

The Prime Cause and Treatment of Cancer

Sorush Niknamian*

Professor, Department of Cell Biology, Molecular Biology and Biomedicine, University of Cambridge, UK.

*Corresponding Author: Sorush Niknamian, Ph.D, Professor, Department of Cell Biology, Molecular Biology and Biomedicine, University of Cambridge, UK. E-mail: so.niknamian@gmail.com

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Introduction

Cancer disease has become the first cause of death in the United States and world-wide. Most Researchers estimate that 595,690 of American people will die from cancer at the end of the year 2017. That means 1,600 deaths/day approximately [1]. Cancer in modern societies is commonly treated with the combination of organ surgery, chemotherapy and radiotherapy which has many side effects and the high possibility of metastasis. Many kinds of diet strategies have been experimented. However, none of them have been particularly effective. According to Otto Warburg hypothesis in the 1930's, the cause of cancer is the change in the metabolism of mitochondrion in human cells. Low oxygen in tissues in combination with high blood glucose will change the cell respiration from aerobic to anaerobic which leads to fermentation type of respiration [2]. Professor Thomas N. Seyfried in 2012 proposed cancer as a metabolic-mitochondrial disease. He has mentioned that all cancer cells' mitochondria are damaged and do not work properly [3]. Mitochondrion is a microorganism living inside our cells and produce energy in the form of Adenosine Triphosphate (ATP) from fats, glucose and glutamine. In order to produce ATP, mitochondria need oxygen through the krebs cycle which is called "Aerobic Respiration". On the other hand, when the amounts of oxygen go under 30% in the tissues and cells, mitochondria cannot respire properly which causes the increase in the amounts of Reactive Oxygen Species (ROS) or better called "Inflammation" inside the cells and causes damage to the mitochondria. According to Professor Thomas N. Seyfried, the main cause of cancer is the increase in ROS inside normal cells. [Thomas N. Seyfried et al., 2012] In 2017 we hypothesized with the sponsorship of the Weston A. Price Foundation (WAPF), that cancer is an Evolutionary Metabolic Disease which we proposed in the 4th International Conference in Medicine and Neurological Diseases which held in Fulda/Germany.

Evolutionary Metabolic Hypothesis of Cancer (EMHC)

The first living cells on Earth are thought to have arisen more than 3.5×109 years ago, when the Earth was not more than

about 109 years old. The environment lacked oxygen but was presumably rich in geochemically produced organic molecules, and some of the earliest metabolic pathways for producing ATP may have resembled present-day forms of fermentation. In the process of fermentation, ATP is made by a phosphorylation event that harnesses the energy released when a hydrogenrich organic molecule, such as glucose, is partly oxidized. The electrons lost from the oxidized organic molecules are transferred via NADH or NADPH to a different organic molecule or to a different part of the same molecule, which thereby becomes more reduced. At the end of the fermentation process, one or more of the organic molecules produced are excreted into the medium as metabolic waste products. Others, such as pyruvate, are retained by the cell for biosynthesis. The excreted end-products are different in different organisms, but they tend to be organic acids. Among the most important of such products in bacterial cells are lactic acid which also accumulates in anaerobic mammalian glycolysis, and formic, acetic, propionic, butyric, and succinic acids. The first cell on the earth before the entrance of the bacteria did contain nucleus and used the fermentation process to produce ATP for its energy. Then an aerobic proteo-bacterium enters the eukaryote either as a prey or a parasite and manages to avoid digestion. It then became an endosymbiont. As we observe, the fermentation process used the glucose or even glutamine to produce ATP, but the aerobic process used the glucose, fat and protein to produce more ATP than the previous one. The symbio-genesis of the mitochondria is based on the natural selection of Charles Darwin. Based on Otto Warburg Hypothesis, in nearly all cancer cells, the mitochondrion is shut down or are defected and the cancer cell do not use its mitochondrion to produce ATP. This process of adaptation is based on Lamarckian Hypothesis of Evolution and the normal cells goes back to the most primitive time of evolution to protect itself from apoptosis and uses the fermentation process like the first living cells 1.5 billion years ago. Therefore, cancer is an evolutionary metabolic disease which uses Glucose and Glutamine as the main food to produce ATP and Lactic Acid.

In another study, we hypothesized that increasing the ROS in a cell can cause damage to the DNA of the mitochondrion and also Nucleus DNA, but another reason behind turning the

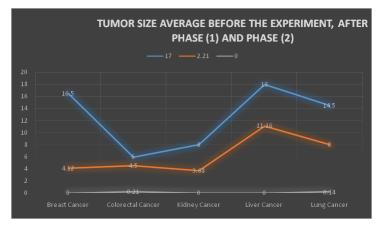
normal cell into cancer cell is the chaos caused by the increasing of inflammation inside each cell and increasing the intracellular ROS. These chaos causes some abnormal messaging between the DNA of the nucleus to stop the apoptosis and turning the oxidative phosphorylation to the fermentation in cytosol. Normally by damaging to the mitochondria, the cell should apoptosis. however; the nucleus sends wrong messages to stop the apoptosis and do fermentation process in cytosol to survive the cell. Even some normal left mitochondria would be shut down and stop the oxidative phosphorylation. This is the main and the real reason how increasing intracellular inflammation [5].

With all the information above, the basic way to stop respiration of cancer cells is to stop feeding them. The main source of fuel in cancer cells are Glucose and Glutamine. The basic diet for weakening or even killing these cells is Ketogenic Diet (KD). Ketogenic diet is a kind of regime which uses high fat content and low carbohydrate. This diet changes the metabolic state into the condition called Ketosis. After several days, fat becomes your body's primary energy source which causes an increase in the levels of compounds which is called "ketones" in the blood [6]. In general, a ketogenic diet used for weight loss is about 60-75% of calories as fat, with 15-30% of calories from protein and 5-10% of calories from carbs. However, when a ketogenic diet is being used therapeutically for the treatment of cancer, the fat content may be significantly higher that is up to 90% of calories, and the protein content lower [7].

Presently, limited researches do seem to show that a ketogenic diet may reduce tumor size and rate of progression in certain cancers. One of the few documented published case studies was performed on a 65-year-old woman with brain cancer. Following surgery, she received a ketogenic diet. Meanwhile, the tumor's progression slowed. But, 10 weeks after returning to a normal diet, she experienced a significant increase in tumor growth [8]. Similar case reports examined the reactions to a ketogenic diet in two young girls who were undergoing treatment for advanced brain cancer. Researchers observed that glucose uptake was decreased in the tumors of both patients. One of the girls reported that her quality of life had improved and remained on the diet for 12 months. During that time her cancer showed no further progression [9]. One study monitored tumor growth in response to a high-carb versus a ketogenic diet in 27 patients with cancer of the digestive tract. Tumor growth increased by 32.2% in patients who received the high-carb diet, but decreased by 24.3% in the patients on the ketogenic diet [10, 11]. In another study, three out of five patients on a ketogenic diet combined with radiation or chemotherapy experienced complete treatment. More interesting, the other two participants found the disease progressed after they stopped the ketogenic diet [12].

We had organized a research in the Violet Cancer Institute (VCI) on 54 patients with 6 different types of cancer. The aim of this research is to figure out the effectiveness of the Sorush Cancer Treatment Protocol (SCTP) which is based on the Evolutionary Metabolic Hypothesis of Cancer (EMHC) and introducing the Specific Ketogenic Diet (SKD) plus Intravenous Ozone Therapy (IOT) in Phase (1) on 54 cancer patients (Ozone has been found to be an extremely safe medical therapy, free from side effects. In a 1980 study done by the German medical society for ozone therapy, 644 therapists were pulled regarding their 384,775 patients, comprising a total of 5,979,238 ozone treatments administered. There were only 40 cases of side effects noted out of this number which represents the incredibly low rate of 0.000007% and only 4 fatalities. Ozone has thus proven to be the safest medical therapy ever devised. [16]), and combination of Hyperbaric Oxygen Therapy with vitamin/mineral and herbal supplementation beside the SKD and IOT in Phase (2) of this research on the remained 31 cancer patients. Based on the researches from 1928-2016 and the experimentation of cancer treatments and protocols on cancer patients, we have reached a treatment and decided to test it on 54 voluntaries cancer patients in the first stage of their disease. In this treatment we used a 5-day water fasting state, the Specific Ketogenic Diet (SKD) designed by ourselves and Intravenous Ozone Therapy (IOT) in the duration of 90 days (Phase 1) and another 90 days (Phase 2) with the entrance of Hyperbaric Oxygen Therapy (HBO2T) and several supplements which we have been effective in previous studies on cancer patients. We have used the measurement of saliva PH, the MRI device and statistical methods to test the shrinkage of the tumors. After Phase (1) of this research on 54 patients the average percentage decrease in the tumors was 58% and after Phase (2) on 31 remained cancer patients the average percentage decrease in the tumors was 98.8%. The average saliva PH in the fasting state of the cancer patients improved from acidic to alkaline as well. In conclusion, we have reached an effective cancer treatment based on SCTP by the usage of SKD, IOT, HBO2T and several supplements.

Figure 1. The blue line shows the tumor size before the experiment begins. The orange line shows the tumor size after Phase (1) of the treatment. The white line shows the shrinkage and improvement of the cancer patients tumors after 180 days.



There was an obvious improvement of cancer tumor decrease, lifestyle, saliva PH and we did not observe any side effects and cachexia in any of the patients [4].

We named this methodology used in this treatment the Sorush Cancer Treatment Protocol (SCTP). This protocol includes: 5-day water fasting, SKD, IOT, HBO2T, natural Vitamin/Mineral Supplements, Herbal Supplements, 3000-5000 mg ascorbic acid, probiotic foods and cottage cheese mixed with flaxseed oil. The total calorie intake by the patients should be 1200-1500 Cal/Day. The dietary restrictions in this protocol are: Dairy, Industrial Vegetable oils, Margarines, Alcohol, Gluten,

Soy Products (Miso and Natto are exceptional), Artificial Sweeteners as well as Stevia, Fruit juices and any types of sugar including Honey. The duration of the SCTP should not exceed 6 months to reduce the possibilities of ketoacidosis occurrence in patients.

The most important note in this protocol is that we banned the consumption of industrial vegetable oils and margarine. In our research which we published with the help of Miriam Kalamian EdM, MS, CNS, we concluded that Omega-6 Linoleic acid from vegetable oils increases oxidative stress in the body of humans, contributing to endothelial dysfunction and heart

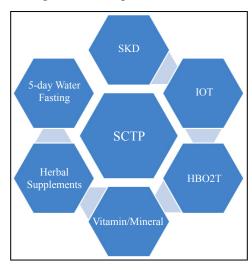


Figure 2. SCTP procedure in brief.

disease. The consumption of these harmful oils which are high in mega-6 polyunsaturated fats results in changing the structure of cell membrane which contribute to increasing inflammation and the incidence of cancer [13]. We replaced MCT by Extra Virgin Coconut Oil which we found to have more advantages in the Ketogenic Diet for the treatment of cancer [14]. Furthermore; we have prohibited the consumption of any soy products in our protocol since there is many evidences show the relation between the consumption of Soy and cancer incidences specially breast cancer [15].

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