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RESEARCH ARTICLE

COMPUTATIONAL STUDY ON PROTEIN INTERACTIONS OF CDKN2C AND ITS IMPACT ON CANCER STUDIES

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CDKN2C protein; CCND1; Osteosarcoma; Multiple myeloma; G1 phase arrest; co-expression

Abstract

CDKN2C is a tumour suppressor that suppress bone cancers like osteosarcoma and multiple myeloma by regulating cell cycle. Deletions or mutations in the gene can lead to tumour progression and worst survival however over expression leads to radiation sensitivity, G1 phase arrest and trigger apoptosis in osteosarcoma. Study on CDKN2C protein and its role in bone cancer is right now gaining evidence and few of the proteins like CDKN1C, CDKN2D, CCNL2 show primary level interactions and gene CDK4 is found to be co-expressed with CCND1 through prediction analysis and observed co-expression of homologs in other species through STRING database tool.

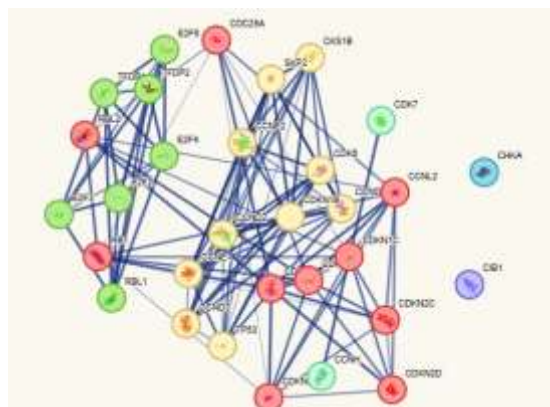
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Introduction:-

Limited radiosensitivity to osteosarcoma is one of the major posing challenge and studies of Qiujian Lian et al 2024 the protein CDKN2C increases radiosensitivity by causing G1 phase cell cycle arrest and subsequent apoptosis. G1 phase arrest is mediated by deregulates the expression of phosphorylated RB protein. Deletions at chromosome 1 is identified in 7 to 4% of the multiple myeloma cancers and CDKN2C presence at 1p32.3 has been identified as one of the potential targets for deletion in multiple myeloma patients (Qiujian Lian et al 2024). One of the examples is homozygous deletion of the gene (Cox C et al., 2005). Using high resolution mapping the deletion of the gene is identified in about 40% of the multiple myeloma cell lines (Kulkarni MS et al., 2002, Dib A et al., 2006).

CDKN2C deviation in expression is known to be recorded with so many cancers but for the first time reports on its upregulation in expression in SCLC at both gene and protein level is done by Guo-Sheng Li et al., 2022 and CDKN2C expression is related to immune microenvironment and it can serve as one of the important prognostic marker in immune therapy. Immune checkpoint inhibitor related treatment is one of the promising treatments for SCLC but however its benefits is limited to few of the patient's due to lack of availability in biomarkers (Esposito G et al 2020).

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CDKN1C, CDKN2D, CCNL2 are the few among the proteins that show primary level of interactors with CDKN2C and whereas CDKN1C, E2F4 are the second level of interactors of protein CDKN2C through proteins CCNL2.

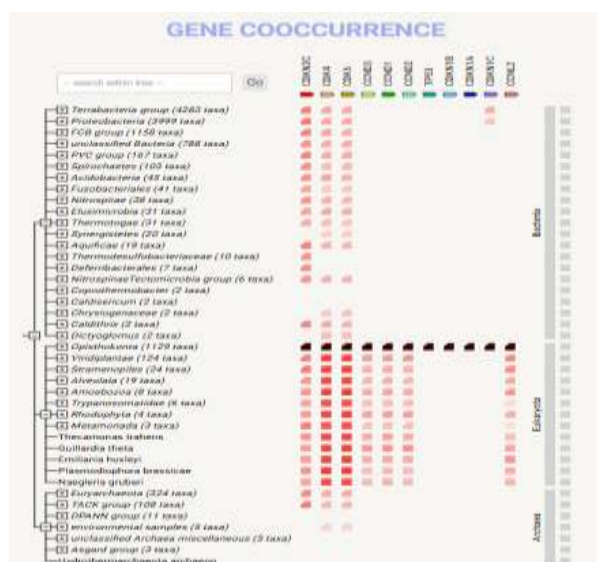


Figure:2 Study of Gene Co-occurrence of protein CDKN2C among different taxa

k means clustering algorithm and working principle:-

k means clustering algorithm is an unpredicted machine learning methodology that clusters data based on predefined number (K) of distinct non overlapping groups with feature similarity. The goal of the algorithm is to minimise the variation and to group the defined points with total similarity in to one group and also to be distinct entirely when marked with another group in a cluster. The algorithm assigns data using Euclidean distance of the data points to the nearest centroid and recalculates the mean of the assigned points to the centroid and the process repeats with initial randomly chosen centroids until the assigned cluster stabilise and indicates convergence as same group.

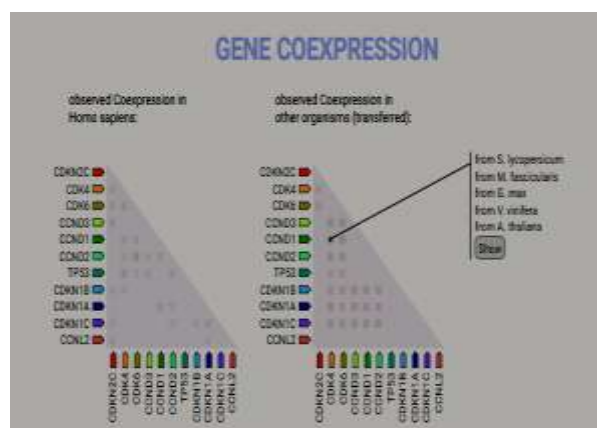


Figure: 3 Gene Co-expression studies of CDKN2C protein in homo sapiens and other related organisms.

Based on Predicted association, CDK4 and CCND1 observed co-expression of homologs in other species through STRING database tool.

Study of transcripts for the gene RANKL using AUGUSTUS tool using FASTA sequence collected from NCBI database:-

AUGUSTUS is a popular open-source software tool used for eukaryotic gene prediction, and utilizes a Generalized Hidden Markov Model (GHMM) to predict gene structures based on sequence alone or incorporates evidence from RNA-SeqESTs, and protein alignments through performing ab initio prediction. It is available both as via web interface (WebAUGUSTUS) or local installation through Git hub.

RANKL: Comparison of RANKL transcripts and protein sequence between Homo sapiens (GenBank: AF019047.1) and mus musculus {(GenBank: AF019048.1) curated from NCBI database using mRNA sequence as the genome input} selecting options both strands and only complete genes with alternative transcripts as few instead of many or none.

Homo sapiens:-

```
ATGCGCCGCGCCAGCAGAGACTACACCAAGTACCTGCGTGGCTCGGAGGAGATGGGCGGCGGCCCGG
GAGCCCCGCACGAGGGCCCCCTGCACGCCCCGCCGCCCTGCGCCGACCAGCCCCCGCGCCTCC
CGCTCCATGTTCTGTTGGCCCTCCTGGGGCTGGGGCTGGGCCAGGTTGTCTGCAGCGTCGCCCTGTTCTTC
TATTTTCAGAGCGCAGATGGATCCTAATAGAATATCAGAAGATGGCACTCACTGCAATTTATAGAATTTT
GAGACTCCATGAAAATGCAGATTTTCAAGACACAACCTCTGGAGAGTCAAGATACAAAATTAATACCTG
ATTTCATGTAGGAGAATTAACAGGCCTTTCAAGGAGCTGTGCAAAAGGAATTACAACATATCGTTGGA
TCACAGCACATCAGAGCAGAGAAAGCGATGGTGGATGGCTCATGGTTAGATCTGGCCAAGAGGAGCA
AGCTTGAAGCTCAGCCTTTTGTCTCATCTCACTATTAATGCCACCGACATCCCATCTGGTTCCCATAAAG
TGAGTCTGTCTCTTGGTACCATGATCGGGGTTGGGCCAAGATCTCCAACATGACTTTTAGCAATGGAA
AACTAATAGTTAATCAGGATGGCTTTTATTACCTGTATGCCAACATTTGCTTTTCGACATCATGAACTT
CAGGAGACCTAGCTACAGAGTATCTTCAACTAATGGTGTACGTCATAAAACCAGCATCAAAATCCCA
AGTTCTCATACCCTGATGAAAGGAGGAAGCACCAAGTATTGGTCAGGGAATTCTGAATTCATTTTAA
TTCCATAAACGTTGGTGGATTTTTTAAGTTACGGTCTGGAGAGGAAATCAGCATCGAGGTCTCCAACC
CCTCCTTACTGGATCCGGATCAGGATGCAACATACTTTGGGGCTTTTAAAGTTCGAGATATAGATTGA
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mus musculus:-

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ATGCGCCGGGCCAGCCGAGACTACGGCAAGTACCTGCGCAGCTCGGAGGAGATGGGCAGCGGCCCGG
GCGTCCCACACGAGGGTCCGCTGCACCCCGCGCCTTCTGCACCGGCTCCGGCGCCGCCACCCGCCGCC
TCCCGCTCCATGTTCTTGGCCCTCCTGGGGCTGGGACTGGGCCAGGTGGTCTGCAGCATCGCTCTGTTT
CTGTACTTTTCGAGCGCAGATGGATCCTAACAGAATATCAGAAGACAGCACTCACTGCTTTTATAGAAT
CCTGAGACTCCATGAAAACGCAGATTTGCAGGACTCGACTCTGGAGAGTGAAGACACACTACCTGACT
CCTGCAGGAGGATGAAACAAGCCTTTTCAGGGGGCCGTCGAGAAGGAAGTGAACACATTGTGGGGCC
ACAGCGCTTCTCAGGAGCTCCAGCTATGATGGAAGGCTCATGGTTGGATGTGGCCAGCGAGGCAAGC
CTGAGGCCCAGCCATTTGCACACCTCACCATCAATGCTGCCAGCATCCCATCGGGTCCCATAAAGTC
ACTCTGTCCTCTTGGTACCACGATCAGGCTGGGCCAAGATCTCTAACATGACGTTAAGCAACGGAAA
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ACTAAGGGTTAACCAAGATGGCTTCTATTACCTGTACGCCAACATTTGCTTTCGGGCATCATGAAACATC
GGGAAGCGTACCTACAGACTATCTTCAGCTGATGGTGTATGTCGTTAAACCAGCATCAAAATCCCAA
GTTCTCATAACCTGATGAAAGGAGGGAGCACGAAAACTGGTCGGGCAATTCTGAATTCCACTTTTAT
TCCATAAATGTTGGGGGATTTTCAAGCTCCGAGCTGGTGAAGAAATTAGCATTACAGGTGTCCAACCC
TTCCCTGCTGGATCCGGATCAAGATGCGACGTACTTTGGGGCTTCAAAGTTCAGGACATAGACTGA

Amino acid sequence:-

Homo sapiens:-

MRRASRDYTKYLRGSEEMGGGPGAPHEGPLHAPPPAPHQPPAASRSMFVALLGLGLGQVVCVSLFFYF
RAQMDPNRISEDGTHCIYRILRLHENADFQDTTLESQDTKLIPDSCRRIKQAFQGA VQKELQHIVGSQHIRAE
KAMVDGSWLDLAKRSKLEAQPF AHLTINATDIPSGSHKVSLSWYHDRGWAKISNMTFSNGKLIVNQDGF
YYLYANICFRHHETSGDLATEYLQLMVYVTKTSIKIPSSHTLMKGGSTKYWSGNSEHFHYSINVG GFFKLRS
GEEISIEVSNPSLLDPDQDATYFGAFKVRDID

Mus Musculus:-

MRRASRDYGKYLRSSSEEMGSGPGVPHEGPLHPAPSAPAPAPPPAASRSMFLALLGLGLGQVVCVSLFLYF
RAQMDPNRISEDSTHCFYRILRLHENADLQDSTLESED TLPDSCR RMKQAFQGA VQKELQHIVGPQRFSGA
PAMMEGSWLDVAQRGKPEAQPF AHLTINAASIPSGSHKVTLSWYHDRGWAKISNMTLSNGKLRVNQDGF
FYYLYANICFRHHETSGSVPTDYLQLMVYVTKTSIKIPSSHTLMKGGSTKNWSGNSEHFHYSINVG GFFKLRS
AGEEISIQVSNPSLLDPDQDATYFGAFKVRDID

Study of Transcripts in protein Matrilin 3 in Homo sapiens present on chromosome 2 using AUGUSTUS online tool : NCBI Reference Sequence: NG_008087.1:

Nucleotide Transcripts:-

GTGTTTGCAAGAGCAGACCCTTGGACCTGGTGTTTATCATTGATAGTTCTCGTAGCGTACGGCCCCTGG
AATTCACCAAAGTGAAAAC TTTTGTCTCCCGGATAATCGACACTCTGGACATTGGGCCAGCCGACACG
CGGGTGGCAGTGGTGA ACTATGCTAGCACTGTGAAGATCGAGTTCCA ACTCCAGGCCTACACAGATAA
GCAGTCCCTGAAGCAGGCCGTGGGTGCAATCACACCCTTGTC AACAGGCACCATGTCAGGCCTAGCCA
TCCAGACAGCAATGGACGAAGCCTTCACAGTGGAGGCAGGGGCTCGAGAGCCCTCTTCTAACATCCCT
AAGGTGGCCATCATTGTTACAGATGGGAGGCCCCAGGACCAGGTGAATGAGGTGGCGGCTCGGGCCC
AAGCATCTGGTATTGAGCTCTATGCTGTGGGCGTGGACCGGCGAGACATGGCGTCCCTCAAGATGATG
GCCAGTGAGCCCCTAGAGGAGCATGTTTTCTACGTGGAGACCTATGGGGTTCATTGAGAACTTTCTCTC
TAGATTCCAGGAAACCTTCTGTGCGCTGGACCCCTGTGTGCTTGGAACACACCAGTGCCAGCACGTCT
GCATCAGTGATGGGGAAGGCAAGCACTGTGAGTGTAGCCAAGGATACACCTTGAATGCCGACAA
GAAAACGTGTT CAGGTGAGGCCTGTGTAGGGGGCCGTGACTGA

Protein sequence:-

VCKSRPLDLVFIISSRSVRPLEFTKVKT FVSRIIDTLDIGPADTRVAVVNYASTVKIEFQLQAYTDKQSLKQ
AVGRITPLSTGTMSGLAIQTAMDEAFTVEAGAREPSSNIPKVAIIVTDGRPQDQVNEVAARAQASGIELYAV
GVD RADMASLKMMASEPLEEHVFYVETYGVIEKLSSRFQETFCALDPCVLGTHQCQHVCISDGEGKHHCE
CSQGYTLNADKKTCSGEACVGGRD

Conclusion:-

By this study we can conclude study of CDKN2C protein interactions and coexpression analysis can help in prediction of biomarkers like CDK4, CCND1 for targeting the cell cycle progression in cells with deletions of CDKN2C gene.

References:-

1. Qiujian Lian, Haonan Zhao, Bingxuan Wang, Ping Ling, Jingyan Li, Peijun Dai, Junyong Ge, Xu Su, Zhiwei Wang, Suchi Qiao (2024). Enhancing radiosensitivity in osteosarcoma via CDKN2C overexpression: A mechanism involving G1 phase arrest mediated by inhibition of CDK4 expression and Thr172 phosphorylation, Biochemical and Biophysical Research Communications, Vol. 735,150840, <https://doi.org/10.1016/j.bbrc.2024.150840>.
2. Cox C, Bignell G, Greenman C, et al. (2005). A survey of homozygous deletions in human cancer genomes. Proc Natl Acad Sci U S A; 102; 4542–7. <https://doi.org/10.1073/pnas.0408593102>

3. Kulkarni MS, Daggett JL, Bender TP, Kuehl WM, Bergsagel PL, Williams ME (2002) . Frequent inactivation of the cyclin-dependent kinase inhibitor p18 by homozygous deletion in multiple myeloma cell lines: ectopic p18 expression inhibits growth and induces apoptosis. *Leukemia*; 16;127–34.<https://doi.org/10.1038/sj.leu.2402328>
4. Dib A, Peterson TR, Raducha-Grace L, et al. (2006). Paradoxical expression of INK4c in proliferative multiple myeloma tumors: bi-allelic deletion vs increased expression. *Cell Div*;1: 23.<https://doi.org/10.1186/1747-1028-1-23>
5. Li GS, Chen G, Liu J, Tang D, Zheng JH, Luo J, Jin MH, Lu HS, Bao CX, Tian J, Deng WS, Fu JW, Feng Y, Zeng NY, Zhou HF, Kong JL. Clinical significance of cyclin-dependent kinase inhibitor 2C expression in cancers: from small cell lung carcinoma to pan-cancers. *BMC Pulm Med*. 2022 Jun 24;22(1):246. <https://doi.org/10.1186/s12890-022-02036-5> PMID: 35751045; PMCID: PMC9233395.
6. Esposito G, Palumbo G, Carillio G, Manzo A, Montanino A, Sforza V, Costanzo R, Sandomenico C, La Manna C, Martucci N, et al.(2020) Immunotherapy in small cell lung cancer. *Cancers (Basel)*;12(9):2522. <https://doi.org/10.3390/cancers12092522>. [DOI] [PMC free article] [PubMed] [Google Scholar].