td>
<td>Surgical excision; alive 1 yr later; uncooperative, re fuses check-up of</td>
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by inhalation of spore-containing dust, the disease commonly disseminates during primary infections. The most frequent type of dissemination is self-limited and benign; in infants, however, the dissemination can be disastrous. Histoplasmic metastatic lesions (the primary focus is almost, without exception, in the lung) have been found in practically every organ; spleen and liver are the most common, and adrenal and brain the most consequential sites.

Blastomycosis shares the pathogenesis with histoplasmosis; soil-borne spore-containing dust is inhaled and a pulmonary primary complex is formed. The skin is by far the most common site of peripheral dissemination, with prostate, joints, and meninges competing as sites of importance.

Fig 2.-Granulation tissue (case 2) with multinucleated giant cells replacing breast tissue (hematoxylin-eosin, x300). Inset, Budding yeast cell of B dermatitidis. Note broad-based daughter cell (Grocott stain, x 1,000).

Fig 3.-Granulomatous reaction (case 3) in periphery of caseous focus of breast, with abundant epithelioid cells and multinucleated giant cells (hematoxylin-eosin, x250). Inset, Three cells of H capsulatum from necrotic area (Grocott stain, x 1,300).

The influence of steroid therapy, in the activation of histoplasmosis and blastomycosis is undeniable but, compared with cryptococcosis, much less prominent. There is no evidence that any of the three diseases is transmitted from man to man, or directly from animal to man. Pigeons (in cryptococcosis) and blackbirds and bats (in histoplasmosis) play a part in connection with the presence of Cryptococcus neoformans and H capsulatum, respectively, in soil, but the exact role of these winged animals is not established. The mechanism of how B dermatitidis reaches the soil is unknown.

It is difficult to determine why mycotic mastitis has not been described more often. This may be due to lack of suspicion or to failure to recognize the causative agents in...
Fig 4.-Epithelioid cell granulomas (case 4) with giant cells in lymph node (hematoxylin-eosin, x 250). Inset, Granuloma with central necrosis (hematoxylin-eosin, X300).

tissue sections or on culture. The use of special stains, especially the Grocott-methenamine silver stain, is indispensable for detection of fungi in tissue. As cases of granulomatous mastitis (simulating carcinoma) are found clinically more often,9,10 mycotic mastitis should become more frequently recognized.

References