Epigenetics: Pharmacology and Modification Mechanisms Involved in Cardiac, Hepatic and Renal Disease

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ABSTRACT

For a long time scientists have tried to describe disorders are due to genetic as well as environmental factors. In the past few years, revolution in technology that has made it possible to decipher the human genome. Epigenetics explains the capability gene expression regulation without modifying the genetic sequence. Epigenetic mechanisms are rooted changes in molecules, or nuclear characteristics that can alter gene expression without altering the sequences of DNA, i.e. DNA methylation, histone modification, and non-coding RNAs. Learning of the fundamental epigenetic modification allowing gene expression as well as cellular phenotype are advanced that novel insights into the epigenetic control of cardiovascular disease, hepatic disease, as well as chronic kidney disease are now emerging. From a half of century ago, in human disease the role of epigenetics has been considered. This subject has attracted many interests in the past decade, especially in complicated diseases like cardiovascular disease, hepatic disease as well as chronic kidney disease. This review first illustrates the history and classification of epigenetic modifications and the factors (i.e. genetic, environment, dietary, thought process and lifestyle) affecting to the epigenetics mechanisms. Likewise, the epigenetics role in human diseases is think out by targeting on some diseases and at the end, we have given the future perspective of this field. This review article provides concepts with some examples to describe a broad view of distinct aspects of epigenetics in biologic and human diseases.

Keywords: - Epigenetics, DNA methylation, Histone modifications, microRNAs and Gene expression and Disease.

Introduction:

In case of modern medicine genetics field has become an essential part in the 50 years therefore the structural model of DNA is illustrated by Watson and Crick. Advances have been made in identifying specific mutations related with some human diseases like cystic fibrosis as well as Huntington's chorea. Additional advances such as Human Genome Project, are giving new opportunities to develop better diagnostic tools and targeted gene therapies. The Epigenetics term itself describes, as 'epi' means above or over in Greek, but it was Conrad Waddington who, in 1940, provided the first broad and operational definition of epigenetics it is nothing but the unusual interactions in genes and their products, that bring the phenotype into being. Conrad Waddington in the 1950s states that "an epigenetic feature is a stably heritable phenotype results in changes in a chromosome without changing the sequences in the DNA".

Epigenetics and Epigenomics:

The epigenetic definitions state that the whole DNA content is exactly similar in somatic cells of one species, though gene expressions patterns have number of differences in various cell types that can be inborn clonally. Epigenetics have an tremendous effect on medicine. Epigenetics is nothing but the study of heritable alteration in gene functions that do not changes the DNA sequence but it can provide a one more layer of transcriptional control that control how genes are explicit.

The term "epigenetics" has been defined for decades as a alteration in the state of expression of a gene (or a feature) that does not contain a mutation but that is nonetheless rooted (at least through a mitotic division) in the absence of the signal or event that begins the change, the newer term "epigenomics" for our knowledge never been formally defined "Epigenomics" on the genome-wide level search
communities has been widely accepted operationally to indicate studies that focus on the analysis of DNA methylation, histone modifications, and noncoding RNAs. An Epigenetics mechanism describes that enable cells to respond quickly to environmental alteration and provide a link between genes and environment. Variation in the epigenetic modification of genes can describe a larger part of the phenotypic fluctuations observed in humans than differences in genotype alone. The current work focuses on epigenetic mechanisms of epigenetics in disease condition, for normal growth, development and homeostasis epigenetic processes are essential. A combination of stochastic events including both genetic as well as environmental factors (e.g., diet, smoking, obesity and stress), which, at the molecular level, cause alteration in gene expression and the aging process of organism. For example, aging of the brain is accompanied by changes of gene expression encoding proteins involved in stress responses, inflammatory responses and neuropeptide metabolism while the elderly heart has an altered transcription profile that leads to disability of cardiac function.

In the past decade various studies have strongly implicated epigenetic mechanisms in controlling the gene expression changes regulating several aging-related diseases like cancer and cardiac failure and in promoting the gene expression changes are responsible for the aging process of different tissues. Therefore, during aging process cells more prone to the transcriptional changes responsible for aging-related diseases due to the alteration of epigenetic mechanism. In this review, we will discuss the possible role of epigenetics in controlling the onset of two aging-related diseases like cancer and cardiovascular disease.

**Mechanism of Epigenetics**

The major epigenetic controlling mechanisms of mammalian cells include:

1. **DNA methylation**
2. posttranslational histone modifications (PTMs) and
3. **RNA based mechanisms** including those controlled by the better characterized small non-coding RNAs (snRNAs).

MicroRNAs (miRNAs) and long non-coding RNAs (lncRNAs) are newly showing their potential as biomarkers.

**1. DNA methylation**

The DNA methylation is nothing but fairly stable epigenetic changes which consist of the covalent binding of a methyl group to the 5' carbon of cytosine. The CpG-rich islands contain 40% of genes and up to 70% of all CpG dinucleotides in the genome are methylated. Regulation of gene expression by changing the transcriptional machinery accessibility to DNA is the significant function of DNA methylation.

High stability of DNA methylation status and it not only serves as a special epigenetic memory of specific cells over a all periods in the cell cycle but also regulate the gene expression and histone codes activity. Acceleration of DNA methylation process accelerated at CpG sites and mediated by DNA methyl transferase enzymes such as DNMT1, DNMT3a, and DNMT3b. S-adenosyl methionine act as an important methyl group donor inside the cell. Folic acid and B12 play the important roles in re-methylation or the attraction of de-methylated form of S-adenosyl methionine via active and passive mechanisms. In the initial years of life level of nutrition could impact on the DNA methylation pattern and level of gene expression in adulthood.
2. Histone modifications:

Histone modification regulating the gene expression via chromatin remodeling and it includes acetylation, methylation, phosphorylation, ribosylation, ubiquitylation, sumoylation, and citrullination. Acetylation is has mediated through five families of mammalian histone acetyltransferase enzymes.\(^{17}\)

Histone are that are essential components of nucleosomes structure in eukaryotic cells it is nothing but a small (11-22 kDa) nuclear proteins hence DNA wraps around an octamer of the core histones H2A, H2B, H3, and H4, it helps in constructing the fundamental unit of chromatin.\(^{18}\) Due to PTMs, histone proteins play an essential role in the regulation of chromatin which control the binding of other proteins called readers. This evidence gives an idea that extracellular histones contribute to human disease because they are cytotoxic for a wide array of human cells and human tissues\(^{19}\) by performing several in vitro and in vivo experiments that showed after binding to phospholipids circulating histone can damage cell membranes, disruption of Ca\(^{2+}\) influx is produced and it leads to the release of other intracellular mediators that results in cell death.\(^{20}\)

3. miRNAs:

miRNAs are a class of conserved, short (18–24nt). At the posttranscriptional level non-coding RNAs that have ability to control the gene expression in distinct biological networks. In recent years miRNAs have potential to be used as biomarkers.\(^{22}\) Recent studies is proposing epigenetic biomarkers mainly based upon circulating miRNAs as biomarkers of various disease conditions [e.g., cardiac disease, Neurodegenerative and psychological disorders, chronic kidney disease and liver disease, among others]. miRNAs are considered as a member of the noncoding RNAs about 17 to 25 nucleotides that can carried out a various biological activities. About 60% of protein-coding genes in human is regulated by the miRNAs,\(^{23}\) and these miRNAs are adjusted epigenetically adjusted by methylation in CpG islands or histone modifications or both of them.\(^{24,25}\)
Factors affecting epigenetics

There are number of factors which impact on epigenetics but mainly five factors which alter the epigenetic mechanism includes

i. Genetic factors
   - Heredity: - The transfer of genes from one generation to the next generation. You inherit your parents' genes is known as heredity. It is helps to make you the person you are today: for example - short or tall, with black hair or blond, with brown eyes or blue.

ii. Dietary factors:
   - Diet also responsible for epigenetic changes. Dietary components have potential to overcome the disease and promote overall individuals health.

iii. Environmental factors:
   - Environmental factors such as nutrition, behavior, stress, physical activity, working habits, smoking and alcohol consumption. Environmental and lifestyle factors may alter the epigenetic mechanisms.

iv. Thought process:
   - A life of happiness and positive thoughts change future generations. Our thoughts and actions can alter the epigenetic mechanism.

   - It’s been shown just thinking about something can cause your brain to release neurotransmitters, chemical messengers that allow it to communicate with parts of itself and your nervous system.

V. Lifestyle:

   - Various lifestyle factors such as nutrition, behavior, stress, physical activity, working habits, smoking and alcohol consumption.

   - These factors leave epigenetic marks on our DNA that leads to the alteration of gene expression.

Epigenetics in Cardiovascular Disease

Recent data associated DNA methylation, as measured in peripheral blood leukocytes, with clinical cardiovascular disease found lower DNA methylation content in peripheral blood leukocytes from patients with atherosclerotic cardiovascular disease. From the aging study it has been found that lower LINE-1 methylation in peripheral blood leukocytes is a predictor of incidence and mortality from ischemic heart disease and stroke.

Global methylation measures gives average estimates of methylation across the genome do not have the resolution needed to target individual genes or gene sequences responsible for cardiovascular disease.

i. Heart failure:

   - Genetic predisposition and multiple environmental factors results in heart failure. In a patient with HF, Movassagh and colleagues reported that three angiogenesis-related genes were discriminatory methylated. In the last stage of due to the altered epigenomics of the three genes may reflect common epigenetic pathways in vasculature and heart remodeling.

   - In case of heart failure histone modifications and Micro-RNAs also have significant importance in epigenetics.

ii. Hypertension:

   - One of the biggest risk factor for cardiovascular disease development is hypertension. Hypertension is not effectively overcome in most of the patients which leads to many cardiovascular complications. Complex interactions between different environmental factors and genes results in hypertension and epigenetic mechanism is implemented to treat the hypertension.

   - Number of factors causing hypertension such as autonomic, renal, vessel diameter/compliance and neuroendocrine.

iii. Congenital Heart Disease

   - Congenital heart disease (CHD) represents a large proportion of significant birth defects, occurring in approximately 8 per 1000 live births. Monogenic causes have been identified but most CHD is considered to be multifactorial with phenotypic differences noted between individuals in the same family or even with the same genetic defect (e.g. trisomy 21 where only ~50% of individuals are born with CHD. A possible role for epigenetics in the etiology of CHD could be the dysregulation of DNA methylation during different stages of embryonic growth that could lead to the inappropriate silencing of tissue-specific gene expression and a resulting increased risk for cardiac malformation.
Epigenetics in liver Disease

Alteration in epigenetics consequently promote a number of human diseases for example fetal, and liver diseases alcohol spectrum disorders, various types of cancer, neuropsychiatric disorders. The second largest organ in the body is liver which performs a various biological functions. The common liver diseases are alcohol related and non-alcohol related liver diseases, viral-mediated liver diseases (Hepatitis A, B and C), fatty liver disease, Chronic liver diseases which needs long-term therapy. Effective treatment and prevention is necessary to reduce the mortality and economic burden caused by the liver diseases worldwide. Epigenetic regulatory mechanisms offer new therapeutic potential to treat these diseases.

i. Hepatocellular Carcinoma

In the world, 250,000 to 1 million annually are only because of hepatocellular carcinoma (HCC) is the fifth most common cancer in the world that causes. There are number of risk factors for hepatocellular carcinoma including diabetes, and obesity, dietary exposure to aflatoxins, hepatitis B virus (HBV) and hepatitis C virus (HCV) infections, smoking, alcohol. Epigenetic changes causes HCC metastasis, DNA methylation, histone modification, invasion, and dissemination encompass noncoding RNAs regulation and all these changes are associated with initiation and progression of hepatocellular carcinoma.

ii. Liver Fibrosis:

Recently it is well documented that cellular phenotype is at least in part under control of chromatin configuration at key regulatory genes, with the help of epigenetic mechanisms. ‘epigenetics’ describes that reversible changes in gene expression that can be inherited via cell division that do not involve alterations of DNA sequence. Epigenetic mechanisms regulate the chromatin structure, modifications and the initiation of transcription and their co-factors that dictate the rate at which a gene is actively transcribed. Here, we consider the epigenetic mechanisms involved in the pathogenesis of liver fibrosis, as well as examine recent advancements, approaches and strategies used in the field of epigenetic for the treatment of liver fibrosis.

iii. Alcoholic Liver Disease:

In recent studies, it has been documented how epigenetic factors can impact on the function and expression of several genes. Alcohol consumption is responsible for 3.8% of global mortality and 4.6% of disability-adjusted life-years (DALYs) lost due to premature death. From the meta-analysis it has been found that risks of mortality is increased from liver cirrhosis among men and women drinking 12-24 g of ethanol per day. Due to the excessive consumption of alcohol it causes exposes the liver cells exposed to a elevated levels of ethanol, which can lead to a class of clinical conditions identified as alcoholic liver disease (ALD).

Identification of epigenetic factors can be important not only for the screening of individuals at risk but also for the study of the epigenetic modification mechanisms underlying ALD.

Epigenetics in kidney diseases

From the emerging evidence recommend that epigenetic changes are involved in the pathogenesis of diseases predisposing to chronic kidney disease (CKD). DNA hypermethylation has been associated with predisposition to, and progression of, atherosclerosis. A recent study DNA methylation levels in more than 14,000 genes between African-American end stage renal disease patients and his panic diabetes it shows that DNA methylation differences were related with disease treatment. By changing the DNA methylation and histone acetylation epigenetic modulations of gene expression that might predispose to hypertension.

Based on the number of experimental studies it describes that local, gene-specific alterations in DNA methylation may play an essential role in the treatment of chronic kidney disease. From some evidence it was recommend that RNA interference is mainly essential for the development and progression of kidney disease.

i. Renal cancer:

Over the last three decades renal cancer rising steadily and currently, it is 9th most common cancer in Europe. The diagnosis of renal cell cancer depends upon early detection of the tumor. Sometimes, clinical signs and symptoms are often not useful in an early diagnosis of cancer, therefore the classic symptoms like haematuria, palpable flank mass and pain and are generally related with advanced stage of the disease. It reported that more malignant renal cancer is generated mainly due to the alterations of DNA methylation in the precancerous kidney cortex tissue. Recently, it is admitted that DNA methylation is the interplay between genetics and epigenetics that leads to tumour genesis and the development of cancer. This interplay have recently been elegantly demonstrated by who identified mutations in genes that encode enzymes, alter chromatin structure and transcriptional control and therefore which demethylate or methylate key lysine residues of histones and it leads to the development of renal cell cancer.

Epigenetic mechanisms can be involved in the treatment of these neoplasms as well.

ii. Chronic Kidney Disease (CKD)

There is considerable evidence showing the role of epigenetic mechanisms in the development of chronic kidney disease. Difference in DNA methylation pattern between normal and diseased kidneys identified in renal fibroblasts is noticed in previous study. Fibroblast multiplication is regulated by DNA methylation and leads to renal fibrogenesis. For example- the study of experimental murine models treated with 5-azacytidine, it results in protection from renal fibrosis. In also demonstrated that hypermethylation of RAS protein activator like-1 was related with the renal fibrogenesis. Various histone modifications in the human kidney have not been assessed with reference to chronic kidney disease.

Future Prospective:

In an individual’s lifetime DNA methylation and histone modifications might help to describes how environmental and lifestyle factors can impose aberrant gene expression patterns. For example, in the pathogenesis of cardiac lesions like, among others, atherosclerosis, arrhythmias, thrombosis and left ventricular dysfunction, DNA and histones play essential roles by reducing the DNA-histone complex may constitute a viable goal for prevention of various cardiac complications. New approaches have brought different clinical trials and have been explored recently targeted for the treatment of cardiovascular disease, liver disease and renal disease. However, a detailed genetic background evaluation helps the physicians to design the best therapeutic approach for each disease. For example - microRNAs involved in the pro-inflammatory switch in macrophages and Kupffer cells will be more efficient and then a steroid-based treatment and will bare less side effects. Moreover, it has to be considered that epigenetic factors can be regulated by environmental factors, i.e. diet and medications, and since represent an ideal genetic therapy target.
Discussion:
At the transcriptional and post-transcriptional levels or at the translation level and post-translational modifications epigenetic mechanisms can influence the gene activity. Potentially broad spectrum effects of epigenetic mechanism can be affected by both genetic and environmental factors could results in more varieties of cell differentiations, morphogenesis, variability, and adaptability of an organism.\(^8\) The epigenetics covers the DNA modifications, histones and DNA-binding proteins, which are important in regulating changes in chromatin structure without any alteration in the sequence of nucleotide in DNA. Also, these alterations could be transferred from one generation to the next generation. The overall studies have shown that exercise and lifestyle modifications including nutritional habits are first protective measures for decreasing the risk of disease.\(^8\) In the pathogenesis of cardiovascular disease, hepatic disease and renal disease Non-coding RNAs were found to be essential and it also offers the possibility of operating as diagnostic and prognostic biomarkers.\(^9\) To improve our understanding of the involvement of lncRNA in regulating gene expression changes underlying heart, liver as well as kidney failure further studies on epigenetics is needed.\(^8\)

Conclusion:
The epigenetics controlling the alteration in gene expression which influence on many different human cardiac, hepatic and renal diseases. Epigenetics helps to explain the complex interactions between genes and environment and understanding of disease by providing a new conceptual framework. Over the life course epigenetic factors potently influence epigenetic variations and changes in gene expression. Epigenetic mechanism in various diseases plays a significant role in controlling the altered gene expression, which is also critical for disease development. These epigenetic modifications may serve as signatures to diagnose the disease and also may provide a promising scope to develop new drug targets to treat these diseases.

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