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Epidemiological, clinical and biological profile of neuromeningeal cryptococcosis among people living with HIV in Kinshasa, Democratic Republic of Congo

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Abstract

Neuromeningeal cryptococcosis (NMC) is one of the most frequent opportunistic infections (OI) in Human Immunodeficiency Virus (HIV) infection. In Kinshasa, the latest data on cryptococcosis were published in 1996. The objective was to describe the epidemiological, clinical and biological profiles of NMC in HIV-infected people living in Kinshasa. This is a descriptive study based on the medical records of patients who attended three clinics in Kinshasa between January 1st 2011 and December 31st 2014. Only the medical records of HIV-infected people presenting the NMC were reviewed. During the 4 year-period of the study, 261 HIV-positive patients presented to the clinics for neuromeningeal syndrome, including 23 with NMC. The global prevalence of NMC was 8.8% for the three clinics. The mean age was 42.8 ± 9.5 years, with male predominance (65.2%). The main symptoms were headache (73.9%), neck stiffness (60.9%), fever (47.8%), and coma (47.8%). Biological records were as follows: median CD4 cell count was 79 cells/mm³; cerebrospinal fluid (CSF) was clear for 56.5% of the cases with predominance of neutrophils in 73.9%. The outcome was fatal in 34.8% of cases. The prevalence and therapeutic outcome of NMC show that it constitutes a non-negligible OI in Kinshasa, especially in HIV-infected people at the AIDS stage. As HIV-infected people with severe immunosuppression are the most affected by NMC, active preventive measures should benefit this vulnerable category of people.

Introduction

Neuromeningeal cryptococcosis (NMC) is one of the most frequent opportunistic infections (OI) in term of morbidity and mortality in advanced stages of HIV infection [1]. The causal agent of NMC is a fungus belonging to the species complex *Cryptococcus neoformans/C. gattii*. Rarely met until the 1980s, this infection has since seen its incidence increase considerably with the emergence of the AIDS pandemic [2]. Despite the availability of antiretroviral (ARV) drugs against HIV infection, the

mortality rate of NMC within 10 weeks after the start of antifungal therapy varies from 20% to 50% in sub-Saharan Africa [3,4]. A survey conducted in 17 European countries by the European Confederation of Medical Mycology (ECMM) found that cryptococcosis was associated with HIV infection in 77% (435/565) of cases [5]. Also, among People Living With HIV (PLWHIV) in Ivory Coast, Kassi Kondo [6] estimated prevalence of NMC at 3.6%; in Togo, this prevalence was estimated at 2.9% [7]. In the Democratic Republic of Congo (DRC), the Demographic and Health Survey 2013-2014 (EDS DRC II) estimated that 1.2% of the general population aged 15-49 are infected with HIV and, according to projections based on Spectrum 5.41, the number of PLWHIV in the DRC was estimated to be 381,187 by the end of 2016, of which only 121,762 would be on ARV Treatment (ART) and 20,854 would die [8]. Among these PLWHIV, more than half do not have access to ART and are thus exposed to all the complications related to the natural history of the disease, including OI [9]. In addition to HIV infection, other immunosuppressive situations have been associated with an increased risk of contracting cryptococcosis [10]. In the DRC, the hospital frequency of cryptococcosis ranged from 19% in 1994 to 11% in 2013 [11,12]. In Kinshasa, the latest available data on cryptococcosis dates back to 1996, yet the extent of HIV infection in this population is constantly increasing [13]. The lack of an up-to-date epidemiological and clinical data on this fungal disease among HIV patients in the DRC in general and specifically in Kinshasa motivated the initiation of the present work. The objective of this study was to describe the epidemiological, clinical characteristics and the profile of biological analyses during NMC among PLWHIV who attended clinics in Kinshasa.

Methods

Study frame: this is a descriptive study based on the medical records of HIV-positive patients who attended three hospitals located in different districts of Kinshasa, between January 1st 2011 and December 31st 2014. Among these hospitals, the

University Clinic of Kinshasa (CUK) was the only tertiary level hospital with an active line of 124 HIV-positive patients. The 2 other hospitals were the Kinshasa Reference General Provincial Hospital (HPGRK) and the Monkole Hospital Center (CH Monkole) with a significant active patient population of 1717 and 1138 PLWHIV, respectively.

Inclusion criteria: the present study took into account the HIV-positive patients in an exhaustive way, whether they were followed regularly or not. Were included in the study, patients who consulted with a clinical presentation of meningoencephalitis, particularly those with NMC confirmed by the direct test using India ink and having been hospitalised in the selected hospitals during the period of study.

Data collection: the data were collected using a survey form. The following variables were considered in this study: socio-demographic (sex, age and marital status), clinical (motive for consultation, medical history, physical examination data, duration and outcome of hospitalisation) and biological (CD4 cell count as well as direct India ink staining, biochemistry, cytology and Sabouraud culture of cerebro-spinal fluid) variables.

Statistical analyses: the data were entered on an excel spreadsheet and then exported to SPSS 26.0 (statistical package for the social sciences 26.0) for descriptive analysis. Quantitative variables were presented as proportions (%), means and standard deviations. Qualitative variables were given in the form of proportions (%).

Ethical considerations: this work has been performed in strict compliance with ethical rules by guaranteeing the anonymity and the safety of patients. The data collected were kept and manipulated by the research team alone.

Results

Epidemiological characteristics

Frequency of neuromeningeal cryptococcosis: out of 261 PLWHIV included in the study, 23 presented

with NMC, an overall prevalence of 8.8%. The relative prevalence was higher at HPGRK with 15 cases out of 98 patients included (15.3%) followed by CUK (6 out of 65 patients included, 9.2%) and CH Monkole (2 out of 98 patients included, 2%).

Sociodemographic data: in this study, men were more affected than women (65.2%) with a sex ratio M/F of 1.9. The mean age of patients was 42.83 ± 9.5 years with an age range from 31 to 62 years. The 35 to 44 age group was the most affected, at 34.8%. The highest frequency of the disease was observed among married, 39.1% (Table 1).

Clinical data: headache (73.9%), fever (47.8%) and coma (39.1%) were the most common motives for consultation. Alcohol uptake (26.1%) and hypertension (13%) were the most commonly reported antecedents in these patients. The mean duration of hospitalisation was 15.3 ± 10.4 days with extremes of 2 and 36 days; the majority of patients had stayed either 1 to 7 days or 15 to 21 days in the hospital (30.4% each). After treatment, survival rate was 60.9%, 3.4% were lost to follow-up, and the outcome was fatal in 34.8% (Table 2). After physical examination, 52.2% of patients were lucid, 60.9% had stiff neck. Facial paralysis, monoparesis of a member, hemibody hypotonia and presence of cutaneous-plantar reflexes were all found in 8.7% of cases (Table 3). Antifungal therapy consisted mainly of fluconazole alone in 82.6% (19/23) of the cases, at a dosage of 400-800 mg/day during the 14 days of the attack phase and then 200 mg/day during the maintenance phase to immune restoration (CD4 cell count > 100 cells/mm³) under ART.

Biological data: the median CD4 cell count in the peripheral blood was 79 (66-105) cells/mm³ with a range of 33 to 218 cells/mm³. The majority of patients had a CD4 cell count between 50 and 199 cells/mm³ (81.8%). In most cases, CSF was clear (56.6%) with predominantly neutrophilic cytorachia (73.9%) and proteinorachia was greater than 45 mg/dl in 6 out of 7 patients from whom proteinorachia data were obtained (Table 4).

Discussion

The objective of this study was to describe the epidemiological, clinical and biological characteristics of NMC in PLWHIV who attended clinics in Kinshasa. The prevalence of NMC was 8.8% among PLWHIV followed during the four-year period of study. While cryptococcosis is in the process of disappearing completely in developed countries, it still constitutes a burden during HIV infection in African countries [14]. The prevalence from the present study is higher than the 4.6% reported by Mwamba *et al.* in 2013 in Lubumbashi in a prospective study looking for cryptococcal antigen in the blood and the CSF of PLWHIV at the AIDS stage [15]. It is also higher than that reported by Kouakou *et al.* in Ivory Coast in 2016 (2.49%) [16] and that observed in Morocco (1.53%) in a retrospective study of PLWHIV covering a period of 7 years [17]. However, it is lower than that reported in a study conducted in 1996 at the CUK and HPGRK, which are 2 of the 3 sites of the present study, by Situakibanza *et al.* [18] who found cryptococcosis to be responsible for 12.5% of fever cases in PLWHIV. This situation could be explained by the prevalence of HIV infection in the population, estimated at 1.6% in Kinshasa; the inaccessibility of ART for more than half of PLWHIV, as described in the EDS-RDC II 2013 -2014 [8]; and the delay in the diagnosis of HIV infection and its related OI. In addition, at that time, the DRC had not yet adopted the test and treat strategy; ARV treatment was conditioned by certain eligibility criteria that were difficult to meet at the time of diagnosis. It was not until 2015 that the DRC adopted the test and treat strategy [19].

The average HIV-positive young adult male, aged 42.8 ± 9.5 years and married, was the most commonly affected by NMC. These results are linked to the vulnerability to HIV for the young population, who are the most sexually active, a major risk factor for NMC [20]. Our results are consistent with those reported in the literature [20]. Our observation revealed a male predominance (65.2%) with a sex ratio M/F of 1.9.

This male predominance is thought to be due to genetic susceptibility linked to a weak phagocytic power and the weak resistance of the male macrophages to *C. neoformans* compared with those of women [21]. In addition, oestrogen may have a protective effect against cryptococcosis [22]. Clinically, headache, fever and loss of consciousness were the most common complaints. However, 52.2% of our patients were afebrile, and had a mild meningeal syndrome limited to neck stiffness in 60.9% of cases. Although the NMC clinical signs are not very specific, fever and headache are often present [20]. In the current study and elsewhere, the predominance of headache and fever has been reported during NMC [15], as well as seizures and motor deficits [23]. The involvement of the cranial nerves, in particular, the facial and oculomotor nerves has also been found [24]. These motor deficits are suggestive of cerebral cryptococcoma (focal lesions) that should be investigated by a brain scan.

From a biological point of view, the CSF was predominantly clear (56.7%) with hyperproteinorachia in 26.1% of cases and, paradoxically, a predominance of neutrophils polymorphonuclear (73.9%). This neutrophilic predominance differs from the literature, which reports the predominance of lymphocytes or a mixed population [25]. In some studies, concomitant Gram and India ink staining of CSF revealed co-infection with *Streptococcus pneumoniae* [26]. However, in our study no bacteriological data could be obtained due to lack of availability of the tests. Apart from cytorachia and proteinorachia, our biological results are corroborated by those reported by Chadli in Morocco [17]. CSF culture, a reference test in the diagnosis of NMC due to its high sensitivity and specificity close to 100%, was not performed on the CSF samples taken from these patients, nor was antigen testing undertaken due to a lack of technical means [1,25]. NMC occurred at a very advanced immunodepression stage with a CD4 cell count <200 cells/mm³ in 81.9% and for the rest of the cases the CD4 cell count was measured between 200 and 349/mm³. These results are in

line with data from the literature reporting that PLWHIV south of the Sahara are mostly detected at a time when the CD4 cell count is <200 cells/mm³ [23]. Some authors have also noted the unusual occurrence of NMC at the third stage of immunodepression (CD4 cell count between 200 and 349 elements/mm³) of HIV disease. Since *Cryptococcus gattii* is present in DRC and is thought to preferentially affect subjects with a good immune status, these cases observed in moderately immunodepressed patients would partially support the presence of this species in the African environment [10]. The viral load, one of the best parameters of the follow-up for the PLWHIV, was not realised in the present study patients for the same reasons evoked above.

Most of the patients received fluconazole alone, 400 to 800 mg/day during the 14 days of the attack phase and then 200 mg/day during the maintenance phase until immune restoration (CD4 cell count > 100 cell/mm³) was reached under ART as the main treatment. Other antifungal treatments such as amphotericin B and 5-flucytosine [1] were not widely available in the three clinics. NMC is associated with high and early mortality. Here, 34.8% of our patients died within a mean hospital stay of 15.3 ± 10.4 days. The majority of deaths were observed between 1-7 days. This high mortality can be correlated to the limits of the technical platform for the management of the NMC, in terms of diagnosis, treatment and consultation time. These results are corroborated by other authors [27]. It is important to note that the medium-term prognosis for NMC in HIV depends on the restoration of cell-mediated immunity through ART, the availability of effective antifungals and the permanence of a well-trained caregiver.

Conclusion

Among the OI occurring during HIV infection, NMC remains a concern due to its prevalence and outcome in Kinshasa. It mainly affects the young adult male with advanced immunodepression. As the clinical signs are not very specific and

conventional biology sometimes does not contribute much, the simultaneous realisation of cytorachia and bacteriology of the CSF should be of great value in diagnostic orientation. In addition, the systematisation of an active search for cryptococcal antigens in any HIV-positive patient with very advanced immunodepression, without ignoring the importance of adherence to ART in the primary prevention of cryptococcosis, are ways to be encouraged when caring for these vulnerable people.

What is known about this topic

- *In recent years, the epidemiology of NMC among PLWHIV has been greatly allayed by the availability of triple antiretroviral therapy in some countries;*
- *The NMC develops most often in the field of dysimmunity, mostly related to HIV and this, at the AIDS stage;*
- *Cytorachia during NMC is often lymphocytic predominance or mixed, with high proteinorachia and hypoglycorachia.*

What this study adds

- *In the major clinics in Kinshasa (DRC), the prevalence of neuromeningeal cryptococcosis among people living with HIV is 8.8%. For a treatment based mainly on fluconazole alone, the therapeutic outcome is fatal in 34% of cases;*
- *The neuromeningeal cryptococcosis clinic is more marked by headaches (73.9%), which in the majority of cases have led to a diagnosis. In addition, neck stiffness, fever and coma completed the diagnosis;*
- *Neuromeningeal cryptococcosis was also found in people living with HIV with a CD4 count > 200 cells/mm³ (8.7% of cases). In addition, a predominantly neutrophil-dominant leukocyte pattern in the CSF was found in most patients (73.9%), suggesting possible bacterial co-infection.*

Competing interests

The authors declare no competing interests.

Authors' contributions

Bive Zono: conception and design, analysis and interpretation, drafting the article. Erick Kamangu: critical revision of the article, final approval of the article. Hippolyte Situakibanza: conception and design, critical revision of the article, final approval of the article. Evelyne Amaela: collection data, final approval of the article. Ben Bepouka: collection data, final approval of the article. Marcel Mboula: analysis and interpretation, final approval of the article. Jean-Marie Kayembe: critical revision of the article, final approval of the article. Georges Mvumbi: critical revision of the article, final approval of the article. Marie-Pierre Hayette: critical revision of the article, final approval of the article. All the authors have read and agreed to the final manuscript.

Tables

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Table 3: clinical data (2)

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References

1. World Health Organization. Guidelines for the diagnosis, prevention and management of cryptococcal disease in HIV-infected adults, adolescents and children, March 2018. Genève **Google Scholar**
2. Park BJ, Wannemuehler KA, Marston BJ, Govender N, Pappas PG, Chiller TM. Estimation of the current global burden of cryptococcal meningitis among persons living with HIV/AIDS. *Aids*. 2009;23(4): 525-530. **PubMed** | **Google Scholar**
3. Bicanic T, Muzoora C, Brouwer AE, Meintjes G, Longley N, Taseera K *et al*. Independent Association between Rate of Clearance of Infection and Clinical Outcome of HIV-Associated Cryptococcal Meningitis: Analysis of a Combined Cohort of 262 Patients. *Clin Infect Dis*. 2009;49(5): 702-709. **PubMed** | **Google Scholar**
4. Lessells RJ, ChB M, Portia Mutevedzi DC, Newell ML. Poor long-term outcomes for cryptococcal meningitis in rural South Africa. *SAMJ S Afr Med J*. 2011;101(4): 251-252. **PubMed** | **Google Scholar**
5. Viviani MA, Cogliati M, Esposto MC, Lemmer K, Tintelnot K, Valiente MFC *et al*. Molecular analysis of 311 *Cryptococcus neoformans* isolates from a 30-month ECMM survey of cryptococcosis in Europe. *FEMS Yeast Res*. 2006;6(4): 614-619. **PubMed** | **Google Scholar**
6. Kassi K. Diversité génétique et sensibilité aux antifongiques d'isolats cliniques et environnementaux de *Cryptococcus* à Abidjan , Côte d'Ivoire . To cite this version: HAL Id: tel-01628628 Spécialité: Sciences Chimiques et Biologiques Présentée par KASSI KO. thèse, Université Montpellier. 2017. **Google Scholar**
7. Kossivi A, Doreen N, Damelan K, Vinyo K, Mensah GK, Komi A *et al*. Infections Opportunistes Du Vih/Sida En Milieu Hospitalier Neurologique Au Togo. *African J Neurol Sci*. 2015;34(2): 352-354. **Google Scholar**
8. Programme National de la Lutte contre le SIDA et les IST. PNLs Rapport Annuel 2015. Kinshasa.
9. Programme National de la Lutte contre le SIDA et les IST. Guide de prise en charge du VIH en République Démocratique du Congo. 2016. Kinshasa.
10. Raiss C. Quand le cryptocoque colonise l'étude digestif: Encore 3 cas. *Animal Genetics*. 2014;39(5).
11. Onusida A. Maladies opportunistes liées au VIH En bref. Collection Meilleurs pratiques de l'Onusida. 1999;12.

12. Kivukuto Mutendela J, Mashupe S, Bihehe Masemo D, Mitima Kashosi T. HIV screening among patients followed for cryptococcal meningitis in developing countries: Data from Bukavu in the Democratic Republic of Congo. *African J Microbiol Res.* 2014;8(7): 721-723. **Google Scholar**
13. UNAIDS and AIDS info. Country factsheets. Democratic Republic of Congo; 2018. Accessed October 2, 2019.
14. Gangneux JP, Bougnoux ME, Hennequin C, Godet C, Chandenier J, Denning DW *et al.* Estimation du poids épidémiologique des infections fongiques graves en France. *J Mycol Med.* 2016;26(4): 385-390. **Google Scholar**
15. Mwamba C, Kapend L, Moutschen M, Kasamba E, Vaira D, Clumeck N. Programme National Multisectoriel De Lutte Contre Le Sida (PNMLS). 2^{ème} Conférence Nationale Sur Le VIH/SIDA en République Démocratique Du Congo. Lubumbashi; 24-26 Juin 2013.
16. Kouakou GA, Ello NF, Kassi NA, Keita M, Doumbia A, Mossou C *et al.* Fluconazole 1200 mg ou 800 mg dans le traitement de la cryptococcose neuroméningée en Côte d'Ivoire. *J Mycol Med.* 2017;27(1): 72-78. **PubMed | Google Scholar**
17. Chadli S, Aghrouch M, Taqarort N, Malmoussi M, Ouagari Z, Moustoufi F *et al.* Neuromeningeal cryptococcosis in patients infected with HIV at Agadir regional hospital, (Souss-Massa, Morocco). *J Mycol Med.* 2018;28(1): 161-166. **PubMed | Google Scholar**
18. Situakibanza H, Kapita B, Mbendi M, Odio W, Mbula M, Muvova D *et al.* Etiologie de la fièvre au cours du SIDA: à propos de 64 cas. *Panorama Médicale.* 1996;1(5): 842-845.
19. World Health Organization. Guideline on when to Start Antiretroviral Therapy and on Pre-Exposure Prophylaxis for HIV. September 2015;1-76. **Google Scholar**
20. Lau C, Muula AS. HIV/AIDS in Sub-Saharan Africa. *Croat Med J.* 2004;45(4): 402-414. **PubMed | Google Scholar**
21. McClelland EE, Hobbs LM, Rivera J, Casadevall A, Potts WK, Smith JM *et al.* The Role of Host Gender in the Pathogenesis of *Cryptococcus neoformans* Infections. *PLoS One.* 2013;8(5): 1-7. **PubMed | Google Scholar**
22. Levitz SM, Farrell TP, Maziarz RT. Killing of *Cryptococcus neoformans* by human peripheral blood mononuclear cells stimulated in culture. *J Infect Dis.* 1991;163(5): 1108-1113. **PubMed | Google Scholar**
23. Bissagnene E, Ouhon J, Kra O, Kadio A. Aspects actuels de la cryptococcose neuroméningée à Abidjan. *Med Mal Infect.* 1994;24(Suppl 2): 580-585. **Google Scholar**
24. Millogo A, Ki-Zerbo GA, Andonaba JB, Lankoandé D, Sawadogo A, Yaméogo I *et al.* La cryptococcose neuroméningée au cours de l'infection par le VIH au Centre hospitalier de Bobo-Dioulasso (Burkina Faso). *Bull la Soc Pathol Exot.* 2004;97(2): 119-121. **Google Scholar**
25. Cryptococcose: Association Française des Enseignants de Parasitologie et Mycologie ANOFEL Sommaire. Université Médicale Virtuelle Francophone. 2014. **Google Scholar**
26. Saleem F, Fasih N, Zafar A. *Cryptococcus neoformans* and *Streptococcus pneumoniae* co-infection in posttraumatic meningitis in a patient with unknown HIV status. *J Pak Med Assoc.* 2015;65(10): 1122-1124. **PubMed | Google Scholar**
27. Bergemann A, Karstaedt AS. The spectrum of meningitis in a population with high. *An International Journal of Medicine.* 1996;89(7): 499-504. **Google Scholar**

Table 1: socio-demographic variables of patients

Variable	Female n (%)	Male n (%)	Total n (%)
Sex			
	8 (34.8)	15 (65.2)	23 (100)
Age group (year)			
25 – 34	2 (8.7)	3 (13.0)	5 (21.7)
35 – 44	3 (13.0)	5 (21.7)	8 (34.8)
45 – 54	1 (4.3)	5 (21.7)	6 (26.1)
55 – 71	2 (8.7)	2 (8.7)	4 (17.4)
Civil status			
Married	2 (8.7)	9 (39.1)	11 (39.1)
Single	1 (4.3)	5 (21.7)	6 (26.1)
Divorced	2 (8.7)	1 (4.3)	3 (13.0)
Widower	3 (13.0)	0	3 (13.0)

Table 2: clinical data (1)

Variable	Female n (%)	Male n (%)	Total n (%)
Motif for consultation			
Headache	6 (26.1)	11 (47.8)	17 (73.9)
Fever	3 (13.0)	8 (34.8)	11 (47.8)
Coma	3 (13.0)	6 (26.1)	9 (39.1)
Other	5 (21.7)	3 (13.0)	8 (34.8)
Vomiting	0	4 (17.4)	4 (17.4)
Convulsions	1 (4.3)	2 (8.7)	3 (13.0)
Antecedents			
Alcohol	1 (4.3)	5 (21.7)	6 (26.1)
High blood pressure	2 (8.7)	1 (4.3)	3 (13.0)
Tobacco	0	2 (8.7)	2 (8.7)
None	5 (21.7)	7 (30.4)	12 (52.2)
Duration of hospitalisation (days)			
1 - 7	4 (17.4)	3 (13.0)	7 (30.4)
8 - 14	2 (8.7)	3 (13.0)	5 (21.7)
15 - 21	0	7 (30.4)	7(30.4)
>21	2 (8.7)	2 (8.7)	4 (17.4)
Outcome			
Alive	4 (17.4)	10 (43.5)	14 (60.9)
Dead	3 (13.0)	5 (21.7)	8 (34.8)
Sight loss	1 (4.3)	0	1 (4.3)

Table 3: clinical data (2)

Physical examination	Female n (%)	Male n (%)	Total n (%)
State of consciousness			
Lucid	4 (17.4)	8 (34.8)	12 (52.2)
Coma	4 (17.4)	7 (30.4)	11 (47.8)
Meningeal sign			
Neck stiffness	5 (21.7)	9 (39.1)	14 (60.9)
Kernig	3 (13.0)	6 (26.1)	9 (39.1)
Brudzinski	2 (8.7)	5 (21.7)	7 (30.4)
Facial paralysis	1 (4.3)	1 (4.3)	2 (8.7)
Motor deficiency			
Monoparesis	0	2 (8.7)	2 (8.7)
Hemiplegia	0	1 (4.3)	1 (4.3)
Hypotonia of the hemicorps	0	2 (8.7)	2 (8.7)
Skin and plantar reflex	0	2 (8.7)	2 (8.7)

Table 4: biological data of patients

Variable	Female n (%)	Male n (%)	Total n (%)
CD4 cell count (cells/mm³)			
<50	0	2 (8.7)	2 (8.7)
50 – 199	7 (30.4)	12 (52.2)	19 (82.6)
200 – 349	1 (4.3)	1 (4.3)	2 (8.7)
CSF appearance			
Clear	6 (26.1)	7 (30.4)	13 (56.6)
Cloudy	1 (4.3)	4 (17.4)	5 (21.7)
Not reported	1 (4.3)	4 (17.4)	5 (21.7)
Cytorachia (elements/mm³)			
Neutrophil predominance	5 (21.7)	12 (52.2)	17 (73.9)
Lymphocyte predominance	2 (8.7)	1 (4.3)	3 (13.0)
Mixed	2 (8.7)	1 (4.3)	3 (12.5)
Proteinorachia (mg/dl)			
>45	2 (8.7)	4 (17.4)	6 (26.1)
15 – 45	1 (4.3)	0	1 (4.3)
<15	0	0	0
Not reported	5 (21.7)	11 (47.8)	16 (69.6)