Something about miasms

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It is necessary to distinguish between miasms and miasms. We have known about Miasms as scenarios for the development of a painful process since the time of Hahnemann. Let's leave them alone for now and try to deal with the miasma with a small letter. What are they, those same miasms according to Schimmel?

To clearly see the whole picture, try testing them in several generations of the same family. The author had a similar experience. Testing of miasms was carried out through pseudo-transparent markers:

1. The sum of DNA and chromosomes or
2. Intox III and chromosomes, which made it possible to clarify the localization of the miasms.

Rather, to identify only miasms associated with DNA and chromosomes, distorting hereditary information and causing DNA damage. It turned out that miasms are distributed in the family group strictly according to the laws of distribution of hereditary properties. In children, only the miasms of their parents were found. So, we can assume that these miasms, which are fragments of RNA and DNA of microorganisms, attached to the genome of our ancestors and passed down from generation to generation, are the very aggravating factor that is called "bad heredity" or "family karma".

Then a legitimate question arises, is there anything you can do about it? It turns out you can. The author proposes miasmatic therapy using grafting techniques.

Having determined with the help of hereditary nosodes the stage of development of the disease (A.E. Kudaev's scheme) and choosing a technique, we proceed to collecting problems that require solution. Each subsequent task of this stage is interconnected with the previous one and is distributed in this way:

1. Chakras (if the inverted nosode of the chakras was used at the first stage of treatment, at this stage, the chakras may not be tested at all);
2. Meridians;
3. DNA;
4. Intox III (sometimes together with Intox II, if you have purchased DNA damage);
5. Chromosomes;
6. Miasms;
7. Resoplexes;

Testing of subsequent tasks through the previous ones turns out to be more reliable and tied to the place. A kind of chain of relationships and influence is formed. In this form, it is presented to the body for a solution. The resulting native US becomes our training drug for UFS, helping it to eliminate damage in the hereditary apparatus.

In virtual testing, 1 ball of the drug completely compensates for both KMH and the entire complex of tasks.

The only weak point of the Schimmel miasms is that they are clearly not enough. Special research work is needed to more fully identify all possible foreign inclusions in DNA, which may have not only biological, but also toxic chemical, and

informational nature. The more complete and comprehensive the miasm scale is, the more effective miasmatic therapy will be. With a preventive purpose, such therapy should be carried out to all who marry and who are going to have children. And then a new generation will appear, born without hereditary burdens.

It should not be forgotten that this therapy session will fully manifest its effect on the body not by the end of the drug intake, but after 6-10 months, which are necessary to restore the correspondence between the corrected genetic apparatus and the renewal of the biochemical components of the cell.

Very interesting information comes to light during testing Schimmel miasms through ONOM miasms. It is possible to determine which Schimmel miasms correspond to each classical miasm.

For example, syphilitic miasm correlates with anthrax, the causative agents of syphilis, smallpox and even herpes. Here the conclusion suggests itself that by getting rid of miasms with a small letter, we will be able to eliminate all the scenarios for the development of diseases. As a result of treatment, all miasms disappear, except for the most ancient Miasms: syphilis, gonorrhea and psora, which in the preparations of hereditary nosodes corresponds to tuberculinum. These three "monsters" are almost always present, being in a state of constant interaction with each other. There is a certain pattern alternating periods of their least and greatest impact on the body. It was reflected in the scheme of the development of the disease according to A.E. Kudaev. The circuit works, and even very well. This means that there is something completely objective, real, which provides these Miasms with "eternal" existence in our inner world.

In the works of E.V. Alekseeva proves the existence in human blood of the originally ancient representatives of life on earth: a unicellular algae, a flagellar microorganism and an imperfect fungus, mimicking erythrocytes. This evolutionarily fixed microflora has become a part of ourselves. She lives in us, feeds on us and interacts with each other. A change in our internal environment leads to an imbalance in this microcosm, when one of the microorganisms gets the opportunity to develop rapidly by suppressing the growth of the other. And if we assume (hypothetically) that it is these microorganisms that give life to Miasms - "monsters" called syphilis, gonorrhea and psora, then the dynamism and constancy in testing hereditary nosodes becomes clear. If this is so, then to fight them is the same as to fight with yourself - stupid, dangerous and ineffective. It is much more practical and important to use the knowledge of their interaction, reflected in the scheme of A.E. Kudaeva to improve the quality of treatment.

Conclusion: miasmatic therapy using Schimmel miasms and grafting techniques allows to repair DNA damage, which provides a qualitative leap in treatment and an increase in the step of therapy.

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