RESEARCH NOTE

The status and health burden of neurocysticercosis in Mbulu district, northern Tanzania

Beda John Mwang'onde1*, Mwita John Chacha2,3 and Gamba Nkwengulila2

Abstract

Objective: The objective of this study was to assess the extent and health burden of neurocysticercosis in the general community of the Mbulu district, northern Tanzania. About 1051 randomly selected participants were screened for human cysticercosis. The Cysticercus Western Blot IgG and Computed Tomography scan were used to detect infection by cysticerci. The DALYs was used to assess the community’s health burden vis-a-vis neurocysticercosis.

Results: The sero-prevalence of HCC was 16.27%. About 76% of 25 selected human cysticercosis sero-positives had neurocysticercosis suggestive lesions on CT scan and 74% had a history of epilepsy. Epilepsy caused 2.8 years of life lost and 2.2 healthy years of life lost due to disability per 1000 person-years in Mbulu. The average DALYs imposed due to neurocysticercosis and epilepsy were 3.0 and 3.9 per 1000 person-years, respectively. Neurocysticercosis is a serious public health concern in northern Tanzania.

Keywords: Human cysticercosis, Neurocysticercosis, Status, Mbulu district

Introduction

Neurocysticercosis (NCC) refers to infection of the brain by the larval form (cysticerci) of the pork tapeworm, Taenia solium. It is a form of human cysticercosis (HCC) and is the most important parasitic infection of the central nervous system (CNS) in humans [1]. NCC is unique in that the living parasite is well tolerated in the human brain [2]. Symptoms and clinical signs primarily result from death of the organism and accompanying inflammatory reactions in the human CNS [2]. The cysticerci tend to inhibit humoral and cellular immune responses as well as inflammatory reactions and cytokines, particularly IFN-γ and IL-2, and to a lesser degree IL-4 production [3, 4]. Live cysticerci also secrete cysteine and serine proteases that deplete CD4+ cells (T helper cells) that send signals to other types of immune cells, including CD8 killer cells which destroy and kill the infection [3].

The exposition of these molecules provides insights into the mechanisms by which T. solium cysticerci evade host immunological attack and are able to survive long periods of time [3]. Nevertheless, even if one develops neurological symptoms in early stages, it is difficult to link it with NCC. Another downside to this is that epilepsy in developing countries like Tanzania is highly associated with witchcraft, familial traits and/or a punishment by the spirits for wrong doing in the community. Ignorance on the epidemiology of NCC exacerbates public health and socioeconomic burden of the affected communities. This study highlights the status and health burden of NCC in Mbulu district for decision science.

Main text

Materials and methods

Study area

The study was carried out in Mbulu District (3.80°–4.50°S, 35.00°–36.00°E) as part of the bigger study on the status and epidemiology of T. solium cysticercosis in southern and northern highlands of Tanzania between March 2013 and January 2014. The district has a population of 320,279 and an average household size of 6.0
The district lies at an estimated altitude of 1000–2400 m above sea level and its climate ranges from semi-arid to sub-humid with an annual rainfall of < 400 and > 1200 mm [6]. Relative humidity ranges from 55 to 75% and mean annual temperatures range from 15 to 24 °C (Ibid.).

**Study design**
This was a cross-sectional study. Blood samples were collected from randomly selected and consenting individuals of different age groups and both sexes in 25 purposefully selected villages. The village executive officers in consultation with the principal investigator conducted meetings for sensitization in every village/hamlet selected for sampling. The principal investigator explained the purpose of the study. Any member of the village/hamlet aged between 0 and 80+ [7, 8] was eligible for participation. To select the study participants, each of the willing village members picked a folded piece of paper on which the words “Yes” or “No” were written. Those who picked “Yes” signed an informed consent form to participate in blood sampling. Children were sampled upon consent from their parents. Strict phlebotomy procedures were followed.

**Sample collection and laboratory analysis**
About 3 mls of cephalic venous blood was collected from each of the consenting participants using sterile plain vacutainer tubes. The samples were then centrifuged at 4800 rpm for serum extraction and the sera were aliquoted and stored in cryogenic vials at −20 °C before use. The presence of cysticercus antibodies from sera was determined using the Cysticercus Western Blot (CYSWB) IgG (LDBIO Diagnostics 69009 Lyon—France) within 21 days of sample collection [9]. Twenty five participants were randomly selected from 48 positive participants with 5–6 well defined molecular bands on CYSWB IgG test for contrasted cerebral computed tomography (cCT scan) for NCC.

**Computed tomography (CT) scan**
The presence of cysticerci in the CNS (brain) from the cysticercosis positive participants was confirmed in a hospital by the use of single slice scanner at one tube rotation cCT scan (TOSHIBA CT Scan x-ray machine; Generator Model: TSX-003, T/No. CXXG-006, S/No. A8572114; Tube S/No. A8572162; Toshiba Medical Systems Corporation, 1998). CT scan detects ring-enhancing lesions (stage II) and multiple ring enhancing lesions (stage III) in NCC infected patients [10]. At the hospital each of the participants underwent strict medical procedures before undergoing a cCT Scan. Positive cases were referred to a neurosurgeon who prescribed medication for management of the condition.

**Estimation of health burden**
The health burden of neurocysticercosis (NCC) was estimated using the serological screening and CT scan results and the Disability Adjusted Life Years (DALYs) [7, 11, 12] in respect of the Mbulu district population [5]. The incidence was obtained from positive cases per each age group over total populations of the specific age group multiplied by 1000.

Data entry and validation was carried out using MS-Excel 2010 version (Ms Corp., Redmond, WA, USA). Descriptive analysis on HCC sero-prevalence was carried out by using Statistical Package for Social Sciences version 19.0 (SPSS Statistic 19, 2010 IBM). The unequal variance t-test (Welch’s t-test) was used to assess variations of infection between different age groups and gender, \[ \alpha = 0.05. \]

**Results**

**Prevalence and burden of human (neuro)cysticercosis**
Sera were collected from 1051 individuals and the participant ratio of female to male was 1:1.7. The mean age of participants was 34.61 ± 18.4 with a range of 3–95 years. About 16.3% (171) of the sampled population in Mbulu district tested positive for HCC Abs with 75% (128) of the positive cases being males. Fifty nine percent (101) of the positive cases displayed the lower MW band (6–8 kDa) indicating an active/live cysticerci infection. The prevalence levels of human cysticercosis between age groups was not statistically different (\[ p = 0.229. \]) However, notable was that the \[ \geq 80 \] years age group was the least infected (\[ p = 0.004. \]) and the 0–4 age group had no positive case (Table 1).

| Table 1 | Seroprevalence of human cysticercosis based on age groups and sex in Mbulu District, Northern Tanzania, 2013–2014 |
|---------|---------------------------------------------------------------------------------------------------------------|
| Age group | Number tested | Total positive cases | % positive | % positive males | % positive females |
| 0–4     | 5             | 0                    | 0          | 0             | 0                     |
| 5–14    | 128           | 14                   | 10.9       | 7.03          | 3.9                  |
| 15–29   | 334           | 44                   | 13.2       | 12.0          | 1.2                  |
| 30–44   | 284           | 53                   | 18.7       | 14.4          | 4.3                  |
| 45–59   | 168           | 37                   | 22.2       | 15.5          | 6.7                  |
| 60–69   | 75            | 14                   | 18.7       | 8             | 10.7                 |
| 70–79   | 37            | 8                    | 21.6       | 13.5          | 8.1                  |
| 80+     | 20            | 1                    | 5          | 5             | 0                     |
Of the 25 HCC sero-positives examined for NCC by CT scan, 76% (19) had one to several calcifications suggestive of NCC. Of the 19 individuals with NCC suggestive lesions, 26.3% (5) did not have any history of epilepsy but recurrent headaches. One participant among the 5 who was found with a single calcified cyst on the left frontal lobe also had multiple rings suggestive of toxoplasmosis in the brain. Seventy four percent [14] of the 19 with lesions had a history of epilepsy for a period of 2–10 years. Of the 6 with no signs of calcified cysts in the brain, 66.7% (4) had been suffering from epilepsy for 2–26 years and on further examination of one of them with frequent unprovoked seizures, it was revealed that the patient was also infected with Brucella spp. The remaining 2 (33.3%) were suffering from recurrent headaches for the past 2–5 years by the time they were examined (Fig. 1).

Generally, epilepsy caused 2.8 years of life lost (YLLs) and 2.2 healthy years of life lost due to disability (YLDs) per 1000 person-years in Mbulu. The YLLs was 2.8 for males as well as for females. The YLDs (0.8) for males was significantly lower than that for females (1.5) (p = 0.001). The DALYs due to epilepsy was 3.9 per 1000 person-years, representing the disease burden in Mbulu district. The DALYs imposed by NCC on rural communities in Mbulu was estimated to be 3.0 per 1000 person-years.

Discussion

The findings of this study show that the prevalence and the burden caused by NCC related epilepsy is considerably high in Mbulu district. The proportion of those exposed to infection by T. solium cysticerci in humans, based on circulating anticysticercal antibodies (Abs) is as high as one person in every six members of the community in the district. Furthermore, cCT scan results from selected sero-positive individuals showed that a significant proportion of them had NCC suggestive lesions with over 56% of them being epileptic. Even though the results of this study cannot be generalized, the number of sero-positive individuals with confirmed NCC suggestive lesions is a strong pointer to the contribution of the disease to epilepsy syndrome in the district. This prevalence level in the northern part of Tanzania is similar to that reported in the southern highlands of the country [13]. On the other hand, the observed prevalence in the present study was higher than that reported in Rwanda (2.8%) [14] and in Mozambique (12.1%) [15]. Furthermore, the observed prevalence is significantly less than that reported in Burundi (31.5%) [16]. While all HCC results compared in here are from the same region i.e. Africa south of the Sahara, the observed differences might be attributed to differences in the target populations studied, methods used for diagnosis/surveillance and the prevalent risk factors in each particular area.

In the present study, the prevalence of HCC was significantly higher in males than females. In Mbulu district the population of males in 2012 was higher than females [5]. The participant ratio for females vis-a-vis males was 1:1.7. Besides, Mbulu district is dominated by agro-pastoral communities. Under such a mode of life, communities’ roles between males and females are distinctly well known [17]. Yet, in such communities, women before menopause hardly participate in social functions and therefore might not suffer the same exposure levels like their husbands. The prevalence of NCC lesions established by this study was alarming! The high prevalence of NCC might be contributed by high preference of T. solium cysticerci in CNS [1].

The transmission of HCC/NCC might be attributed to the life style of the Mbulu community [13]. Some households do not share toilets with their daughters and mothers-in-laws thus heads of households prefer defecating
out in the bush. In this area, pigs are kept under freerange systems thus facilitating the cycle of this disease.

The burden caused by NCC related epilepsy was considerably high in Mbulu district when compared to the WHO estimates, 29% [18]. The differences in proportions of NCC associated epilepsy might be due to research approaches in establishing the prevalence. For example studies by Bhattarai et al. [12], Praet et al. [19] and Carabin et al. [20], where WHO rely under this case, were based on systematic reviews and meta-analysis data. Such studies are very important in making policy decisions although they are limited by selection of studies, limitations in executions of design and methods used to obtain primary data and heterogeneity. Bhattarai et al. also used stratified data from urban areas where the risk of acquiring *T. solium* eggs is relatively lower compared to rural areas where sanitation and hygiene standards may be questionable. This could be the reason that the proportion of NCC associated epilepsy in developing countries was estimated at 29%. The current study was cross-sectional. Data were obtained from the general communities limited to a district. However, the study was wider population because it focused on agro-pastoral systems thus facilitating the cycle of this disease.

As CNS syndromes of unknown origin is kept on increasing in cysticercosis endemic regions, wider and broad studies from different countries and regions in order to control *T. solium* cysticercosis are of paramount importance. It must also be noted that diagnosis challenges for HCC might be a limiting factor towards achieving this goal.

**Limitations of the study**

The results of this study may not be generalized to a wider population because it focused on agro-pastoral communities limited to a district. However, the study mainly depended on exposure seroprevalence and suggestive lesions diagnosed via CT Scan as confirmatory test.

**Authors’ contributions**

Conceived and designed the experiments: BJM, MCJ, GN. Performed the experiments: BJM, MCJ. Analyzed the data: BJM. Wrote the paper: BJM, MCJ, GN. All authors read and approved the final manuscript.

**Author details**

1 Department of Biosciences, Soloman Mahlangu College of Science and Education, Sokoine University of Agriculture, P.O. Box 3038, Morogoro, Tanzania. 2 Department of Zoology and Wildlife Conservation, University of Dar es Salaam, P.O. Box 35064, Dar es Salaam, Tanzania. 3 Department of Zoology and Wildlife Conservation, Wellcome Trust Fellow, P.O. Box 35064, Dar es Salaam, Tanzania.

**Acknowledgements**

Authors wish to thank Misses. Suzan Gabriel and Teddy Marceli of the Mbulu District Hospital for collecting and processing the blood samples during field work. We thank the Haydom Lutheran Hospital officials and technical staff particularly Dr. Naaman for conducting CT scanning. Further thanks go to the Mbulu District officials for accepting the study to be conducted in their district.

The authors would like to acknowledge the United States Department of Defense, Defense Threat Reduction Agency (DTRA), Cooperative Biological Engagement Program (CBEP) for their support to develop this manuscript. While DTRA/CBEP did not support the research described in this publication, the Program supported the presentation of this research in an international forum and supported grantsmanship training related to the development of this manuscript. The contents of this publication are the responsibility of the authors and do not necessarily reflect the views of DTRA or the United States Government.

**Competing interests**

The authors declare that they have no competing interests.

**Availability of data and materials**

Data and materials used to come up with this manuscript are available at the Department of Zoology and Wildlife Conservation, College of Natural and Applied Sciences, University of Dar es Salaam, Tanzania.

**Consent for publication**

Not applicable.

**Ethics approval and consent to participate**

The ethical clearance for this study was provided by the National Institute for Medical Research (NIMR), Tanzania, NIMR/HQ/R.8a/Vol. IX/1297. Parents and other participants aged above 18 years signed an assent form to participate in the study. Participants aged 18 years and above signed an informed consent form on their own, while for the participants aged less than 18 years, their parents signed the form on behalf.

**Funding**

The research leading to the writing of this paper was carried out within the framework of the "African Research Consortium for Ecosystem and Population Health: Expanding the Frontiers in Health", Africa One, which was funded by the Wellcome Trust (WT-087535MA), and the University of Dar es Salaam hosted and facilitated the project. The study design and collection, analysis, and interpretation of data were carried out by authors. Development of the manuscript was supported by the United States Department of Defense, Defense Threat Reduction Agency (DTRA), Cooperative Biological Engagement Program (CBEP).

**Publisher’s Note**

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Received: 3 October 2018 Accepted: 8 December 2018 Published online: 13 December 2018

**References**

1. Markell EK, John DT, Krotski W. Modern pathology Markell and Voge’s medical parasitology. 8th ed. Philadelphia:W.B. Saunders; 1999.
2. Sharma R, Bargotra R, Tandon VR, Gupta SK, Verma SSJ. Neurocysticercosis: current vitae. JK Sci. 2005;7(1):1–4.
3. Molinari JL, Tato P. Molecular determinants of host-parasite interactions: focus on parasite. In: Singh G, Prabhakar S, editors. *Taenia solium* cysticercosis from basic to clinical science. Wallingford: CAB International; 2002. p. 25–33.
4. Flisser A, Correa D, Evans C. *Taenia solium* cysticercosis: new and revisited immunological aspects. In: Singh G, Prabhakar S, editors. *Taenia solium* cysticercosis from basic to clinical science. Wallingford: CAB International; 2002. p. 15–24.
5. NBS. National Accounts of Tanzania Mainland (2001–2011); 2012. http://www.nbs.go.tz/. Accessed 03 June 2013.
6. MDC. Mbulu District profile, updated fourth quarter development report; 2013. http://www.citypopulation.de/php/tanzania-admin.php?adm2id=2103. Accessed 21 Feb 2014.

7. Murray CJ, Lopez AD. Burden of Disease: DALYs, Global Burden of Disease 2004, Updated 2008:40–51.

8. World Health Organization (WHO). Global burden of disease and injury series. Glob Burd Dis. 1997;36–200.

9. Jayashi CM, Gonzalez AE, Castillo R, Rodríguez S. Europe PMC funders group validity of the enzyme-linked immunoelectrotransfer blot (EITB) for naturally acquired porcine cysticercosis. Vet Parasitol. 2017;199:42–9.

10. Joshi DD, Bista PR, Ito A, Yamasaki H, Zoonoses N. Present situation of porcine taeniasis and human. Southeast Asian J Trop Med Public Health. 2007;38(suppl 1):1–2.

11. Ding D, Hong Z, Wang WZ, Wu JZ, De Boer HM, Philpko L, et al. Assessing the disease burden due to epilepsy by disability adjusted life year in rural China. Epilepsia. 2006;47(12):2032–7.

12. Bhattarai R, Helene C, Christine B. The burden of cysticercosis. Nov Asp cysticercosis neurocysticercosis. 2013;(3):59–76.

13. Mwanjali G, Khamia C, Kakoko DVC, Lekule F, Ngowi H, Johansen MV, et al. Prevalence and risk factors associated with human Taenia solium infections in Mbozi District, Mbeya Region, Tanzania. PLoS Negl Trop Dis. 2013;7(3):e2102.

14. Newell E, Vyunjimana F, Geerts S, Van Kerckhoven I, Tsang V, Engels D. Prevalence of cysticercosis in epileptics and members of their families in Burundi. Trans R Soc Trop Med Hyg. 1997;91:389–91.

15. Zoli A, Shey-Njila O, Assana E, Nguekam JP, Dorny P, Brandt J, et al. Regional status, epidemiology and impact of Taenia solium cysticercosis in Western and Central Africa. Acta Trop. 2003;87(1):35–42.

16. Nsengiyumva G, Druet-Cabanac M, Ramanankandrasana B, Bouteille B, Nsizabira L, Preux PM. Cysticercosis as a major risk factor for epilepsy in Burundi, East Africa. Epilepsia. 2003;44(7):950–5.

17. Flintan F. Changing nature of gender roles in the dry lands of the horn and east Africa: implications for DRR programming; 2011. https://www.preventionweb.net/publications/view/24271, Accessed 16 Dec 2013.

18. Bhattarai R, Bucke CM, Carabin H, Proaño JV, Flores-Rivera J, Corona T, et al. Estimating the non-monetary burden of neurocysticercosis in Mexico. PLoS Negl Trop Dis. 2012;6(2):e1521.

19. Praet N, Speybroeck N, Manzano D, Berkvens D, Nforninwe DN, Zoli A, et al. The disease burden of Taenia solium cysticercosis in Cameroon. PLoS Negl Trop Dis. 2009;3(3):1–8.

20. Carabin H, Cowan L, Nash T, Willingham AL. Estimating the Global Burden of Taenia solium Cysticercosis/Toeniosis; 2006. http://www.who.int/foodsafety/publications/foodborne_disease/burden_sept06/en/index2.html. Accessed 6 July 2013.