



BLASTOCYSTOSIS : ETHIOLOGY, BIOLOGY AND PREVALENCE

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ABSTRACT

Blastocystosis is a protozoic invasion which clinical course displays as a healthy carrier or an enterocolitic syndrome. The infection is famous as Zierd-Garavelli disease. Residing the small bowel and colon *Blastocystis hominis* is a typical anaerobic and opportunistic infection. It is presented in several morphologic forms : vacuol, unvacuol, multivacuol, amebic and cystic ones. Transmission factors are food, drinks, hands and dust contaminated with cysts. Toxoallergic reaction causes the pathologic findings and leads to nonspecific inflammation of colonic mucosa. The enzymes produced by Blastocystic organisms cleave secretory Ig A of host and provide persisting in the human body. The disease appears to follow the course same as enterocolitis in immunocompromized people (eg. AIDS , leucaemia, diabetes). It is characterized with eosinophilly, stool leucocytosis and occult bleeding. Treatment regimens include Metronidazol, Tinidazole, Paromomycin, Biseptol and Furazolidon. There is possibility of self- limiting of the infection.

Key words: blastocystosis, opportunistic infection, protozoonosis, metronidasole, apoptosis

Blastocystosis is a protozoic disease presenting with diarrhoea or without symptoms. It is widespread opportunistic parasitosis which has a strong influence on the symptoms of the patients affected with AIDS and viral enteral infection.

Short historical data. The first morphological description of the parasite was done by Alexeieff (1911) as a microorganism with the name of Blastocystosis enterocola. In 1912 it was identified by E. Brunt as saprophytic fungus which is harmless to the human and was named Blastocystosis *Hominis*. In 1967 Zierdt C.H. et al describe it as protozoic. From 1976 the majority of authors considered it as a typical nonobligated (opportunistic) protozoic organism from subdivision Sarcodina (Amebi). Later Zierdt (1991) shows the role of *B. hominis* in enteral diseases and includes it to the enteral parasites. Because of this reason Blastocystosis is also notorious as Zierdt - Garavelli's disease. Based on

molecular and genetic research Cavalier-Smith, T. (1998) suggests *B. Hominis* to be qualified to a new class Blastocystea, subtype Opalinata, part of subkingdom Heterokonta, kingdom Chromista.

Ethiology and biology. The source of the disease is the parasite *Bl. hominis*, described above. In taxonomic plan the place of the parasite has not been determined yet. The genetic research determine genetic heterogeneity of the kind. Blastocystic isolates in different people show different number chromosomes (9-13). Noel et al. (2006) found out through genetic analysis an isolate of *Bl. ratti* which qualified to *Bl. hominis*. Stensvold, C.R. et al. (2007) suggest standardization of the subtypes. To avoid mistakes for now it is assumed that with *B. hominis* to be signed the parasite isolated from humans and respectively *Bl. species* the one from animals (birds and mammals).

B. hominis inhabits the small bowel and colon of the human body and the animals. In humans *B. hominis* inhabits the cecum, proximal part of colon transversum and at some degree in the distal part, where the parasite is found in the vacuolar, unvacuolar, granular, multivacuolar, amebic and cystic forms.

In human stool samples usually are observed unvacuolar, multivacuolar and cystic forms. If the parasite is cultivated in nutritious artificial environment vacuolar, granular and amebic forms are found.

Granular form is a spherical body with diameter of 4-200 μm . It is wrapped with bilaminar membrane and is characterized with a big central vacuole (that takes up to 75% of the cell's capacity), rounded by a thin cytoplasmic strip with cell organelles and 1-4 nuclei. The granular form has a spherical body as well with diameter of 10-80 μm . It lacks vacuole. It is usually observed in old cultures. The content consists of a big number closely situated granules which are believed to be the result of the parasite metabolism and are most likely food reserve. Both of the forms reproduce through binary cell-division.

Multivacuolar trophozoite is round or egg-like form with small measures - from 5-8 μm . It possesses a wide outer casing. In the cytoplasm there is a big number of small transparent vacuoles. It has not been determined whether they are independent formations or have connections between them forming a net.

Unvacuolar forms can be found only within material taken by colonoscopy. They have measures around 5 μm and also possess 1-2 nuclei and lack vacuoles. They only have cell membrane.

Amebic form was discovered by Tann and Zierdt (1973). It is round and is measured from 2,6-7,8 μm . It has 1-2 nuclei and 1-2 pseudopods. It does not have cell membrane and outer casing. It lacks central vacuole, Golgi complex and mitochondria. It is considered to be pathogenic form.

Cystic form was discovered by Mehlhorn (1988) in stool of the patient affected by AIDS. It is round with measures 3,7-5 μm (sometimes 5-10 μm). They are covered with thick outer membrane. It has 2-4 nuclei, in its cytoplasm there is a big quantity of glycogen and lipid drops. It is considered to be the invasive form of the parasite.

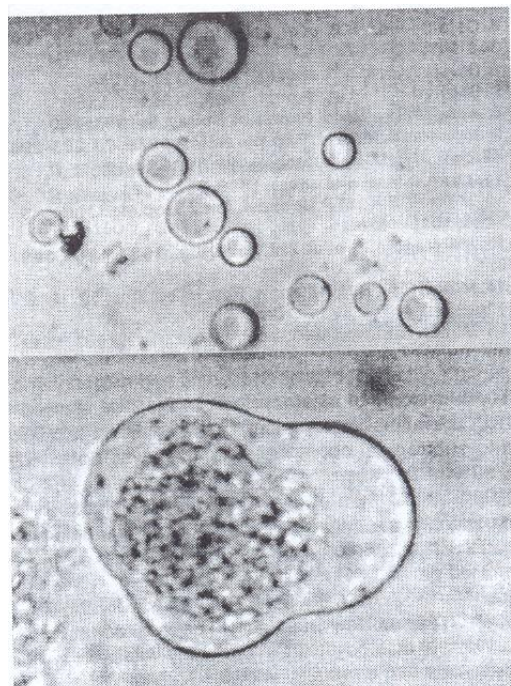


Fig.1. a,b *B. hominis* in stool probe (after R.Kurdova et al.,1995)



Fig. 2. *B. hominis* (after W. Peters & H.M.Gilles, 1989) (coloured with Iodine, x 1800)

The biological cycle of the parasite has not been studied enough. Thick-wall cysts get into stool of final host and contaminate the outside environment. They are responsible for the outer transmission along the stool to oral way. The infection represents swallowing of thick-wall cysts, which invade the mucosa of the small bowel. (fig.3). They excyst and rupture under the influence of the gastric juice and intestinal enzymes and transform into unvacuolar forms. After frequent asexual reproduction (mitosis) vacuolar forms are formed. It is presumed that the vacuolar form transforms into multivacuolar one. The litter passes through the phases of the pre-cyst, schizogony and encysting into thick-wall cysts, securing the autoinvasion. After multiplication of the vacuolar form amebic one is formed which increases in number, transforms in pre-cyst and after schizogony encysts into thick-wall cyst, the last is excysting into the stool. A possibility of encysting of vacuolar forms is assumed. When multivacuolar forms are cultivated vacuolar ones are formed.

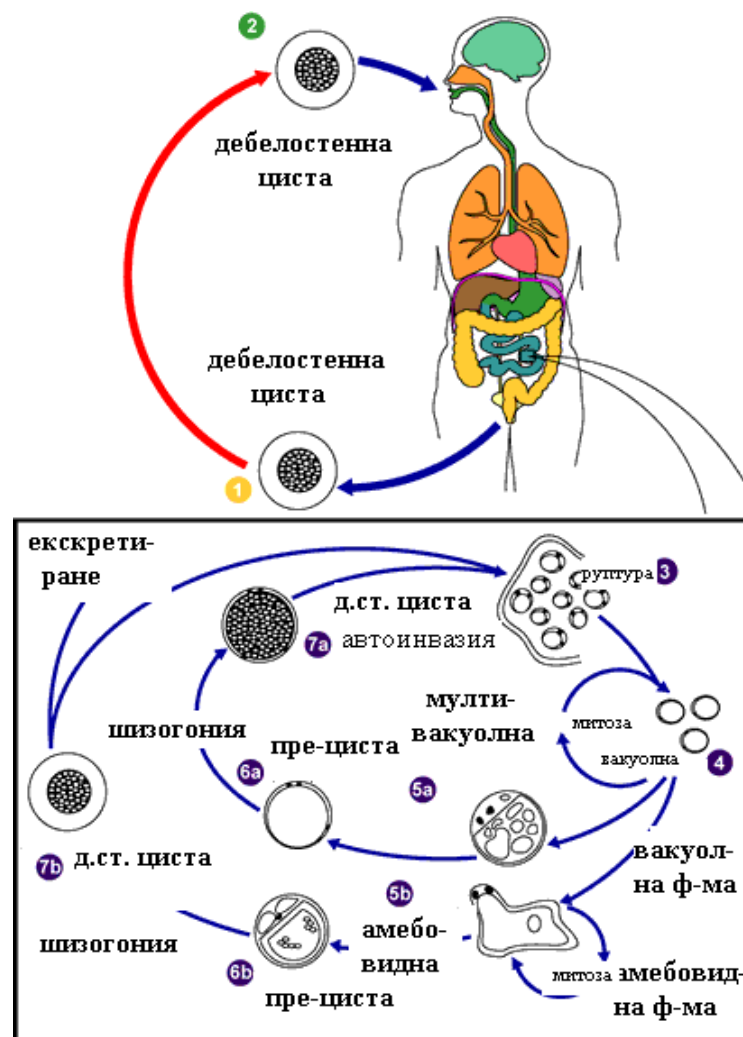


Fig.3. Biological cycle of *Blastocystis hominis* (no Singh M., Suresh K., Ho.L.C.et al.- *Parasitol.Res*, 1995; 81: 449)

Legend: to fig.3: 1. Separation of thick-wall cysts through stool in outer environment; 2. Infection by swallowing of thick-wall cysts and invasion of the intestinal mucosa; 3. Rupture of the cysts; 4. Asexual reproduction (mitosis) and formation of vacuol forms and their multiplication; after mitosis the vacuol forms transform into amoeba ones and their multiplication; 5-a. Multivacuolar form; 5-b. amoeba form; 6-a. pre-cyst; 6-b. pre-cyst; 7-a. After schizogony - the thick-wall cysts is responsible for the autoinvasion; 7-b. After schizogony - the thick-wall cyst excreting into the stool.

Epidemiology. Blastocystosis is an intestinal protozoanosis. The source of the infection is the human (sick or healthy carrier). Most likely a natural tank are the animals (birds, mammals including the monkeys). The mechanism of infection is stool to oral. It is possible a sexual way of infection. Factors of infection are fecal polluted with cysts water and food (fruits, vegetables), hands, dust etc.

There have been registered sporadic cases, group and family infections and epidemiologic explosion. The parasite affects both sexes, but more often children and older people. Most often are infected people in their active ages (30-40 yrs), in comparison

with children. Frequently victims of the infection are the homosexual people and the older people. Often the blastocystosis is detected among tourists who have been to tropical countries or come from countries with warm climate "traveling diarrhoea". It is also observed seasonal prevalence of the disease related to the climatic factors (increase of the frequency and intensity of the carriers in the months before the monsoons).

The cysts are resistant in outer environment. They do not destroy in distilled water and also in nutritious circles consisting parasiticide drugs. The rest of the forms quickly disintegrate in unpleasant conditions.

Usually the parasite is found in warm countries where the frequency of the invasion varies from 30 to 50%. In the developed countries of America and Europe this per cent is lower - 1,5-18%. Higher frequency is observed among the population of economic undeveloped regions and in people traveling to tropic countries.

In Bulgaria the first research and epidemiological surveys were done by R. Kurdova and co-authors (1995). They found

B. hominis in 64 (9.90%) among all of the 646 people from Bulgarian cruise that had frequently flown to tropic and subtropic countries. There have been diagnosed 7(12,28%) patients with blastocystosis related to AIDS. In 2006 in our country 277 (0,12%) people infected with *B. hominis* were found. Still the blastocystosis is not diagnosed from the medico-diagnostic laboratories in 11 regions of the country (2,3,4,12).

