

LIST OF
PUBLICATIONS

LIST OF PUBLICATIONS

- 1) **Anukriti**, Dhasmana A., Uniyal S., Somvanshi P., Bhardwaj U., Gupta M., Haque S., Lohani M., Kumar D., Ruokolainen J., Kesari KK., (2019) Investigation of Precise Molecular Mechanistic Action of Tobacco-Associated Carcinogen ‘NNK’ Induced Carcinogenesis: A System Biology Approach. Genes, 10, 564; doi:10.3390/genes10080564. PMID: PMC6723528. (**Impact factor : 3.759**)
- 2) **Anukriti**, Dhasmana A., Hobani YH., Farasani A., Habibullah M., Alshammary FL., Khan S., Haque S., Lohani M. (2020) Tobacco Smoke Carcinogens Induce DNA Repair Machinery Function Loss: Protection by Carbon Nanotubes. (Impact Factor: 1.583).

OTHER PUBLICATIONS

- 1) **Anukriti**, Uniyal S, Dhasmana A, Gupta M, Kesari KK, Jamal QMS and Lohani M. System Network Biology Approaches in Exploring of Mechanism Behind Mutagenesis. Networking of Mutagens in Environmental Toxicology, Environmental Science, https://doi.org/10.1007/978-3-319-96511-6_6.
- 2) Dhasmana A., Uniyal S., **Anukriti.**, Kashyap VK., Somvanshi P., Gupta M., Bhardwaj U., Jaggi M., Yallapu MM., Haque S., Chauhan SC., (2020) Topological and system-level protein interaction network (PIN) analyses to deduce molecular mechanism of curcumin. Scientific Reports 10(1):12045; DOI: 10.1038/s41598-020-69011-0. (**Impact Factor: 4.120**)

CERTIFICATES




3rd International Conference
On
“*Nutraceuticals and Chronic Diseases-2018*”
(INCD-2018)


Certificate of Appreciation


This is to certify that *Anukriti* has **delivered oral presentation / presented poster / participated** at the Third International conference on “*Nutraceuticals and Chronic Diseases*” (INCD-2018).

Conference accredited from Uttarakhand Medical Council for 10.0 Hours.

Jointly organized by Swami Rama Himalayan University (Dehradun, India), International Society of Nutraceuticals & Chronic Diseases and University of Tennessee Health Science Center (USA), in Dehradun, India, from 14th-16th September, 2018.


Patron
Bharat B. Aggarwal


Organizing President
Subhash C. Chauhan


Organizing Chairman
Sunil Saini



MIT School of
**Bioengineering Sciences
 & Research**
 Breathe Life into Engineering Career!



**MIT-ADT
 UNIVERSITY**
 PUNE, INDIA
 A Leap Towards The World Class Education

3rd International Conference on Recent Trends in Bioengineering (ICRTB 2020)

January 31 - February 01, 2020

CERTIFICATE

This is to certify that Dr/Mr/Ms

Anukriti

from Swami Rama Himalayan University presented a poster/paper/Model entitled

Investigation of systems network pattern of NNK and BaP (Tobacco carcinogens) induced cell perturbation; guarding by carbon monoxide
 in ICRTB 2020 organised by MIT School of Bioengineering Sciences & Research at MIT-ADT University, Pune, India from January 31

- February 01, 2020. The organizing committee acknowledges your efforts and participation.

Renu Vyas

Dr. Renu Vyas
 Convener
 Organizing Committee ICRTB 2020

Vinayak Ghalsas

Prof. Vinayak Ghalsas
 Director
 MIT School of Bioengineering Sciences & Research

Mangesh Karad

Mangesh Karad
 Executive President
 MIT ADT University

Vishwanath D. Karad

Hon. Prof. Dr. Vishwanath D. Karad
 Founder President, MIT Group of Institutions
 President, MIT ADT University



Certificate of Participation



This is to certify that

Anukriti

has actively participated in

14th Uttarakhand State Science and Technology Congress 2019-2020
27th- 29th February, 2020

presented a research paper (Oral/Poster) entitled

Investigation of systems network pattern of NNK (Tobacco carcinogen) induced cell cycle perturbation; Guarding by carbon nanometerals
under the discipline *carbon nanometerals*

Biotechnology, Biochemistry & Microbiology

at Uttarakhand Council for Science and Technology, Vigyan Dham, Dehradun.

Ashutosh

Dr Ashutosh Mishra
Organizing Secretary
14th USSTC
Council for Science and Technology
Uttarakhand

Rajendra

Dr Rajendra Dobhal FNASc
Chairman - 14th USSTC
Director General
Council for Science and Technology
Uttarakhand





3rd International Conference
On
“Nutraceuticals and Chronic Diseases-2018”
(INCD-2018)

Certificate of Appreciation

This is to certify that*Anukrite*..... has delivered *Oral Presentation/ Presented Poster in the category of Young/Budding Scientist* and scored.....*Third*..... at the Third International conference on **“Nutraceuticals and Chronic Diseases” (INCD-2018)**.

Conference accredited from Uttarakhand Medical Council for 10.0 Hours.

Jointly organized by Swami Rama Himalayan University (Dehradun, India), International Society of Nutraceuticals & Chronic Diseases and University of Tennessee Health Science Center (USA), in Dehradun, India, from 14th -16th September, 2018.

Patron
Bharat B. Aggarwal


Organizing President
Subhash C. Chauhan

Organizing Chairman
Sunil Saini

FULL LENGTH
PUBLISHED
PAPER

Article

Investigation of Precise Molecular Mechanistic Action of Tobacco-Associated Carcinogen ‘NNK’ Induced Carcinogenesis: A System Biology Approach

Anukriti ¹, Anupam Dhasmana ¹, Swati Uniyal ², Pallavi Somvanshi ³ , Uma Bhardwaj ⁴, Meenu Gupta ⁵, Shafiul Haque ⁶, Mohtashim Lohani ⁷, Dhruv Kumar ⁸, Janne Ruokolainen ⁹ and Kavindra Kumar Kesari ^{9,*}

¹ Department of Biosciences, Himalayan Institute of Medical Sciences, Swami Rama Himalayan University, Dehradun 248016, India

² School of Biotechnology, Gautam Buddha University, Greater Noida 201310, India

³ Department of Biotechnology, TERI School of Advanced Studies, 10, Institutional Area, Vasant Kunj, New Delhi 110070, India

⁴ Research and PhD Cell, Swami Rama Himalayan University, Dehradun 248016, India

⁵ Department of Radiotherapy, Cancer Research Institute, Himalayan Institute of Medical Sciences, Swami Rama Himalayan University, Dehradun 248016, India

⁶ Research and Scientific Studies Unit, College of Nursing and Allied Health Sciences, Jazan University, Jazan 45142, Saudi Arabia

⁷ Department of EMS, College of Applied Medical Sciences, Jazan University, Jazan 45142, Saudi Arabia

⁸ Amity Institute of Molecular Medicine & Stem Cell Research, Amity University, Noida, Uttar Pradesh 201303, India

⁹ Department of Applied Physics, Aalto University, 00076 Espoo, Finland

* Correspondence: kavindra.kesari@aalto.fi; Tel.: +358-4514-72943

Received: 14 June 2019; Accepted: 24 July 2019; Published: 26 July 2019



Abstract: Cancer is the second deadliest disease listed by the WHO. One of the major causes of cancer disease is tobacco and consumption possibly due to its main component, 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK). A plethora of studies have been conducted in the past aiming to decipher the association of NNK with other diseases. However, it is strongly linked with cancer development. Despite these studies, a clear molecular mechanism and the impact of NNK on various system-level networks is not known. In the present study, system biology tools were employed to understand the key regulatory mechanisms and the perturbations that will happen in the cellular processes due to NNK. To investigate the system level influence of the carcinogen, NNK rewired protein–protein interaction network (PPIN) was generated from 544 reported proteins drawn out from 1317 articles retrieved from PubMed. The noise was removed from PPIN by the method of modulation. Gene ontology (GO) enrichment was performed on the seed proteins extracted from various modules to find the most affected pathways by the genes/proteins. For the modulation, Molecular COMplex DETection (MCODE) was used to generate 19 modules containing 115 seed proteins. Further, scrutiny of the targeted biomolecules was done by the graph theory and molecular docking. GO enrichment analysis revealed that mostly cell cycle regulatory proteins were affected by NNK.

Keywords: NNK; cancer; systems biology; protein–protein interaction network; topological analysis; gene ontology

1. Introduction

Cancer is one of the major non-communicable diseases [1] and is accountable for millions of deaths per year worldwide. According to the World Health Organization (WHO), cancer is the second major cause of morbidity, with an estimate of 9.6 billion deaths in 2018 [2]. Cancer is a multistage process caused by aberrations in the cellular processes. Cancer is not only caused by mutation in any single gene but also by the accumulation of mutations in multiple genes, a phenomenon described as ‘oncogene addiction’ [3]. According to the WHO, there are mainly three reasons that lead to these aberrations, with tobacco consumption heading the list, which is single-handedly responsible for around 22% of deaths by cancer globally [4].

Currently, we have immense information on how tobacco consumption has direct implications in cancer, specially lung, head and neck, stomach, liver, and pancreatic cancers [5,6]. 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanone (NNK) is one of the main components in tobacco that plays a major role in the causation of cancer [6]. NNK and its derivative, 4-(methylnitrosamino)-1-(3-pyridyl)-1-butanol (NNAL), binds with the DNA and forms DNA adducts, the resultant of which may lead to genetic mutations followed by the deregulation of normal cellular processes [6,7].

NNK is not just responsible for causing cancer but also holds serious implications in other diseases as well. Earlier studies have shown that NNK has significant impact on steatohepatitis [8], Alzheimer’s disease [9], and tuberculosis [10], for example.

In this study, attempts have been made to exploit the system biology approach for an investigation of the overall impact of tobacco-generated carcinogen NNK on molecular systems.

Systems biology is an interdisciplinary field, which is a combination of mathematics, computer science, and biology [11]. Systems biology holds importance as it helps in getting a holistic view of the connections of biomolecules. It provides an anti-reductionist approach towards the involvement of different biomolecular components in a variety of biological systems. It focuses on the development of interactome of the affected targets and then analyzing it by using mathematical models [12]. Graph theory is the core assessment tool for the topological analysis of the interactome that aims to identify hub biomolecular targets based on their clustering coefficients, degrees, and betweenness centralities. Whereas, clustering and gene ontology (GO) enrichment analysis categorize the biomolecules on the basis of functions to procure more promising insight into complex networks. This study aims to find the most probable biomolecular targets of tobacco-associated carcinogen NNK along with its interactions and associated pathways that get perturbed by various cellular mechanisms and lead to cancer development. The most probable key targets of NNK are identified on the basis of their bottleneck scores and based on their thermodynamic interactions with NNK calculated by molecular docking simulations.

2. Materials and Methods

The full methodology scheme is mentioned in Figure 1.

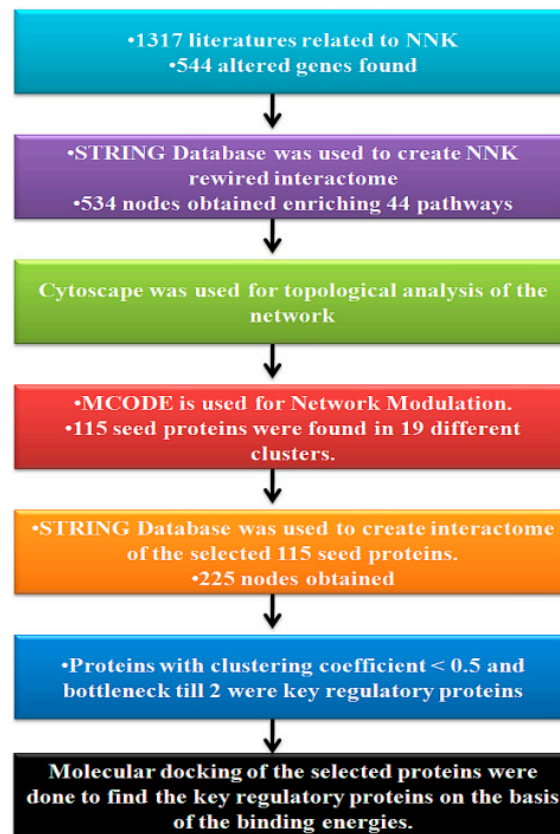


Figure 1. Schematic diagram of the adopted methodology.

2.1. Construction and Visualization of Protein–Protein Interaction Network

In total, 544 biomolecular targets were found to be affected by NNK in approximately 1320 studies using PubMed. A protein–protein interaction network was developed using the STRING database version 10.5 [13]. The network was evidence-based and developed with the highest confidence level score of 0.9, having 50 interactors in the first as well as second shell.

2.2. Protein–Protein Interaction Network (PPIN) Analysis

The Cytoscape Software (version 3.6.1) program [14] was used for the analysis of the protein–protein interaction network (PPIN) to generate protein interaction networks. Network analyzer, an in-built plugin of Cytoscape, was used to analyze the topological properties of an NNK modulated PPIN [15]. The topological properties of any network provide a deep insight into complex biological networks [16,17]. The topological analysis also helps in reducing noise in the data and offers reliable information regarding the network [18]. Node properties, like degree distribution, shortest path length, average clustering coefficient, betweenness centrality, and closeness centrality, were also analyzed.

2.3. Protein Interaction Network Modular Analysis and Pathway Enrichment

Clusters or modules are closely connected nodes in a network that come together and form a dense sub-network [19]. The analysis of clusters or modules helps in attaining detailed information about PPIN. Molecular COMplex DETection (MCODE) is a plug-in available in the Cytoscape software program, which was used for the cluster analysis. The clusters are scored on the basis of size and density—a high score means a big and dense cluster—while gene ontology (GO) serves the purpose of validating the cluster that belongs to a specific function [20]. Thus, the enrichment of clusters helps in enriching the pathways by providing an additional number of external genes that are not present in the dataset. GO analysis provides detailed information about the biological process underneath that

cluster. For GO functional enrichment analysis, ClueGO (version 2.5.1) plug-in of Cytoscape software was used [21]. The analysis was done using a threshold p value < 0.05 . A two-sided hypergeometric test was used for the statistical analysis along with the Bonferroni correction method, in case applied.

2.4. Molecular Docking Analysis

Molecular docking is one of the most preferred methods to find the orientation of two molecules when they form a complex. Docking simulation also explores the thermodynamic stability of the complexes by providing information regarding the binding energy and inhibition constant (K_i) value and helps in finding the best binding modes or orientations of a ligand with its biomolecule. The docking parameters used were based on the studies published by [22]. Autodock 4.0 MGL suite [23] was used for docking simulations. The simulations were performed on AMD E1-6015 APU processor, CPU 1.4 GHz and 4 GB RAM of Hewlett-Packard (HP) machine.

3. Results

3.1. Construction of the Network

In total, 544 biomolecular targets were found to be affected by NNK interaction through a literature survey. A protein–protein interaction network (PPIN) was developed using the STRING database version 10.5. The network developed at the 0.9 confidence level score and 50–50 interactors in the first and second shell comprised of 534 nodes and 2909 edges. The average node degree was 10.09 and average local clustering coefficient was 0.501. The PPI enrichment p value was $< 1 \times 10^{-16}$. Figure 2 represents the NNK rewired protein–protein interaction network having 534 nodes and 2909 edges. Figure 3A,B depicts the number of biomolecular targets involved in various processes and pathways, respectively.

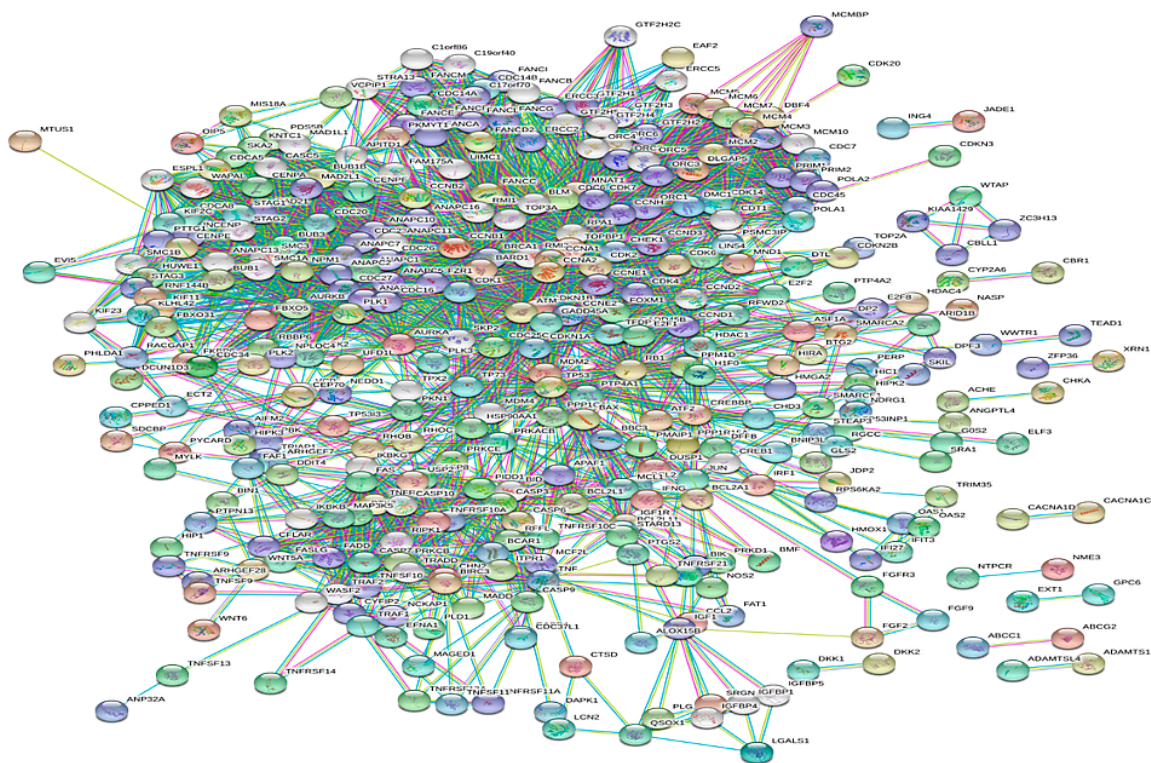
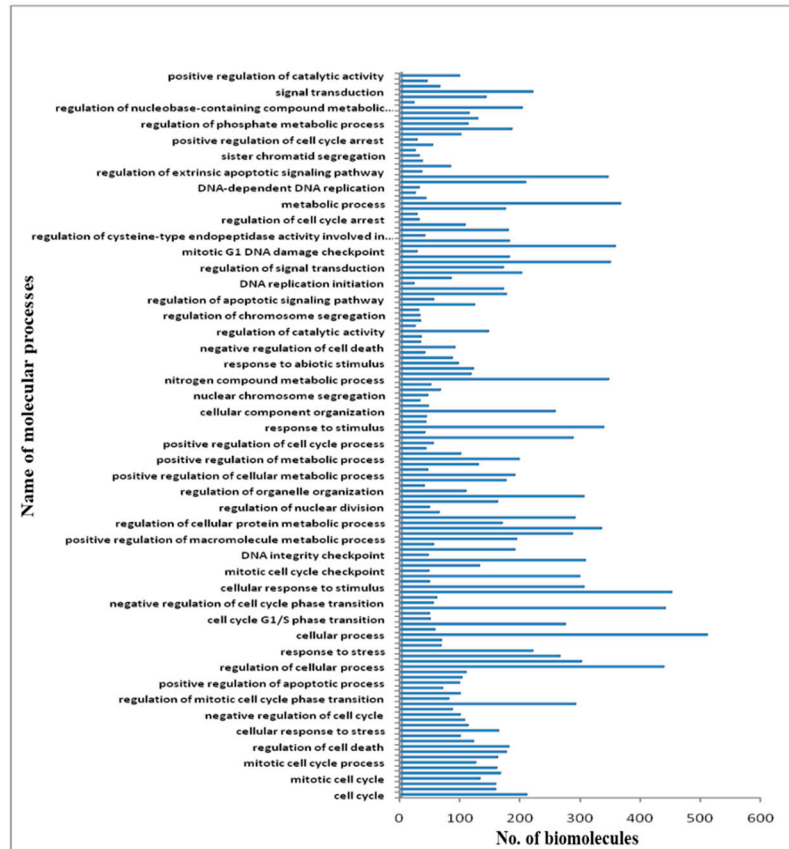
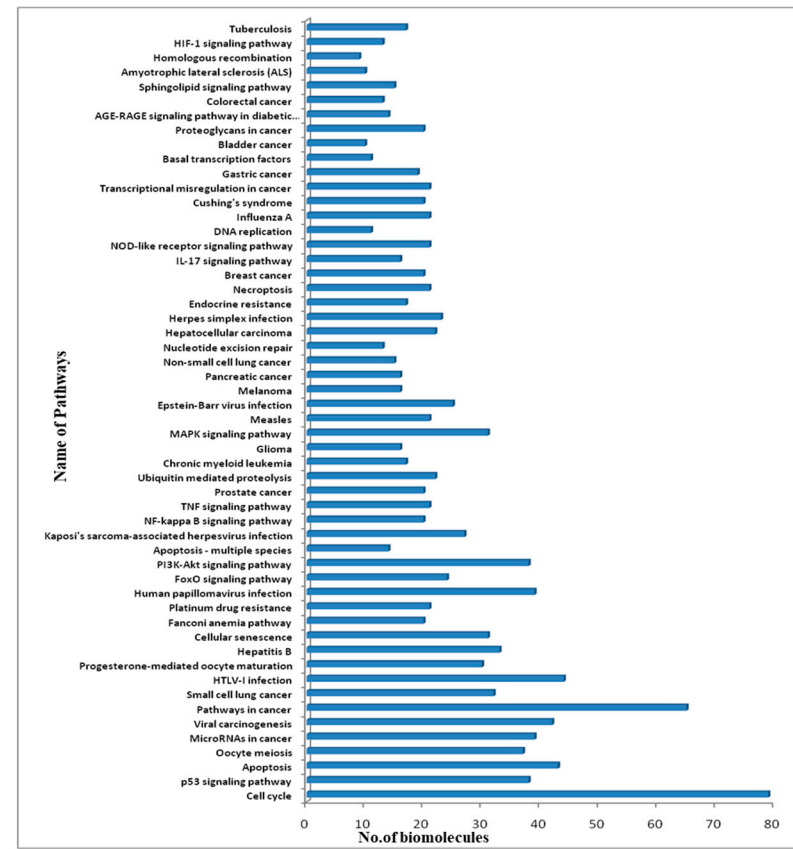


Figure 2. STRING generated NNK rewired protein–protein interaction network with 534 nodes and 2909 edges.



A



B

Figure 3. (A) Processes enriched by NNK rewired PPIN. (B) Pathways enriched by NNK rewired PPIN.

3.2. Topological Properties of the Network

The protein–protein interaction network developed was further analyzed using Cytoscape software. The topological properties of the PPIN calculated with the help of Network Analyzer plug-in were the shortest path length average, neighborhood connectivity distribution, clustering coefficient, node degree distribution, betweenness centrality, closeness centrality, etc. The shortest path length in any network depicts the shortest communication mode between two nodes. Figure 4 shows a graphical representation of path length distribution 2971. This means that at the 2971-unit path length, the information is being passed on at the highest frequency. The degree of a node describes the connectivity of a node with other nodes; it is the total number of links, which are either reaching or starting from that node [24]. The node degree distribution (Figure 5) is one of the most important topological properties of a network. It fits a power law that indicates the presence of hubs in the network [25]. The nodes, which lie close to the power line and have a higher degree, can be considered as the hubs. The average neighborhood connectivity distribution stands for the average number of neighbors, which was observed as 15,940 in this study (Figure 6). Figure 6 shows the average connections of each node with its neighbors. The above parameter helps in understanding the density of a network. The clustering coefficient of a network depicts the tendency of a graph to be divided into clusters [26]. The local clustering coefficient is the number of edges around a particular node, whereas the average clustering coefficient is the clustering coefficient of the whole network [25], and in this study, the average clustering coefficient in this network was found to be 0.597 (Figure 7).

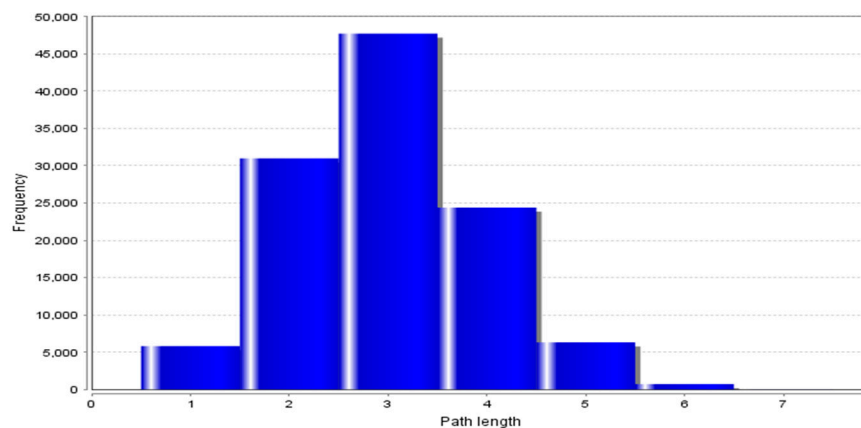


Figure 4. Characteristic path length distribution: 2971.

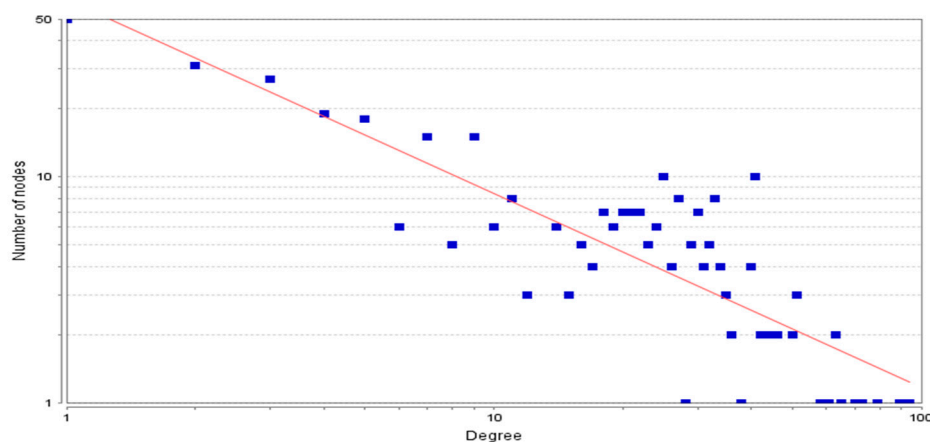


Figure 5. Node degree distribution following power law fitting $y = 61,323x^{-0.861}$ (R-squared 0.722).

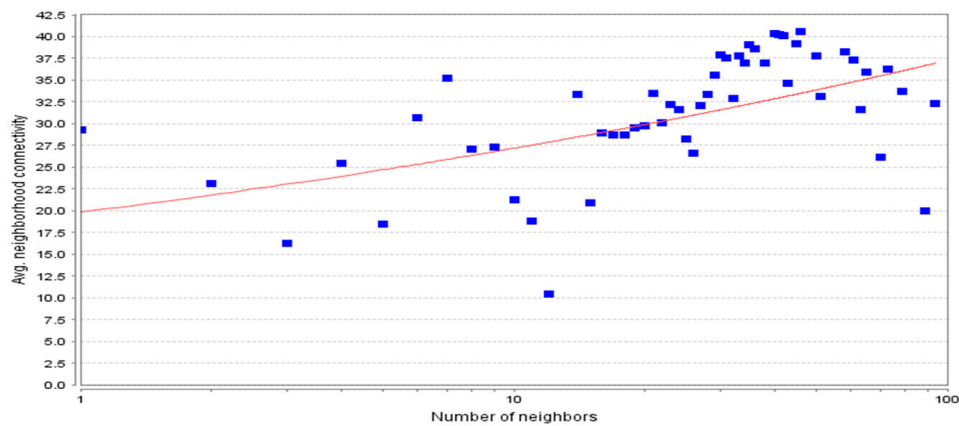


Figure 6. Average (Avg.) neighborhood connectivity distribution following power law fitting $y = 19,838x^{0.137}$ (R-squared 0.253). The average number of neighbors is 15,940.

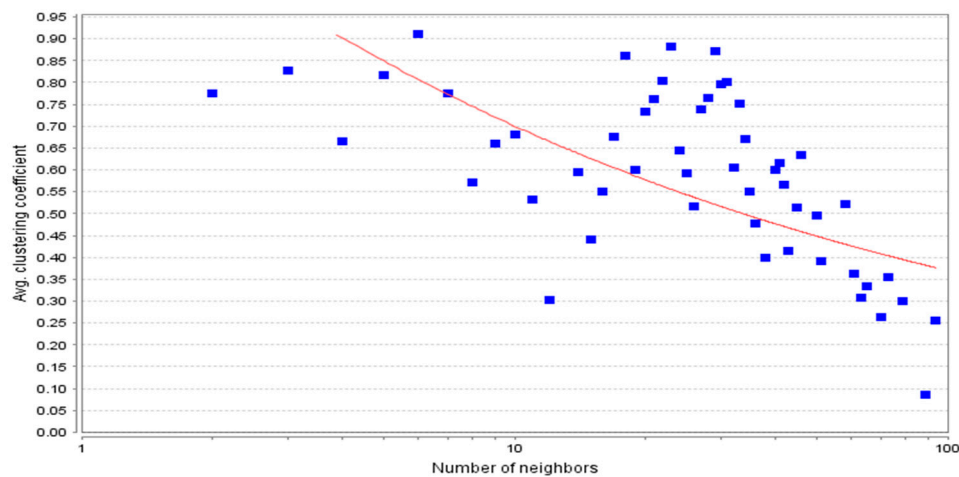


Figure 7. Average (Avg.) clustering coefficient following power law fitting $y = 1.326x^{-0.277}$ (R-squared 0.329). Clustering coefficient distribution of 0.597.

3.3. Clustering and GO Enrichment Analysis

For modular analysis and pathway enrichment, the MCODE plug-in of the Cytoscape was used. Modules or clusters were created from the network and were scored on the basis of their size and density. A high score depicts a denser and tighter cluster. Formation of clusters also helps in the reduction of the noise and getting a better understanding of the genes involved in the clusters. Figure 8 shows the clusters generated by MCODE. The nodes in red color represent the seed proteins, and the yellow nodes are the connectors. The seed proteins are the proteins that were reported earlier to get either upregulated or downregulated by the action of NNK and connector proteins are the proteins that are associated with the seed proteins in the transfer of information, but are not reported to have any direct relation with NNK. The seed proteins were checked in all the clusters and are presented in bold in Table 1, while the remaining proteins (non-highlighted) are the connectors. The cluster that was ranked first had the highest score of 29,862, with 30 nodes and 433 edges. Moreover, during analysis, it was found that cluster 1 had 19 seed proteins and 11 connectors and the overall analysis of the entire cluster found 115 seeds and 88 connectors.

Table 1. Generated clusters with MCODE scores, seeds (bold letters), and connector (non-bold letters) proteins.

Cluster	Score	Nodes	Edges	Seeds	Connectors	Node IDs
1	29.862	30	433	19	11	PLK1 , CASC5 , BUB3 , CDCA8 , STAG1 , BUB1B , CCNB2 , CCNB1 , BUB1 , ESPL1 , CENPF , KIF2C , CDCA5 , MAD2L1 , CDK1 , PDS5B , WAPAL , APITD1 , SKA2 , RAD21 , SMC1A , KNTC1 , CDC20 , CENPA , CENPE , SMC3 , STAG2 , MAD1L1 , AURKB , INCENP
2	19.3	41	386	22	19	CDKN1A , FBXO31 , CDC6 , CDK2 , RNF144B , KLHL42 , CCNA1 , DBF4 , CDC7 , MCM4 , POLA2 , PRIM2 , CDC34 , POLA1 , SKP2 , ORC3 , MCM2 , MCM5 , ORC2 , ORC6 , MCM6 , MCM3 , ORC4 , CHEK1 , AURKA , PTTG1 , CDC45 , PRIM1 , MCM10 , CDT1 , ORC1 , HUWE1 , ORC5 , RBBP6 , NEK2 , MCM7 , ANAPC16 , ANAPC13 , CDKN1B , RPA1 , CCNA2
3	14.652	47	337	9	38	ERCC2 , E2F1 , FBXO5 , CDC23 , ANAPC5 , FANCE , FANCC , FANCG , BLM , ANAPC4 , ANAPC1 , BRCA1 , RB1 , FANCF , FANCL , C17orf70 , CCNE1 , C19orf40 , FANCD2 , FANCI , ANAPC10 , FANCM , FANCB , GTF2H5 , CCND2 , FZR1 , CDC26 , RMI1 , TOP3A , FANCA , GTF2H4 , ANAPC11 , CDC27 , ANAPC2 , GTF2H2 , ANAPC7 , GTF2H1 , STRA13 , CDC16 , CCNH , C1orf86 , ERCC3 , ATM , CCND3 , BARD1 , GTF2H3 , CCND1
4	10.762	22	113	13	9	FAS , RIPK1 , CASP8 , IKBKKG , BID , HDAC1 , TNFRSF10B , TRADD , TP53 , TFDP1 , TRAF2 , TNFSF10 , FADD , FASLG , TNFRSF10A , CASP10 , E2F2 , TOPBP1 , IKBKB , CDK4 , CCNE2 , CDK6
5	7	7	21	7	0	BBC3 , BCL2A1 , BCL2L11 , BCL2 , MCL1 , PMAIP1 , BCL2L1
6	5	5	10	2	3	IGFBP1 , LGALS1 , IGFBP4 , QSOX1 , IGFBP5
7	4.333	7	13	5	2	NCKAP1 , WASF2 , CYFIP2 , PRKCA , PTK2 , PRKCB , BCAR1
8	4	4	6	4	0	H1F0 , HMG2A , ASF1A , HIRA
9	4	4	6	1	3	KIAA1429 , WTAP , CBLL1 , ZC3H13
10	3.333	4	5	3	1	EAF2 , GTF2H2C , CDK7 , MNAT1
11	3.333	4	5	4	0	OAS1 , IFI27 , OAS2 , IFIT3
12	3.333	7	10	5	2	STAG3 , CEP70 , FKBP6 , SMC1B , NEDD1 , HSP90AA1 , TPX2
13	3	3	3	3	0	MAP3K5 , TRAF1 , BIRC3
14	3	3	3	3	0	PSMC3IP , MND1 , DMC1
15	3	3	3	3	0	OIP5 , MIS18A , NPM1
16	3	3	3	3	0	TNFRSF11A , TNFRSF12A , TNFSF11
17	3	3	3	3	0	RHOB , RHOC , CDC25C
18	3	3	3	3	0	SDCBP , PYCARD , CPPED1
19	3	3	3	3	0	FGF2 , FGFR3 , FGF9

Nodes in “bold text” represent the seed proteins and the nodes in “normal” text represent the connector proteins.

The seed proteins were identified in each cluster. Thereafter, the PPIN was generated using the STRING database with 0.9 confidence level, where 50 interactors in the first and another 50 in the second shell were recorded. The PPIN generated now was further enriched using the ClueGO plug-in.

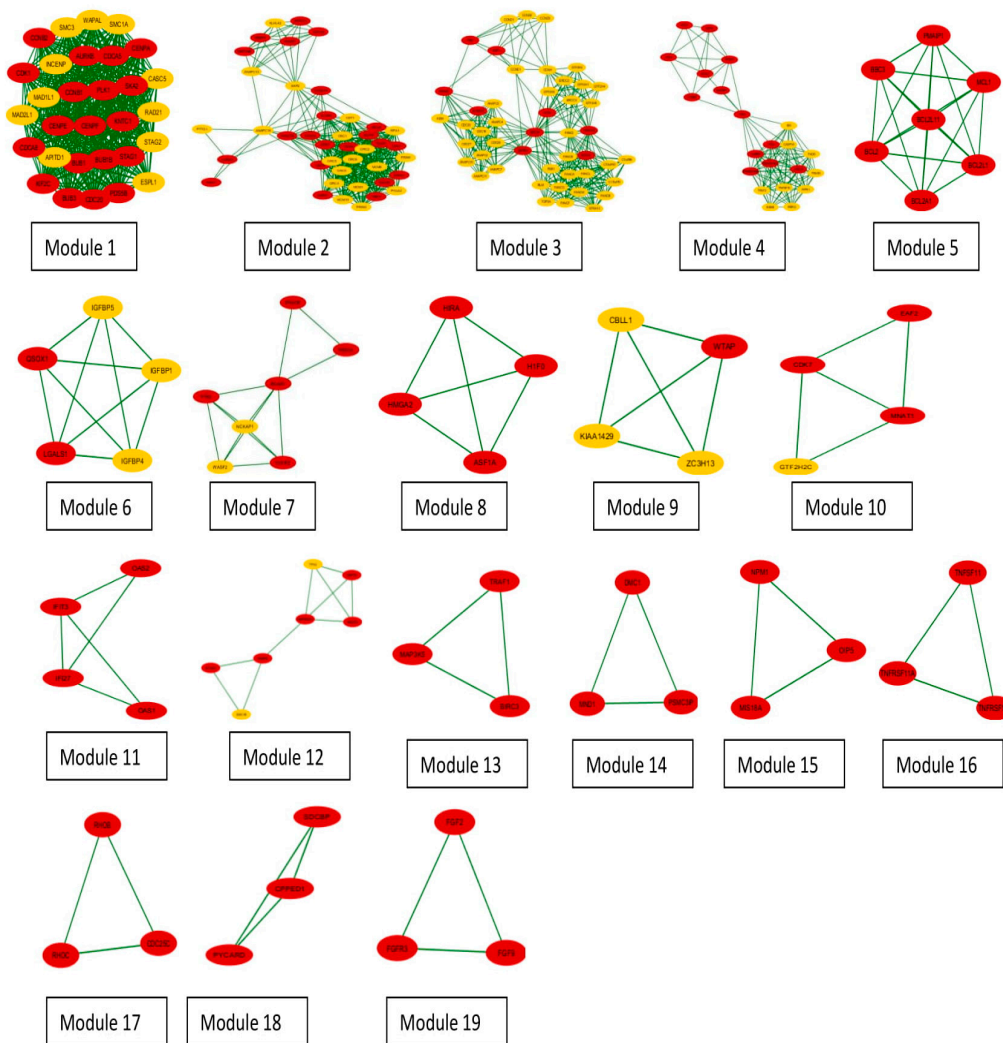


Figure 8. Network modules: red circles show the seed proteins and yellow circles are the connectors.

3.4. PIN Construction and Topological and GO Analysis of Final Selected Seed Proteins

After the modularization process, 115 seed proteins were obtained, and used to further create a PPIN (Figure 9) with 100 connectors using the highest confidence level score of 0.9. The PPIN generated had 213 nodes and 2509 edges, with an average node degree 23.6 and average clustering coefficient of 0.761. The PPI enrichment p -value was less than 1×10^{-16} .

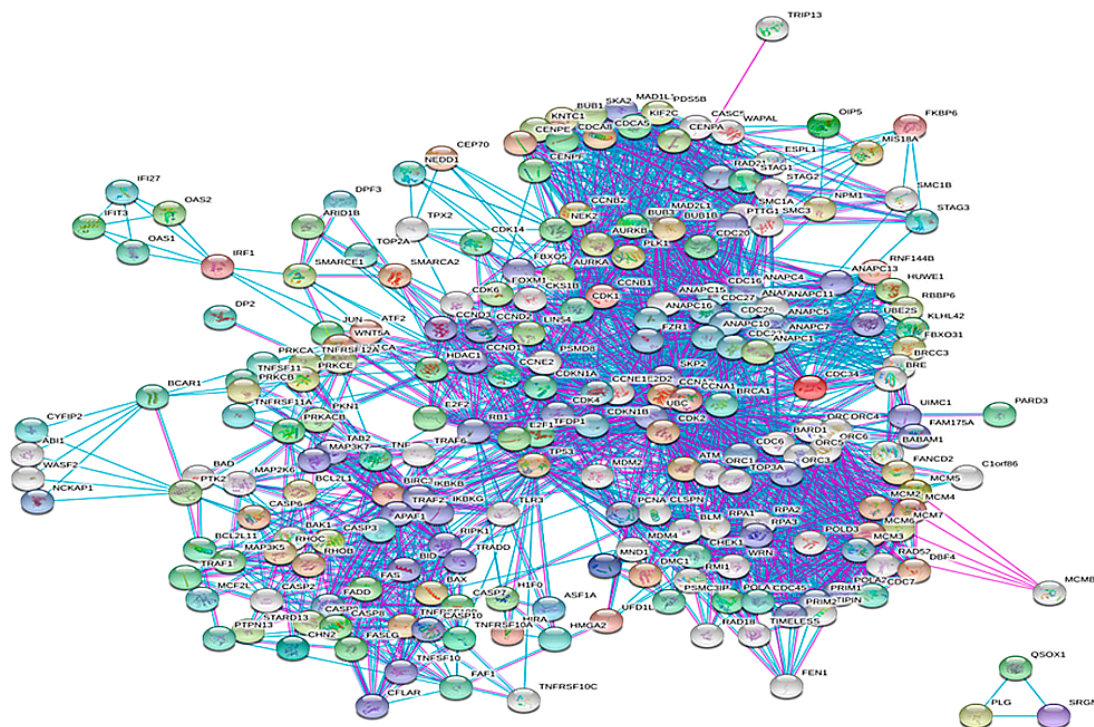


Figure 9. Protein-protein interaction network (PPIN) of final seed proteins.

The network generated was analyzed using Cytohubba, a plug-in of Cytoscape [27]. Moreover, cytohubba analysis of each node of the network was performed and scored based on various parameters, like the degree, closeness centrality, clustering coefficients, betweenness, bottleneck, and stress. The proteins were first sorted on the basis of the clustering coefficient and then on the basis of bottleneck. The proteins with a clustering coefficient less than 0.5 were selected. The proteins having a clustering coefficient more than 0.5 were rejected as this depicts that these proteins were highly clustered and have no further spaces for the attachment of other molecules. The nodes with a clustering coefficient less than 0.5 and high degrees were found to be highly significant. A high degree and clustering coefficient less than 0.5 depict that the nodes are important in various connecting networks and they also have binding spaces available on their surface for other molecules to bind to. Once the nodes were sorted on the basis of clustering coefficients, the next most important parameter was the bottleneck. The proteins with high bottleneck were considered the most critical proteins in any PPIN. The median of the bottleneck scores of selected proteins was calculated which came out to be 3. All the proteins with bottleneck more than or equal to 2 were finally selected. Table 2 enlists the final selected proteins, which were sorted on the basis of bottleneck, with clustering coefficients less than 0.5. CHEK1, showing the highest bottleneck score of 29, was ranked in first position followed by TP53 with a bottleneck of 27. These were also analyzed with ClueGO for the pathway enrichment analysis. Figure 10 shows the pie chart representation of the enriched pathways in the form of groups. In Figure 10, the cell cycle is occupying the maximum area (43.64%) of the pie chart. From this, we can say that the cell cycle is the most enriched pathway, having the maximum number of biomolecules (a detailed graph displaying the sub-pathways in each group along with the number of genes present in it and the proteins involved in each group is enlisted in Table 3, also depicted in Supplementary Figure S1). Figure 11 shows that the maximum number of genes are associated with the cell cycle followed by the cellular macromolecule metabolic process.

Table 2. Key proteins with their bottleneck, clustering coefficient, and degree scores.

Name	Betweenness	Bottleneck	Closeness	Clustering Coefficient	Degree
CHEK1	835.45994	29	116.61667	0.36501	52
TP53	8007.14223	27	126.41667	0.19394	55
BRCA1	2686.48081	23	127.45	0.26346	65
CDK1	3705.28949	19	140.41667	0.32157	85
CDK4	935.26901	14	112.66667	0.42521	35
HSP90AA1	3657.32532	13	101.66667	0.27273	22
RPA2	1523.40238	9	125.78333	0.33978	64
ATM	1803.73545	8	115.36667	0.33718	40
TFDP1	426.33754	6	111.78333	0.46702	34
CDKN1B	1218.04331	4	120.41667	0.44245	50
CASP8	1842.98595	4	90.11667	0.44853	17
PYCARD	796	3	62.18333	0.33333	3
CCNA1	894.78122	3	125.2	0.39548	60
CCNB1	1208.39103	3	127.26667	0.40665	69
RPA1	1657.28981	2	126.45	0.33269	65
CDK2	1887.40725	2	134.2	0.34035	76
CHEK2	126.13416	2	95.95	0.3619	15
BID	577.03969	2	90.78333	0.4269	19
RB1	608.84911	2	107.66667	0.44138	30
PLK1	1284.13385	2	121.25	0.49545	56
CDK7	488.15015	2	108.78333	0.49733	34

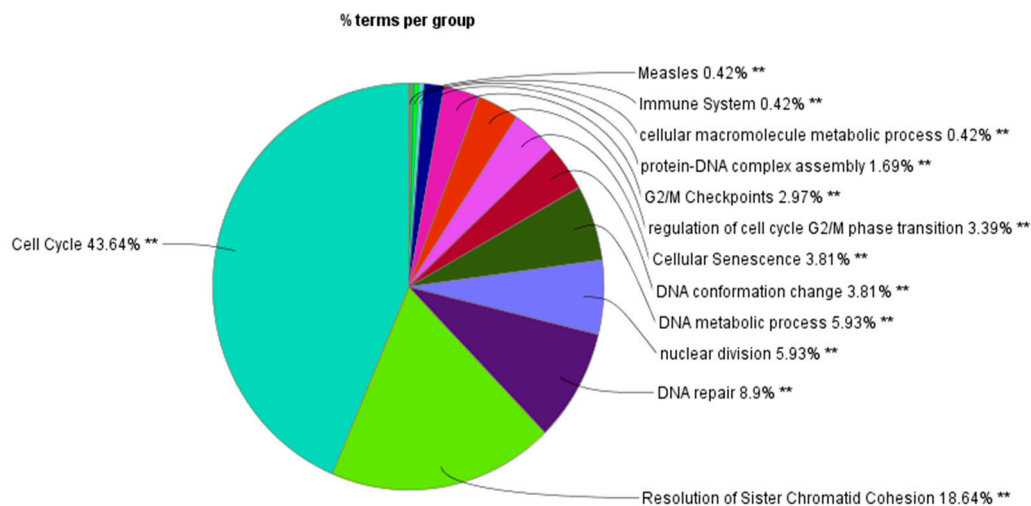
**Figure 10.** ClueGO results of gene ontology (GO) functional enrichment of key proteins.

Table 3. Pathway enrichment of the modulated seed proteins using ClueGO.

Function	Groups	Group Genes
Cell cycle	Group12	ANAPC1 ASF1A ATM AURKA AURKB BARD1 BBC3 BCAR1 BCL2 BCL2A1 BCL2L1 BCL2L11 BIRC3 BRCA1 BUB1 BUB1B BUB3 CASP8 CCNA1 CCNA2 CCNB1 CCNB2 CCNE2 CDC20 CDC25C CDC34 CDC45 CDC6 CDC7 CDCA5 CDCA8 CDK1 CDK2 CDK6 CDKN1A CENPA CENPE CENPF CEP70 CHEK1 CYFIP2 DBF4 DMC1 E2F1 E2F2 EAF2 FANCC FANCD2 FAS FASLG FBXO31 FBXO5 FGF2 FGF9 FGFR3 FKBP6 H1F0 HDAC1 HIRA HMGA2 HSP90AA1 HUWE1 KIF2C KNTC1 LGALS1 MCL1 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MIS18A MNAT1 MND1 NEDD1 NEK2 NPM1 OIP5 PDS5B PLK1 PMAIP1 POLA1 PRKCA PRKCB PSMC3IP PTK2 PYCARD RBBP6 RHOB RHOC RN144B SDCBP SKA2 STAG1 STAG3 TFDP1 TNFRSF10A TNFRSF10B TNFRSF10C TNFRSF12A TOPBP1 TP53 TRAF1
Cellular senescence	Group06	ANAPC1 ASF1A ATM BBC3 BCL2 BCL2L1 BCL2L11 BIRC3 BRCA1 CASP8 CCNA1 CCNA2 CCNE2 CDC20 CDC25C CDCA5 CDK1 CDK2 CDK6 CDKN1A CHEK1 E2F1 E2F2 FAS FASLG FGF2 FGF9 FGFR3 H1F0 HDAC1 HIRA HMGA2 HSP90AA1 MAP3K5 MCL1 MDM2 PMAIP1 PRKCA PRKCB PTK2 TFDP1 TP53 TRAF1
DNA conformation change	Group07	ASF1A ATM BBC3 BCL2L11 BIRC3 CASP8 CCNB1 CDC45 CDCA5 CDK1 CENPA CENPE CENPF DMC1 FANCC FAS FBXO5 H1F0 HDAC1 HIRA HMGA2 HSP90AA1 KNTC1 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MIS18A MNAT1 NPM1 OAS1 OIP5 PMAIP1 POLA1 PRKCA PTK2 PYCARD RHOC TNFSF11 TP53 TRAF1
DNA metabolic process	Group08	ASF1A ATM AURKA AURKB BARD1 BBC3 BCL2 BCL2A1 BCL2L1 BCL2L11 BRCA1 BUB1 BUB1B BUB3 CCNA1 CCNA2 CCNB1 CCNE2 CDC25C CDC34 CDC45 CDC6 CDC7 CDCA5 CDK1 CDK2 CDK6 CDKN1A CENPF CHEK1 DBF4 DMC1 E2F1 EAF2 FANCC FANCD2 FAS FBXO31 FBXO5 FKBP6 H1F0 HDAC1 HMGA2 HSP90AA1 HUWE1 KNTC1 MCL1 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MIS18A MNAT1 MND1 NEK2 NPM1 PDS5B PLK1 PMAIP1 POLA1 PRKCB PSMC3IP PYCARD RBBP6 TFDP1 TNFRSF10A TNFRSF10B TNFRSF10C TOPBP1 TP53
DNA repair	Group10	ANAPC1 ASF1A ATM AURKA AURKB BARD1 BBC3 BCL2 BCL2L11 BRCA1 CASP8 CCNA1 CCNA2 CCNB1 CCNB2 CCNE2 CDC20 CDC25C CDC34 CDC45 CDC6 CDC7 CDCA5 CDK1 CDK2 CDK6 CDKN1A CENPF CEP70 CHEK1 DBF4 DMC1 E2F1 E2F2 EAF2 FANCC FANCD2 FAS FASLG H1F0 HDAC1 HIRA HMGA2 HSP90AA1 HUWE1 MAP3K5 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MNAT1 MND1 NEDD1 NEK2 NPM1 PDS5B PLK1 PMAIP1 POLA1 PRKCA PRKCB RBBP6 TFDP1 TNFRSF10A TNFRSF10B TNFRSF10C TOPBP1 TP53 TRAF1
G2/M checkpoints	Group04	ASF1A ATM AURKA AURKB BARD1 BBC3 BCL2 BCL2A1 BCL2L1 BCL2L11 BRCA1 CCNA1 CCNA2 CCNB1 CCNB2 CCNE2 CDC25C CDC34 CDC45 CDC6 CDC7 CDCA5 CDK1 CDK2 CDKN1A CENPF CHEK1 DBF4 DMC1 E2F1 FANCC FANCD2 FBXO31 FKBP6 H1F0 HDAC1 HMGA2 HSP90AA1 HUWE1 MCL1 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MIS18A MNAT1 MND1 NEK2 NPM1 PDS5B PLK1 PMAIP1 POLA1 PSMC3IP PYCARD RBBP6 TFDP1 TOPBP1 TP53
Immune system	Group01	ANAPC1 BCL2 BCL2L1 BIRC3 CASP8 CDC20 CDC34 CDK1 CDKN1A CENPE CPPED1 CYFIP2 FASLG FBXO31 FGF2 FGF9 FGFR3 HSP90AA1 HUWE1 IFI27 IFIT3 KIF2C MCL1 OAS1 OAS2 PRKCB PTK2 PYCARD QSOX1 RBBP6 RN144B SDCBP TNFRSF11A TNFRSF12A TNFSF11 TP53
Measles	Group00	BBC3 CCNE2 CDK2 CDK6 FAS FASLG OAS1 OAS2 TNFRSF10A TNFRSF10B TNFRSF10C TP53
Resolution of sister chromatid cohesion	Group11	ANAPC1 ATM AURKA AURKB BARD1 BBC3 BCL2 BCL2L11 BIRC3 BRCA1 BUB1 BUB1B BUB3 CASP8 CCNA1 CCNA2 CCNB1 CCNB2 CCNE2 CDC20 CDC25C CDC34 CDC6 CDC7 CDCA5 CDCA8 CDK1 CDK2 CDKN1A CENPA CENPE CENPF CEP70 CHEK1 CYFIP2 DMC1 E2F1 FANCD2 FAS FASLG FBXO31 FBXO5 FKBP6 HMGA2 HSP90AA1 HUWE1 IFI27 KIF2C KNTC1 MAP3K5 MDM2 MNAT1 MND1 NEDD1 NEK2 NPM1 PDS5B PLK1 PMAIP1 PRKCA PRKCB PSMC3IP PTK2 PYCARD RBBP6 RN144B SDCBP SKA2 STAG1 STAG3 TFDP1 TNFRSF10A TNFRSF10B TOPBP1 TP53

Table 3. Cont.

Function	Groups	Group Genes
Cellular macromolecule metabolic process	Group02	ANAPC1 ASF1A ATM AURKA AURKB BARD1 BBC3 BCL2 BCL2L1 BIRC3 BRCA1 BUB1 BUB1B BUB3 CASP8 CCNA1 CCNA2 CCNB1 CCNE2 CDC20 CDC25C CDC34 CDC45 CDC6 CDC7 CDCA5 CDCA8 CDK1 CDK2 CDK6 CDKN1A CENPE CENPF CHEK1 CPPED1 CYFIP2 DBF4 DMC1 E2F1 E2F2 EAF2 FANCC FANCD2 FAS FASLG FBXO31 FBXO5 FGF2 FGF9 FGFR3 FKBP6 H1FO HDAC1 HIRA HMGA2 HSP90AA1 HUWE1 IFI27 LGALS1 MAP3K5 MCM2 MCM4 MCM5 MCM6 MCM7 MDM2 MIS18A MNAT1 MND1 NEK2 NPM1 OAS1 OAS2 PDS5B PLK1 PMAIP1 POLA1 PRKCA PRKCB PSMC3IP PTK2 PYCARD QSOX1 RBBP6 RNFI44B SDCBP STAG1 TFDP1 TNFRSF10A TNFRSF10B TNFRSF10C TNFRSF11A TNFSF11 TOPBP1 TP53 TRAF1 WTAP
Nuclear division	Group09	ANAPC1 ATM AURKA AURKB BRCA1 BUB1 BUB1B BUB3 CCNA1 CCNA2 CCNB1 CCNE2 CDC20 CDC25C CDC6 CDCA5 CDCA8 CDK1 CDK2 CENPE CENPF CHEK1 DMC1 FANCD2 FBXO5 FKBP6 KIF2C KNTC1 MND1 NEK2 PDS5B PLK1 PSMC3IP STAG1 STAG3
Protein–DNA complex assembly	Group03	ASF1A ATM AURKA AURKB BBC3 BCL2 BCL2L1 BIRC3 BRCA1 CASP8 CCNB1 CDC20 CDC34 CDC45 CDK1 CDK2 CENPA CENPE CENPF CEP70 DMC1 FANCC FAS FBXO5 H1FO HDAC1 HIRA HMGA2 HSP90AA1 KNTC1 MCM2 MCM7 MDM2 MIS18A MNAT1 NEDD1 NEK2 NPM1 OAS1 OIP5 PLK1 PMAIP1 PRKCA PTK2 PYCARD RHOC SDCBP STAG1 STAG3 TNFSF11 TP53 TRAF1
Regulation of cell cycle G2/M phase transition	Group05	ANAPC1 ATM AURKA AURKB BARD1 BRCA1 BUB1B BUB3 CCNA1 CCNA2 CCNB1 CCNB2 CDC20 CDC25C CDC7 CDK1 CDK2 CDK6 CDKN1A CENPF CEP70 CHEK1 FBXO5 HMGA2 HSP90AA1 MDM2 MNAT1 NEDD1 NEK2 NPM1 PLK1 PRKCA PRKCB TOPBP1 TP53

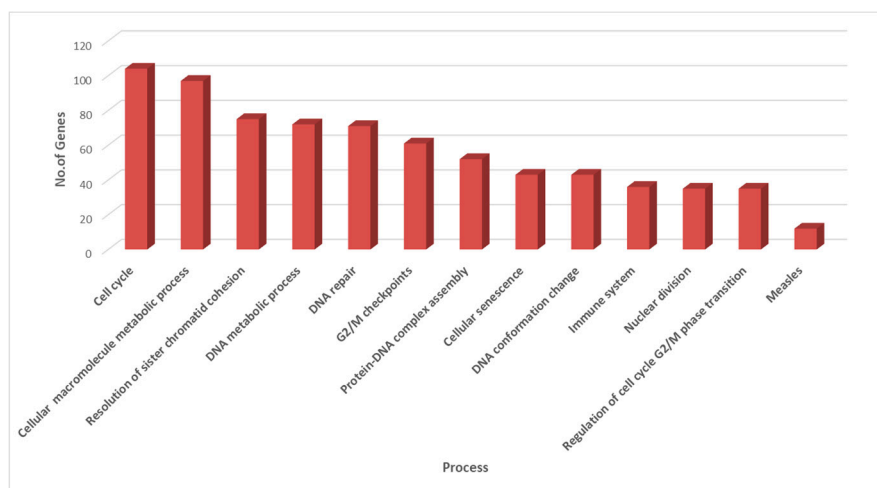


Figure 11. Representation of the pathways enriched and the total number of genes associated with them.

3.5. Molecular Docking Simulation of NNK with Key Proteins

Autodock 4.0 was used for docking the final 30 target proteins with NNK. CDK7 showed the highest binding energy of -5.93 Kcal/Mol followed by CCNA1 (-5.6 Kcal/Mol). The binding energies of the proteins with NNK will further help in refining the results in the selection of the best-suited target proteins of NNK. The more negative the binding energy, the stronger the interaction between the ligand and the protein. The binding energies, K_i values, and the H-bonds formed along with their distances for all the 20 target proteins are listed in Table 4. Figure 12 shows the top three interactions of NNK with its target biomolecules, namely CDK7, CCNA1, and CDKN1B. Table 5 shows the key biomolecular targets of NNK and their role in cell cycle regulation.

Table 4. Final 22 target proteins docked with NNK.

S.No	Protein	Ligand	Binding Energy (Kcal/Mol)	K _i	Binding Residues	H-Bond	Distance
1.	CDK7	NNK	-5.93	45.31 μM	Leu18, Val26, Ala39, Lys41, Ile75, Phe91, Asp92, Phe93, Met94, Glu95, Thr96, Asp97, Leu144, Asp155	CDK7:MET94:N - :NNK:O7	2.86224
2.	CCNA1 (connector)	NNK	-5.60	79.09 μM	Cys97, Gly98, Gln99, Gly100, Val164, Asp165, Thr166, Gly167, Thr168, Leu169, Lys170, Leu173, Tyr218	:GLY98:H - :NNK:O7 :GLY167:H - :NNK:N10 :THR168:H - :ASP165:O :LYS170:H - :NNK:N14 :LYS170:H - :NNK:O15 :NNK:N14 - :THR168:O	1.83311 1.98057 2.11393 2.36928 1.99807 3.02246
3.	CDKN1B	NNK	-5.42	106.27 μM	His573, Lys574, Pro575, Leu576, Glu581, Trp582, Gln583, Glu584	CDKN1B:GLN583:N - :NNK:N14 CDKN1B:GLN583:N - :NNK:O15 :NNK:N14 - CDKN1B:GLN583:O	2.84311 2.79833 2.96186
4.	CASP8	NNK	-5.35	119.75 μM	Lys2224, Tyr2226, Gln2227, Asp2308, Gly2350, Lys2351, Pro2352, Asp2398, Arg2471, Lys2472	CASP8:LYS2351:HZ1 - :NNK:N10 CASP8:ARG2471:HH21 - :NNK:N2 CASP8:ARG2471:HH21 - :NNK:N14 CASP8:ARG2471:HH21 - :NNK:O15 CASP8:LYS2472:HZ1 - :NNK:O7	1.9527 2.48624 2.19123 2.25672 1.82255
5.	CHEK2 (connector)	NNK	-5.35	120.25 μM	Ser228, Gly229, Ala230, Cys231, Gly232, Val234, Lys249, Leu301, Thr367, Asp368	CHEK2:CYS231:N - :NNK:N10 CHEK2:GLY232:N - :NNK:N10 CHEK2:LYS249:NZ - :NNK:N14 CHEK2:ASP368:N - :NNK:O15	3.11295 3.11412 3.20168 2.86381
6.	PLK1	NNK	-5.21	152.98 μM	Lys413, Trp414, Val415, Asp416, Leu490, Asn533, Lys540	PLK1:TRP414:N - :NNK:O7 PLK1:ASP416:N - :NNK:N14 PLK1:ASP416:N - :NNK:O15 PLK1:ASN533:ND2 - :NNK:N10	2.97424 2.94596 2.71702 3.10323
7.	BID (connector)	NNK	-5.13	174.27 μM	Leu21, Phe24, Gly25, Gln28, Leu39, Asp40, Leu42, Gly43, Arg86, Ala89, Arg90, Phe173	:NNK:N14 - BID:GLN28:OE1 BID:PHE24:HA - :NNK:O15 BID:ARG86:HA - :NNK:N10 :NNK:C3 - BID:GLN28:OE1 :NNK:C1 - BID:LEU39:O :NNK:C11 - BID:ARG86:O :NNK:O15 - BID:PHE24 :NNK:N14 - BID:GLN28:OE1 BID:PHE24:HA - :NNK:O15 BID:ARG86:HA - :NNK:N10 :NNK:C3 - BID:GLN28:OE1 :NNK:C1 - BID:LEU39:O :NNK:C11 - BID:ARG86:O :NNK:O15 - BID:PHE24	3.29101 2.94776 2.82375 3.41587 3.00214 3.32997 3.70686 3.29101 2.94776 2.82375 3.41587 3.00214 3.32997 3.70686
8.	HSP90AA1	NNK	-5.10	183.46 μM	Leu48, Asn51, Ser52, Ala55, Asp93, Ile96, Gly97, Met98, Asn106, Phe138, Thr184, Val186	HSP90AA1:ASN51:ND2 - :NNK:N14 HSP90AA1:ASN51:ND2 - :NNK:O15 HSP90AA1:THR184:OG1 - :NNK:O7	3.12471 2.99653 2.69827
9.	BRCA1	NNK	-5.08	187.68 μM	Val1654, Ser1655, Gly1656, Leu1657, Thr1658, Pro1659, Phe1662, Thr1700, Leu1701, Lys1702	BRCA1:SER1655:OG - :NNK:N14 BRCA1:GLY1656:N - :NNK:O7 BRCA1:LEU1657:N - :NNK:O7 BRCA1:LEU1701:N - :NNK:O15 BRCA1:LYS1702:N - :NNK:O15 :NNK:N14 - BRCA1:SER1655:OG	3.19252 2.75464 2.77131 3.00037 2.75018 3.19252
10.	CDK1	NNK	-5.00	217.06 μM	Lys88, Leu91, Asp92, Ile94, Pro95, Pro96, Glu196, Lys200	CDK1:LYS200:NZ - :NNK:N14 CDK1:LYS200:NZ - :NNK:O15 :NNK:N14 - CDK1:ILE94:O	3.0124 2.96559 2.89466
11.	CDK2	NNK	-4.90	255.02 μM	Val29, Glu81, Phe82, Leu83, His84, Ile135, Asn136, Thr137	CDK2:PHE82:N - :NNK:N10 CDK2:HIS84:N - :NNK:N14 CDK2:HIS84:N - :NNK:O15 CDK2:HIS84:ND1 - :NNK:N14 :NNK:O15 - CDK2:ILE135:O	2.88593 2.95506 2.76842 3.18956 2.91961

Table 4. Cont.

S.No	Protein	Ligand	Binding Energy (Kcal/Mol)	K _i	Binding Residues	H-Bond	Distance
12.	CCNB1	NNK	-4.86	274.97 μM	Ile253, Lys256, Tyr257, Glu285, Leu289, Phe294, Gly295, Leu296, Gly297	CCNB1:TYR257:N - :NNK:N10 CCNB1:LEU296:N - : NNK:N14 CCNB1:LEU296:N - : NNK:O15 CCNB1:GLY297:N - : NNK:O15 : NNK:N14 - CCNB1:PHE294:O	3.14377 2.86319 2.81795 2.91169 3.10697
13.	CHEK1	NNK	-4.80	303.50 μM	Val23, Val37, Ile39, Glu55, Asn59, Leu82, Phe149	CHEK1:ILE39:N - :NNK:N10 CHEK1:ASN59:ND2 - :NNK:N14 CHEK1:ASN59:ND2 - :NNK:O15 CHEK1:PHE149:N - :NNK:O15 :NNK:N14 - CHEK1:GLU55:OE2 :NNK:O15 - CHEK1:GLU55:OE2 :NNK:O15 - CHEK1:PHE149:N	2.8265 2.71952 3.02158 3.15867 3.05383 2.67183 3.15867
14.	RPA2 (connector)	NNK	-4.70	358.90 μM	Cys49, Thr50, Ile76, Val77, Asp96, Met97, Tyr125, Phe155, His158, Ile159	RPA2:VAL77:N - :NNK:O15 :NNK:O15 - RPA2:CYS49:O :NNK:O15 - RPA2:VAL77:O :NNK:O15 - RPA2:HIS158:NE2	3.09031 3.06724 3.18087 2.98064
15.	ATM	NNK	-4.48	523.49 μM	Thr2059, Ala2062, Gly2063, Ile2065, Gln2066, Gln2069, Leu2077, Tyr2080, Leu2081, Leu2084, Glu2094, Leu2095, Leu2098	ATM:GLN2066:N - :NNK:O7	3.06063
16.	CDK4	NNK	-4.35	650.98 μM	Val44, Leu54, Pro55, Thr58, Val59, Val62, Ala63, Arg66, Val82, Ile92, Val94	CDK4:PRO55:CD - :NNK:O15 CDK4:VAL59:CA - :NNK:O7	2.9883 2.92285
17.	TFDP1	NNK	-4.20	833.96 μM	Val264, Phe285, Asn286, Phe287, Phe291	TFDP1:PHE287:N - :NNK:O15 :NNK:O15 - TFDP1:PHE287:O	2.78538 3.08744
18.	TP53	NNK	-4.14	927.72 μM	Gln136, Leu137, Ala138, His179, Cys182, Asp184, Asn239, Cys275, Ala276	TP53:LEU137:N - :NNK:O15 TP53:ASN239:ND2 - :NNK:N14 TP53:ASN239:ND2 - :NNK:O15 :NNK:O15 - TP53:CYS275:O	2.93164 3.07617 2.87469 2.82627
19.	RB1	NNK	-4.02	1.14 mM	Val434, Gly435, Gln436, Cys438, Asn505, Leu506, Asp507, Ser508, Gly509, Thr510	RB1:GLN436:HN - :NNK:O15 RB1:SER508:HN - :NNK:O7 RB1:GLY509:HN - :NNK:O7	1.93726 2.36625 1.90514
20.	RPA1 (connector)	NNK	-3.67	2.03 mM	Val375, Asn402, Pro403, Ala408, Tyr409, Arg412, Gly413	RPA1:ARG412:NH1 - :NNK:N14 :NNK:O15 - RPA1:ALA408:O	3.06161 2.78707

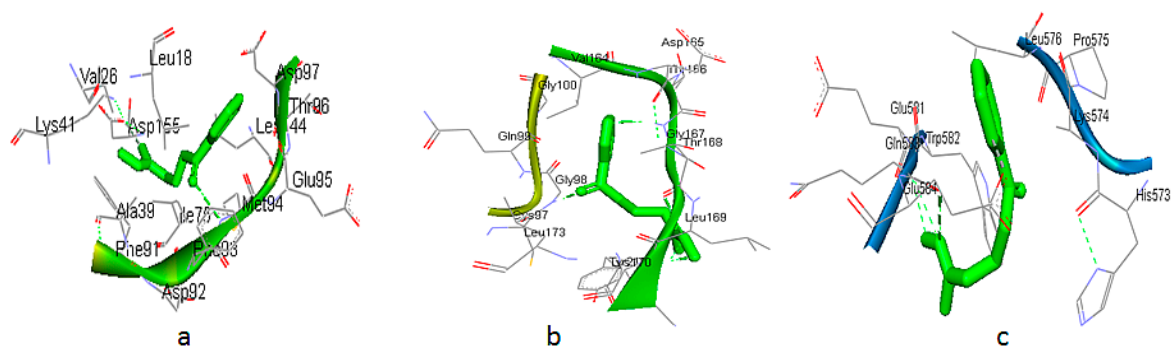


Figure 12. Binding interactions of the top three proteins with NNK: (a) CDK7 (−5.93 Kcal/Mol), (b) CCNA1 (−5.6 Kcal/Mol), (c) CDKN1B (−5.42 Kcal/Mol).

Table 5. Key proteins and their roles in cell cycle regulation.

S.No	Protein	Role in Cell Cycle	References
1.	CDK7	Catalytic subunit of CAK complex which activates the cyclin-associated kinases CDK1, CDK2, CDK4, and CDK6 by threonine phosphorylation, thus regulating cell cycle progression	[28,29]
2.	CCNA1	Controls the cell cycle at the G1/S (start) and G2/M (mitosis) transitions	[30]
3.	CDKN1B	Important regulator of cell cycle progression. Acts either as an inhibitor or an activator of cyclin type D-CDK4 complexes depending on its phosphorylation state and/or stoichiometry.	[31]
4.	CASP8	Responsible for the positive and negative regulation of apoptosis and inflammation	[32]
5.	CHEK2	required for checkpoint-mediated cell cycle arrest in response to p53 defects	[33]
6.	PLK1	Regulates entry and exit from mitosis and cytokinesis. Regulates DNA replication	[34]
7.	BID	induces apoptosis on interaction with BCL2 family proteins	[35]
8.	HSP90AA1	regulates the function of ATM in sensing and repairing the DNA damages	[36]
9.	BRCA1	Has role in DNA damage repair, cell cycle control and transcriptional regulation	[37]
10.	CDK1	Promotes G2-M transition and regulates G1 progress and G1-S transition via association with multiple interphase cyclins.	[38]
11.	CDK2	Essential for transition of cell cycle from G1 to S phase and then from S to G2 phase.	[39,40]
12.	CCNB1	Essential for the control of the cell cycle at the G2/M (mitosis) transition	[41]
13.	CHEK1	required for smooth cellular proliferation by inducing the degradation of cdc25a	[42]
14.	RPA2	Binds and stabilizes single-stranded DNA intermediates. Controls DNA repair and DNA damage checkpoint activation.	[43,44]
15.	ATM	Activates checkpoint signaling upon double strand breaks (DSBs), apoptosis. May function as a tumor suppressor.	[45,46]
16.	CDK4	Hypophosphorylates RB1 in early G ₁ phase. Cyclin D-CDK4 complexes are major integrators of various mitogenic and antimitogenic signals.	[47]
17.	TFDP1	Can stimulate E2F-dependent transcription induces growth arrest.	[48]
18.	TP53	Prevents CDK7 kinase activity when associated to CAK complex in response to DNA damage, thus stopping cell cycle progression.	[49,50]
19.	RB1	Key regulator of entry into cell division that acts as a tumor suppressor. Promotes G0-G1 transition when phosphorylated by CDK3/cyclin-C.	[51]
20.	RPA1	controls DNA repair and DNA damage checkpoint activation	[52]

4. Discussion

Cancer is a global inflammable problem, which pathologically occurs due to the accumulation of mutations in one or many genes, a phenomenon known as “oncogene addiction”. The loss of fidelity during the replication of DNA or the repair of damaged DNA leads to a cell becoming cancerous. Tobacco consumption has a major influence on these aberrations due to its main component, NNK, which has been strongly related with various cancers, mainly lung cancer [53,54]. NNK binds

with various proteins involved in several cellular processes and leads to cancer. Hence, it is vital to fully understand the precise mechanism of the disease development caused by NNK and the interactome analysis of biomolecular target genes/proteins for attaining an overview of the highly complex functional interdependency of these targets for the efficient performance of the cell.

Systems biology is an interdisciplinary field that facilitates an understanding of complex biological network systems. These networks can be metabolic networks, gene regulation network, cell signaling network, or protein–protein interaction network [55]. With the help of system biology, we can analyze huge real-time networks precisely. Significantly, the huge amount of data from research being conducted globally were collected and statistically analyzed to uncover the functions of individual genes and proteins. The results may then be integrated to provide higher-level information [56]. Mathematical modeling and computational simulations help in understanding the internal dynamics of the system or the process and thus help in predicting the future of the process [57]. The networks created represent the genes or proteins with the nodes and their relations or physical connections with the edges [58]. The biological systems are immensely huge and complex networks depicting how one protein is connected with other proteins and how a slight impact on a gene or protein will affect the whole interactome [59]. These highly complicated and tightly packed networks, which govern all biological processes, are also referred to as protein–protein interaction networks (PPINs). All these graphs and networks work on the principle of graph theory. In the current study, we focused on NNK rewired PPIN to identify its bio-molecular targets that regulate the various cellular processes and cause cancer.

To construct PPIN, a literature survey was performed on PubMed database using keywords, like NNK, and around 544 genes extracted from 1317 research articles. The STRING database was used for the generation of the PPIN as it gives freedom to the users to choose the active interaction sources for their network. For this study, we used experimental data sources, like BIND, IntAct, etc., and pathway database interaction sources, like KEGG and reactome. The network generated had 534 nodes, including 100 connectors, at the highest confidence level score of 0.9. It contained 2909 edges that depicted that all the reported nodes were well connected with each other, which were further confirmed by the topological parameter, i.e., the average node degree, which was found to be 10.9. The degree of a node gives the information about the connection of the node with other nodes or simply the number of edges that either enter a node or leave it. All the biological systems are undirected and scale-free so the node degree depicts the average connectivity of each node with other nodes present in the network. The identification of key nodes in any network is not possible just by calculating its degree, hence various other parameters require evaluation, like the betweenness centrality, closeness centrality, and bottleneck scores [60]. Nodes with clustering coefficients less than 0.5 are key nodes because lesser clustering coefficients depict that the protein still have space for interaction with other molecules. The average neighborhood connectivity details provide the information about the density of the network. The shortest path length is important as it gives details about the minimum number of the edges between the two nodes and how fast information can pass on from one node to another node.

The network generated was then divided into modules. Modularization is an important step as it helps in reducing the noise of the data. On modularization, 19 clusters were obtained and each cluster contained seed proteins. Finally, 115 seed proteins were procured from the initial list of 534 proteins. Again, a PPIN was generated using these 115 seed proteins and 100 connectors at the first and second shells, and we obtained 213 nodes and 2509 edges. In addition, 100 connectors were done in order to include some more relevant proteins, which were possibly missed in the above procedure. The topological analysis was done on the basis of clustering coefficient and bottleneck scores. We used the bottleneck score as our final selection criteria as bottlenecks are the significant proteins, which have great functional and dynamic significance in a network [61]. To find the statistically significant cut-off for the proteins to be selected as key targets, the median was calculated on the bottleneck score, which came out to be 2. The enrichment analysis was done to find the pathways and processes that are enriched in the process of the PPIN generation. On the GO enrichment analysis, it was found that

most of the seed proteins fall under the pathways that are related to cell cycle regulation and these results were also reflected on the final selected seed proteins that had a bottleneck score of 2 or more than 2. The cell cycle is a tightly regulated and a precisely timed event that is divided into phases and is monitored by checkpoints [62]. It is a highly orchestrated event, which ensures the integrity and stability of the genome. NNK binds to various receptors and hampers various pathways, leading to the cell becoming cancerous [63]. It not only binds with the receptors but also targets many important genes and proteins, like CDKs and cyclins, that are crucial for cell cycle processes [6]. Table 3 and Figure 10, Supplementary Figure S1, and Figure 11 clearly depict that most of the pathways that were enriched fall under cell cycle regulation at different phases.

Finally, docking studies were performed for the target proteins using NNK as a ligand. Molecular docking is the key tool to check whether the binding of ligand and protein is thermodynamically possible or not. The binding energies of these docking results further helped in refining the targets for NNK. The major targets of NNK found were CDK7, CCNA1, CDKN1B, CASP8, CHEK2, PLK1, BID, HSP90AA1, BRCA1, CDK1, CDK2, CCNB1, CHEK1, RPA2, ATM, CDK4, TFDP1, TP53, RB1, and RPA1, with their binding energies ranging from -5.93 to -3.67 Kcal/Mol. PYCARD and QXOS1 were excluded from this screening because of their low degree and 0 clustering coefficient. In addition to the above key proteins, our present study also revealed five connector proteins, which can be proven as putative biomolecular targets of NNK. These proteins are CHEK2, CCNA1, BID, RPA1, and RPA2. From these docking results, we can easily conclude that a majority of the key targets of NNK belong to the cell cycle regulatory proteome. CDKs and cyclins are the key regulatory proteins of the cell cycle that control the various phases of the cell. Different CDK/cyclin complexes are required for a smooth transition of the cell cycle. Other proteins, like ATM, RB1, and TP53, may act as tumor suppressor proteins and signaling proteins for DNA damage. Table 5 enlists the final key biomolecular targets of NNK and their role in the cell cycle.

5. Conclusions

In conclusion, a NNK rewired PPIN, with the help of various systems biology tools, was explored for the identification of potential targets involved in tobacco induced cancer development. The biomolecules that were suspected for being the most probable targets of NNK were further screened using functional enrichment techniques and then using molecular docking techniques. The results of this study show that the maximum proteins that are the most probable targets of NNK are involved in cell cycle regulation. With these, we can conclude that NNK has a major impact on the biomolecules of the cell cycle regulatory proteome. The present study opens future possibilities to identify new biomarkers for the cancers associated with NNK. This can also help in the pre-symptomatic diagnosis of diseases and in the development of precision medicines.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2073-4425/10/8/564/s1>, Figure S1: Graph of GO enriched pathways depicting the involvement of genes under various sub-graphs.

Author Contributions: Conceptualization, A.D., U.B., M.G., and K.K.K.; methodology, A.D., A.N.; software, A.N., M.L.; validation, A.D., A.N., D.K.; formal analysis, A.D., S.U., P.S., U.B., M.G., S.H. and K.K.K.; investigation, A.N., A.D., S.H.; resources, A.D., K.K.K.; data curation, A.D., A.N., S.U. and P.S.; writing—original draft preparation, A.D., A.N., K.K.K.; writing—review and editing, D.K., M.L., J.R.; visualization, A.N.; supervision, A.D.; project administration, A.D.

Acknowledgments: The authors are grateful to Vijay Dhasmana, Honorable Vice Chancellor Swami Rama Himalayan University, India, Sunil Saini Chairperson Research Committee, Head of Department of Biosciences, other faculty members of Biosciences and all members of Research Committee—Himalayan Institute of Medical Sciences, Swami Rama Himalayan University, India, for providing the motivation and necessary facilities for this study.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Pavlopoulou, A.; Spandidos, D.A.; Michalopoulos, I. Human cancer databases (Review). *Oncol. Rep.* **2015**, *33*, 3–18. [[CrossRef](#)] [[PubMed](#)]
2. Bray, F.; Colombet, M.; Mery, L.; Piñeros, M.; Znaor, A.; Zanetti, R.; Ferlay, J. *Cancer Incidence in Five Continents*; IARC: Lyon, France, 2017; Volume XI.
3. Weinstein, I.B.; Joe, A.K. Mechanisms of Disease: Oncogene addiction—A rationale for molecular targeting in cancer therapy. *Nat. Rev. Clin. Oncol.* **2006**, *3*, 448–457. [[CrossRef](#)] [[PubMed](#)]
4. Forouzanfar, M.H.; Afshin, A.; Alexander, L.T.; Anderson, H.R.; Bhutta, Z.A.; Biryukov, S.; Brauer, M.; Burnett, R.; Cercy, K.; Charlson, F.J.; et al. GBD 2015 Risk Factors Collaborators. Global, regional, and national comparative risk assessment of 79 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990–2015: A systematic analysis for the Global Burden of Disease Study 2015. *Lancet* **2016**, *388*, 1659–1724.
5. Hecht, S.S. Tobacco smoke carcinogens and breast cancer. *Environ. Mol. Mutagen.* **2002**, *39*, 119–126. [[CrossRef](#)] [[PubMed](#)]
6. Haneef, M.; Lohani, M.; Dhasmana, A.; Jamal, Q.M.; Shahid, S.M.; Firdaus, S. Molecular Docking of Known Carcinogen 4-(Methyl-nitrosamino)-1-(3-pyridyl)-1-butanone (NNK) with Cyclin Dependent Kinases towards Its Potential Role in Cell Cycle Perturbation. *Bioinformation* **2014**, *10*, 526–532. [[CrossRef](#)]
7. Xue, J.; Yang, S.; Seng, S. Mechanisms of Cancer Induction by Tobacco-Specific NNK and NNN. *Cancers* **2014**, *6*, 1138–1156. [[CrossRef](#)] [[PubMed](#)]
8. Zabala, V.; Tong, M.; Yu, R.; Ramirez, T.; Yalcin, E.B.; Balbo, S.; Silbermann, E.; Deochand, C.; Nunez, K.; Hecht, S.; et al. Potential Contributions of the Tobacco Nicotine-Derived Nitrosamine Ketone (NNK) in the Pathogenesis of Steatohepatitis in a Chronic Plus Binge Rat Model of Alcoholic Liver Disease. *Alcohol Alcohol.* **2015**, *50*, 118–131. [[CrossRef](#)]
9. Nunez, K.; Kay, J.; Krotow, A.; Tong, M.; Agarwal, A.R. Cigarette Smoke-Induced Alterations in Frontal White Matter Lipid Profiles Demonstrated by MALDI-Imaging Mass Spectrometry: Relevance to Alzheimer’s Disease. *J. Alzheimers Dis.* **2016**, *51*, 151–163. [[CrossRef](#)]
10. Pai, M.; Mohan, A.; Dheda, K.; Leung, C.C.; Yew, W.W.; Christopher, D.J.; Sharma, S.K. Lethal Interaction: The colliding epidemics of tobacco and tuberculosis. *Expert Rev. Anti Infect. Ther.* **2007**, *5*, 385–391. [[CrossRef](#)]
11. Wang, Y.; Zhang, X.S.; Chen, L. Modelling biological systems from molecules to dynamical networks. *BMC Syst. Biol.* **2012**, *6* (Suppl. 1), 2–5. [[CrossRef](#)]
12. Villaverde, A.F.; Ross, J.; Banga, J.R. Reverse engineering cellular networks with information theoretic methods. *Cells* **2013**, *2*, 306–329. [[CrossRef](#)] [[PubMed](#)]
13. Szklarczyk, D.; Morris, J.H.; Cook, H.; Kuhn, M.; Wyder, S.; Simonovic, M.; Santos, A.; Doncheva, N.T.; Roth, A.; Bork, P.; et al. The STRING database in 2017: Quality-controlled protein-protein association networks, made broadly accessible. *Nucleic Acids Res.* **2017**, *45*, D362–D368. [[CrossRef](#)] [[PubMed](#)]
14. Shannon, P.; Markiel, A.; Ozier, O.; Baliga, N.S.; Wang, J.T.; Ramage, D.; Amin, N.; Schwikowski, B.; Ideker, T. Cytoscape: A software environment for integrated models of biomolecular interaction networks. *Genome Res.* **2003**, *13*, 2498–2504. [[CrossRef](#)] [[PubMed](#)]
15. Doncheva, N.T.; Assenov, Y.; Domingues, F.S.; Albrecht, M. Topological analysis and interactive visualization of biological networks and protein structures. *Nat. Protoc.* **2012**, *7*, 670–685. [[CrossRef](#)] [[PubMed](#)]
16. Barabasi, A.L.; Oltvai, Z.N. Network biology: Understanding the cell’s functional organization. *Nat. Rev. Genet.* **2004**, *5*, 101–113. [[CrossRef](#)] [[PubMed](#)]
17. Almaas, E. Biological impacts and context of network theory. *J. Exp. Biol.* **2007**, *210*, 1548–1558. [[CrossRef](#)] [[PubMed](#)]
18. Klemm, K.; Bornholdt, S. Topology of biological networks and reliability of information processing. *Proc. Natl. Acad. Sci. USA* **2005**, *102*, 18414–18419. [[CrossRef](#)] [[PubMed](#)]
19. Cingovska, I.; Kovachev, A.; Gmbh, A.T.; Trivodaliev, K.; Kalajdziski, S. Protein Function Prediction by Clustering of Protein-Protein Interaction Network. In *Proceedings of the ICT Innovations 2011*; Kocarev, L., Ed.; Advances in Intelligent and Soft Computing; Springer: Berlin, Germany, 2012; Volume 150. [[CrossRef](#)]
20. Rajendran, P.; Thangavel, D. Clustering of Microarray Data to Identify Enriched Go Terms of Genes in Severe Asthma Dataset using Gene Enrichment Analysis. *Ind. J. Sci. Tech.* **2016**, *9*, 1–11. [[CrossRef](#)]

21. Bindea, G.; Mlecnik, B.; Hackl, H.; Charoentong, P.; Tosolini, M.; Kirilovsky, A.; Fridman, W.H.; Pagès, F.; Trajanoski, Z.; Galon, J.; et al. Cytoscape plug-in to decipher functionally grouped gene ontology and pathway annotation networks. *Bioinformatics* **2009**, *25*, 1091–1093. [[CrossRef](#)] [[PubMed](#)]
22. Dhasmana, A.; Jamal, Q.M.; Gupta, R.; Siddiqui, M.H.; Kesari, K.K.; Wadhwa, G.; Khan, S.; Haque, S.; Lohani, M. Titanium dioxide nanoparticles provide protection against polycyclic aromatic hydrocarbon BaP and chrysene-induced perturbation of DNA repair machinery: A computational biology approach. *Biotechnol. Appl. Biochem.* **2016**, *63*, 497–513. [[CrossRef](#)] [[PubMed](#)]
23. Morris, G.M.; Huey, R.; Lindstrom, W.; Sanner, M.F.; Belew, R.K.; Goodsell, D.S.; Olson, A.J. AutoDock4 and AutoDockTools4: Automated docking with selective receptor flexibility. *J. Comput. Chem.* **2009**, *30*, 2785–2791. [[CrossRef](#)]
24. Huberman, B.; Alto, P.; Kaneko, K.; Ouyang, Q. *Handbook on Biological Networks*; World Scientific Publishing Co. Pte. Ltd. 5 Toh Tuck Link: Singapore, 2009; Volume 10, ISBN 9789812838797.
25. Junker, B.H.; Schreiber, F. *Analysis of Biological Networks*; A Wiley Interscience Publication: New York, NY, USA, 2008; ISBN 9780470041444.
26. Pavlopoulos, G.A.; Maria Secrier, M.; Moschopoulos, C.N.; Soldatos, T.G.; Kossida, S.; Aerts, J.; Schneider, R.; Bagos, P.G. Using graph theory to analyze biological networks. *BioData Min.* **2011**, *4*, 10. [[CrossRef](#)]
27. Chin, C.H.; Chen, S.H.; Wu, H.H.; Ho, C.W.; Ko, M.T.; Lin, C.Y. Cytohubba: Identifying hub objects and sub-networks from complex interactome. *BMC Syst. Biol.* **2014**, *8* (Suppl. 4), S11. [[CrossRef](#)] [[PubMed](#)]
28. Fisher, R.P. Secrets of a double agent: CDK7 in cell-cycle control and transcription. *J. Cell Sci.* **2005**, *118*, 5171–5180. [[CrossRef](#)] [[PubMed](#)]
29. Bartkova, J.; Zemanova, M.; Bartek, J. Expression of CDK7/CAK in normal and tumor cells of diverse histogenesis, cell-cycle position and differentiation. *Int. J. Cancer.* **1996**, *66*, 732–737. [[CrossRef](#)]
30. Yang, R.; Müller, C.; Huynh, V.; Fung, Y.K.; Yee, A.S.; Koeffler, H.P. Functions of cyclin A1 in the cell cycle and its interactions with transcription factor E2F-1 and the Rb family of proteins. *Mol. Cell. Biol.* **1999**, *19*, 2400–2407. [[CrossRef](#)]
31. Ray, A.; James, M.K.; Larochelle, S.; Fisher, R.P.; Blain, S.W. P27kip1 inhibits cyclin D-cyclin-dependent kinase 4 by two independent modes. *Mol. Cell. Biol.* **2009**, *29*, 986–999. [[CrossRef](#)] [[PubMed](#)]
32. Liccardi, G.; Ramos, G.L.; Tenev, T.; Annibaldi, A.; Legrand, A.J.; Robertson, D.; Feltham, R.; Anderton, H.; Darding, M.; Peltzer, N.; et al. RIPK1 and Caspase-8 Ensure Chromosome Stability Independently of Their Role in Cell Death and Inflammation. *Mol. Cell* **2019**, *73*, 413–428. [[CrossRef](#)]
33. Furusawa, Y.; Yamanouchi, Y.; Iizumi, T.; Zhao, Q.; Mitsuhashi, Y.; Morita, A.; Enomoto, A.; Tabuchi, Y.; Kondo, T. Checkpoint kinase 2 is dispensable for regulation of the p53 response but is required for G2/M arrest and cell survival in cells with p53 defects under heat stress. *Apoptosis* **2017**, *22*, 1225–1234. [[CrossRef](#)]
34. Giráldez, S.; Galindo-Moreno, M.; Limón-Mortés, M.C.; Rivas, A.C.; Herrero-Ruiz, J.; Mora-Santos, M.; Sáez, C.; Japón, M.Á.; Tortolero, M.; Romero, F. G1/S phase progression is regulated by PLK1 degradation through the CDK1/βTrCP axis. *FASEB J.* **2017**, *31*, 2925–2936. [[CrossRef](#)]
35. Renshaw, S.A.; Dempsey, C.E.; Barnes, F.A.; Bagstaff, S.M.; Dower, S.K.; Bingle, C.D.; Whyte, M.K. Three novel Bid proteins generated by alternative splicing of the human Bid gene. *J. Biol. Chem.* **2004**, *279*, 2846–2855. [[CrossRef](#)]
36. Pennisi, R.; Antocchia, A.; Leone, S.; Ascenzi, P.; di Masi, A. Hsp90α regulates ATM and NBN functions in sensing and repair of DNA double-strand breaks. *FEBS J.* **2017**, *284*, 2378–2395. [[CrossRef](#)]
37. Mullan, P.B.; Quinn, J.E.; Harkin, D.P. The role of BRCA1 in transcriptional regulation and cell cycle control. *Oncogene* **2006**, *25*, 5854–5863. [[CrossRef](#)]
38. Timofeev, O.; Cizmecioglu, O.; Settele, F.; Kempf, T.; Hoffmann, I. Cdc25 phosphatases are required for timely assembly of CDK1-cyclin B at the G2/M transition. *J. Biol. Chem.* **2010**, *285*, 16978–16990. [[CrossRef](#)]
39. Neganova, I.; Vilella, F.; Atkinson, S.P.; Lloret, M.; Passos, J.F.; von Zglinicki, T.; O'Connor, J.E.; Burks, D.; Jones, R.; Armstrong, L.; et al. An important role for CDK2 in G1 to S checkpoint activation and DNA damage response in human embryonic stem cells. *Stem Cells* **2011**, *29*, 651–659. [[CrossRef](#)]
40. De Boer, L.; Oakes, V.; Beamish, H.; Giles, N.; Stevens, F.; Somodevilla-Torres, M.; Desouza, C.; Gabrielli, B. Cyclin A/cdk2 coordinates centrosomal and nuclear mitotic events. *Oncogene* **2008**, *27*, 4261–4268. [[CrossRef](#)]
41. Brown, N.R.; Lowe, E.D.; Petri, E.; Skamnaki, V.; Antrobus, R.; Johnson, L.N. Cyclin B and cyclin A confer different substrate recognition properties on CDK2. *Cell Cycle* **2007**, *6*, 1350–1359. [[CrossRef](#)]

42. Goto, H.; Natsume, T.; Kanemaki, M.T.; Kaito, A.; Wang, S.; Gabazza, E.C.; Inagaki, M.; Mizoguchi, A. Chk1-mediated Cdc25A degradation as a critical mechanism for normal cell cycle progression. *J. Cell Sci.* **2019**, *132*, jcs223123. [[CrossRef](#)]
43. Sleeth, K.M.; Sørensen, C.S.; Issaeva, N.; Dziegielewska, J.; Bartek, J.; Helleday, T. RPA mediates recombination repair during replication stress and is displaced from DNA by checkpoint signalling in human cells. *J. Mol. Biol.* **2007**, *373*, 38–47. [[CrossRef](#)]
44. Grudic, A.; Jul-Larsen, A.; Haring, S.J.; Wold, M.S.; Lønning, P.E.; Bjerkvig, R.; Bøe, S.O. Replication protein A prevents accumulation of single-stranded telomeric DNA in cells that use alternative lengthening of telomeres. *Nucleic Acids Res.* **2007**, *35*, 7267–7278. [[CrossRef](#)]
45. Bakkenist, C.J.; Kastan, M.B. DNA damage activates ATM through intermolecular autophosphorylation and dimer dissociation. *Nature* **2003**, *421*, 499–506. [[CrossRef](#)] [[PubMed](#)]
46. Taira, N.; Yamamoto, H.; Yamaguchi, T.; Miki, Y.; Yoshida, K. ATM augments nuclear stabilization of DYRK2 by inhibiting MDM2 in the apoptotic response to DNA damage. *J. Biol. Chem.* **2010**, *285*, 4909–4919. [[CrossRef](#)] [[PubMed](#)]
47. Wang, Z.; Xie, Y.; Zhang, L.; Zhang, H.; An, X.; Wang, T.; Meng, A. Migratory localization of cyclin D2-Cdk4 complex suggests a spatial regulation of the G1-S transition. *Cell Struct. Funct.* **2008**, *33*, 171–183. [[CrossRef](#)] [[PubMed](#)]
48. Zaragoza, K.; Bégay, V.; Schuetz, A.; Heinemann, U.; Leutz, A. Repression of transcriptional activity of C/EBPalpha by E2F-dimerization partner complexes. *Mol. Cell. Biol.* **2010**, *30*, 2293–2304. [[CrossRef](#)] [[PubMed](#)]
49. Guo, A.; Salomoni, P.; Luo, J.; Shih, A.; Zhong, S.; Gu, W.; Pandolfi, P.P. The function of PML in p53-dependent apoptosis. *Nat. Cell Biol.* **2000**, *2*, 730–736. [[CrossRef](#)] [[PubMed](#)]
50. Huarte, M.; Guttman, M.; Feldser, D.; Garber, M.; Koziol, M.J.; Kenzelmann-Broz, D.; Khalil, A.M.; Zuk, O.; Amit, I.; Rabani, M.; et al. A large intergenic noncoding RNA induced by p53 mediates global gene repression in the p53 response. *Cell* **2010**, *142*, 409–419. [[CrossRef](#)]
51. Ren, S.; Rollins, B.J. Cyclin C/cdk3 promotes Rb-dependent G0 exit. *Cell* **2004**, *117*, 239–251. [[CrossRef](#)]
52. Maréchal, A.; Li, J.M.; Ji, X.Y.; Wu, C.S.; Yazinski, S.A.; Nguyen, H.D.; Liu, S.; Jiménez, A.E.; Jin, J.; Zou, L. PRP19 transforms into a sensor of RPA-ssDNA after DNA damage and drives ATR activation via a ubiquitin-mediated circuitry. *Mol. Cell* **2014**, *53*, 235–246. [[CrossRef](#)]
53. Jamal, Q.M.S.; Dhasmana, A.; Lohani, M.; Firdaus, S.; Ansari, M.Y.; Sahoo, G.C.; Haque, S. Binding pattern elucidation of NNK and NNAL cigarette smoke carcinogens with NER pathway enzymes: An onco-informatics study. *Asian Pac. J. Cancer Prev.* **2015**, *16*, 5311–5317. [[CrossRef](#)]
54. Huang, R.Y.; Li, M.Y.; Hsin, M.K.; Underwood, M.J.; Ma, L.T.; Mok, T.S.; Warner, T.D.; Chen, G.G. 4-Methylnitrosamino-1-3-pyridyl-1-butanone (NNK) promotes lung cancer cell survival by stimulating thromboxane A2 and its receptor. *Oncogene* **2011**, *30*, 106–116. [[CrossRef](#)]
55. Boccaletti, S.; Latora, V.; Moreno, Y.; Chavez, M.; Hwang, D.U. Complex networks: Structure and dynamics. *Phys. Rep.* **2006**, *424*, 175–308. [[CrossRef](#)]
56. Koutsogiannouli, E.; Papavassiliou, A.G.; Papanikolaou, N.A. Complexity in cancer biology: Is systems biology the answer? *Cancer Med.* **2013**, *2*, 164–177. [[CrossRef](#)] [[PubMed](#)]
57. Kitano, H. Systems Biology: A brief overview. *Science* **2002**, *295*, 162–1664. [[CrossRef](#)] [[PubMed](#)]
58. Bosley, A.D.; Das, S.; Andresson, T. A Role for Protein–Protein Interaction Networks in the Identification and Characterization of Potential Biomarkers. In *Proteomic and Metabolomic Approaches to Biomarker Discovery*; Issaq, H.J., Ed.; Academic Press: London, UK, 2013; Chapter 21; pp. 333–347. ISBN 9780123947956.
59. Klipp, E.; Liebermeister, W.; Wierling, C.; Kowald, A.; Lehrach, H.; Herwig, R. *Systems Biology: A Textbook*; Wiley-VCH: Weinheim, Germany, 2009; ISBN 978-3-527-31874-2.
60. Liu, J.; Xiong, Q.; Shi, W.; Shi, X.; Wang, K. Evaluating the importance of nodes in complex networks. *Phys. A Stat. Mech. Appl.* **2016**, *452*, 209–219. [[CrossRef](#)]
61. Yu, H.; Kim, P.M.; Sprecher, E.; Trifonov, V.; Gerstein, M. The importance of bottlenecks in protein networks: Correlation with gene essentiality and expression dynamics. *PLoS Comput. Biol.* **2007**, *3*, 713–720. [[CrossRef](#)] [[PubMed](#)]

62. Hustedt, N.; Durocher, D. The control of DNA repair by the cell cycle. *Nat. Cell Biol.* **2017**, *19*, 1–9. [[CrossRef](#)] [[PubMed](#)]
63. Huang, R.Y.; Chen, G.G. Cigarette smoking, cyclooxygenase-2 pathway and cancer. *Biochim. Biophys. Acta Rev. Cancer* **2011**, *1815*, 158–169. [[CrossRef](#)]



© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).