Case 18
A 38 year-old Thai man from Samutprakan

**Chief complaint:** Multiple discrete indurated erythematous to purplish dermal papules on trunk and extremities for one year

**Present illness:**
The patient developed productive cough for 3 months with low grade fever and significant weight loss of 10 kilograms, so he went to visit a pulmologist and chest film revealed bilateral diffuse reticulonodular infiltration. Further investigations were performed e.g. anti-HIV, blood culture for aerobe. A few days later, the blood culture was detected both budding yeasts and gram positive cocci in chain. His positive serology for HIV was definitely confirmed.

Suddenly, he was called for emergency treatment. Numerous encapsulated budding yeasts were also found in his cerebrospinal fluid.

A dermatological consultation was established later to evaluate multiple discrete indurated asymptomatic erythematous to purplish dermal papules on trunk and extremities which lasted for one year without any previous treatments.

**Past history:**
- He has no other underlying diseases.
- He had experienced unprotected sexual intercourse 10 years ago.

**Physical examination:**
- V/S: T 38.6°C, P 140/min, RR 24/min, BP 134/93 mmHg
- HEENT: mildly pale conjunctivae, present of oral thrush
- Lymph node: no lymph node enlargement
- CNS: no stiffness of neck
- Other systems: unremarkable

**Dermatological examination:**
- Multiple discrete indurated erythematous to purplish dermal papules on trunk and extremities
**Histopathology** (S16-16484A, Left arm)

- Perivascular and interstitial inflammatory-cell infiltration of lymphocytes and histiocytes with large vacuolated cytoplasm
- Numerous round to oval spores with thick capsules within some histiocytes
- Increased number of dilated bizarre-shaped thin-walled vessels surrounding preexisting vessels and interstitium
- Increased number of spindle cells between collagen bundles
- **Immunohistochemistry** for HHV-8: scattering positive

**Investigation:**
- CBC: WBC 4,190 (N 93%, L 4%, Mo 2%, Eo 1%) , Hb 10.2, Hct 31.2%, Platelet 346,000
- CD4 T cell 5% (8 cell/mm³)

- CXR: bilateral diffuse reticulonodular infiltration
- Sputum AFB: negative for 3 days
- Blood culture for aerobe: *Granulicatella adiacens*, *Cryptococcus neoformans.*
- BAL culture for aerobe: *Cryptococcus neoformans.*
- CSF culture for aerobe: *Cryptococcus neoformans.*
- BM culture for aerobe: *Cryptococcus neoformans.*
- Skin culture for aerobe: *Cryptococcus neoformans.*

**Diagnosis:** Kaposi sarcoma with disseminated cryptococcosis

**Treatment:**
- IV amphotericin B 0.7 mg/kg/day for 4 weeks then fluconazole 800 mg/day
- IV paclitaxel weekly 2 doses (cessation due to CMV colitis)
- Delayed HAART 6 weeks after treatment of cryptococcosis
- Ceftriaxone 2 gm IV OD for 14 days

**Presenter:** Thiraphong Mekwilaiphan, MD  
**Consultant:** Ploysyne Rattanakaemakorn, MD

**Discussion:**
Kaposi sarcoma is an angioproliferative neoplasm,¹ presents as a slow growing erythematous to purplish papule, plaque, or nodule² on the skin especially on lower extremities³, the mucous membranes lining the mouth, nose, and throat, lymph nodes, or other organs e.g. lung.⁴ ⁵ It is different from other cancers in that lesions may begin in more than one place in the body at the same time.⁶

Those infected with HHV-8 who are most likely to develop Kaposi sarcoma have immune systems weakened by disease or by drugs given after an organ transplant⁶ as well as in this immunocompromised patient which the immunohistochemistry for
HHV-8 is scattering positive.

There are four distinct types of Kaposi sarcoma as shown in table 1.7

<table>
<thead>
<tr>
<th>Type</th>
<th>Population</th>
<th>Clinical</th>
<th>Course</th>
</tr>
</thead>
<tbody>
<tr>
<td>Classic</td>
<td>Older men (50-80 years)</td>
<td>Usually confined to lower</td>
<td>Usually indolent, survival</td>
</tr>
<tr>
<td></td>
<td></td>
<td>extremity</td>
<td>10-15 years</td>
</tr>
<tr>
<td>Endemic (African)</td>
<td>Young black males, 15-40 yrs, and children</td>
<td>Localized nodular lesions or large exophytic, aggressive lesions</td>
<td>Nodules indolent, aggressive lesion survival 3-5 years</td>
</tr>
<tr>
<td>Iatrogenic</td>
<td>Immune suppressed (e.g., renal transplant)</td>
<td>Localized or widespread involvement</td>
<td>May regress when immune suppressants discontinued</td>
</tr>
<tr>
<td>Epidemic (AIDS-related)</td>
<td>Primarily gay men in U.S., Europe; adults in Africa</td>
<td>Head, face, neck, GI and lung most common</td>
<td>Fulminant, survival 1-3 years without effective HIV therapy</td>
</tr>
</tbody>
</table>

AIDS, acquired immunodeficiency syndrome; GI, gastrointestinal system; HIV, human immunodeficiency virus.

The typical histopathology shows slit-like spaces that frequently contain red blood cells. Plasma cells and hemosiderin deposits are also usually apparent. Eosinophilic hyaline globules, 1–7 μm in size, are commonly present and form grape-like agglomerations which are predominantly intracellular. These structures are thought to represent digested erythrocytes, as the neoplastic cells seem to have phagocytic activity.8

For AIDS-related Kaposi sarcoma, the risk of developing Kaposi sarcoma for untreated AIDS patients is estimated to be 300 times greater than that of other immunosuppressed individuals and 20,000 times greater that of the general population9 as well as the more aggressive clinical course.10

Several modalities of treatment have been used for Kaposi sarcoma including chemotherapy, radiation therapy, surgical excision and Highly Active Anti-Retroviral Therapy (HAART) in patients with AIDS-related Kaposi sarcoma.11 The choice of treatment is determined by the stage of Kaposi's sarcoma, its rate of progression, the degree of immune competence and HIV associated diseases.12

Cryptococcosis is a potentially severe infection by Cryptococcus neoformans. that usually occurs in a setting of immunosuppression13, especially in HIV patients with a CD4 count less than 200 cells/mm³,14 mainly affects with pulmonary and resulting and/or meningeal involvements or disseminated infections.15 Clinical manifestations of cutaneous cryptococcosis are varied. Lesions may resemble molluscum contagiosum, or appear acneliform, nodular, herpetiform, cellulitic, or keloid-like.16

The diagnostic evaluation depends on organ involvements which microbiological confirmation is required for definite diagnosis. It can easily found encapsulated unipolar budding yeasts, varying in size under light microscope. For the culture media it can grow in blood agar in aerobe culture and the appropriate culture media is Sabouraud’s dextrose agar.

The current guidelines from the Infectious Diseases Society of America (IDSA) recommend treating HIV patient with at least 2 weeks of induction therapy with a combination of amphotericin B deoxycholate 0.7–1.0 mg/kg per day intravenously plus flucytosine 100 mg/kg per day orally in 4 divided doses followed by fluconazole (400 mg per day orally) for a minimum of 8 weeks and maintenance with fluconazole 200 mg per day orally life-long. Initiation of HAART should be considerably prescribed 2–10 weeks after commencement of initial antifungal treatment17 to avoid cryptococcosis-associated immune reconstitution inflammatory syndrome (C-IRIS) which is fatal condition.18

As the literature review, the coexistent Kaposi sarcoma and cutaneous cryptococcosis is rare. There were not exceed ten report cases which usually found in AIDS patients that histologically conspicuous spindle cell component for Kaposi sarcoma and small aggregates of cryptococcal yeasts. Unfortunately, half of them died during treatment.19, 20

As well as this patient whose CD4 count was only 8 cells/mm³ is another one who had coexistent Kaposi sarcoma and cutaneous cryptococcosis. He was prescribed IV amphotericin B 0.7 mg/kg/day for 4 weeks then oral fluconazole 800 mg/day. Before the initiation
of HAART, the lesions on his feet became ulceration that cannot be distinguished the effect of cryptococcosis or Kaposi sarcoma, so IV paclitaxel was prescribed weekly, concurred with systemic antifungal, and discontinued due to infection of CMV colitis meanwhile HAART was then initiation 6 wks after treatment of cryptococcosis which could be effective in AIDS-related Kaposi sarcoma. Finally, the lesions were remised into post-inflammatory hyperpigmentation and the patient is still alive without recurrence in 3 following months.

References: