Cryptococcal meningitis after transnasal transsphenoidal pituitary microsurgery of ACTH-secreting pituitary adenoma

A case report

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Abstract

Rationale: Microbial infection should be regarded in the differential diagnosis of neurosurgical complications after transnasal transsphenoidal pituitary microsurgery, albeit cryptococcal meningitis is rare. This article will discuss the risk factors of cryptococcal meningitis in patients underwent transnasal transsphenoidal pituitary microsurgery, and summarize the potential origins of infection.

Patient concerns and diagnosis: Here, we report a case of 37-year-old male who had cryptococcal meningitis after transnasal transsphenoidal pituitary microsurgery of a relapsing ACTH-secreting pituitary adenoma.

Intervention: Standard therapy for Cryptococcus neoformans (fluconazole [400mg per day] and flucytosine) was administered and followed by maintenance dose.

Outcomes: The patient had been on treatment for one and a half years during follow-up and reported neurologically well with repeated negative cerebrospinal fluid (CSF) culture until sudden death of heart arrest.

Main lessons to learn: C neoformans can be a possible cause of meningitis in immunocompetent patients after transnasal transsphenoidal pituitary microsurgery. Risk factors, such as pre-existed pulmonary infection and Cushing-associated hypercortisolism, should be stressed. Promising preventive measures may include preoperative routine sputum smear and India-ink stain for screening, preoperative treatment of cryptococcal pneumonia, postoperative antibiotic management, and a more secure skull base reconstruction. Radiation and pharmaceutical treatment may be alternative for recurrent Cushing disease.

Abbreviations: ACTH = adrenocorticotrophic hormone, CSF = cerebrospinal fluid, WBC = white blood cell.

Keywords: ACTH, cryptococcal meningitis, pituitary adenoma, transnasal transsphenoidal pituitary microsurgery

1. Introduction

Microbial infection should be regarded in the differential diagnosis of posttransnasal transsphenoidal neurosurgical complications. Cryptococcus neoformans was once recognized as an opportunistic pathogen among immunocompromised patients. However, meningitis caused by C neoformans, as a result of fungal dissemination from initial pulmonary infection in immunocompetent patients, was rarely seen. Here, we describe a patient who developed cryptococcal meningitis after transnasal transsphenoidal excision of adrenocorticotrophic hormone (ACTH)-secreting pituitary adenoma. This should be the first report of immunocompetent individual who was infected with C neoformans after surgeries with transnasal transsphenoidal approach.

2. Case report

This work was approved by the hospital’s medical ethics committee. Informed consent was obtained in accordance with the institutional guidelines.

A 37-year-old man after transnasal transsphenoidal excision of ACTH-secreting pituitary adenoma 2 years ago was admitted with relapse of hypercortisolism for 4 months. Past history collection revealed negative history other major medical problems. He had no surgical history except the previous surgery for Cushing disease and he was not taking medicines. On admission, baseline inferior petrosal sinus to peripheral ACTH gradient was greater than 2, and inferior petrosal sinus to peripheral gradient after desmopressin activation was greater than 3. Contrast-enhanced pituitary magnetic resonance imaging evidenced hypointensity around the right-half pituitary gland. Thus, a transnasal transsphenoidal reoperation for relapsing pituitary tumor was indicated as elective surgery. Intraoperative leak was detected, and primary reconstruction of sella turcica was done using fascia lata and fibrin glue with fat packing of the sphenoid sinus. However, postoperative cerebrospinal fluid (CSF) leak still occurred and hypercortisolism was not alleviated with serum cortisol >75μg/dL. Lumbar CSF
drainage was placed and CSF was regularly collected for laboratory study. Prophylactic cefuroxime was then administered. The 1st month after surgery, the patient developed intermittent fever between 37 and 39°C. The diagnosis of suppurative meningitis was confirmed based on both the typical turbidity and the microscopic leukocyte counting of CSF. CSF germiculture tests revealed oxacillin-resistant Staphylococcus epidermidis and Acinetobacter baumanii. Antibiotic scheme was modified into 3 courses of meropenem and vancomycin followed by 1 course of cefazidime, and the patient gradually restored normal body temperature and CSF test results returned to normal level.

Second month after surgery, the patient was presented with recurrent hyperpyrexia, neurological examination revealed stiff neck and Lasegue sign. Laboratory CSF study suggested central nervous system infection, with a leukocyte count of 4869/mm³ (92% neutrophils and 8% monocytes), protein 5.58g/L, glucose 0.5mmol/L, and chloride ion 117mmol/L. The patient was diagnosed with central nervous system infection based on the CSF results. But after administrating cefazidime and vancomycin empirically for 2 weeks, the CSF still showed high leukocyte count. Therefore, further microbiological investigation on CSF was conducted, including India-ink stain, which broke through with an encouraging result. Further past history collection regarding the patient’s household environment did not suggest frequent contact with birds. Fungal culture of CSF grew cryptococcus species, and the titer of cryptococcal antigen was 1:8192. Further serum cryptococcal antigen titer was 1:1024. The patient started standard therapy as daily intravenous amphotericin B deoxycholate at 1mg and oral flucytosine at 1g tid. However, there was acute onset of epileptic seizure after administration of amphotericin B deoxycholate. Standard therapy had to be discontinued and alternated to intravenous fluconazole (400mg qd) plus oral flucytosine (1g tid) instead.

Detailed data of CSF biochemistry during antifungal course for postsurgery cryptococcal meningitis was presented in Figure 1. First week of antifungal therapy showed little effect. The patient was yet presented with fever, chill, even delirious status. But 1 week later, the patient’s general condition went better. CSF re-evaluation revealed white blood cell (WBC) count 70/mm³, protein 1.39g/L, and glucose 2.2mmol/L, with negative fungal culture. The patient had been well on this antifungal therapy for 3 months. He then continued with maintenance therapy of oral fluconazole (400mg per day) and flucytosine, without clinical relapse. Repeated CSF smears and CSF culture confirmed microbiologically negative.

One and a half years following the surgery, the patient developed acute onset of cyanosis, with oxygen saturation lower than 70% and blood pressure lower than 90/60mmHg. Complete blood count was abnormal for WBCs as high as 12.6/mm³ with neutrophilic predominance. Bedside chest X-ray revealed extensive plaques in his bilateral lungs. Pulmonary infection was highly suspected. Unfortunately, the patient had a heart arrest the other night, even cardiopulmonary resuscitation and epinephrine failed to retrieve his life.

3. Discussion
To our knowledge, cryptococcal meningitis following transnasal transsphenoidal pituitary microsurgery has not been reported previously. Common surgical complications of transnasal transsphenoidal pituitary microsurgery include pituitary dysfunction, CSF leak, and infection, which mainly comprises of sphenoiditis and meningitis. Transnasal transsphenoidal surgery entails the passage through nasal cavity, which is a flora reservoir for Staphylococcus aureus, Haemophilus influenzae, Streptococcus pneumoniae, Staphylococcus epidermidis, and other streptococcal species. Such a clean-contaminated passage renders those microbes a way to the brain. Organisms predominantly involved in this surgical context are S aureus, coagulase-negative staphylococci, gram-negative bacilli, and enterococci; however, C neoformans is rare condition.
C. neoforms has been recognized as an opportunistic pathogen among immunocompromised patients. Acquired immune deficiency syndrome epidemic has driven cryptococcal disease worldwide. Other underlying associations include the use of immunosuppressant, solid organ transplantation, malignancy, and congenital immunological disorders. Apart from individuals with immunological defects, there have been reports of immunocompetent hosts developing cryptococcal infection. By observation, cryptococcal meningitis among immunocompetent patients tends to have an indolent onset but harsh CSF result, with higher initial WBC count and higher protein level. And our patient, who was presented with fever and stiff neck, just had typical CSF results.

Hypercortisolemia should be a risk factor for postoperative infection. Therefore for patients with Cushing syndrome who are suffering extreme hypercortisolism, the likelihood of developing bacterial or opportunistic infection can be greatest. In our case, this is a 37-year-old male patient with relapsing ACTH-secreting pituitary adenoma. He did not arrive hormonal remission after transnasal transphenoidal surgery, rendering him vulnerable to opportunistic infection. Other risk factors associated with immune compromise, such as malignancy or HIV infection, are not applicable to this patient. Another concern for cause of infection is pulmonary originated pathogens, that is, cryptococcal meningitis might be secondary to asymptomatic cryptococcal pneumonia. Previous report has demonstrated that patients with pulmonary cryptococcal infections usually show no symptoms until the disease disseminates to extrapulmonary sites, most notably to central nervous system. In our case, this partially due to the stress of surgical excision, which bring down the immune system and unblock the fungal infection from disseminating into central nervous system. Direct spread from respiratory tract (nasopharynx) to brain through sphenoid sinus cannot be neglected as well.

This case deserves further reflections on measurements to avoid cryptococcal meningitis after transnasal transphenoidal neurosurgery, especially for patients with Cushing syndrome. One of the major risk factors for neurosurgery-associated infection is persistent CSF leak. Since CSF leak is the prime concern for transnasal transphenoidal pituitary microsurgery, different procedures on skull base reconstruction have been put forward. The combination of fascia lata graft, nasoseptal flap, and dural sealant show long-term effectiveness in secure closure of the cranial base. Reconstruction with bone or bone substitute proved stable.

Relapsing pituitary adenoma accounts for large percentage of pituitary surgeries. But do think twice before you operate on a relapsing pituitary adenoma. There are multiple reasons. First of all, reoperation of pituitary adenoma usually has poor outcome, unlikely to achieve biochemical remission. And the hypointensity foci on preoperative magnetic resonance imaging, which was suspected of recurrent tumor, might prove to be scar tissue by pathology. In such circumstance, positron emission tomography imaging potentially serves as a valuable tool in distinguishing recurrent adenoma, residual pituitary gland, and scar tissue. Also, a group of Cambridge team testified that 11C-methionine positron emission tomography–computed tomography may play a role in detection and treatment of ACTH-secreting tumors in Cushing syndrome.

Apart from surgical reintervention of the ACTH-secreting adenomas, there are alternatives including radiotherapy, adrenal surgery, and pharmaceutical therapy. Progression has been made in the field of drug development, and pasireotide, mifepristone, ketokonazole, metyrapon, and cabergoline have approved effective in managing Cushing disease.

Regarding treatment of cryptococcal meningitis, standard antifungal scheme for C. neoforms has arrived at consensus and proved effective. But amphotericin B can bring the problem of drug intolerance. However, our experience suggests intravenous fluconazole might be an alternate.

This case serves to demonstrate that C. neoforms can be a possible cause of meningitis after transnasal transphenoidal pituitary microsurgery, and physicians must be aware of it. Perioperative management includes routine sputum smear and India-ink stain for pneumonia, preprocessing of cryptococcal pneumonia, empirical antibiotic management, regular and frequent neurological examination, and a more secure skull base reconstruction may secure a more stable postsurgical course.

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