

Surgical Management of a Case of Multiple Hydatidosis

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Abstract: A 14 year old female was admitted to Elshab Medical Teaching Hospital, Khartoum, Sudan. As informed by the father, there was a history of cough for 2 months, chest pain for 2 months and breathlessness for 3 weeks. An intercostals drainage tube was administered to alleviate the condition and to relief pneumothorax. The patient was referred to thoracic surgery unit for thoracotomy of the left side of the chest. The second X-ray and MRI revealed the presence of a large hydatid cyst in the lower lobe of the right lung. The hydatid cyst was situated at the junction of the ventrolateral aspect of the upper lobe and dorsolateral aspect of the lower lobes of the right lung. Because of the deteriorating condition of the patient, thoracotomy was initially performed in the right side of the chest to remove the giant hydatid cyst. A week later, thoracotomy was also performed to remove a ruptured pulmonary cyst from the upper lobe of the left lung, which was found infected with secondary bacterial organism. An incidental cyst was also observed in the liver. Thus, this patient represents a case of multiple hydatid cysts. Conventional bacteriological examination revealed isolation and identification of *Pseudomonas stutzeri* from the infected hydatid cyst. Molecular characterization revealed that the cyst is of genotype 6 (G6) strain of *Echinococcus granulosus* as detected by polymerase chain reaction (PCR)-based assay.

Key words: Hydatid cyst, lungs, thoracotomy

INTRODUCTION

Cystic pulmonary hydatidosis in children or adolescence is very rare although it is commonly observed in adults. The mature worms of this cestode parasite are maintained in a carnivore definitive host whereas the larval stages are harboured by a herbivore intermediate host. The disease is prevalent in Africa including the Sudan, especially in rural areas where animal slaughtering is practiced on farms. In addition, improper disposal of offal from slaughterhouses and the presence of large populations of stray dogs could also contribute towards the endemicity of the disease^[1,2].

The parasite causes serious public health problems in certain parts of the Sudan^[2,3]. Humans can accidentally become infected by ingesting eggs of adult worms and hence, cystic hydatidosis, caused by *Echinococcus granulosus*, is a disease of public health importance. Ten genotypes (strains) of *E. granulosus* have been identified worldwide, designated (G1 to G10). In a previous study, the camel genotype (G6) was believed to be the most prevalent strains of the parasite in the Sudan^[4]. Despite the endemicity of cystic hydatidosis in African and Mediterranean countries, very little information is known in regard to molecular

characterization of the genotype of this cestode parasite. In this study, we present an unusual case of bilateral pulmonary hydatidosis with a ruptured cyst in the right lung and an infected cyst in the left lung. The pulmonary cysts were surgically removed and submitted to the diagnostic laboratory for further parasitological and microbiological examinations, and for identification of the genotype (strain) of the parasite using PCR.

MATERIALS AND METHODS

Patient: A fourteen- year old female was admitted to Elshab Medical Teaching Hospital, Khartoum, Sudan. As informed by the father, there was a history of cough for 2 months, chest pain for 2 months and breathlessness for 3 weeks. An intercostals drainage tube was administered to alleviate the condition and to relief pneumothorax. The patient was referred to thoracic surgery unit for thoracotomy of the left side of the chest. The second X-ray and MRI revealed the presence of a large hydatid cyst in the lower lobe of the right lung. The hydatid cyst was situated at the junction of the ventrolateral aspect of the upper lobe and dorsolateral aspect of the lower lobes of the right lung. Because of the deteriorating condition of the patient, thoracotomy was initially performed in the

right side of the chest to remove the giant hydatid cyst. Hydatid fluid was aspirated aseptically and hydatid cyst was surgically excised from the lung tissue.

Parasitological examination: Conventional parasitological examination was used for identification of the cyst. The cyst fluid containing protoscolices was stained with eosin and examined under the microscope. Definitive diagnosis was made possible by demonstration of free moving protoscolices under the microscope.

Microbiological examination: A loopful of infected materials from the cyst were inoculated in Macconkey media. Further characterization and bacterial identification was made possible using biochemical tests.

Extraction of nucleic acid from the hydatid cyst: The surgically removed pulmonary hydatid cysts were preserved in 70% alcohol and submitted to the molecular diagnostic laboratory for molecular characterization. The cysts were washed thoroughly with nucleic acid free water by centrifugation to remove the 70% alcohol. DNA was then extracted from protoscolices using QIAamp extraction kit as described previously. The DNA concentration was determined by spectrophotometer at 260 nm wavelength. Five µl of the suspended nucleic acid was used in the PCR amplification.

Polymerase Chain Reaction (PCR): The eluted DNA was stored at -20°C until used for PCR amplification. The Polymerase Chain Reaction (PCR) was employed to determine the strains of *E. granulosus* as described by Dogan^[4]. Primers p1 and p2 were used to amplify the 254 bp first PCR product from pig, cattle and camel strains (G5, G6 and G7), respectively. Primers P1 and p3 were used to amplify the internal (semi-nested) 171 bp PCR product, which is specific for camel strain (G6). PCR products were loaded onto 1.5% agarose gel and electrophoresed. The gels were then stained with ethidium bromide and the specific bands were visualized under UV light.

RESULTS

Parasitological examination of the right pulmonary cyst revealed the presence of hydatid cyst. Confirmation of hydatid cyst was made possible by demonstration of viable protoscolices under the microscope. The viability of living protoscolices was tested following surgical removal of the cyst (Fig. 1). However, no protoscolices were seen in association with the infected hydatid cyst of the left lung. Microbiological examination revealed the

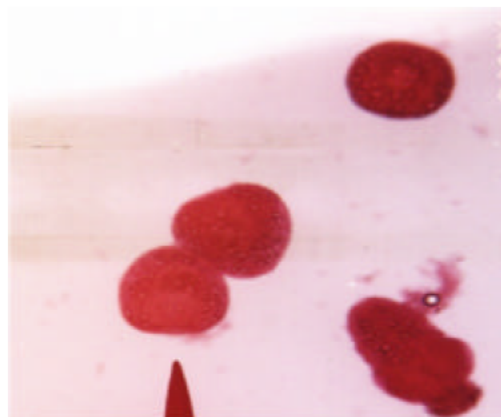


Fig. 1: The viability of living protoscolices derived from hydatid cyst of the infected patient

isolation and identification of gram negative bacilli *Pseudomonas Sp.* Using biochemical reactions, the bacterial organism was further identified as *Pseudomonas stutzeri*. The antibiotic sensitivity test indicated that bacterial growth showed a very large zone of inhibition around tetracycline. Molecular characterization of hydatid cyst-derived protoscolices was made possible using G6-specific PCR-based detection assay. Using a pair of outer primers, the PCR assay produced a 254 bp specific PCR products from DNA samples extracted from hydatid cyst of the infected patient. Using a pair of semi-nested primers, the PCR assay produced a 171-bp nested PCR product from the primary PCR product. The primary and the nested PCR products were visualized onto ethidium bromide-stained agarose gels. Thus, the recovered hydatid cysts were identified as camel strain (G6) of *E. granulosus*.

DISCUSSION

Hydatid lung disease caused by *Echinococcus granulosus* is often asymptomatic and usually benign. Surgical interference is not generally recommended unless the cyst is interacting with the normal function of the affected organ. Hydatid lung disease of adolescence in the African continent is usually caused by *E. granulosus*. Slowly enlarging *Echinococcus* cysts usually remain asymptomatic until their expanding size or their space occupying effect elicits symptoms. The cysts may be discovered as an incidental finding on a routine X-ray chest. Pulmonary hydatid cyst may rupture into pleural cavity, pericardium or the bronchial tree leading to cough, chest pain and haemoptysis. In the present case, the hydatid cyst, located at the lower left lobe of the lung, had ruptured and was invaded by secondary bacterial

infection leading to empyema, which mandates surgery to remove the cyst^[6,7,8]. Following this surgical procedure, the patient returned rapidly to Good health.

Diagnosis becomes difficult once the pulmonary hydatid cyst ruptures spontaneously or following trauma and gets secondarily infected. A more common cause of empyema is the infection of pleural fluid as reported in our case. It is worth mentioning that the general practitioner, who handled the case at the time of admission, did not investigate for hydatid lung cyst before insertion of intercostals drainage tube. This is probably due to the fact that one would not think of the disease in such young age group. Nevertheless, he did not attempt to obtain some information about previous history of contact with dogs. It is well documented that younger patients (children and teenagers) are more prone to be symptomatic and to develop complications from ruptured hydatid cysts. This is because children develop very large cysts in relation to the size of the lung (6-12 cm), due to greater elasticity of the lung. The cyst could also lead to a large residual cavity that takes more time to resolve and is then more likely to develop infections. It is well documented that the bronchial tree in younger patient is smaller than in adults and hence expulsion of membranes and particles is compromised.

It is worth mentioning that very little information is available in regard to genotypes of *E. granulosus* strain in the Sudan. Previous studies showed that the camel strain (G6) is the most prevalent strain of the parasite in humans and animals. Two cases of cattle strain of *E. granulosus* (G5) were reported sporadically in a survey of the disease in Khartoum State. The camel strain (G6) has never been reported in south of Sudan. In this study we report on isolation of G6 strain of *E. granulosus* in a child from Southern Sudan for the first time. From an epidemiological point of view, the patient is originally from Malakal area of the upper Nile Province of the Southern Sudan. Thus, it is probably that this strain (G6) of the parasite is circulating in southern Sudan. However, additional data are needed to confirm this assumption. In this study we would also like to point out the importance of a good history, especially of contact with dogs in the patients who develop respiratory complaints spontaneously or following mechanical trauma to chest. Such patients should be investigated on the lines of pulmonary hydatid cyst so as to facilitate rapid diagnosis and subsequent successful treatment. An incidental hydatid cyst was also seen in the dorsal aspect of the live during MRI examination. However, surgery was not recommended as the health condition of the patient is

deteriorating. The patient was treated with oral albendazole at a dose rate of 10 mg Kg⁻¹ body weight. Following surgical treatment and medicinal treatment, the patient health condition improved rapidly

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