

Electron microscopy and 3D nano-organization of virus-like and surface structure of *Entamoeba*, *Candida* and *Escherichia* species

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Protozoa, yeast and bacteria can be considered as attractive models for different organisms to analyze cytoplasmic and surfaces structure as well as cellular processes of eukaryotes and prokaryotes [1-3].

For electron-microscopic (EM) study of virus-like symbiotic strains of *Entamoeba histolytica*, *E. moshkovskii*, *Candida guilliermondii* (Cg strains NP-4), *Escherichia coli* (serogroup O₁₂₄) and their nano-particles it has been using the methods of negative contrasting, cryo- and ultrathin sections of computer morphometric and stereometric analysis (Video-test program, structure-5, nanotechnology) both.

The virus nano-particles of *Entamoeba* are shown to have stick-type form: their length is 180- 200 nm, diameter - 60-70 nm, surface square - 11036 nm². They are in the cytoplasm of *Entamoeba* solely and with rosette type. Ultra structural pathological changes of viruses-symbionts were established under γ -ionizing radiation in cellular membrane structures with thickness size of 9-10 nm and in nucleoids both, and also in formation of electron density round particles with diameter of 5-35 nm in virions, eliminating from the individual virus-like particles: diameter of the big round particles "O1" of 35 nm, surface square - 885 nm², diameter of the small "O2" - 8 nm, surface square - 45 nm² (Fig. 1). The length of biomolecular ribonucleoprotein helical-like nano-particles and crystal body in the vegetative and cyst forms of *Entamoeba* is of 300 nm, diameter - 40 nm. Morphometric scanning EM analysis of the *C. guilliermondii* after x-ray radiation allow to consider the relation of square of destructive structures to the total square cell of Cg is equal to 26,2773 : 73,723 (Fig. 2).

The application of direct EM method allows visualization of the surface of enteropathogenic *E. coli*: there is a large number of fimbria. Computer stereometric EM analysis of these fimbria points out that their diameter is 8-15 nm (Figs. 3a, 3b).

Thus, the results obtained with EM morphometric and stereometric analysis of nano-particles of virus-symbionts of protozoa, yeast and bacteria allow considering the effectiveness of comparative EM approach during the investigation of radiation effects as well as of bio-nanoparticles.

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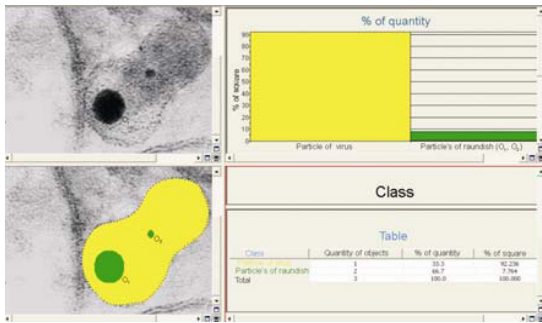


Figure 1. Transmission EM picture of virussymbionts of *Entamoeba*. Computer morphometrical analysis of the “Video-test, structure-5, nanotechnology” was used under the program.

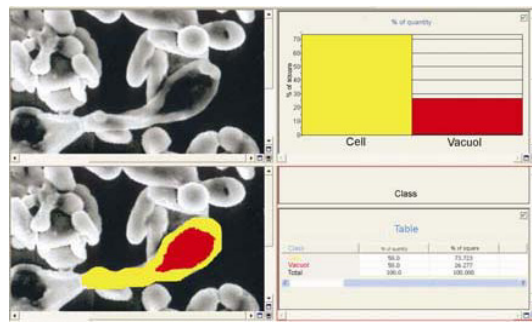


Figure 2. Scanning EM picture of *Candida guilliermondii* after x-ray radiation. Computer morphometrical analysis of the “Video-test, structure-5, nanotechnology” was used under the program.

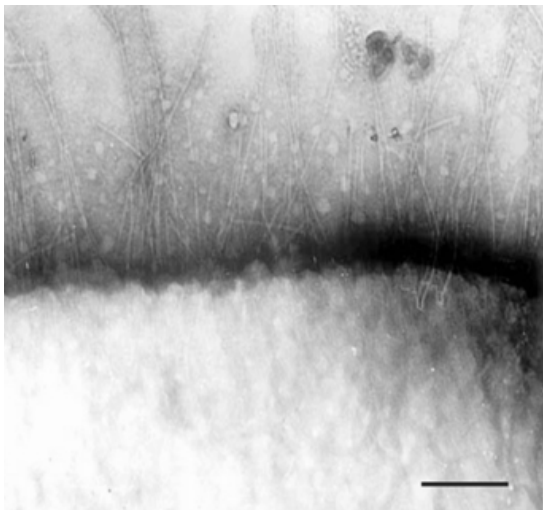


Figure 3a. Transmission EM picture of fimbri of *E. coli* (serogroup O124) (negative contrast). Bars: 100nm.

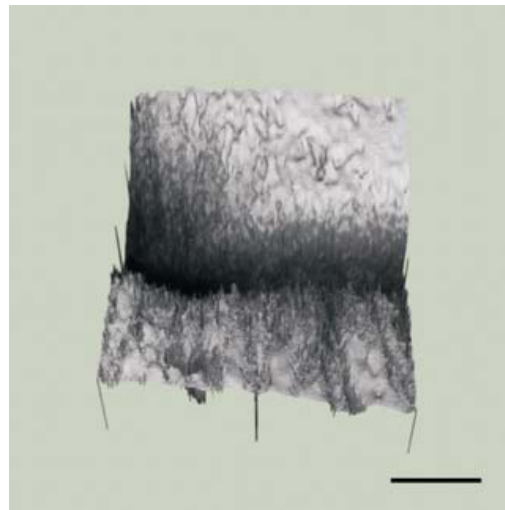


Figure 3b. Fimbri of *E. coli* (serogroup O124). 3D imaging analysis with program “Video-test, structure-5, nanotechnology” Bars: 100nm.