Meningoencephalitis, communicating hydrocephalus, drug reaction with eosinophilia and systemic symptoms (DRESS) and dengue infection in an HIV patient, Sri Lanka

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Abstract

Drug reaction with eosinophilia and systemic symptoms (DRESS) is a potentially life threatening, rare drug hypersensitivity reaction. Here, we present a case of meningoencephalitis, communicating hydrocephalus in a person living with HIV (PLHIV) who developed DRESS syndrome and dengue infection.

Key words: Meningoencephalitis, Communicating Hydrocephalus, Drug reaction with eosinophilia and systemic symptoms (DRESS), Dengue infection, HIV, Sri Lanka.

Introduction

Drug reaction with eosinophilia and systemic symptoms syndrome (DRESS) is a potentially life threatening, rare drug hypersensitivity reaction which results in a skin eruption, eosinophilia and systemic symptoms due to involvement of internal organs such as liver, lungs, kidney, central nervous system (CNS), lymphatic system and heart. It usually has a long latency period such as 2 to 8 weeks. Co-trimoxazole (trimethoprim-Sulfamethoxazole) is a known drug which can cause DRESS. (1) Co-trimoxazole is frequently used as pneumocystis pneumonia (PCP) prophylaxis in HIV positive patients who have CD4 counts below 350 cells/μl. INAH prophylaxis of 300mg daily for tuberculosis given for 6 months is started in all people living with HIV prior to starting antiretroviral therapy (ART) as per WHO guidelines. There is no available literature regarding the incidence of DRESS among HIV positive patients. In 2018, estimated HIV prevalence in Sri Lanka was 0.02%. (2) Meningoencephalitis in PLHIV can be the result of a viral, bacterial, parasitic, fungal or HIV unrelated pathologies. HIV is not known to infect neurons, but causes neuronal death due to indirect ways. Therefore, distinct from other types of viral encephalitis, HIV does not itself acutely present with meningoencephalitis. (3) DRESS syndrome can also have central nervous system complications such as meningoencephalitis. (1)  

Case report

Forty years old male was diagnosed to have HIV infection in Jan 2019. He was asymptomatic on presentation and following exclusion of opportunistic infections and active tuberculosis (TB) he was commenced on INAH prophylaxis for TB and co-trimoxazole prophylaxis for PCP in February 2019. He was started on tenofovir, emtricitabine and
efavirenz (TDF+FTC+EFV) regimen 2 weeks after starting INAH prophylactic therapy. Four weeks after starting Co-trimoxazole, he developed fever with chills and rigors and generalized pruritic macular papular rash (Figure 1) and elevated liver functions were noticed. But he was not dyspneic, jaundiced or pale. His blood pressure was 120/70 mmHg. All systemic examinations were normal. He was hospitalized and all the drugs were withheld due to the reaction. While in the hospital, his ALT was rising from 44 IU/l to 2,666 IU/l; AST to 271 IU/l; total bilirubin was 66.5 IU/l and direct bilirubin increased to 63.1 IU/l. His eosinophil count rose to 9.3%. His total WBC was 6.87 x 10^3 cells/μl (reference range 4-10) with an absolute eosinophil count of 0.68 x 10^3 cells/μl (reference range 0.02-0.5). He was clinically diagnosed to have DRESS syndrome by the consultant dermatologist. He was treated with oral chlorpheniramine, prednisolone 30mg twice daily tapering off dose, betamethasone local application and was monitored for hepatic encephalopathy. He was continuing to have daily high fever spikes for initial 12 days without any evidence of underlying infections. He was discharged after 19 days of hospital stay following settling of his fever and normalization of liver transaminases. He was restarted on TDF, FTC, raltegravir (RAL) regimen and it was withheld one month later due to increasing liver transaminases. Even though there were few attempts to reintroduce ART, it failed due to his rising liver enzymes.

He was again hospitalized in September 2019 with fever, persistent erythematous rash and low platelets. He was managed as dengue fever with his positive dengue NS1 antigen and dropping platelet count. He was discharged after 10 days of the hospital admission once he became stable. Two days after the discharge he was readmitted with fever, photophobia, headache, and confusion with a GCS of 13/15 without any focal neurological signs. Fundal examination was normal. Respiratory system was normal with no crepitations or rhonchi. Blood pressure was 116/86 mmHg and pulse rate was 120/min. Abdominal examination was normal. A full opportunistic infection screening was repeated (table 1, table 2, table 3). Non-contrast computer-tomography (NCCT) brain was normal at admission. Lumbar puncture was done (table 2) and revealed neutrophil pleocytosis and gram positive diplococci. Antimeningetic doses of antibiotics IV merapenem 2g 8hrly and IV vancomycin 500mg 6hrly was given. Empirically oral valgancyclovir 900 mg was started. Patient’s conscious level improved initially with antibiotics, but it was deteriorated on third day rapidly and the high fever spikes were continued. Urgent MRI brain revealed communicating hydrocephalus and basal meningeal enhancement without any focal lesions. External ventricular drainage was inserted and CSF was sent for investigations (table 2) to diagnose underlying cause for having non resolving meningoencephalitis with communicating hydrocephalus. There were fluctuations in his serum sodium which was initially 133mmol/l and went down to 120 mmol/l but later it became normalized with hypertonic saline. His serum potassium level remained normal. His urinary sodium level, potassium level, urine osmolality, serum osmolality and urine to serum osmolality ratio was normal. His blood urea and creatinine level remained normal with 0.83 mg/dl an eGFR of 110 ml/min/1.73 m2. Persistently his GCS was less than 8, pupils being sluggishly reactive, 4mm in size with bilateral extensor plantar reflex and in decerebrate position.

The repeat CSF report became positive for Toxoplasma antibodies IgM and IgG CSF Toxoplasma titre were not done due to unavailability. Although confirmatory test, CSF T.gondii PCR was lacking, it was decided to give a trial of anti-toxoplasmosis therapy. He was started empirically on pyrimethamine 200mg loading dose, followed by 50 mg oral daily, intravenous clindamycin 600mg 6 hourly, folinic acid 10 mg orally. Sulfadiazine was replaced by clindamycin due to increased risk of drug reactions. Though his fever subsided the day following the antibiotics, there was no clinical improvement. The repeat MRI revealed periventricular enhancement even though the hydrocephalus settled, patient subsequently succumbed to death.
Table 1: HIV related investigations

| Investigations                  | Pre ART screening of Opportunistic infections January 2019 | In ward opportunistic screening September to October 2019 |
|--------------------------------|------------------------------------------------------------|-----------------------------------------------------------|
| CD4                            | 262 cells/μl                                              | 31 cells/μl                                               |
| Viral load                     | 29100 copies/μl                                           |                                                           |
| CMV IgM-                       | Negative                                                  | Equivocal                                                 |
| CMV IgG                        | Negative                                                  | Positive                                                  |
| S.CMV PCR                      | Negative                                                  |                                                           |
| S.cryptococcal - Ag            |                                                           | Negative                                                  |
| HBsAg                          | Negative                                                  | Negative                                                  |
| Hepatitis C antibodies         | Negative                                                  | Negative                                                  |
| Hepatitis A IgM                | Negative                                                  |                                                           |
| Hepatitis A IgG                | Positive                                                  |                                                           |
| EBV IgG                        | Positive                                                  |                                                           |
| Serum Toxoplasma IgM           | Negative                                                  | Positive                                                  |
| Serum Toxoplasma IgG           | Negative                                                  | Positive                                                  |
| Sputum TB gene Xpert culture   | Negative                                                  |                                                           |
| Sputum TB culture              | Negative                                                  |                                                           |
| VDRL/TTPA                      | Negative                                                  | Negative                                                  |

Figure 1. Generalized, symmetrical, pruritic, macular papular rash

Figure 2. MRI brain. Showing sub-acute meningitis complicated with communicating Hydrocephalus and basal meningeal enhancement without any focal lesions

Figure 3. NCCT brain, showing marked hydrocephalus and brain edema

Table 2: CSF investigations

| CSF            | Result                              |
|----------------|-------------------------------------|
| Color          | Colorless                           |
| Appearance     | Slightly cloudy                     |
| Protein        | 57 mg/dl                            |
| Glucose        | 46 mg/dl                            |
| WBC            | 513 mm3                              |
| Neutrophils    | 80%                                 |
| Lymphocytes    | 20%                                 |
| RBC            | 73                                   |
| Organisms      | Gram positive diplococci seen       |
| CSF culture    | No growth                           |
| CSF cryptococcal Ag | Negative                  |
| EBV IgG        | Detected                            |
| VDRL/TTPA      | Negative                            |
| TB Gene Expert | Not detected                        |
TB culture | Negative
---|---
CMV IgM | Equivocal
CMV IgG | Negative
CMV PCR | Not detected
VZV | No detected
Enterovirus | Not detected

**Table 3: Inward investigations**

| Investigation | In ward investigations done during dengue admission (Sept 2019) | In ward investigations done during last medical admission (Oct 2019) |
|---|---|---|
| CXR | Normal | Normal |
| 2D echo | Not done | No infective endocarditis |
| Coagulation profile INR APTT PT | Normal | 0.93 (normal) 22 (normal) 10.6 sec (normal) |
| Blood picture | Features compatible with viral infection | no features of microangiopathic hemolytic anemia, no malignant cells |
| AST | 568 IU/l | 51 → 71U/l |
| ALT | 827 IU/l | 44 → 2,666 U/l |
| Total Bilirubin | Normal | 66.5 |
| Direct Bilirubin | Normal | 63.1 |
| Albumin | Normal | 26 |
| Globulin | Normal | 29 |
| CRP | 10 | 45 mg/ |
| ESR | 97mm | |
| Platelets | 131 X 103 | 155 X 103 |
| Blood Culture | Negative | Negative |
| Urine culture | Negative | Negative |

**Discussion**

Management of HIV infection had drastically improved during the last decade due to advent of ART and standardized and prompt management of opportunistic infections (OIs). But the use of ART and medications for OIs can be limited because of the adverse effects and hypersensitive reactions caused by these medications. This patient was diagnosed to have DRESS possibly due to co-trimoxazol. This was considered since there was the latency period between drug exposure and onset of symptoms in DRESS syndrome when compared to INAH and EFV. DRESS syndrome is described to have a long latency period of 2 to 8 weeks after exposure to a possible drug. DRESS syndrome was first described to be caused by sulphonamide containing drugs but later antiepileptic drugs such as phenytoin, carbamazepine, lamotrigine, phenobarbital and allopurinol were frequently reported causes.(1) This case points to an instance where the clinicians could not continue the patient on ART due to possible DRESS syndrome and development of subsequent opportunistic infections as a result of severe immune deficiency.

Drug reaction with eosinophilia and systemic symptoms is a drug induced, rare, but potentially life-threatening hypersensitivity reaction which manifests with skin rashes, hematological abnormalities like eosinophilia, atypical lymphocytosis, lymphadenopathy and internal organ involvement (Lung, Liver, Kidney). (4)

As the diagnosis of DRESS is based on clinical judgment with high degree of suspicion, the European Registry of Severe Cutaneous Adverse Reactions (RegiSCAR) has developed a scoring system, by which DRESS can be classified as definite, probable or possible disease. (5) According to this classification, our patient who had been exposed to high risk medications, namely, Cotrimoxazole, Isoniazide and Efaviranz two to four weeks prior to the onset of symptoms, had fever > 38.5 °C, eosinophilia of >9% with an absolute eosinophil count of 0.68 x 10³ cells/μl (reference range 0.02-0.5), rash involving > 50% of body surface area, involvement of one organ (liver) and disease duration of more than 15 days had “probable DRESS” with a score of 5.

The patient could not be continued on ART as his liver transaminases started rising drastically, despite being attempted twice with least hepatotoxic regimens of TDF, FTC and RAL combination. One possible reason for elevation of liver enzymes with reintroduction of ART could be due to FTC and RAL which are known to cause drug induced hepatitis as well. Another reason could be drug hypersensitivity reaction due to RAL which is a rare cause for drug hypersensitivity reactions. (6)
The patient developed the next episode of fever with positive dengue NS1 antigen when the patient was off ART and was on a tapering off dose of prednisolone for DRESS. He had the persistent erythematous rash following the diagnosis of DRESS syndrome. Even though DRESS can have a relapsing course, with the positive dengue NS1 antigen and thrombocytopenia, the patient was treated as for dengue fever. Even though Dengue NS1 antigen has a specificity of 89.1% in diagnosing dengue fever, the dengue epidemic in the country and the rising platelet count with settling fever established the diagnosis of dengue fever in the first admission in September. (7) On the other hand missed opportunity of diagnosing possible opportunistic infections with 6 days of hospital admission and attribution of whole picture to dengue fever is a diagnostic dilemma in this case.

Two days following discharge the patient was readmitted with fever, photophobia, headache and confusion. There were no signs to suggest meningism. The initial NCCT being normal and CSF report with slightly high proteins and neutrophil pleocytosis with gram positive diplococci made the diagnosis of bacterial meningitis and commenced on meningitic doses of antibiotics, Intra venous meropenum intravenous vancomycin and intravenous acyclovir was empirically started. However, patient’s condition progressively deteriorated despite treatment. Communicating hydrocephalus and basal meningeal enhancement without any focal lesions were clearly evident from the subsequent MRI scan, which suggests poor prognosis. Hence it can be inferred that the initial disease diagnosis was suboptimal.

In an HIV infected patient presenting with CNS symptoms, the most important factor in differentiating the diagnosis is the degree of immune suppression of the patient. (8) Having CD4 cell count of 31 cells/μl signify a marked immunosuppression and most likely diagnoses are AIDS associated tumors and opportunistic infections. The leading considerations are Toxoplasma encephalitis, primary CNS lymphoma, progressive multifocal leukoencephalopathy, HIV encephalopathy, CMV encephalitis and cryptococcal meningitis. CNS tuberculosis infection also must be considered in our setting where TB is common.

Ventricular enlargement, meningeal enhancement in MRI scan with slightly elevated CSF proteins with positive CMV IgG suggested CMV encephalitis although it was very rare and commenced on IV gancyclovir on day 7 of admission. Even though, CSF CMV PCR report is negative possibility of CMV encephalitis could not be excluded.

Toxoplasmosis being the commonest OI in the central nervous system (CNS), was suspected in this patient as the patient was not on co-trimoxazole prophylaxis and it was suspected to be the culprit drug for his DRESS syndrome. Absence of the typical clinical and radiological findings and the initial negative toxoplasma IgG and very high liver transaminase and bilirubin levels with impending liver failure and DRESS syndrome made us reluctant to start toxoplasma treatment empirically.

The patients with toxoplasma encephalitis are uniformly seropositive for anti-toxoplasma IgG and the absence of it makes the diagnosis unlikely. Further, it usually presents with focal neurological signs due to multiple ring enhancing lesions giving rise to mass effect localized in the parietal and frontal lobes, in the thalamus or basal ganglia or in the corticomedullary junction. (9) Multiple ring enhancing lesions with mass effect are visible after injection of gadolinium T1-weighted MRI in about 90% of cases, rarely toxoplasma can present without focal neurological signs with diffuse encephalopathy. (10) However, when there is diffuse encephalitis, the progression is rapid, and the outcome is fatal and is a rare entity and there is limited literature available with very few case reports. (11)

Common causes for hydrocephalus in HIV/AIDS patients are primary normal pressure hydrocephalus, subarachnoid hemorrhage and meningitis. However, hydrocephalus due to toxoplasmosis, tuberculous meningitis and cryptococcal meningitis is very rare with few, reported cases in literature. (12) The mechanism of development of hydrocephalus in toxoplasma is defined as either compression of cerebro spinal fluid (CSF) pathway by
surrounding parenchymatous space occupying lesion or blockage of CSF flow within the ventricular system and ependymal canal by an exudate from necrotizing ependymitis. Development of hydrocephalus in a patient with diffuse toxoplasma encephalitis was not been reported so far.

Even though toxoplasma encephalitis usually occurs as a reactivation of latent infection, this patient’s toxoplasma IgG at the time of HIV diagnosis was negative. However, both toxoplasma IgM and IgG became positive in blood and CSF later on around day 10 of last admission. CSF toxoplasma titres were not done due to unavailability. Although CSF T.gondii PCR was lacking, it was decided to give a trial of anti-toxoplasmosis therapy. Therefore, the patient was started on the alternative regimen suspecting diffuse toxoplasma encephalitis with oral pyrimethamine, oral clindamycin with leucovorin because the definitive treatment cannot be administered due to impending DRESS syndrome.

Even though, the fever settled with the treatment, the clinical improvement was minimal due to ongoing brain damage as evident on repeat MRI, with periventricular enhancement and hydrocephalus.

Patient might have developed systemic inflammatory response syndrome (SIRS) at the end stage as it fulfilled more than two criteria such as fever for more than 38°C, rapid heart rate of 120/min with blood pressure of 116/86 mmHg. He had deep irregular breathing which was rapid and abnormal hematological indices. But his septic screening remained negative. Although his serum sodium level decreased 120 mmol/L which was corrected by hypertonic saline. His serum potassium level remained normal and there were no features of syndrome of inappropriate ADH secretion.

Cryptococcal meningoencephalitis can be excluded by negative blood and CSF findings. Even though TB investigations were normal, TB meningitis cannot be totally excluded due to several factors. Severely immunocompromised state, on long term steroids for DRESS and poor sensitivity of CSF Gene xpert and TB culture (sensitivity of 10 to 60%).(13) Any other fungal meningencephalitis can not be totally excluded due to lack of testing facilities.

Although atypical presentation of toxoplasma encephalitis is another high probability. The reason for atypical presentation could be the long term steroid therapy and administration of high dose steroids at the time of diagnosing dengue and bacterial meningitis. But high index of suspicion should always be in mind when treating severely immune compromised patients with fever and CNS symptoms as toxoplasmosis is one of the commonest CNS opportunistic infections. However, the possibility of drug reactions and possible liver failure made it reluctant of starting toxoplasma treatment empirically with misleading dengue fever and delaying of definitive diagnosis, finally lead to the unavoidable brain damage and death of this patient.

References
1. Roujeau JD.Drug reaction with eosinophilia and systemic symptoms (DRESS), UpToDate 2019.
2. National STD/AIDS Control programe. Elimination of Mother to Child transmission of HIV and syphilis, National Validation report, Sri Lanka. 2. National STD/AIDS Control programe, Ministry of Health, Colombo: s.n., August, 2019.
3. Wendel KA, McArthur JC. Acute meningoencephalitis in chronic human immunodeficiency virus (HIV) infection: putative central nervous system escape of HIV replication. Clinical infectious diseases. 2003 Oct 15;37(8):1107-11.
4. Mockenhaupt M. Drug reaction with eosinophilia and systemic symptoms (DRESS). UpToDate. 2019.
5. Kardaun SH, Sidoroff A, Valerye-Allanore L et al. Variability in the clinical pattern of cutaneous side-effects of drugs with systemic symptoms: Does a DRESS syndrome really exist? Br J Derm. 2007;157(609).
6. CDC guideline for use of Antiretroviral agents in adults and adolescents living with HIV 2018.
7. Vaishali N Solanke, Mohan G Karmarkar PRM. Early dengue diagnosis: Role of rapid NS1 antigen, NS1 early ELISA, and PCR assay. Trop J Med Res. 2015;18(2):95–9.
8. Koralnik IJ. Approach to HIV-infected patients with central nervous system lesions. UpToDate.
9. Miller RF, Hall-Craggs MA, Costa DC, Brink NS, Scaravelli F, Lucas SB, Wilkinson ID, Ell PJ, Kendall BE HM. Magnetic resonance imaging, thallium-201 SPET scanning, and laboratory analyses for discrimination of cerebral lymphoma and toxoplasmosis in AIDS. Sex Transm Infect. 1998;74(4):258.
10. Image M ring enhancing lesions with mass effect are visible after injection of gadolinium in this T1-weighted image. Clin Infect Dis. 1992;15(2):211.

11. Frangoise Gray, Romain Gherardi, Elizabeth Wingate, Jeffrey Wingate, Gilles F6nelon, Andr Gaston AS, Poirier and J. Diffuse “encephalitic” cerebral toxoplasmosis in AIDS. J Neurol. 1989;236:273–7.

12. Djientcheu VDP, Njamnshi AK, Ongolo-Zogo P, Dongmo L, Eloundou JN, Rilliet B, et al. Hydrocephalus: A rare presentation of central nervous system toxoplasmosis in the acquired immunodeficiency syndrome. African J Neurol Sci. 2004;23(2).

13. British HIV Association guidelines for the treatment of TB/HIV Co-infection. HIV Medicine. British HIV Association guidelines. United Kingdom. 2011;12:517-537