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CBL-interacting protein kinases (CIPKs) in Chickpea: Genome-wide identification, structure and expression analysis under abiotic stresses and development

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Abstract: Calcineurin B-like proteins (CBL)-interacting protein kinases (CIPKs) by interacting with CBLs regulate developmental processes, hormone signalling transduction and mediate stress responses in plants. Although the genome of chickpea is available, information of CIPK gene family has been missing in chickpea. Here, a total of 22 CIPK encoding genes were identified in chickpea and characterized by *in silico* methods. We found a high structural conservation in chickpea CIPK family. Our analysis showed that chickpea CIPKs have evolved with dicots from common ancestors, and extensive gene duplication events have played an important role in evolution and expansion of CIPK family in chickpea. Most chickpea CIPK proteins localize in cytoplasm and nucleus. Promoter analysis revealed various cis-regulatory elements related to plant development, hormone signaling and abiotic stresses. Expression analysis indicated that CIPKs are significantly expressed in a spectrum of developmental stages, tissue/organs that hinted their important role in plant development. Several CIPK genes had specific and overlapping expressions in different abiotic stresses and seed development stages, suggesting the important role of CIPK family in abiotic stress signaling, and seed development in chickpea. Thus, this study provides the avenue for detailed functional characterization of CIPK family in chickpea and other legume crops.

Keywords: CIPK, Calcium, Chickpea, Structure, Phylogeny, Expression, Stress, Development.

1. Introduction

South Asia is the major producer of the world's second most important food legume chickpea. Importantly, India is the largest producer of chickpeas and is credited for about 70% of world's chickpeas production. India contributes an estimated production of 5.9 million tonnes (mt) annually [1]. Chickpea is an important dietary source for vegetarians due to vital nutritive constituents in its seeds, including 20–30% crude protein, 40% carbohydrate, and 3–6% oil [2]. In addition, chickpea seeds are rich in minerals, such as calcium, magnesium, potassium, phosphorus, iron and zinc [3]. The chickpea production is severely affected by various stresses. Consequently, a huge gap is developed between its demand and supply. Abiotic stresses alone account for an estimated 40-60% global chickpea production losses annually. Drought causes major damage and accounts for about 50% of chickpea yield loss. Temperature fluctuations and soil salinity combined are responsible for about 25% of chickpea yield loss [4]. The chickpea yield loss due to drought, cold and salinity, respectively costed approximately 1.3 billion, 186 million and 354 million US dollars, which have economically dented several chickpea-producing countries [5]. These stresses have an adverse effect on flower set, pollen viability, pod set/abortion and retention. As all these crucial developmental stages essentially determine seed number, a negative impact on these stages significantly hampers chickpea yield. Due to

continuous climate change, severe and frequent challenges of drought in arid and semi-arid areas where chickpea is traditionally cultivated are predicted [6] and that can be detrimental for overall productivity of chickpea. Thus, identification and utilization of important stress related genes in biotechnological programmes to generate improved chickpea varieties is need of the hour.

Environmental cues, such as biotic and abiotic stresses are known to elicit the increase in cytosolic Ca2+ with specific spatio-temporal features. The spatio-temporal accumulation of Ca2+ generates specific "Ca2+ signature" in the form of spikes, waves and oscillations. The stimulus specific Ca2+ signature is decoded by Ca2+ sensors and downstream effectors towards a response [7]. Several Ca2+ sensors have been identified and characterized in plants, including calmodulin (CaM) and CaM-like proteins (CMLs) [8], Ca2+ -dependent protein kinases (CDPKs) [9], and calcineurin B-like proteins (CBLs) [1]. Among these, CBLs are a unique group of Ca2+ sensors and to determine their functional identity, a family of plant specific serine/threonine kinases; CBL-interacting protein kinases (CIPKs) functions as important downstream signaling component [10]. Both CBLs and CIPKs are encoded as multi-gene families in higher plants, for example: 10 CBL and 26 CIPKs members have been identified in Arabidopsis, and 10 CBL and 30 CIPKs in rice [11]. Large numbers of CBL and CIPK members in higher plants constitute a complex and sophisticated signaling network. For instance, In Arabidopsis, each CBL interacts with multiple CIPKs and vice-versa [12], consequently, some CBLs share a common CIPK partner and some CIPKs are regulated by a common CBL. Such specific and overlapping patterns of CBL-CIPK interactions may provide functional specificity and synergism to CBL-CIPK signalling networks.

Structurally, CBLs are typical Ca2+ sensor proteins with four EF-hand domains which are responsible for Ca2+ binding. On the other hand, CIPKs harbours several functionally distinct domains. All CIPKs consist of a conserved catalytic kinase domain at the N-terminal and a regulatory domain at the C-terminal [10, 12]. A typical of a functional kinase protein, CIPK kinase domain contains an ATP binding site and an activation loop. The regulatory domain contains FISL/NAF and PPI motifs which are responsible for the interaction with CBL and type 2C protein phosphatases, respectively [13, 14]. The function of CBL-CIPK pathways could be regulated by pattern of gene expression, Ca2+ binding affinity, protein stability and protein-protein interactions [7]. CBL-CIPK networks have been implicated in diverse functions that regulate plant response to biotic stress [15–17], abiotic stress [1, 18, 19], nutrient deficiency [1, 20, 21] metal toxicity [22, 23] and plant development [24–27]. Majority of the knowledge about CBL-CIPK signaling has developed from the research with model plant Arabidopsis thaliana. Information about CBL-CIPK networks and their role is scarce in important legume crop chickpea. Though, CBL family has been identified in chickpea [28], identification and characterization of CIPK family is missing. A comprehensive gene expression profiling of the CIPK family will help in understanding CBL-CIPK functions in chickpea. Information obtained from expression analysis will encourage the utilization of crucial genes for genetically engineering the chickpea plant towards better stress tolerance and development.

With this rationale, we have identified the CIPK gene family in chickpea. Phylogenetic analysis and chromosomal localization have provided insight into the evolution and expansion of chickpea CIPK family. Analysis of gene and domain structure ensured the authenticity and integrity of identified genes. Homology modelling helped to understand the three-dimensional structure of chickpea CIPKs. In-silico analysis revealed various stress, hormone and development related cis-regulatory elements in CIPK promoters. Expression profiling using various datasets in public repositories suggested involvement of the CIPK family in biotic and abiotic stress signaling, and seed development in chickpea.

2. Materials and Methods

2.1. Identification of CIPKs in the chickpea genome

The chickpea genome submitted by Varshney et al. (2013) was downloaded from the NCBI and explored to identify CIPK encoding genes. Rice and Arabidopsis thaliana CIPK proteins were retrieved from Uniprot (Swiss-Prot), and homology search was performed using BLAST tool (E-value =10-6) against the chickpea proteome. Significant hits were selected on the basis of >=50% identity, and >=100 amino acid length alignment. Further, the HMM sequence of CIPK-NAF domain was extracted from Pfam (http://pfam.xfam.org/) database, and BLAST search was done (E-value =10) against the chickpea proteome. Furthermore, both the sets of putative candidates were mixed, and redundant sequences were removed using CD-HIT [29]. The domain analysis was performed by using a standalone version of the InterproScan [30]. The gene attributes such as gene ID, protein ID, CDS, size of amino acid and chromosomal coordinates were extracted from NCBI web server.

2.2. Phylogenetic analysis

To examine the evolutionary relationship between CIPKs in chickpea and other species, Multiple Sequence Alignment (MSA) was performed with the amino acid sequences of CIPKs from four different plant species e.g., Arabidopsis thaliana, Oryza sativa, Glycine max, and Cicer arietinum) using ClustalW [31] at default settings in MEGA X version 10.1.8 [32]. The neighbour-joining method was used to construct the phylogenetic tree and bootstrap values were calculated in 1000 replicates to determine the phylogenetic relationship among the CIPKs. iTOL [33] webserver was used to mark the different clades of CIPKs with different colours and shapes for better visualization.

2.3. Gene structures, motif organization and domain prediction

Gene Structure Display Server (GSDS) program (http://gsds.gao-lab.org/) was used to compare the CDS sequences with their corresponding genomic DNA sequences in order to investigate the coding sequences and intron structure. Motif organization of CIPK proteins was examined via the Multiple Expression motifs for Motif Elicitation (MEME) tool [34] with default parameters; site distribution - zero or one occurrence per sequence; motif discovery mode - classic; motif length 6-50; and the top ten most enriched motifs were selected based on lowest E-values. The identification of domains was performed by a standalone package of InterProScan [30]. The coordinates of the essential domains and active sites were extracted and used as input in Illustrator for Biological Sequences [35] for the visualization.

2.4. Gene nomenclature, chromosomal location and gene duplication

The names of CIPK genes in chickpea were assigned according to their closest orthologous relationship with Arabidopsis CIPK genes in the phylogenetic tree. The information of the chromosome coordinates was obtained from NCBI. Their localization was displayed in different chromosomes using TBtools [36]. To search for all duplicated gene pairs within the chickpea genome, the protein sequence of chickpea was used to run the all-versus-all local BLASTP with parameters of E-value 1e-5, max target sequences 5, and m6 format output. MCScanX software package [37] was used to analyse the segmentally duplicated regions of CIPK genes of chickpea. The genes and the intra-species collinear gene pairs were mapped to the eight chromosomes of chickpea using the family_circle_plotter.java script. The protein sequences of each duplicate gene pair were aligned by CLUSTALW. The alignment file in FASTA format and the CDS sequences of the

corresponding genes were used to calculate the non-synonymous (Ka) and synonymous (Ks) substitution values by the PAL2NAL server [38].

2.5. In silico promoter analysis

For the identification of cis-regulatory elements in the promoters of chickpea CIPKs, 2000 bp upstream sequences of the coding region of genes were extracted from NCBI and used as input in the PlantCARE (http://bioinformatics.psb.ugent.be/webtools/plant-care/html/) tool.

2.6. Subcellular localization and physicochemical properties of CaCIPK proteins

The full-length protein sequences of all the CIPKs of chickpea were used as input to predict their subcellular localization using the CELLO program [39]. The locations were displayed in different parts of the cell by Biorender software (https://biorender.com/). The online tool Compute pI/MW of Expasy [40] was used to calculate the molecular weight (MW) and isoelectric point (pI) of CaCIPKs.

2.7. Protein-Protein Interaction Network Construction for CaCBLs -CaCIPKs

To elucidate the interaction network between CIPK and CBL proteins in chickpea, the amino acid sequences of 22 CIPKs, and 9 CBLs from the study of Meena et al., 2015 were used as input in STRING (http://string-db.org/). At STRING, the interaction network can be constructed using low confidence value of 0.15, medium confidence of 0.4, high confidence of 0.7 and highest confidence of 0.9. Experimental data of interacting CBL and CIPK proteins in Arabidopsis were constructed using the confidence value > 0.4. Homologous proteins of the determined interactive Arabidopsis proteins in chickpea were identified by reciprocal best BLASTP analysis.

2.8. Protein tertiary structure prediction

The tertiary structures of the 22 CaCIPK proteins were predicted with the Phyre2 web portal (http://www.sbg.bio.ic.ac.uk/phyre2). Phyre2 uses advanced remote homology detection methods to build 3D models for protein sequences [41]. All the proteins were modelled with 100% confidence by the single highest scoring template model.

2.9. Expression analysis using RNA-seq data

To generate the genome-wide expression profiles of CaCIPK genes in different tissues and developmental stages, RNA-Seq data was extracted from NCBI Sequence Read Archive; SRA number SRP121085. RNA-seq data for different seed stages in two distinct desi chickpea varieties (JGK3 and Himchana 1) was extracted from SRA number SRP072563 and SRP072564. Data for three abiotic stresses (desiccation, cold and salinity stress) in root, and shoot tissue of ICC4958 chickpea variety was extracted from SRA number SRP034839.

The raw reads downloaded from SRA at NCBI were processed using FASTP [42] to remove the adapter, poly-N, short and low-quality reads. The reference genome of chickpea was downloaded from the NCBI genome web server. The HISAT2 [43] tool was used for building the index of the reference genome, and for mapping of filtered reads onto the genome. The alignments were assembled into potential transcripts using StringTie [44] and the transcript abundance was calculated as fragments per kilobase of transcript per million reads (FPKM) values. For differential expression analysis, three biological replicates of each treatment and control were analysed and fold change expression was

calculated by the ratio of average FPKM of test samples and average FPKM of control samples. The 'pheatmap' package of R was used to generate the heatmaps of the expression data using the logarithm of normalized expression values for the tissue study and the logarithm of fold change for the remaining studies.

3. Results and Discussions

3.1. Identification and sequence analysis of CaCIPK genes

A total of 39 putative CIPK sequences were obtained from homology search with Arabidopsis and rice CIPKs. Further HMM profile search against chickpea proteome revealed 38 putative CIPK sequences. After combining both sets, followed by manual curation a total of 26 unique CIPK sequences were obtained. Domain analysis revealed that the necessary domains (e.g. PPI and NAF domain) were absent in four sequences, therefore, they were removed from the list. Finally, a total of 22 non-redundant CIPK encoding genes were found in the chickpea genome. Previously, 26 CIPK members have been reported in Arabidopsis [45] and 33 in rice [46]. The number of CIPKs in chickpea is comparable with wheat (20 members), tomato (22 members) and canola (23 members) [47–49].

The length of 22 CaCIPK proteins varied from 418aa (CaCIPK16) to 503aa (CaCIPK12) with average molecular weight of 51.16 kDa. Most of the CaCIPK proteins (except CaCIPK3 and CaCIPK11) were found to have an isoelectric point (pI) greater than 7 (Table 1).

Table 1: Various features of the chickpea CIPK family.

			Transcript					Protein			
Gene Name	NCBI ID	Chrom osome	Start	End	Identifier	Intron count	CDS length	Identifier	Length (aa)	Isoelectric point (pI)	Protein wt. (kDa)
CIPK1	LOC101499928	Ca1	25702213	25709285	XM_004488130.3	11	1347	XP_004488187.1	448	8.02	50.34
CIPK2	LOC101488582	Ca2	27180279	27182763	XM_004490648.3	0	1368	XP_004490705.1	455	9	51.9
CIPK3	LOC101491417	Ca2	33011673	33018054	XM_004491146.3	13	1326	XP_004491203.1	441	6.59	50.44
CIPK4	LOC101504371	Ca4	32563304	32564889	XM_004497948.3	0	1305	XP_004498005.1	434	9.12	49.29
CIPK5	LOC101513526	Ca6	53224442	53226557	XM_012717669.2	0	1353	XP_012573123.1	450	8.26	51.83
CIPK6	LOC101511702	Ca7	1293373	1294939	NM_001309656.1	0	1335	NP_001296585.1	444	9.11	50.34
CIPK7	LOC101496895	Ca1	8839486	8841422	XM_004486556.3	0	1347	XP_004486613.1	448	9.14	50.77
CIPK8	LOC101511605	Ca2	23829223	23836498	XM_027331998.1	14	1425	XP_027187799.1	474	8.03	53.78
CIPK9	LOC101493574	Ca1	36737555	36744930	XM_027334254.1	14	1398	XP_027190055.1	465	8.93	52.14
CIPK10	LOC101498077	Ca5	26642943	26645602	XM_004500288.3	0	1392	XP_004500345.1	463	8.76	52.43
CIPK11	LOC101499591	Ca5	26665837	26667956	XM_012715825.2	0	1320	XP_012571279.1	439	6.49	48.88
CIPK12	LOC101506657	Ca1	3314537	3316964	XM_012712534.2	0	1512	XP_012567988.1	503	7.14	56.66
CIPK13	LOC101489428	Ca5	28720785	28723545	XM_004500487.3	0	1380	XP_004500544.1	459	9.04	51.78
CIPK14	LOC101488926	Ca2	27205251	27207496	XM_004490649.3	1	1308	XP_004490706.1	435	8.67	48.82
CIPK15	LOC101507945	Ca6	37592424	37596217	XM_004506380.3	0	1395	XP_004506437.1	464	8.82	52.36
CIPK16	LOC101510050	Ca4	44315144	44317032	XM_004498761.3	0	1257	XP_004498818.1	418	8.94	47.54
CIPK17	LOC101510187	Ca7	3876530	3882455	XM_012717926.2	11	1344	XP_012573380.1	446	8.43	50.8
CIPK18	LOC101492060	Ca3	21042046	21046759	XM_004492487.3	0	1392	XP_004492544.1	463	8.41	52.63
CIPK19	LOC101507330	Ca7	6850555	6859747	XM_012718074.2	13	1341	XP_012573528.1	446	9.2	50.84

CIPK20	LOC101506991	Ca1	3307254	3309153	XM_012712535.2	0	1362	XP_012567989.1	453	8.96	52.14
CIPK21	LOC101512343	Ca7	2026868	2030503	XM_012717974.2	13	1359	XP_012573428.1	452	8.17	51.07
CIPK22	LOC101493164	Ca3	21099833	21101565	XM_004492489.3	0	1296	XP_004492546.1	431	8.89	48.89

This indicates that most CaCIPK proteins function in similar microenvironments. Similar structural features and functional pI had been previously reported in CIPKs of other plants such as P.mume [50], C.annum [51] and B.napus [49]. To gain insights into the homology of the CaCIPK proteins, the sequence identity and similarity was calculated by the SIAS tool (http://imed.med.ucm.es/Tools/sias.html). The analysis showed that different CaCIPKs have 46.13 to 81.87% sequence similarity among themselves. Four protein pairs; CaCIPK15/18, CaCIPK5/7, CaCIPK1/17 and CaCIPK2/10) showed a high degree of identity i.e., 76.67%, 76.39%, 76.01% and 73.84%, respectively (Figure S1). Even the most divergent protein pair CaCIPK8 and CaCIPK22 shared 35.49% identity (49.18% similarity). These findings suggest that plant CIPKs are highly conserved in terms of sequence and structure, which hints towards their similar mode of action.

3.2. Gene and domain structure

The evolution of gene families is often reflected by their gene structure [52, 53]. A large variation in the number of introns was found in CIPK genes in chickpeas with number of introns ranging from 0 to 14 (Figure 1A). Out of 22 CaCIPK genes, only seven had more than two introns. Thus, CaCIPKs could be classified into two groups: i) intron-poor subgroup with zero (CaCIPK2, -4, -5, -6, -7, -10, -11, -12, -13, -15, -16, -18, -20, -22) or one (CaCIPK14) intron, and ii) intron-rich subgroup with greater than 10 introns (CaCIPK1, -3, -8, -9, -17, -19, -21). Division of CIPKs into two subgroups was also supported by the division of clade in the phylogenetic tree. Similar pattern of intron-rich and intron-poor CIPK genes has been reported in different plant species, including Arabidopsis [54], rice [55], soybean [56], wild sugarcane [57] and wheat [47]. These findings suggest that CIPK gene family is structurally conserved across plant kingdom.
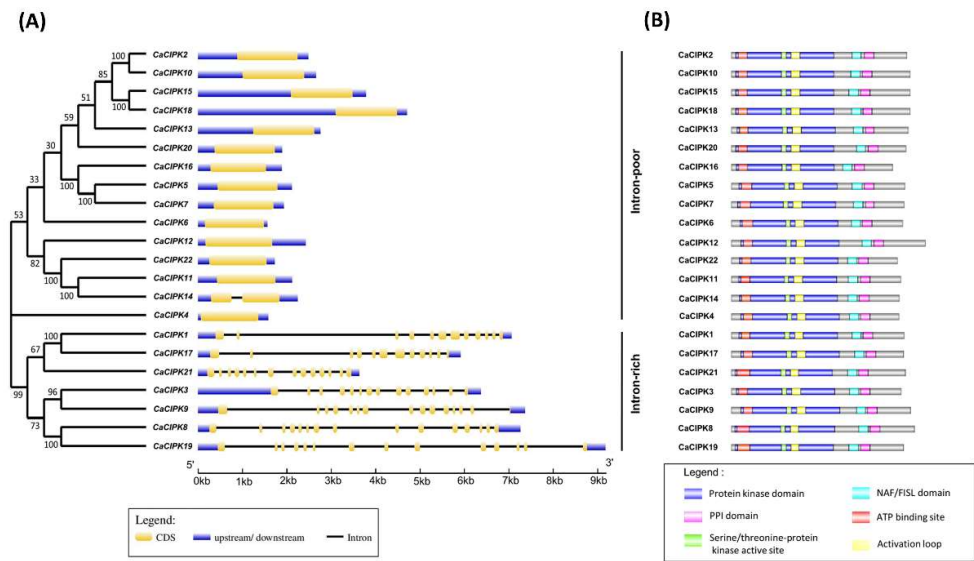

Figure 1. Structural features of chickpea CIPK family. (A) Exon-intron organization is shown for 22 CaCIPK genes. The values in the phylogenetic tree (left side) represent

bootstrap values. (B) Protein structure of chickpea CIPK family is showing conserved protein kinase, NAF and PPI domain along with some important sites present at the N-terminal e.g. ATP binding site, Serine/threonine-protein kinase active site and activation loop.

The domain structure analysis revealed that all the CaCIPK proteins possess three essential domains, at N-terminal a kinase domain, and at C-terminal regulatory NAF and protein phosphatase interaction (PPI) domain (Figure 1(B)). Kinase domains contain an ATP binding site and an activation loop. During CIPK activity, the stabilization of substrates at the active site is regulated by the phosphorylation of activation loop [58]. The activation loop has been found between the conserved 'DFG' and 'APE' amino acid residues. However, in our analysis, few variations were observed in the conserved short motifs. Alignment of 22 CaCIPK proteins showed that the glycine residue of 'DFG' was changed to asparagine (DFN) in CaCIPK1, whereas the alanine of 'APE' was modified to serine (SPE) in CaCIPK6 and CaCIPK18 (Figure S2). Importantly, sites important for phosphoregulation of activation loop i.e., serine, threonine, and tyrosine [59] were found to be conserved in all CaCIPK proteins except CaCIPK6 where serine was modified to cysteine. Further investigations are required to assess an effect of these changes on the function of the activation loop. In CIPKs, NAF motif mediates the interaction between CIPK and CBL, and FISL motif keeps the kinase inactive under normal conditions, hence act as autoinhibitory domain [13, 14]. PPI domain is required for interaction of CIPKs with protein phosphatase 2C (PP2C) [60]. In CaCIPK proteins, a total of 10 conserved motifs were identified by using the MEME tool. Out of those, motif 7 was annotated as NAF domain due to the presence of conserved asparagine-alanine-phenylalanine residues [13], whereas motif 8 which is located just after the motif 7 was designated as the PPI domain as it contains important arginine and phenylalanine residues. All motifs except motif 6 and 10 were present in 22 CaCIPK proteins (Figure S3). Motif 6 was absent in CaCIPK4, whereas motif 10 was absent in the subgroup of intron-rich CaCIPKs. This may explain the presence of CaCIPK4 in a separate phylogenetic clade. The similar motif composition of proteins of the same clade was also reported in tomato and saccharum [48, 57]. Furthermore, the sequential arrangement and the size of motifs in all the CaCIPK proteins were similar, which hints towards structural conservation and a close phylogenetic relationship, as previously reported in Prunus mume [50]. The sequence logo of different motifs is depicted in Table S1.

3.3. Phylogenetic analysis of CaCIPK family

A total of 133 CIPK protein sequences from four species: Arabidopsis thaliana (26), Oryza sativa (33), Glycine max (52), and Cicer arietinum (22) were used to construct the phylogenetic tree to explore the evolutionary relationship among the CIPKs. Based on high bootstrap values, the tree was divided into two major groups: group I and II. Two groups were further sub-divided IA, IB, and IIA, IIB (Figure 2).
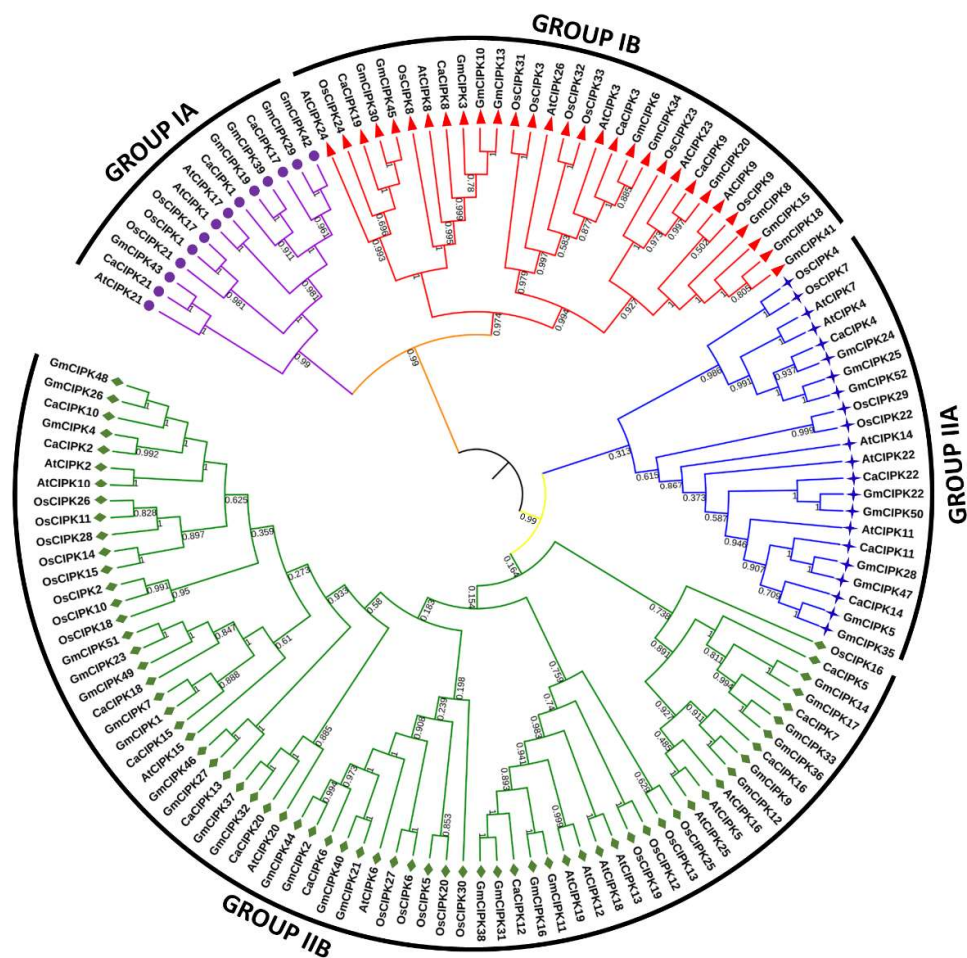
Figure 2. Phylogenetic relationship among CIPKs from different plants. The evolutionary relationship from Arabidopsis thaliana (At), Oryza sativa (Os), Glycine max (Gm) and Cicer arietinum (Ca) constructed using Neighbor-Joining method is shown via phylogenetic tree. Labels above the nodes represent bootstrap values calculated from 1000 replicates. The Group IA, IB, IIA and IIB are indicated by violet, red, blue and green branch lines respectively.

Interestingly, all intron-rich CaCIPKs were clustered in group I, and intron-poor CaCIPKs were clustered in group II. This indicates evolutionary conservation among intron-rich and intron-poor CIPKs in chickpea, and thus CIPKs may have evolved as two groups in chickpea. Similar phylogenetic pattern has been observed for CIPKs in other plants, such as Zea mays [61] and Prunus mume [50]. Remarkably, CaCIPKs were found to be close to CIPKs from Arabidopsis and soybean, but distantly placed from rice CIPKs. This suggests that CIPKs might have evolved separately in dicots and monocots. Group IIB contained most members of CaCIPKs. Group IA includes CaCIPK1, -17, -21, group IB includes CaCIPK3, -8, -9 and -19, Group IIA includes CaCIPK4, -11, -14 and -22 and Group IIB includes CaCIPK2, -5, -6, -7, -10, -12, -13, -15, -16, -18 and -20.

3.4. Chromosomal location and gene duplication of CaCIPK genes

All the CaCIPK genes were mapped onto the seven out of eight chromosomes of chickpea. None of them was localized on chromosome 8. All the genes were unevenly

distributed on the seven chromosomes (Figure 3). Chromosome 1 contains the maximum number of genes viz. CIPK1, -7, -9, -12, -20. Chromosome 2 and 7 harbours four genes each i.e., CIPK2, -3, -8, -14 and CIPK6, -17, -19, -21 respectively. Chromosome 5 contained three genes (CIPK10, -11, -13) that were located very close to each other. The other chromosomes comprised only two CIPKs each. The uneven distribution pattern of CIPKs in chickpea is consistent with previous reports of CIPKs in tomato, pepper, sorghum, rice, apple and woody plant [48, 50, 51, 55, 62, 63].
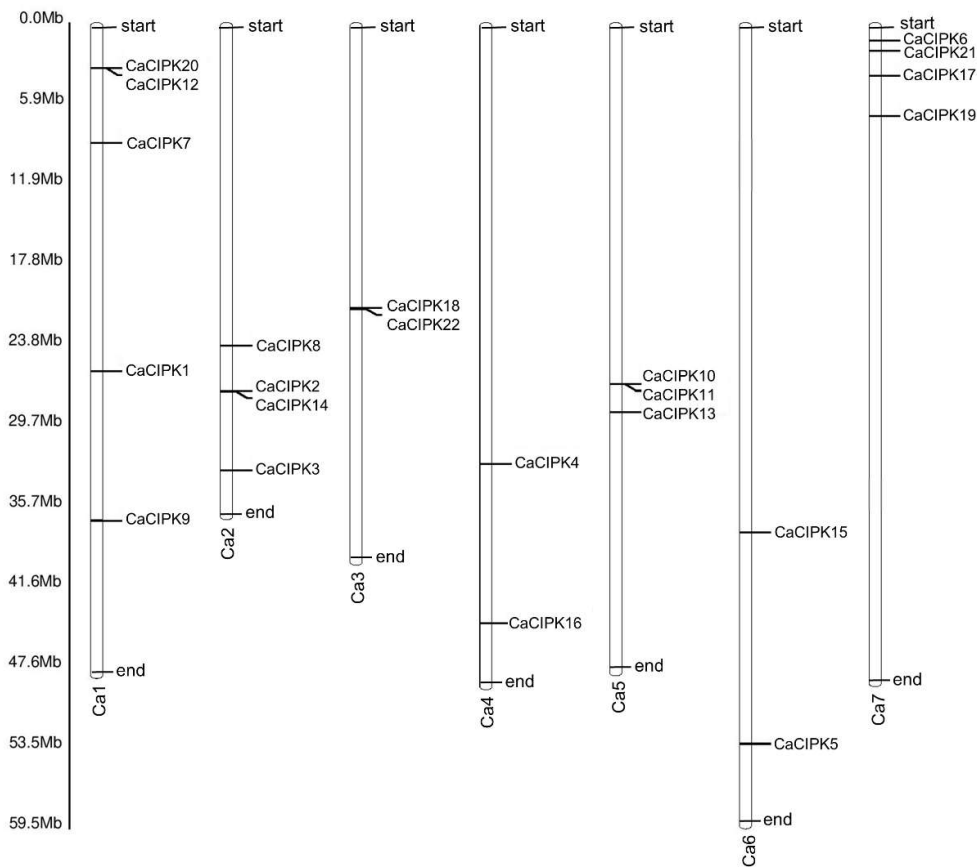

Figure 3. Chromosomal locations of chickpea CIPK genes. The 22 CIPK genes of chickpea were mapped to different chromosomes using TBtools.

CaCIPK family has undergone gene duplication as nine gene pairs showed segmental duplications which include CaCIPK1/17, CaCIPK2/10, CaCIPK2/18, CaCIPK5/7, CaCIPK10/18, CaCIPK11/14, CaCIPK11/22, CaCIPK14/22 and CaCIPK15/18 (Figure S4). In addition, four gene pairs (CaCIPK12/20, CaCIPK2/14, CaCIPK18/22, CaCIPK10/11) showed tandem duplications. Similarly, CIPK gene family has been found to exhibit significant gene duplication in plant species like Arabidopsis, rice and Vitis vinifera [64–66]. This suggests that gene duplication has been the major driving force behind evolution and expansion of CIPK gene family in chickpea and other plants. A ratio of Ka (non-synonymous)/Ks (synonymous substitution) = 1 signifies neutral selection (drift), Ka/Ks < 1 indicates purifying selection and Ka/Ks > 1 implies positive selection (adaptive evolution) [67]. In our study, the ratio of Ka to Ks for CaCIPKs was calculated which ranged from 0.0068 to 0.1675. Thus, Ka/Ks was found to be less than 1 for all segmentally duplicated gene pairs which suggests that all duplicated genes of CaCIPK family had undergone purifying selection on the whole genome duplication (WGD) (Table S2).

3.5. Cis-regulatory elements in CaCIPK promoters

Various cis-regulatory elements were found to be unevenly distributed on CaCIPK promoters (Figure 4).
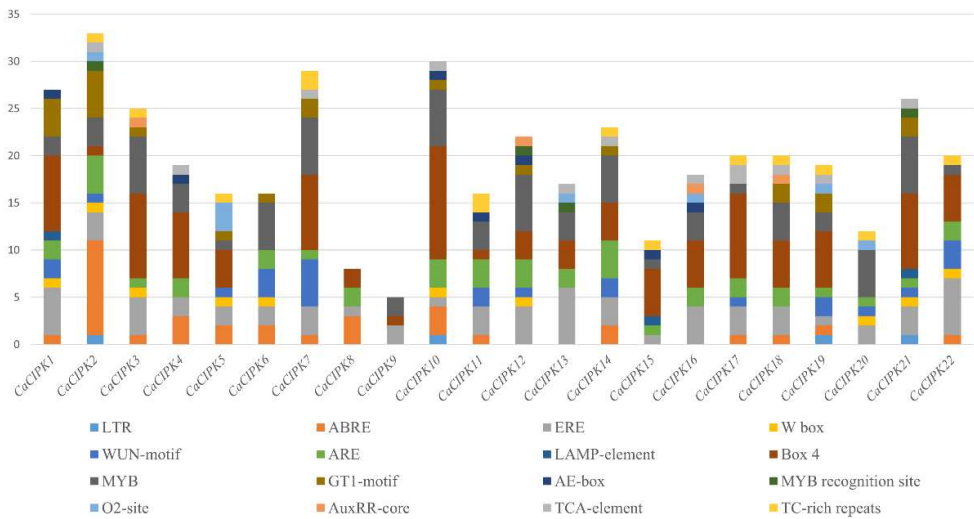

Figure 4. Different cis-regulatory elements in the 2 kb upstream region of 22 CaCIPK genes are illustrated by different colours in the bar chart. The X-axis represents the name of genes and Y-axis represents the number of cis-elements in each promoter. The names of cis-regulatory elements are mentioned below the graph.

These cis-elements have been found to be associated with different stress responses, including ARE (anaerobic induction), MYB (drought stress), LTR (low-temperature responsiveness), WUN-motif (wound-responsiveness), MYB recognition site (drought stress), responsive to phytohormones, such as ERE (ethylene responsive element), ABRE (abscisic acid responsiveness), W-box (WRKY transcription factor binding site), TCA-element (salicylic acid responsiveness), AuxRR-core (auxin responsive element), and, O2-site (cellular development), TC-rich repeats (defence and stress responsiveness) and Box4, GT1-motif, LAMP-element (light responsiveness) [68–74] . Apart from these, an oxidative stress responsive element ERE [75] was found to be present in all CaCIPKs. W box has a role in both biotic and abiotic stress [76], and was present in CaCIPK1, -2, -3, -5, -6, -10, -12, -20, -21, -22. MYB was present in all the CaCIPK genes except CaCIPK8. O2 site involved in zein metabolism and circadian motif [75], was present in only six CaCIPK genes viz. CaCIPK2, -5, -13, -16, -19, -20. Another well characterized cis-element, ABRE involved in abiotic stress and ABA responsiveness [77, 78] was found in the promoter of 15 CaCIPK genes, including CaCIPK1, -2, -3, -4, -5, -6, -7, -8, -10, -11, -14, -17, -18, -19, -22. Besides, LTR-motif was one of the least common elements and only four CaCIPK genes (CaCIPK2, -10, -19, -21) contained this motif (Table S3). Overall, presence of different cis-regulatory elements in CaCIPK promoters indicate their role in abiotic and biotic stress responses, phytohormone signaling and plant development.

3.6. Subcellular localization and structure prediction

In our study, the majority of CaCIPK proteins were found to localize in the cytoplasm, and only six CaCIPK proteins were localized in the nucleus (Figure 5). Among 22 CIPKs, five proteins namely CaCIPK2, -6, -10, -13, -19 were found to be localized both in the nucleus and cytosol.
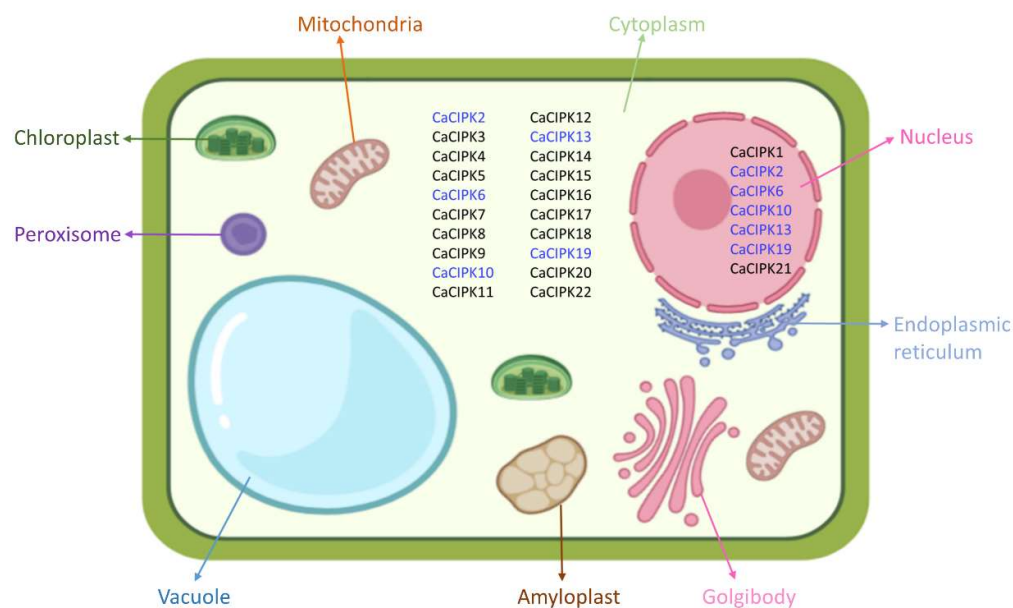
Figure 5. Subcellular localization of chickpea CIPK proteins predicted using the CELLO program. Genes located in both cytoplasm and nuclei are indicated by blue font.

Till now, the subcellular localization of few CIPKs in Arabidopsis has been reported and they have been localized in cytoplasm and nucleus [79, 80]. However, subcellular localization of the majority of CIPKs in other plants such as rice, maize and soybean has not been determined yet. In general, CBLs recruit CIPKs to plasma membranes or tonoplasts to perform their different functions. For example, AtCBL10 recruits AtCIPK24 at the tonoplast to regulate the vacuolar homeostasis of Na+ [81]. AtCBL1 and AtCBL9 recruit AtCIPK1 to the plasma membrane whereas, AtCBL2 recruit to the tonoplast [82, 83] Interestingly, AtCBL1 and AtCBL9 also recruit AtCIPK23 at the plasma membrane [84]. Therefore, future investigation of CaCBL-CaCIPK interactions under different conditions will reveal their exact subcellular localization.

The three- dimensional structures of 22 CaCIPK proteins were modelled with 100% confidence by the single highest scoring template (Figure 6).
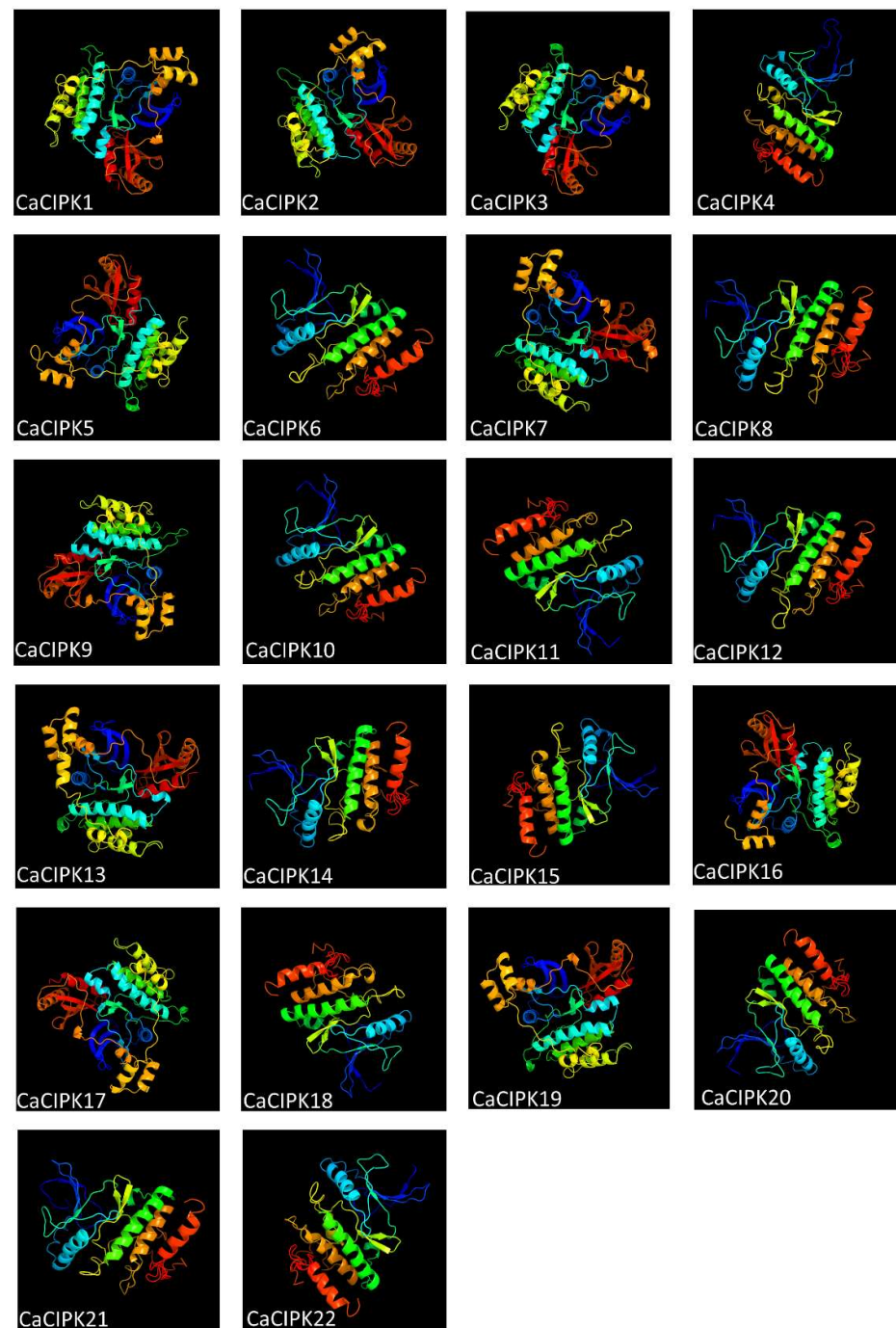
Figure 6. Predicted tertiary structures of chickpea CIPK proteins by PHYRE2.

Majority part of the models were built based on two templates – c6c9dB (Serine/threonine-protein kinase MARK1) and c5ebzF (inhibitor of nuclear factor kappa-b kinase subunit alpha) on the basis of raw alignment score which takes into account sequence and secondary structure similarity, inserts and deletes. CaCIPK16 showed the maximum coverage (95%) whereas CaCIPK12 showed the least coverage (50%). The identity of the template model c4czuC which belonged to CIPK23 with other CIPKs varied from 52% to 85%

(Table S4). All the CaCIPK proteins were found to have comparable numbers of α-helices and β-sheets, ranging from 16 to 21 and 14 to 17. The 3D structure of CIPKs have not been fully explored in other plants, however in Prunus mume 3D structures of PmCIPK proteins were predicted which shared the highest identity with the hypothetical protein c6c9dB [50], similar to CaCIPK proteins.

3.7. Interaction patterns between CBL and CIPK proteins in chickpea

CIPKs are generally activated by interaction with CBLs, and then perform different functions. Thus, it is crucial to determine interactions and functional complexes of CBLs and CIPKs in chickpea. Therefore, in silico analysis was performed to analyse the CBL and CIPK interactions in chickpea. A combined score of co-expression, experimentally determined interaction and automated text mining was used to predict the strength of interaction. A score of less than 0.7 was taken as weak, whereas a score greater than 0.7 was considered as strong interaction. CaCBL1 showed strong interaction (thicker lines) with CaCIPK2, -3, -4, -6, -8, -9, -12, -14, -16, -17, -19, -22 and exhibited weak (thinner lines) interactions with CaCIPK18 and CaCIPK20 (Figure 7).
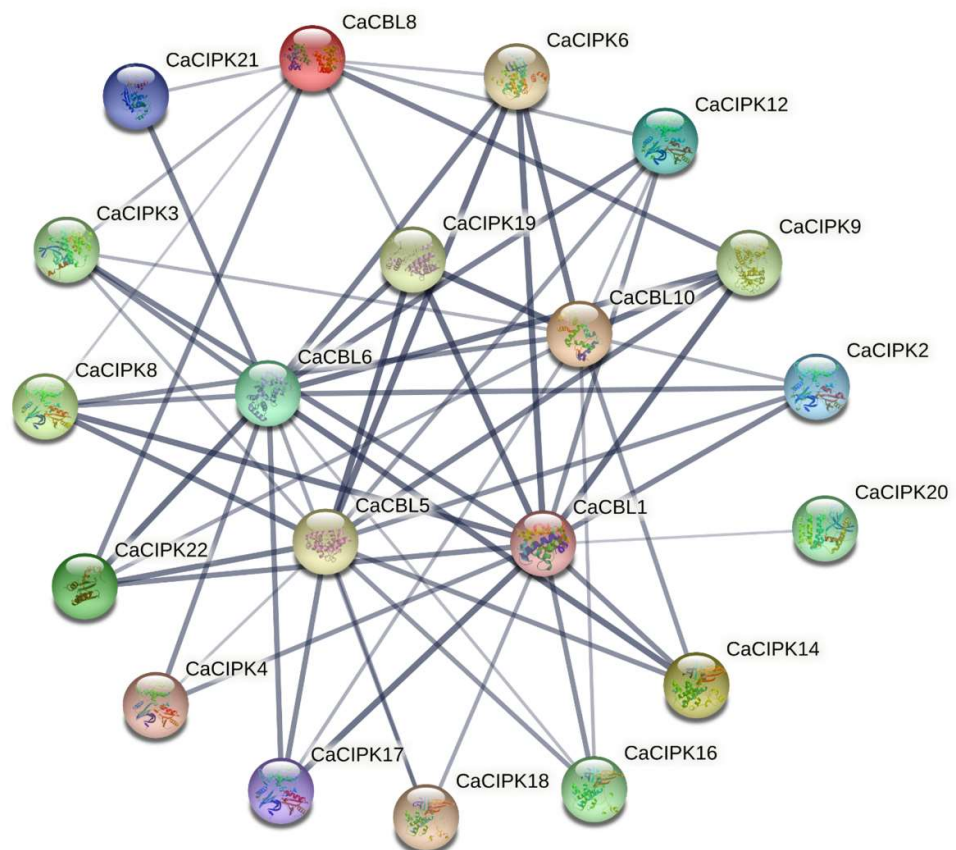

Figure 7. Interaction networks of CaCIPKs based on A.thaliana orthologues in the STRING database. The thicker and thinner lines represent strong and weak interaction respectively.

CaCBL5 showed strong interactions with nine CaCIPKs, including CaCIPK2, -6, -8, -9, -14, -16, -17, -19, -22 and weak interactions with four CaCIPKs (CaCIPK3, -4, -12 and -

18). CaCBL6 was found to interact with 14 CaCIPKs, including CaCIPK2, -3, -4, -6, -8, -9, -12, -14, -16, -17, -18, -19, -21, -22CaCBL8 interacted with eight CaCIPKs (CaCIPK3, -6, -8, -9, -12, -19, 21, -22) among which it showed strong interactions with only two CaCIPKs (CaCIPK9 and CaCIPK22). Lastly, CaCBL10 interacted with 11 CaCIPKs among which it showed strong interactions with CaCIPK6, -8, -9, -14, -19 and weak interactions with CaCIPK2, -3, -12, -16, -17, -22 (Table S5). Interestingly, CaCIPK9 was found to interact strongly with all the CBLs. This interaction pattern suggests that each CaCIPK may interact with multiple CaCBLs and vice versa. Such interactions may determine functional specificity or overlap of chickpea CBL-CIPK complexes and subcellular localization of CaCIPKs in condition specific manner. Similar interaction patterns of CBL-CIPK complexes have been found in different plant species, including Arabidopsis [12] and rice [65]. This suggests that CBLs and CIPKs make diverse complexes to display functional specificity and synergism across plant species.

3.8. Expression profile of CIPK genes in different developmental stages

The expression analysis of CaCIPKs was carried out in 27 tissues of chickpea belonging to different stages i.e. germination stage (radicle, plumule, embryo), seedling stage (epicotyl, primary root), vegetative stage (root, petiole, stem, leaf), reproductive stage (nodules, flowers, buds, pods, immature seeds), and senescence stage (yellow leaf, immature seeds, mature seeds, seed coat, and nodules) (Figure 8).
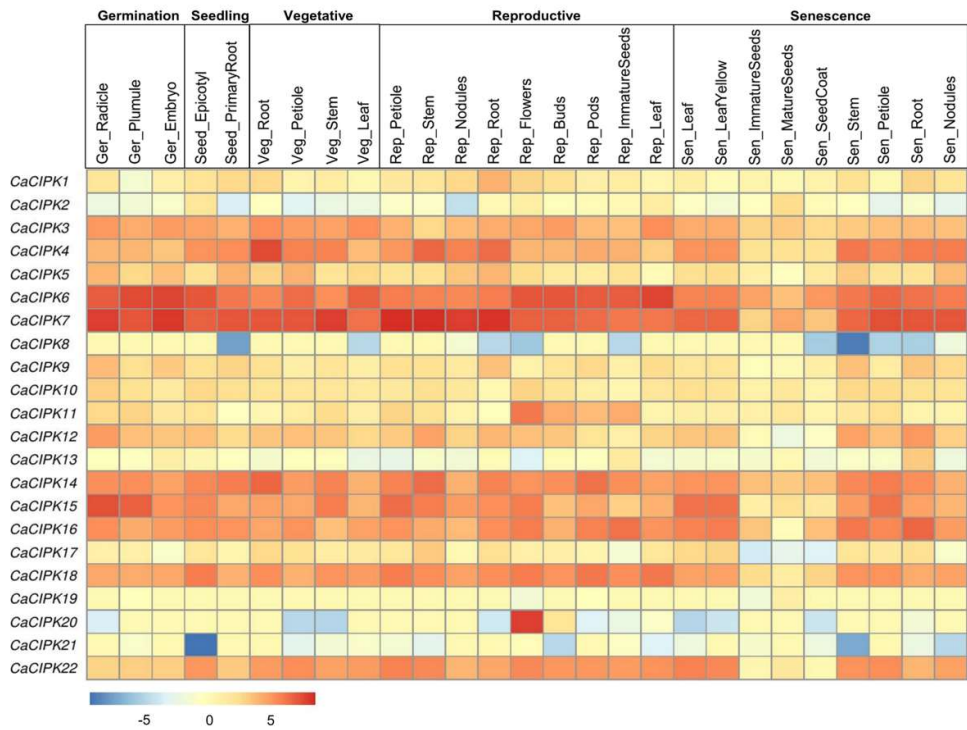
Figure 8. Expression profiles of different CIPK genes in different developmental stages of chickpea. The heatmap represents the expression pattern of CaCIPK genes in different tissues of developmental stages such as germination, seedling, vegetative, reproduction and senescence. The genes are named on the left and different tissues/developmental stages are labelled at the top. Scale bar represents the normalised $\log_2$ FPKM values.

The results indicate that most CaCIPK genes show significant expression in multiple tissues and developmental stages. CaCIPK3, -4, -6, -7, -14, -15, -16, -18 and -22 were found

to have ubiquitously high expression in all the tissues. Whereas CaCIPK2, -8, -13, -17 and -21 showed low expression in almost all the tissues. This expression pattern indicates that CIPKs might be involved in regulation of a wide array of processes during different stages of plant development in chickpea. (Table S6). Similar expression pattern has been observed for CIPKs in plants such as Arabidopsis, rice and wheat [47, 65] AtCIPK19 has been found to express highly in pollen grains and pollen tubes and analysis of atcipk19 mutant and overexpression in plants revealed that AtCIPK19 is required for pollen tube growth and polarity [24]. AtCIPK6 and its chickpea ortholog have been shown to regulate root development via controlling auxin transport in Arabidopsis [26]. Tomato SlCIPK2 is specifically expressed in floral organs and through interaction with different SlCBLs and transcription factors regulates stamen development and stress tolerance [85]. Also, Manihot esculenta CIPKs; MeCIPK16 and MeCIPK20 were predominantly expressed in flowers [86]. These findings suggest that CIPKs are key regulators of plant development.

3.9. Expression profile of CIPK genes in abiotic stress

To investigate the possible involvement of chickpea CIPKs in abiotic stress signaling, their expression pattern was analysed in root and shoot under three major abiotic stresses, desiccation, salinity and cold. Several CIPK genes were found to differentially express both in root and shoot (Figure 9).
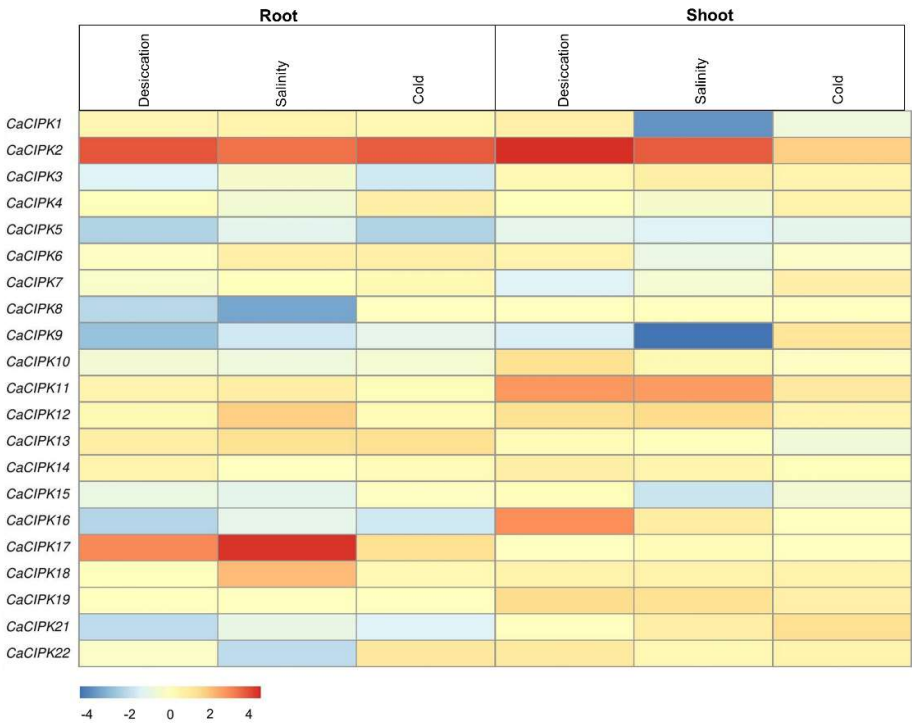
Figure 9. Expression profiles of CIPK genes of chickpea in three abiotic stress – cold, desiccation and salinity in root and shoot. The genes are named on the left and condition/tissue are labelled at the top. The scale bar represents the log_2 Fold Change based on FPKM values.

Seven genes, CaCIPK2, -4, -6 -11, -12, -13, and -18 were found to be commonly upregulated in all three abiotic stresses, whereas five genes CaCIPK3, -5, -9, -16 and -21 were commonly downregulated in root (Table S7). Notably, CaCIPK8 was upregulated in cold

stress but commonly downregulated in desiccation and salinity stresses. Similarly, eight genes CaCIPK2, -10, -11, -12, -14, -16, -18 and -19 were commonly upregulated in multiple stresses in shoot. Only a single gene CaCIPK5 was commonly downregulated in all stresses in shoot. CaCIPK7 and -9 were commonly downregulated in desiccation and salinity but upregulated in cold stress. Similarly, CaCIPK1 and -15 were commonly downregulated in salinity and cold stresses but upregulated in desiccation. CaCIPK2 was ubiquitously expressed in all the stresses, both in root and shoot. These results suggest that several CaCIPK genes may have an important role in abiotic stress signaling in chickpea. Previously, CIPKs from different plant species have been implicated in abiotic stress responses. For example, salt overly sensitive (SOS) pathway is well characterized in Arabidopsis, where membrane localized AtCBL4 (SOS2) interact and activate AtCIPK24 (SOS2), which in turn regulated membrane localized Na+/H+ antiporter (SOS1) to confer salt tolerance [87]. In addition, AtCIPK8, the homolog of AtCIPK24 interacts with AtCBL10 and activates SOS1 leading to salt tolerance [88]. AtCIPK21 is readily induced under salt and osmotic stress. AtCBL2 and AtCBL3 interact with AtCIPK21 and target it to the tonoplast. Analysis of AtCIPK21 overexpression and null mutant plants showed that it is a positive regulator of salt and osmotic stress tolerance in Arabidopsis [89]. Similarly, OsCIPK23 and ZmCIPK8 have been known to positively regulate drought tolerance in rice and maize respectively [90, 91].

3.10. Expression profile in different stages of seed development

Optimum development of seeds leads to their production with sufficient quantity as well as quality, thereby it determines the yield. To understand the role of CIPKs in chickpea seed development, expression profile was generated with mature leaf as control and seven different seed stages, representing early-embryogenesis (S1), mid-embryogenesis (S2), late-embryogenesis (S3), mid-maturation (S4-S5), and late-maturation (S6-S7), in two desi cultivars: JGK3 (large seeded) and Himchana1(small seeded) (Figure 10).

Figure 10. Expression profiles of CIPK genes in different stages of seed in chickpea in two cultivars: JGK3 and Himchana1. The genes are named on the left and stages/cultivars are labelled at the top. The scale bar represents the log_2 Fold Change based on FPKM values.

Few CaCIPK genes, including CaCIPK2, -11, -13 were ubiquitously expressed during all the seed stages in both the chickpea varieties however, level of expression varied (Table S8). CaCIPK2 expressed highly during S5-S7 in JGK3 whereas, during S4 in Himchana1. CaCIPK11 showed high expression during S1-S5 in both the varieties and during S7 in Himchana1. Similarly, CaCIPK13 showed significant expression during S1-S5 however, the level of expression was higher in JGK3 than Himchana1. In contrast, CaCIPK18 and -21 showed significant expression during S1-S5, but expression was higher in Himchana1. Remarkably, CaCIPK6 and -16 were upregulated during S1-S4 but downregulated during S5-S7 in both the varieties. CaCIPK10 was upregulated during S4-S5 in both the varieties but upregulated during S6 only in JGK3. Two CaCIPK members, CaCIPK12 and -17 were significantly downregulated during all seed stages in both JGK3 and Himchana1. These findings suggest the crucial role of CIPKs in chickpea seed development. Some CIPKs might be involved in regulating all the seed development stages in both the varieties, whereas some members might regulate specific seed stages in both varieties or any specific variety. Thus, CIPKs could play an important role in determining the chickpea yield.

Very few studies have analysed the role of CIPKs in seed development till date. The role of CIPKs in seed development has been proposed in plant species like rice and Phaseolus vulgaris. Along with CBLs, several rice CIPK genes were differentially expressed during five stages of seed development [65]. In Phaseolus vulgaris, PvCIPK1, -2, -3 and -5 were expressed only in small and mid-size developing seeds but show no expression in large developing seeds [92].

4. Conclusions

CIPKs have been studied at the genome-wide scale in diverse plant species, however, in-depth study of the CIPK gene family was missing in important legume crop chickpea. Therefore, in this study, genome-wide identification and characterization of the CIPK gene family was carried out in chickpea which unearthed a total of 22 CIPK genes. Gene and protein structure analysis indicated structural conservation among chickpea CIPKs and homology with other plant species. Phylogenetic analysis suggests that chickpea CIPKs have evolved from common dicot ancestors and gene duplication is the major driving force behind their evolution and expansion. Subcellular analysis showed that the CIPK proteins are majorly located in the nucleus and cytoplasm. In-silico interaction analysis revealed various specific and overlapping functional complex formations between CBLs and CIPKs which could be tested functionally in future. Expression analysis during various developmental stages indicated that CIPKs are expressed in a wide range of tissue/organs and could play an important role in their development. Promoter and expression analysis of the CIPK gene family strongly suggest their role in abiotic stress signaling and seed development in chickpea. Thus, this study provides a useful platform for detailed functional characterization of the CIPK family in chickpea and other legume crops.

Supplementary Materials: Table S1: Sequence logo of different motifs identified through MEME; Table S2: List of segmentally duplicated gene pairs; Table S3: List of cis-regulatory elements identified in the promoter regions of CaCIPK genes; Table S4: The confidence, coverage and sequence identities of the homologous relationship of the CaCIPKs; Table S5: Type and strength of interactions between CBL and CIPK proteins in chickpea; Table S6: Log2FPKM values of different tissues belonging to different developmental stages; Table S7: Log2 Fold Change values of CaCIPKs in response to abiotic stress; Table S8: Log2 Fold Change values of CaCIPKs in various stages of seed development; Figure S1: Percentage of identity and similarity between CaCIPKs; Figure S2:

Alignment of 22 CaCIPKs for domain identification; Figure S3: Identification of motifs through MEME; Figure S4: Duplication of chickpea CIPK genes performed by MCScanX is shown via Circos plot.

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