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RESEARCH ARTICLE

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INTERACTIONS BETWEEN METABOLOMES: A VIEW INTO HEALTH AND ILLNESS

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Abstract

The concept of metabolic regulations deals with the varied and innumerable metabolic pathways that are present in the human body. A combination of such metabolic reactions paves the way to the proper functioning of different physiological and biological processes. Dealing with the adversities of a disease, engineering of novel metabolic pathways showcases the potential of metabolic engineering and its application in the therapeutic treatment of diseases. A proper and deeper understanding of the metabolic functions in the human body can be known from simulated yeast models. This gives a brief understanding about the interactions between the molecular set of metabolomeand its complexity.

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Introduction:-

As a central hub or an area concentrated with majority of metabolic reactions, it can be understood from the studies of the gut that systemic metabolism in human is not just regulated by their genes and their personal dietary habits but also by gut microbes [1]. If the gut microbiota present in the body is in a state of intestinal dysbiosis, certain micro-organisms like *E.coli*can be engineered and modelled metabolically to improve the functioning and growth of the indigenous microbiome [2]. The role of microorganism is widely being known and explored in the recent years due to their exploitable advantages and disadvantages that are meant to be kept in check [3]. Apart from different cohorts and divisions of microbiota present, a decent understanding and knowledge about the gut microbiota present in the human digestive system is required to evaluate, explore and treat the different diseases related to the human intestine tract [4]. A proper balance in the growth and bioactivity of different intestinal flora is required for the homeostasis of the human biological system [5]. A metabolic pathway is basic for every disease or any biological function that takes place in the body [6]. So study about these metabolic pathways and identifying the metabolites involved in them as markers helps in easy diagnosis and treatment of different diseases [7]like Non-small cell lung cancer and Anaplastic large cell lymphoma (ALK), Alpha-fetoprotein (AFP) - Liver cancer andgerm cell tumours, Beta-2 — microglobulin (B2M) - Multiple myeloma, chronic lymphocytic leukemia, and some lymphomas and problems faced by the organism [8].

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Role Of Metabolites In Precision Medicine

It is known from statistics that out the many people who are being treated with a disease only few of them are responding to the treatment and some are not. As an example when it comes to the radiotherapy and immunotherapy for the treatment of different cancers, only a handful people are being cured. This is because not every individual's body responds in the same way. So this is where the concept of precision medicine comes in to picture [9]. With the required biomarkers and companion (diagnostic tests used as a companion to a therapeutic drug to determine its applicability to a specific patient), researchers based on the patients disease progression and other key factors can stratify patients in to subsets [10]. This facilitates in the better prediction of disease outcomes so that appropriate treatment regimes can be formulated for the different sub groups identified. In turn, stratified medicine can give rise to precision medicine, where treatment is tailored for each patient according to their medical history, results from other tests, their response to medication and other clinical features [11]. In this respect there are a set of plant derives secondary metabolites like Vinblastine [12], Capsaicin [13], Curcumin[14] which are of medicinal impotence. These metabolites can be harnessed by modifying their respective metabolic pathways so that these metabolites are produced in large amounts. As a result they can be commercially produced in large amounts and aid in the treatment of different diseases [15].

Gut Microbiota - Impacting Flora In Human Body

The fluctuation seen in the growth of different gut microbiota can be due to host genotypes, physiological status, diet, drugs, and living conditions. The system formed as a combination of both the gut microbiota and the human system is called a "Superorganism" [16]. Based on the effects they induce, gut microbiota can be divided into 3 groups: - 1) beneficial bacteria 2) conditional pathogenic bacteria 3) pathogenic bacteria. As per the growth and functioning of different classes of gut microbiota listed above it results in the different diseases and ill effects of the gut and organs concerned to it like the liver and the gall bladder.

SCFA And Gut Microbiota

In general the food that enters into the digestive system is partly digested by the digestive enzymes and partly by the gut microbiota. The complex carbohydrates that entering the human gut is fermented into SCFA (small chain fatty acids) via the gut microbiota which further promotes the process of intestinal gluconeogenesis, and the formation of lipids [17]. This SCFA produced is known to play certain significant role in host organisms by improving the intestinal functioning, increases the resistance against pathogenic microorganisms, fight tumours, maintaining the electrolyte balance of the host and they also provide energy to the host epithelial cells [19]. Anotherintriguing factor about the gut microbiota is found out through a study that the peroxisome proliferation receptor- γ (PPAR- γ) signal induced by them is the one responsible for maintaining homeostasis. The compound that is responsible for the transduction of PPAR- γ is butyrate which is mainly produced by the metabolism of Clostridia. Butyrate also decrease the production of TGF- β 1 and IL-6, increases the activity of cytokines (anti-inflammatory) and by inducing the T cells enhances body immunity through anti-inflammatory effects [19].

It was further known that the *Bifidobacteriaceae* in the intestine of the mice have started to increase in number after the treatment with oligofructose weakened the weight gain, fat accumulation and ameliorated metabolic disorders induced due to high fat diet in mice [20]. *Akkermansiamuciniphila* is microbewhose abundance in the gut is closely related with the health of the host. It majorly survives on the intestinal mucin as the only carbon and nitrogen source with its main metabolite being propionate (SCFA) and its intestinal abundance is around 1-3%. These bacteria with its metabolite are seen to have their effect in the inflammatory responses of obese and diabetic patients, improve adverse symptoms such as insulin resistance and glucose tolerance [21].

Gut Microbiota- A Regulative Biome For Many Diseases

A comprehensive study on gut microbiota can give us a idea about different diseases on which the gut microbiota can have their effect [22]. If seen every disease has its own specific microbial markers for the targeted treatment of diverse diseases. In this point of view Louis et al. found out that, in a weightloss problem conducted the Firmicutes/Bacteroidetes was high in obese patients [23] and the Akkermansiaan intestinal microbiota abundance was found in successful weight loss patients [24]. Additionally it was also found that the Lactobacillus additives maintain homeostasis and reduced body weight considerably [25]. Similarly when it comes to liver diseases and liver cirrhosis, compared to healthy individuals the significant increase in the number of Enterobacteriaceae, Enterococcus species and Proteus species were found in patients with liver cirrhosis [26]. Seen at the pathogenesis of gastrointestinal diseases the microorganisms like EnteritoxigenicB. fragilis induced inflammatory responses in

colorectal cancer (CRC) mouse models [27]. In this disease model it was also found out that the colon epithelial regeneration was hindered to an extent due to the low availability or absence of *Bifidobacterium*[28].

Conclusion:-

From above mentioned strategies using metagenomics it is clear that it has become a powerful technology in analyzing the gut microbiota and in understanding its relationship with host. But there are some limitations. It is not an easy task to know the expression of microbial systems and it also requires higher sequence coverage. The time and cost are also considerable constraints for limitations. Among above all limitations mentioned above getting highly purified and high quality DNA samples is important because there may be 50% of human contaminants in DNA sample selected

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