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# **Recent Advancements in the Study of Cancer**

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# ABSTRACT

A brand-new, developing field of study called clinical bioinformatics combines the fields of omics science, information technology, bioinformatics, and clinical informatics. Despite the obstacles that existed for physicians applying genetic tests, such as their low tolerance for uncertainty, negative attitudes about their responsibility for genetic counselling and testing, and lack of familiarity with the ethical issues raised by testing, clinical physicians still needed to be informed and open to advancements in omics technology at the beginning of the 20th century. In the fight against cancer, understanding epigenetic modifications, particularly the acetylation and deacetylation of histones, is becoming more and more important. The ultimate goal of cancer research is the organisation and analysis of the data, which is always expanding, as well as the creation of new methods for diagnosis or therapy. This article reviews the several efficient bioinformatics methods for cancer research. The essential principles and precepts of the important and popular bioinformatics techniques are introduced.

Keywords: Cancer Research, Interactome, Epigenetics, Bioinformatics, SNP, And Cheminformatics.

## Introduction:

Epigenetic modifications are primarily responsible for a subset of illnesses known as cancer. Epigenetic modifications are changes in genes that affect gene expression and function. The epigenetic mechanism's identification required a lot of investigation. Histone acetylation and deacetylation are among these and play a critical role in the development of cancer. The essential enzymes and regulators in this process, histone deacetylases (HATs) and histone deacetylases (HDACs), have been the focus of substantial study. Since epigenetics is a hallmark of cancer, a substantial amount of data is being generated through a wide range of biological testing and laboratory procedures. In order to provide greater clarity and better decision-making, big data analysis techniques are thus applied to research and development in the field of cancer research. One such strategy is .In order to address problems in the biological sciences, this multidisciplinary area combines information technology (IT) and all branches of the life sciences. There has been great advancement in the development of bioinformatics as a platform for cancer research. The most important bioinformatics technique is omics, which also includes genomes, proteomics, transcriptomics, metabolomics, etc. In this article, we tried to explain the role of bioinformatics and some of its potential applications in the study



# Benefits of Organoid Use in Cancer Research

The unique, biological platform known as the three-dimensional (3D) organoid system is based on cell culture and is physiologically relevant. An organoid is a miniature organ grown in vitro in three dimensions, displaying accurate microanatomy. Organoids are developed and passaged in a basement membrane matrix, which aids in their capacity for self-renewal and differentiation, starting with just one to a few cells extracted from tissue or cultured cellsWith the emergence of the science of stem cell biology in the early 2010s, the method for producing organoids has significantly advanced. Numerous kinds of cancer tissues and cells share the properties of stem, embryonic stem cells (ES cells), or induced pluripotent stem cells (iPS cells) that enable them to produce an organoid in vitro. As a result, ES cells have been used by cancer researchers[35,36].

In recent years, the 3D organoid system has emerged as a powerful tool for fundamental research with the potential to be used to customised therapy. This technique uses living tissue fragments from biopsies, surgically resected specimens, or even frozen tissues to passage dissociated main structures to produce secondary 3D organoids.

#### Method

Starting with straightforward search phrases like "bioinformatics," "cancer research," and "cancer bioinformatics," we used Google Scholar and PubMed (https://www.ncbi.nlm.nih.gov/pubmed) to locate publications that matched our search parameters. The documents on each page were analysed and arranged in accordance with their relevancy, importance, year, and substance. Manual duplicate detection was used to remove them from the final list. Our paper is organised as follows: We begin by providing a summary of some important bioinformatics methods, then we examine their importance in cancer research and go into how to employ them. A conclusion then follows.

## **Microbiome Research in Cancer**

The human body is home to many microorganisms. These microorganisms consist of fungus, viruses, and bacteria. Of all of them, bacteria have the closest connection to the human body. Bacteria can exist anywhere in the human body, including the respiratory system, oral cavity, and digestive tract. There may be 1000 different varieties of bacteria in the digestive tract, and there are more than 100 trillion distinct bacterial cells there. The term "intestinal flora" refers to the general population of different microorganisms found in human intestines. The phrases "microbiota" and "microbiome" have also gained popularity recently[34].

#### Study of the interactome and pathways:

The interactome refers to the network of genes and proteins in a cell[8]. Studies on protein-protein interactions (PPIs) reveal the underlying molecular pathways of the illness and identify its exact aetiology. This makes it possible to design hub genes for the gene responsible for the growth of cancer. For the purpose of determining networks' therapeutic usefulness and networking, it is imperative to understand networks correctly in light of all these elements. By using Interactome and pathway analysis, researchers can identify several genes that were differentially expressed in clinical samples and help in the identification of corresponding biomarkers[9].

#### **Biological information structure**

The prediction and analysis of the structures of biological molecules like DNA, RNA, and proteins is known as structural bioinformatics. By comparing database structures and validating them, this method allows the function of molecules to be deduced from their sequence or structural details[10]. As a result, homology modelling has gained widespread acceptance as a technique for clarifying the theoretical model of molecules, particularly proteins. The use of this for receptor-drug interaction studies would enable further validation

#### Drug discovery and cheminformatics:

Designing and producing efficient treatment components for cancer can be challenging. Cheminformatics is a modern discipline that examines the intricate structures of chemical compounds to identify those with the potential to be medicinal molecules. Pharmaceutical companies and medical researchers can produce potential cancer medicines with the use of these drug discovery techniques[11]. Adsorption, distribution, metabolism, and excretion (ADME), lipinski's rule analysis, and additional wet-lab biological testing, where only actual trials are conducted, are however necessary for further validation of compounds found through docking, dynamics, and quantitative structure-activity relationship (QSAR) studies. Drug development is a crucial area, says QSAR[12]For clinical testing and drug validation, ADME testing is essential[13], and Lipinski's rule determines if a molecule is orally active or not using a set of rules[14].

S.No	Method	Softwares/databases
1	GWASs (genome-wide association studies)	METAL[16], PLINK/GPLINK[15], GWAMA[17], and
		MANTRA[18].
2	Analysis using phylogenetics	Clustalw/X[19], Phylip[20], MEGA[21], BEAST[22],
		PAUP[23].
3	Study of the interactome and pathways	IntAct[24], PANTHER[25], KEGG[26], STRING[27],
		BioGrid[28].
4	Biological information structure	Modelers SWISS-MODEL[29], Phyre2[30], PDB[31], and
5	Drug discovery and cheminformatics	BioVia DS visualisation (Biovia, 2016), Schrodinger[33],
		Patchdock[34],

Table 1: Overview of the important bioinformatics methods and list of available softwares/databases

#### Artificial intelligence battles cancer

Partners of the World Economic Forum are transforming cancer care in India by utilising cutting-edge technology like artificial intelligence (AI) and machine learning. For instance, AI-based risk profiling can assist in cancer screening, resulting in early diagnosis of common malignancies like breast cancer. In situations when imaging specialists might not be available, AI technology can potentially be utilised to examine X-rays and spot tumours. The Centre for Fourth Industrial Revolution of the World Economic Forum India seeks to speed cancer interventions, including these two.

# **Discussion:**

The results of this study outline how bioinformatics is used in the study of cancer. In addition to what we've already mentioned, research indicates that emerging disciplines like systems biology and precision medicine are creating areas that will have a big impact on the development of cancer research. A large dataset of gene expression data might be generated utilising novel sequencing techniques, which could save costs and time spent on research while enhancing its results.

#### **References:**

1. Zaidi SK, Young DW, Montecino M, Lian JB, Stein JL, van Wijnen AJ, et al. Architectural epigenetics: mitotic retention of mammalian transcriptional regulatory information. Mol Cell Biol. 2010;30:4758–66.

2. Hirschhorn JN, Daly MJ. Genome-wide association studies for common diseases and complex traits. Nat Rev Genet. 2005;6:95–108.

3. Easton DF, Eeles RA. Genome-wide association studies in cancer. Hum Mol Genet. 2008;17:R109–15.

4. He Q, He Q, Liu X, Wei Y, Shen S, Hu X, et al. Genome-wide prediction of cancer driver genes based on SNP and cancer SNV data. Am J Cancer Res. 2014;4:394–410.

5. Lake JA, Moore JE. Phylogenetic analysis and comparative genomics. Trends Biotechnol. 1998;16:22-3.

6. Brown D, Smeets D, Szekely B, Larsimont D, Szasz AM, Adnet P-Y, et al. Phylogenetic analysis of metastatic progression in breast cancer using somatic mutations and copy number aberrations. Nat Commun. England; 2017;8:14944.

7. Somarelli JA, Ware KE, Kostadinov R, Robinson JM, Amri H, Abu-Asab M, et al. PhyloOncology: Understanding cancer through phylogenetic analysis. BiochimBiophysActa. 2017;1867:101–8.

8. Vidal M, Cusick ME, Barabási A-L. Interactome Networks and Human Disease. Cell. 2011;144:986–98.

9. Coulombe B. Mapping the Disease Protein Interactome: Toward a Molecular Medicine GPS to Accelerate Drug and Biomarker Discovery. J Proteome Res. 2011;10:120–5.

10. Chandra N, Anand P, Yeturu K. Structural bioinformatics: deriving biological insights from protein structures. Interdiscip Sci. Germany; 2010;2:347-66.

11. Begam BF, Kumar JS. A Study on Cheminformatics and its Applications on Modern Drug Discovery. Procedia Eng. 2012;38:1264–75.

12. Gini G. QSAR Methods. Methods Mol. Biol. 2016. p. 1–20.

13. Vugmeyster Y, Harrold J, Xu X. Absorption, Distribution, Metabolism, and Excretion (ADME) Studies of Biotherapeutics for Autoimmune and Inflammatory Conditions. AAPS J. Boston; 2012;14:714–27.

14. Pollastri MP. Overview on the Rule of Five. CurrProtocPharmacol. 2010; Chapter 9: Unit 9.12.

15. Purcell S, Neale B, Todd-Brown K, Thomas L, Ferreira MAR, Bender D, et al. PLINK: A Tool Set for Whole-Genome Association and Population-Based Linkage Analyses. Am J Hum Genet. 2007;81:559–75.

16. Willer CJ, Li Y, Abecasis GR. METAL: fast and efficient meta-analysis of genomewide association scans. Bioinformatics. 2010;26:2190-1.

17. Mägi R, Morris AP. GWAMA: software for genome-wide association meta-analysis. BMC Bioinformatics. 2010;11:288.

18. Morris AP. Transethnic meta-analysis of genomewide association studies. Genet Epidemiol. 2011;35:809-22.

19. Thompson JD, Higgins DG, Gibson TJ. CLUSTAL W: improving the sensitivity of progressive multiple sequence alignment through sequence weighting, position-specific gap penalties and weight matrix choice. Nucleic Acids Res. 1994;22:4673–80.

20. Retief JD. Phylogenetic analysis using PHYLIP. Methods Mol Biol. 2000;132:243-58.

21. Kumar S, Nei M, Dudley J, Tamura K. MEGA: A biologist-centric software for evolutionary analysis of DNA and protein sequences. Brief Bioinform. 2008;9:299–306.

22. Drummond AJ, Rambaut A. BEAST: Bayesian evolutionary analysis by sampling trees. BMC Evol Biol. 2007;7:214.

23. Wilgenbusch JC, Swofford D. Inferring evolutionary trees with PAUP. CurrProtocBioinforma. 2003;Chapter 6:Unit 6.4.

24. Kerrien S, Alam-Faruque Y, Aranda B, Bancarz I, Bridge A, Derow C, et al. IntAct—open source resource for molecular interaction data. Nucleic Acids Res. 2007;35:D561–5.

25. Mi H, Muruganujan A, Casagrande JT, Thomas PD. Large-scale gene function analysis with the PANTHER classification system. Nat Protoc. 2013;8:1551.

26. Kanehisa M, Goto S. KEGG: Kyoto Encyclopedia of Genes and Genomes. Nucleic Acids Res. 2000;28:27-30.

27. Szklarczyk D, Morris JH, Cook H, Kuhn M, Wyder S, Simonovic M, et al. The STRING database in 2017: quality-controlled protein–protein association networks, made broadly accessible. Nucleic Acids Res. 2017;45:D362–8.

28. Chatr-aryamontri A, Oughtred R, Boucher L, Rust J, Chang C, Kolas NK, et al. The BioGRID interaction database: 2017 update. Nucleic Acids Res. 2017;45:D369–79.

29. Biasini M, Bienert S, Waterhouse A, Arnold K, Studer G, Schmidt T, et al. SWISS-MODEL: modelling protein tertiary and quaternary structure using evolutionary information. Nucleic Acids Res. 2014;42:W252-8.

30. Kelley LA, Mezulis S, Yates CM, Wass MN, Sternberg MJE. The Phyre2 web portal for protein modelling, prediction and analysis. Nat Protoc. 2015;10:845–58.

31. Berman HM, Westbrook J, Feng Z, Gilliland G, Bhat TN, Weissig H, et al. The Protein Data Bank. Nucleic Acids Res. 2000;28:235-42.

32. Eswar N, Webb B, Marti-Renom MA, Madhusudhan MS, Eramian D, Shen M-Y, et al. Comparative protein structure modeling using Modeller. CurrProtocBioinforma. 2006;Unit-5.6.

33. Bhachoo J, Beuming T. Investigating Protein-Peptide Interactions Using the Schrodinger Computational Suite. Methods Mol Biol. 2017;1561:235–54.

34. Schneidman-Duhovny D, Inbar Y, Nussinov R, Wolfson HJ. PatchDock and SymmDock: servers for rigid and symmetric docking. Nucleic Acids Res. 2005;33:W363-7.