Allele mining, evolutionary genetic analysis of *TaHKT1;5* gene and evaluation of salinity stress in selected lines of wheat.

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ABSTRACT

This study reports the novel allelic diversity in *HKT 1;5* gene (*High Affinity Potassium Transporter*) in bread wheat and its phylogenetic relationship among the paralogs/orthologs of bread wheat and its wild relatives. *HKT 1;5* gene is a known and pivotal gene associated with the salinity tolerance in plants upon the discrimination of K^+ over Na^+ in leaves without change in Na^+ concentration in root. This gene was sequenced in a diverse collection of bread wheat, durum wheat, wild relatives, and ditelosomic lines. Sequence characterization in bread wheat led to the identification of four alleles, which could be distinguished by number of SNPs. Sequence comparison between Triticeae monocot bread wheat and dicot *Arabidopsis thaliana* revealed that the *HKT1;5* gene is conserved at exonic regions, however presence of transposable elements especially in intronic regions is further intriguing towards evolutionary relatedness with repeated elements. Two paralogous or major alleles observed in *Triticum monoccum* and *Aegilops tauschii* were further categorized as sub-alleles based on their SNPs comparison. The gene was not present in *T. urartu* in accordance with existing evidence, while in *A. speltoides* with few base pairs insertion at exon1 region caused a frameshift mutation with an altered amino acids and genomic database mining unveiled additional alleles in this species. Ditelosomic lines with 4DL and 4DS chromosomes revealed a higher similarity with bread and durum wheat respectively. Phylogenetic studies of *HKT1;5* orthologs from different Poaceae species revealed the occurrence of five different ortholog groups with taxonomic consistency. Phenotyping based on salinity stress distinguished the saline tolerant lines from sensitive lines in comparison with reference lines and possible allelic comparison was made.

KEY WORDS: *HKT* transporter, *Triticum*, Diversity, Evolution, Phylogenetics, Phenotype, Ditelosomics, SNPs, Transposable Elements, Allelic Variant.

INTRODUCTION

Salinity or sodicity marks 6% of the world's land covering around 400 million hectares of cultivated and uncultivated land (FAO Land and Plant Nutrition Management Service, 2008). Soil salinity imposes ion toxicity, osmotic stress, nutrient (N, Ca, K, P, Fe, Zn) deficiency and oxidative stress on plants. It is one of the important stresses resulting in the reduction of growth and yield of different crops including wheat. In several wheat-producing countries of the Indian subcontinent (e.g. India and Pakistan) and the Middle Eastern region (Iran, Egypt, and Libya), salinity affects up to 10% of the total wheat belt, but in some regions, such as Western Australia, salinization affects more than half of wheat farms (Colmer et al. 2006, Honghong et al. 2014). Salt concentrations beyond 40 mM or electrical conductivity greater than 4 dS/m in soil can cause significant reduction in spikelets per spike, delayed spike emergence and reduced fertility, which results in poor grain yields (Shrivastava and Kumar 2015) and reductions in wheat grain yield (Kalhoro1 et al., 2016). Adverse of effect of salinity during developmental stages of wheat and strategies to overcome problems associated with salinity stress has been recently reviewed (Sabagh et al., 2021). The role of salinity tolerance concerning *HKT1;5* gene and its translocation events were shown based on the available research evidence (James et al., 2011, Byrt et al., 2007) (Fig. 1A, 1B). Salinity tolerance and causes of sensitivity of plants is depicted with illustration (Fig. 1C).

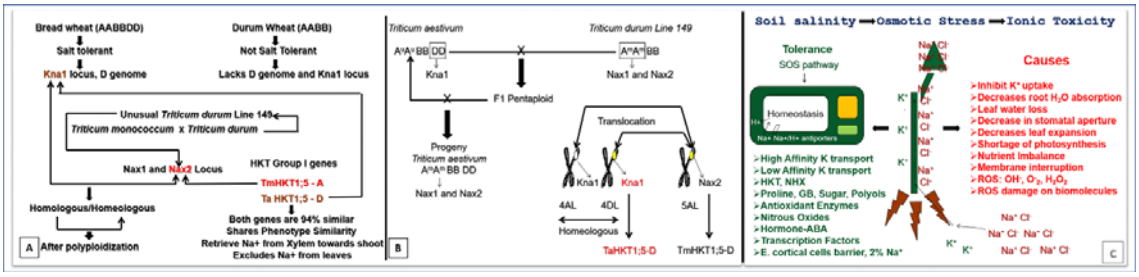

Figure 1. A). Genes behind the salinity tolerance in wheat. B). Reciprocal translocation of *Kna1* and *Nax2* locus. Schematic representation. Based on sources: James et al., 2011, Byrt et al., 2007. C) Schematic representation of salinity tolerance and sensitiveness in plants.

Improving salinity tolerance is an important breeding target in wheat breeding programs. A major genetic mechanism contributing towards salinity tolerance of most crops including the cereals rice (*Oryza sativa*) and wheat (*Triticum aestivum* and *Triticum monococcum*) is Na^+ exclusion from leaves. High-affinity potassium (K^+) transporter (*HKT*) protein family have been reported to play a major role in this trait (Ren *et al.* 2005; James *et al.* 2006; Munns *et al.* 2012; Munns and Gilliham, 2015; Campbell *et al.* 2017; Xu *et al.* 2018). Several *HKT* genes, including *HKT1;1*-like, *HKT1;3*-like, *HKT1;4*-like, and *HKT1;5*-like, have been identified and mapped to wheat homoeologous chromosome groups 2, 6, 2, and 4, respectively (Huang et al. 2008). *HKT 1;5* gene associated *Kna1* locus was reported for salinity tolerance in plants due to the discrimination of K^+ over Na^+ in leaves with net Na^+ loading in root xylem than net uptake (Gorham et al., 1990). The wheat sodium transporters *TmHKT1;5-A* and *TaHKT1;5-D* (Genbank:

DQ646342) are encoded by genes underlying major shoot Na^+ exclusion loci *Nax2* (Chromosomal location: 5AL, ancestrally 4AL) and *Kna1* (Chromosomal location: 4DL) from *Triticum monococcum* (Tm) and *Triticum aestivum* (Ta), respectively (Byrt et al., 2007). Role of *TmHKT1;5-A* in improving grain yield on saline soils by 25% has been reported (Munns et al 2012). The presence of HKT gene in several plant species has been reported (Waters e al., 2013). Comparative genomics suggested that there is no synteny of the *HKT1;5* gene between *O. sativa* and bread wheat unlike other *HKT* genes (Huang *et al*., 2008) and only a single form of the *HKT* gene family exist in the model dicot species *A. thaliana* (Mayer *et al*., 1999, Uozumi *et al*., 2000) and divergence of this gene must have occurred early during evolution of monocots and dicots (Waters e al., 2013). Allele mining pertain to natural variations is useful strategy for the identification of better salinity tolerant variety from natural populations (Baxter et al., 2010). Efforts have taken to genetically characterize the *HKT1;5* gene to better understand the molecular evolutionary basis of diversity of salt tolerance among bread wheat and its wild relatives. Therefore, understanding the allelic diversity, natural variations, and divergence of *HKT1;5* gene would be an asset and provides the genetic basis of the evolution of this gene. In this study, using a diverse wheat material and multifaceted genetic approaches, we demonstrate that *HKT1;5* gene has multiple allelic variants present within the wheat germplasm pool and its wild progenitor species.

MATERIALS AND METHOD

Plant materials and DNA extraction

The plant materials included 16 bread wheat and a durum wheat cultivar, two ditelosomic lines were included (Ditelosomic lines DTGH09_1559 possessing 4DL and lacking 4DS chromosome, DTGH09_1556 possessing 4DS and lacking 4DL chromosome) (Supplementary Table 1). A total of 18 selected wild accessions represented the diploid progenitors of the three genomes of bread wheat, *T. monococcum*, *T. uratu*, *A. speltoides* and *A. tauschii* (Supplementary Table 2). After attaining four leaf stage, DNA was extracted from collected leaves of each cultivar using the CTAB method (Murray and Thompson, 1980).

NCBI GenBank Submission and retrieval of sequences from public databases for Phylogeny

For this study, we have generated 29 sequences so far; among them, 7 were about 3 kb each and are complete coding sequences exclusively from wheat. The remaining 22 sequences were about 1 kb each and are from different progenitor species, ditelosomics and some varieties of wheat as well. For phylogenetic analysis, public database was searched for *HKT1;5* gene sequences; apart from the sequences generated in this study. Therefore, additional sequences were retrieved from Public Data

Bases such as NCBI (https://www.ncbi.nlm.nih.gov/nuccore/) and URGI (https://urgi.versailles.inra.fr/Species/Wheat), for comparative genomic analysis. SIFT analysis were carried out using existing SNPs derived VCF file of *T. aestivum HKT1;5* and further comparisons made using TraesCS4D02G361300 from ENSEMBL.

Primer design, Polymerase Chain Reaction and Sequencing

A and D genome specific primers for the *Triticum aestivum HKT1;5-D* (Genbank: DQ646342) and *Triticum monococcum HKT1;5* (Genbank: DQ646339) genes were designed using the reference sequence available at NCBI (Byrt *et al*., 2007). Primers allowed amplifying the gene from start to stop codon within a single Polymerase Chain Reaction (PCR) in genome specific manner. Alternatively, additional UTR anchoring, and internal primers were used to amplify different gene fragments that overlapped to obtain the entire length of the gene (Supplementary Table 3). The primers designed from *T. monococcum* (A^m) were also amplified a gene in *A. tauschii* and *T. aestivum* 4DL or *A. tauschii* (D^t genome) specific primers were also amplified a gene in *T. monococcum*. The PCR amplification was performed in 20 µl reaction volume consisting of 50 ng of template DNA, 0.5µl of dNTP (2.0mM each), 0.1 µl of Phusion Hifidelity Polymerase (2U/ µl) (Thermo Fisher Scientific Co.), 0.4 µl of 1pM Forward primer, 0.4 µl of 1pM Reverse primer, 0.6 µl of DMSO (100%), 4 µl of HF5X buffer, 0.4 µl of $MgCl_2$ (50mM), and 12.1 µl of Sigma H_2O. PCR was performed using Mastercycler® pro (Eppendorf) using the following cycling program: Initial denaturation at 98°C for 40 s, 35 cycles of 12 s denaturation at 98°C, 45 s annealing at 64°C, 1.4 mins extension at 72°C and final extension at 72°C for 5 mins. PCR conditions and cycles were adjusted based on primer pair, amplicon length, GC content and annealing temperature etc. Subsequently, amplified PCR fragments were resolved on 1.2% agarose gel pre-stained with GelRed™ (Biotium Inc.) at 90V for 90 minutes. PCR fragments were visualized using a UV illumination-based documentation unit. The visualized bands were cut from the agarose gels using clean scalpel and purified using Sigma Aldrich GenElute agarose gel purification kit. Additionally, amplified fragments were purified using a Qiagen QIAquick PCR purification kit (Cat. No. 28106) following the manufacturer's instructions. The purified PCR fragments were sequenced through Macrogen Co., South Korea. The primers designed from *T. monococcum* (A^m) were also amplified a gene in *A.tauschii* and also *A. tauschii* (D^t genome) specific primers were also amplified in *T. monococcum*.

Identification of SNPs for allele mining

The quality of chromatograms was reviewed based on PHRED quality scores using FinchTV (Geospiza Inc., USA) chromatogram viewer software. Chromatogram ab1 extension files were assembled using reference sequence (Byrt et al., 2007) with CodonCode aligner program (https://www.codoncode.com/aligner/). Alternatively, clear chromatogram regions were copied as text and converted to FASTA format as an input for the CAP3 assembly program and assembled to display the entire gene (Huang and Madan, 1999). Subsequent sequence alignment of the gene from different

cultivars was performed using Clustal Omega with default parameters (Sievers *et al*., 2011). The presence of SNPs was manually detected with multiple sequence alignment, based on the alignment SNPs derived different alleles were designated. Whole gene from selected lines of bread wheat was sequenced and used for SNPs/allele prediction and relevant analysis, while in durum wheat, wild progenitors and ditelosomics lines region spanning around 1kb at 5'UTR and/or Exon1 was exclusively sequenced focused to detect UTR or coding sequence variations and for related studies.

Identification of Repeated Elements

A search for identification of transposon and its related repeated elements in whole *HKT* gene of *T. aestivum* and *A. thaliana* was done using Repbase Censor server (https://www.girinst.org/censor/index.php). A predicted complete non-autonomous mariner transposon element was subjected for BLASTN analysis with IWGSC-URGI server (https://wheat-urgi.versailles.inra.fr/Seq-Repository/BLAST).

Phylogenetic analysis

The sequences were aligned using Clustal Omega (Sievers *et al*., 2011) or Clustal X and exported as multiple sequences aligned FASTA format (Larkin et al., 2007). Phylogenetic analysis of the gene sequences and relative divergence using RelTime were carried out with MEGA11 (Molecular Evolutionary Genetic analysis software) and gaps/missing parameters were excluded (Tamura et al., 2021). The Maximum Likelihood method with Jukes Cantor (JC) model and 1000 bootstrap resampling was used to reconstruct the phylogenetic tree of the *TaHKT1;5-D* gene sequences generated in this study along with the reference and orthologous gene sequences retrieved from public databases. Global alignment of the coding sequences was done using LAGAN alignment toolkit (Brudno *et al*., 2003). Genetic analysis was done using DNAsp for Gamma statistics for gene flow estimates of haplotypes (Nei 1973), DeltaST (Nei, 1982), Nst (Lynch, 1990), Fst (Hudson *et al*., 1992) (Rozas J, 2017).

Insilco analysis of protein and 3-D modeling

Corresponding putative proteins of *A. thaliana HKT1* gene and *T. aestivum HKT1;5-D* gene tertiary structure protein modeling was done using HHPhred (Söding *et al*., 2005) using protein templates from Protein Data Bank (http://www.rcsb.org/pdb/home/home.do). Visualization and annotation of generated protein models were done with Rasmol (Sayle and Milner-White, 1995) and compared with the primary structure of the protein. Secondary structure was predicted using HNN secondary structure prediction method (Guermeur, 1997). Documentation of protein physico-chemical parameters including instability index was calculated using Protparam tool at Expasy server http://web.expasy.org/tools/protparam/protparam-doc.html (Gasteiger *et al*., 2005).

PETRIPLATE BASED PHENOTYPIC TRAITS MEASUREMENT AND ANALYSIS

Completely Randomized Design Experiment

Seeds of the bread wheat cultivars C306, HDR-77, Oasis F86 and WK1204 were surface sterilized with 5% of sodium hypochlorite for 5 minutes and were subsequently exposed to five salinity treatments: Control (Water), 50 mM, 100 mM, 150 mM, 200 mM salt solution. For each cultivar, 15 seeds were used for each of the five salinity treatments in each experiment including controls. The 15 seeds were placed between the layers of cellulosic filter papers. During the experiment, Petri plates were covered with lid to reduce the excess evaporation and for maintaining a homogenous ambience. The average temperature of the experimental condition was 25.13°C (range 23-27.8°C) during the day and 14.29°C (range 12-18°C) during the night with 8 hours of photoperiod. Salinity or control treatment was performed with three replicates through a completely randomized design. To compensate for the evaporation, an equal amount of water was given to all samples periodically.

Phenotypic measurements: Estimation of germination traits

Phenotypic traits viz. Final Germination Percentage (FGP), Germination Energy Percentage (GEP), Radical Length (RL), Number of Radicles (NoR), Coleoptile Length (CL), Relative Water Content (RWC), Fresh Weight (FW), Dry Weight (DW) were measured. Germination was counted at every one-day interval until the 7th day when the germination was ceased. Each day was assigned as T; hence time-frequency interval was 24 hours from T1 to T7. Measurement of seedlings morphological parameters viz., seedling coleoptiles and radicle lengths were measured using a millimeter ruler. Final Germination Percentage (FGP) was calculated using following formula: FGP = (Number final germinated seeds/ Total number of seed tested) x 100. Germination Energy (GE) was calculated using following formula according to Ruan et al., 2002. GE (%) = (Number of germinated seeds at 4th after sowing/ Total number of seed tested) x 100. Seed Vigor Index (SVI) was calculated according to the formula described by Abdul Baki and Anderson (1973). SVI= Radicle length (cm) × Germination (%). Relative Water Content (RWC) was calculated according to Sumithra et al., 2006. RWC % = 100 x ((FW – DW) / FW). Fresh weight was measured when the germination got ceased and dry weight was observed after drying at 70°C in hot air oven. Further germination traits such as Mean Germination Time (MT) Mean Germination Rate (MR) Coefficient variation of germination time (CVt), Synchrony (Z) was calculated according to Ranal et al., 2009, Ranal et al., 2006. Statistical Analysis: Analysis of variance (ANOVA) was carried out and Least Significant Difference (LSD) with Bonferroni correction was estimated using R (R Core Team, 2014).

GREEN HOUSE BASED PHENOTYPIC TRAIT EXPERIMENT

Green house experiments were conducted in temperature controlled green house at CIMMYT, El Batan, Mexico. A completely randomized design experiment at greenhouse was conducted with 4 salinity concentrations (50mM, 100mM, 150mM, 200mM) and a control (water without salt). Soil content of the pot was about 3kg with 1:2:1 proportions of Sand: Soil: Organic matter) homogenously for all pots. Each pot was filled with gravel stones (3-5mm) from bottom up to 5cm to facilitate excess water drainage. Totally 90 pots were filled as designed experiment consisted of 6 varieties with 3 replicates for 4 concentration of salt stress with a control.

Pot water holding capacity (PWC) measurements

For measuring water holding capacity 10 random pots were chosen and soil+ pot's net weight was measured without adding water. Thereafter the pots were filled completely with excess water. The pots were then left for an hour to drain excess water and pots net weight was measured twice a day (morning and afternoon)**.** Likewise, pots were measured continuously for 4 days. One-way ANOVA analysis was done resulted with the P Value 1.28E-26, $P<0.001$. Water holding capacity of pots for each day depended upon the evaporation rate and ambient temperature. This experiment was useful to select a pot with an appropriate and adequate soil without excess water and salt leakage to minimize the errors due to deviation of physico-chemical properties in a controlled ambient.

Preparation of seedlings

Five seeds of bread wheat lines viz., C306, HDR-77, Kharchia, OasisF86, WK1204 and a durum wheat line DWK26 were placed in Petri plate with appropriate amount of water to achieve germination. Seedlings of each line were allocated to blocks of pots as mentioned below with appropriate genotype and salt concentration labelling. After a week, thinning was done to leave a single seedling per plot. Five pots were allocated for the concentrations of 0 (control), 50mM, 100mM, 150mM and 200mM for each variety and three replicates were maintained. Therefore, in total 90 pots were completely randomized inside the greenhouse to have a homogenous ambient.

Phenotypic traits measurement

The phenotypic traits including Plant Height (PH) (cm), Number of Tillers (NT), Number of Effective Tillers (NoET), Flag Leaf Length (FLL), Total Number of Spikes (TS), Spike Length (SL) were measured. Flowering was measured 50 days after sowing (DAS) from control and salt stressed genotypes. Statistical Analysis: Analysis of variance (ANOVA) was carried out and Least Significant Difference (LSD) with Bonferroni correction was estimated using R (R Core Team, 2014).

RESULTS

HKT1;5 gene sequence characterization

Amplification of salinity tolerance gene *TaHKT1;5-D* was done in diverse wheat lines (Fig. 2). Amplification was achieved in all 16 lines of bread wheat. Subsequently, 12 selected lines were selectively sequenced (based on preference with salinity tolerance, sensitiveness and/or with other interesting features such as drought tolerance etc) comprising four alleles in bread wheat. Gene sequence of bread wheat varieties, progenitor species, durum wheat and ditelosomic lines was submitted at GenBank (Table 1).

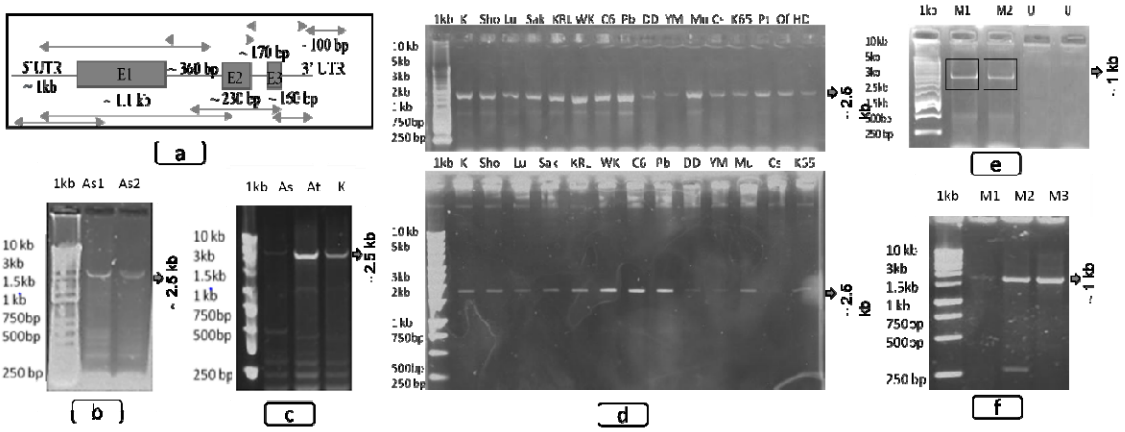

Figure 2. a) Schematic diagram of *HKT1;5* gene structure and approximate size of UTRs, Exons, and Introns. b) *HKT1;5* gene amplified in *A. speltoides.* c) *HKT1;5* gene amplified in *A. speltoides*, *A. tauschii* and bread wheat variety Kharchia d) Top gel: Amplification of *HKT1;5* gene in diverse *T. aestivum* varieties, K: Kharchia, Sho: Shorowaki, Lu: Lu26, Sak: Sakha, KRL: KRL1-4, WK: WK1204, C6: C306, Pb: Pasban 90, DD: Dharwar dry, YM: YMI#6, Mu: Munal#1, CS: Chinese spring, K65: Kharchia65, Pt: Pastor, Of: Oasis F86, HD: HDR-77. Bottom Gel: Purified fragments showing the intact banding pattern of the first 13 samples from top gel. e) *HKT1;5* gene amplified in *T. monococcum*, but not amplified in *T. urartu.* f) *HKT1;5* gene amplified in *T. monococcum*.

Subsequent sequencing and allele mining detected three SNPs, among them two SNPs at 830th and 832nd positions located at 5'UTR and third one located at 2106th position of first intron with reference to Byrt et al., 2007. The gene is located at chromosome 4DL between 507964629 bp to 507967781 bp (TraesCS4D02G361300) that revealed with BLASTN at wheat ENSEMBL. We found four alleles were designated as A, B, C and D among 12 geographically diverse bread wheat cultivars and allele 'C' is

prevalent (Table 1). Variant Effect Predictor revealed that 67% upstream variants (Genomic positions: 507965458 bp, 507965460 bp) and 33% intron variants (Genomic position: 507966734 bp) without amino acids change as expected for corresponding SNPs positions at 830, 832 and 2106 on the sequence of *TaHKT1;5* gene. However, a missense variant (506th Serine/Glycine) with deleterious mutation (SIFT score: 0.05) was observed in exon 3 as an inter-homoeologous variant (Genomic position: 507967636) from *TaHKT1;5* gene between chromosome 4DL to 7BS (ENSEMBL: BA00457515 SNP) comparison. *TmHKT1;5* gene at IWGSC-URGI BLASTN search revealed that 99% identity (0 gaps) and 100% query coverage with nucleotide database of *T. monococcum* (TGAC_WGS_monococcum_v1_contig_910810).

Table 1. NCBI GenBank accession numbers with allele description of *TaHKT1;5* gene from diverse collection of *T. aestivum* cultivars and wild relatives.

Wheat/wild relatives	Allele	GenBank	Size (bp)	SNPs/Allele Description
T. aestivum WK1204	A	KU184266	3073	g.[830T>C;832T>C;2106;T>C]
T. aestivum Dharwar Dry	B	KU212871	2826	g.[830T>C;832C>T;2106;T>C]
T. aestivum Munal#1	B	KU184267	2438	g.[830T>C;832C>T;2106;T>C]
T. aestivum Dharwar Dry	C	KU212872	2826	g.[830T>C;832T>C;2106;C>T]
T. aestivum OasisF86	C	KU212870	3068	g.[830T>C;832T>C;2106;C>T]
T. aestivum HDR-77	C	KU212869	3122	g.[830T>C;832T>C;2106;C>T]
T. aestivum C306	C	KU242566	1066	g.[830T>C;832T>C;2106;C>T]
T. aestivum Kharchia	C	KU212875	3074	g.[830T>C;832T>C;2106;C>T]
T. aestivum Sakha8	C	KU242567	965	g.[830T>C;832T>C;2106;C>T]
T. aestivum YMI#6	C	KU242568	1047	g.[830T>C;832T>C;2106;C>T]
T. aestivum Shorowaki	C	KU242569	965	g.[830T>C;832T>C;2106;C>T]
T. aestivum Lu-26	C	KU242570	1035	g.[830T>C;832T>C;2106;C>T]
T. aestivum Kharchia 65	C	KU212873	2883	g.[830T>C;832T>C;2106;C>T]
T. aestivum Kharchia 65	D	KU212874	2883	g.[830C>T;832T>C;2106;C>T]
A. tauschii CWI94958SH	T1	KU253614	958	g.[724G>T]
A. tauschii CWI94956SH	T2	KU253615	1009	g.[724T>G]
T.monococcum PI272558	M1	KU253607	994	g. [234 A>G; 745C>G; 832C>T]
T. monococcum PI272558	M2	KU253608	994	g. [234 G>A; 745C>G; 832T>C]
T. monococcum PI428150	M1	KU253609	966	g. [234 A>G; 745C>G; 832C>T]
T. monococcum PI326317	M1	KU253610	936	g. [234 A>G; 745C>G; 832C>T]
T. monococcum PI264935	M1	KU253611	672	g. [234 A>G; 745C>G; 832C>T]
T. monococcum PI272560	M1	KU253612	978	g. [234 A>G; 745C>G; 832C>T]
T. monococcum PI272560	M3	KU253613	978	g.[234 A>G; 745 G>C; 832T>C]
T.monococcum PI264935	N1	KY110746	942	N/A
T.monococcum PI272560	N1	KY110747	893	N/A
A.speltoides CWI94201SH	As2	KY110745	632	N/A
Ditelosomic DTGH09_1566	DS2	KY110743	902	N/A
Ditelosomic DTGH09_1559	DS1	KY110744	879	N/A
T. durum DWK26	B-Td1	KY110748	1007	N/A

Identified SNPs reflect the homozygotic nature of alleles in all the varieties with an exception in varieties Dharwar Dry (alleles 'B' and 'C') and Kharchia 65 (alleles 'C' and 'D') whereby a heterozygotic nature of the allele was observed suggesting the existence of an alternative variant of the gene *TaHKT1;5-D*. To check the allelic diversity in wild relatives, we used species-specific primers developed for *T. monococcum* and *T. aestivum* 4DL or *A. tauschii*. These primers successfully amplified *TaHKT1;5-D* gene in *T. monococcum, A. speltoides* and *A. tauschii.* The species-specific primers from *T. monococcum* amplified a native *TmHKT1;5-A* gene itself (M allele), also an ortholog in *A. tauschii* (Amplified, though not sequenced as M allele related *A.tauschii* (named as putative *At_HKT*) allele sequence directly retrieved from public database). Likewise, *T. aestivum* 4DL or *A. tauschii* specific primers also amplified a native *AtHKT1;5-D* gene (Allele 'T') and an ortholog from *T. monococcum* (N allele). These results highlight the existence of two major alleles 'M' and 'N' in *T. monococcum*. Interestingly, we could not amplify any orthologs in *T. urartu*. Based on SNPs within the M allele, three sub-alleles were detected, designated as M1, M2, M3, while major allele N exhibited only a single variant (N1) (Table 1). Also, two major alleles were identified in *A. tauschii*. The first allele was designated as 'T' and consists of two sub-alleles, T1 and T2 and the second allele (of *A.tauschii* related to M allele of *T. monococcum*) was derived from a public database IWGSC-URGI (https://wheat-urgi.versailles.inra.fr/Seq-Repository/BLAST) and named as putative *At_HKT*.

The comparison between major alleles (M and N from *T. monococcum*, T and *At_HKT* from *A. tauschii)* exhibited the polymorphism at 5% level, while the comparison within sub-category of major alleles called minor alleles showed 0 to 0.2% polymorphism. The phylogenetic tree, constructed based on maximum likelihood with 1000 bootstrap, clearly revealed some distinct clusters of major and minor alleles (Fig. 3a). In 5' UTR region, a continuous stretch of adenine was observed at higher number in *A. tauschii* gene (*At_HKT* allele) from 554 bp onwards, whereas in *T. aestivum* and *T. monococcum* the repetition of adenine was lower. One variant from *A. tauschii* (*At_HKT*) is a major source of the 4DL chromosome bearing *HKT 1;5* gene to *T. aestivum* as it shown close phylogenetic relationship each other. (Fig. 3b). Distance matrix of these alleles from corresponding species is shown in supplementary table 4. The alignment around exon1 between wheat and *A. tauschii* indicates the vicinity of evolutionary relatedness between At_HKT allele and alleles of wheat, while T allele (T1, T2) lies farther (Fig. 3c). Relative divergence time concerning *HKT1;5* gene is recent for Triticeae members in Pooideae subfamily in Poaceae than model grass *Brachypodiuum distachyon* and other species belongs to the subfamily of Oryzoideae and Panicoideae as shown in Fig. 3d.

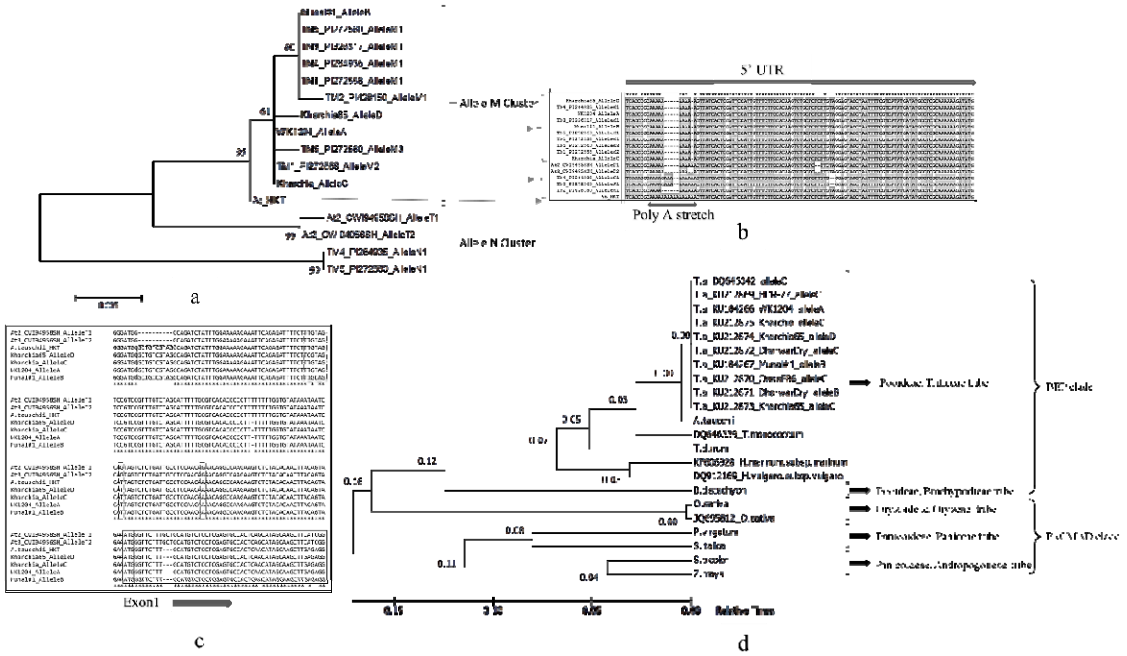

Figure 3. a) Phylogenetic comparison of M and N alleles from *T. monococcum* to *T. aestivum* and *A. tauschii* alleles. b) Alignment showing Poly A stretches in 5' UTR and nucleotide comparison among the species/sequences. c) Interspecific SNPs and insertion/deletion mutations between *A. tauschii* and *T. aestivum* alleles. Large box indicates exon1 start codon and an insertion. Dotted box indicates the difference between *A. tauschii* alleles T1, T2 and *A. tauschii* allele '*At_HKT*' sequence. Shaded box indicates SNPs difference from *T. aestivum*. d) *HKT1;5* gene based Relative Divergence time among specific tribes in Poaceae.

Concerning *A. speltoides*, amplification was achieved in some accessions accession, it perhaps due to accumulated mutations in and around primer binding sites. The *HKT1;5* gene partial sequence generated from *A. speltoides* (Allele 'As2') revealed an alteration of amino acids in primary structure of protein due to the insertion of four nucleotides (CGCG) caused a frameshift mutation at exon1 suggesting a distinct allelic variant (Genbank: KY110745, Fig. 4a). Besides, a BLASTN search for additional alleles with top score sequences from IWGSC-URGI (https://urgi.versailles.inra.fr/Species/Wheat), revealed the existence of four more alleles in *A. speltoides* with available genomic resource. Hence, there is a need of more sequencing effort in natural accessions/populations of this species to excavate further alleles. None of the database sequences showed a frameshift mutation with CGCG insertion at exon1 as 'As2' allele of *A. speltoides*. The BLASTN of the sequences (https://urgi.versailles.inra.fr) revealed that the complete gene sequence was present only on homeolog chromosome 4D. To better understand the number of gene variants on chromosome 4D, we used two ditelosomic lines, the first one called DTGH09_1559 possessed 4D chromosome long arm (L) but lacking 4D chromosome short arm (S) (Allele 'DS1') and the second one namely DTGH09_1556 (Allele 'DS2') was the opposite of DTGH09_1559 i.e. possessing 4D chromosome short arm (S) but lacking 4D chromosome long arm (L). The alignment of sequenced

homeolog from DTGH09_1559 with 4DL (-4DS) (GenBank: KY110744) revealed a 99.89% identity with *T. aestivum* (GenBank: KU184266), while it was 93.25% identical to *T. durum* (GenBank: KY110748). In contrast, the homeolog from DTGH09_1556 4DS (-4DL) (GenBank: KY110743) revealed a 98.89% identity to *T. durum* while it was 93.08% identical to *T. aestivum*. Both ditelosomic lines DTGH09_1559 and DTGH09_1556 have 93.78 % identity to each other. Protein sequence alignment among bread wheat, durum wheat and two ditelosomic lines corresponding to exon1 is shown in Fig. 4b, the nucleotide alignment with an insertion and deletion is shown in Fig 4c. This indicated that the homeolog from DTGH09_1556 4DS (-4DL) has similarity with an ortholog of durum wheat and might have amplified from 'A' or 'B' genome rather than from the D genome of ditelosomic line DTGH09_1556 4DS (-4DL). BLASTN has further confirmed that DTGH09_1556 4DS (-4DL) gene sequence was located on chromosome 4B long arm (4BL) as it showed 100% sequence identity with the scaffold TGACv1_scaffold_320406_4BL from bread wheat.

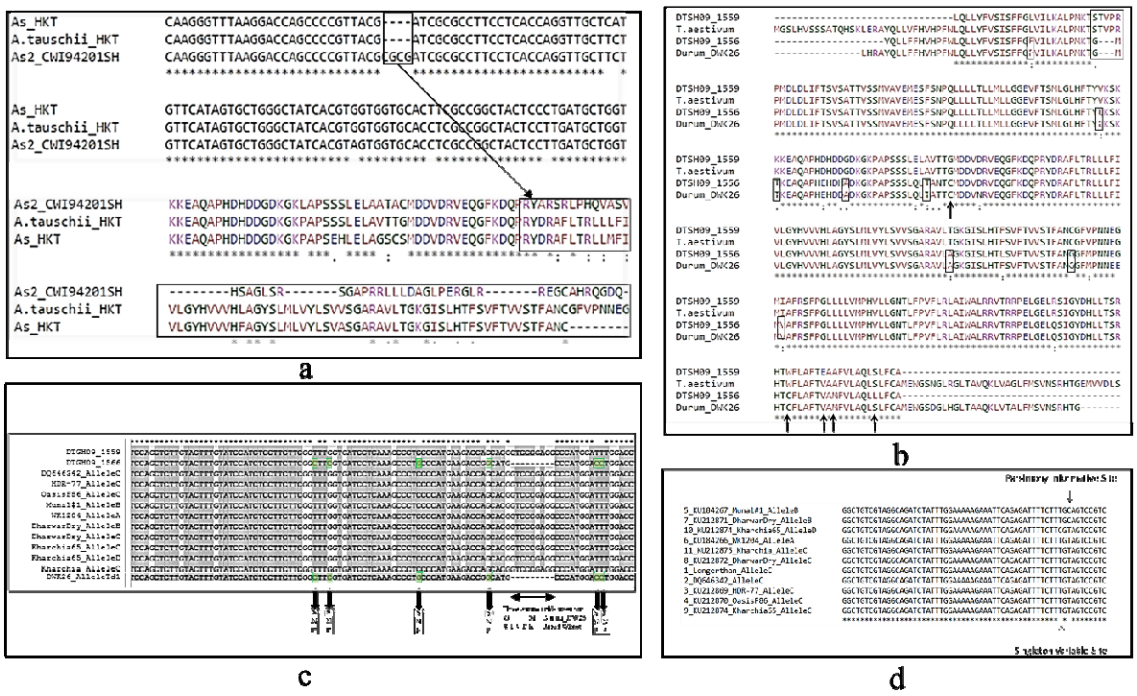

Figure 4: a) Inter specific SNPs insertion deletion variation between *A. tauschii* and *A. speltoides*. Box indicating the 4bp insertion caused frameshift with altered protein. b) Amino acids alignment corresponding to the region of exon1 of *HKT1;5* gene among bread wheat, durum wheat and diltelosomic lines. Small box indicates conservative amino acid changes. Large box indicates insertion deletions. Arrow indicates non conservative amino acid changes. c) Nucleotide alignment of *HKT 1;5* gene exon1 part among *T. aestivum*, *T. durum* and diltelosomic lines. d) Parsimony informative site and singleton variant site in *T. aestivum* alleles.

Overall, the coding sequences of *TaHKT1;5-D* against the survey sequences of all bread wheat revealed the presence of an entire genic region on 4DL, while remnants or portions of gene regions on 4BL, 4BS and 4DS chromosomes. A non-autonomous mariner transposon element was found in intron 1 of the gene on all A, B, and D chromosomes with >3120 BLASTN hits. These results clearly indicate the mobile nature of the gene through transposition events either as a whole or part of the gene, hence multiple variants of *HKT1;5* gene in bread wheat and its progenitors is highly plausible. The presence of huge variation in sequence alignment between *T. durum* cultivar DWK26 and *T. aestivum* indicated variation between the amplified gene from 4BL and 4DL respectively. The amino acid alignment evidencing that *HKT1;5* gene sequence of durum wheat was highly comparable with diltelosomic line 1556 (+4DS, -4DL), while *T. aestivum* was highly comparable with DTGH09_1559 (+4DS, -4DL). In *A. tauschii* the presence of two variants (*At_HKT* and T allele) revealed that these sequences sharing proximity among ditelosomic line 1559 (+ 4DL and - 4DS), bread wheat and *A. tauschii*, while Ditelosomic line 1556 (+4DS and -4DL) was differing with its own SNPs from others. The trend suggested that two paralogous variants existed in *A. tauschii*. Besides, ditelosomic bread wheat lines showed the existence of 2 paralogous variants, which revealed that one was amplified from DT line 1559 (+4DL -4DS) while another from 1556 (+ 4DS, -4DL). Both the alleles from *A. tauschii* (*At_HKT* and T allele) had proximity with the sequence of DT line 1559 rather than DT1556.

Genetic diversity analyses of the *HKT1;5* alleles in bread wheat

There were 2301 invariant sites, and 3 variant sites detected. Among the variant sites, 2 were singleton variant sites (at position 830th in 5' UTR and at 2106th position in Intron1) and 1 parsimony informative site (position 832nd of 5' UTR) with two variants. The variant sites at 5' UTR is shown in Fig. 4d. Diversity analyses showed low polymorphism and recombination rate for this gene among the bread wheat lines concerning these four alleles in *TaHKT1;5-D* (Table 2). Gene flow and genetic differentiation showed a similar trend with least effective migrants (0.13) F_{ST} (0.65) among the four alleles represents lesser migrants among the defined population/genotypes. It further elucidates the least frequency of recombination in this gene owing to autogamous nature and lesser allelic variation within bread wheat lines. The conservative nature of the sequences with a low number of mutations has resulted in low haplotype diversity of the four alleles. The observed total nucleotide difference was also significantly lower (Table 3).

Table 2. Polymorphic sites and DNA polymorphism. h-Number of haplotypes, hd-Haplotype diversity, SD-Standard deviation, θ- 4Nμ, where N is effective population size and μ is mutation rate per site per generation, π- Nucleotide diversity, η –Total number of mutations, S-Segregating sites, R-Recombination, NR- No Recombination, k-Average number of nucleotide differences, Sq-Sequence.

H	**Hd**	**Variance**	**SD**	π	**θ from η**
4	0.694	0.02162	0.147	0.00036	0.00048
θ from S	**Variance θ (NR)**	**SD θ (NR)**	**Variance θ (R)**	**SD θ (R)**	**θ from π**
0.00048	0.0000001	0.00032	0.0000001	0.00028	0.00036
θ from S	**θ from η**	**k**	**θ from S/Sq**	**Variance θ (NR)/Sq**	**Variance θ (R)/Sq**
0.00048	0.00048	0.833	1.104	0.545	0.406

The percentage of nucleotides was 22.96% (A), 27.73% (T/U), 26.83% (C), and 22.48% (G) derived from the alleles studied corresponding to 2304 positions. The C to T transition mutation (pyrimidines) substitution pattern was higher (50.59%) than other type of transition mutations. Besides, the transversion mutations (purine to pyrimidine or vice versa) were lower between the ranges of 0.04 to 0.05 %. Among the bread wheat cultivars, SNPs were not observed in coding region of the gene, hence Ks/Ka (synonymous/non-synonymous mutation) analysis was not performed. Tajima D test (D: -0.93613) with negative value indicates purifying selection due to low frequency of variations along this gene among the bread wheat cultivars and previse population size expansion. However, Tajima D test was significant and thus increasing sample size may provide a better clue.

Table 3. Genetic diversity, differentiation and gene flow analysis from B and C Alleles of *TaHKT1;5-D* gene. Hd-Haplotype Diversity, Kt- Average Nucleotide Differences, PiT- Nucleotide diversity, χ^2- Chi Square test, Hst- Haplotype based statistics, Kst- Sequence based statistics, Z*- Rank Statistics, Snn- Near Neighbour Statistics, Gst- Differentiation of population, GammaSt- Gamma Statistics, Nst- N statistics, Fst- Fixation index, Nm- Effective number of migrants, PM- Permutation test with 1000 replicates, * 0.05<P.

Genetic Diversity						
Alleles	**Sequences**	**Segregating sites**	**Haplotypes**	**Hd**	**Kt**	**PiT**
A, B, C, D	9	3	4	0.6944	0.8333	0.00036
Genetic differentiation						
Estimated	χ^2	**Hst**	**Kst**	**Kst***	**Z***	**Snn**
Value	9.77	0.15	0.27	0.12	3.84	0.43
PM test	0.13	0.10	0.027*	0.16	0.11	0.18
Gene flow						
	Haplotype data		**Sequence Data**			
Estimated	**Gst**		**DeltaSt**	**GammaSt**	**Nst**	**Fst**
Value	0.319		0.00008	0.31624	0.65908	0.65909
Nm	0.53		0.54	0.54	0.13	0.13

Phylogenetic analysis of the *HKT 1;5* gene

The phylogenetic tree based on all generated gene sequences and additional Triticeae member sequences from public databases revealed the presence of five orthologous groups with each group consisting of closely related species or from closely related chromosomes in case of polyploid species (Fig. 5). The ditelosomomic line DTGH09_1559 clustered together with the *T. aestivum* and *A. tauschii* cultivars. The alleles from *T. monococcum* and *A. speltoides* deviated from the orthologous group I and were designated as orthologous group II. Alleles from ortholog group III existed with III sub-groups of bread wheat and durum wheat samples and have proceeded from A, B and D putative genome donors from ortholog group II. The four alleles of *A. speltoides* retrieved database comes under three ortholog groups with other related species. The orthologous group IV consisted of alleles from *A. sharonensis*, *A. speltoides*, *Hordeum spp* and *B. distachyon*. This group appeared as a distinct group from the ortholog groups I-III. Group V consists of other Poaceae members used in this study apart from Triticeae and Brachypodieae tribes.

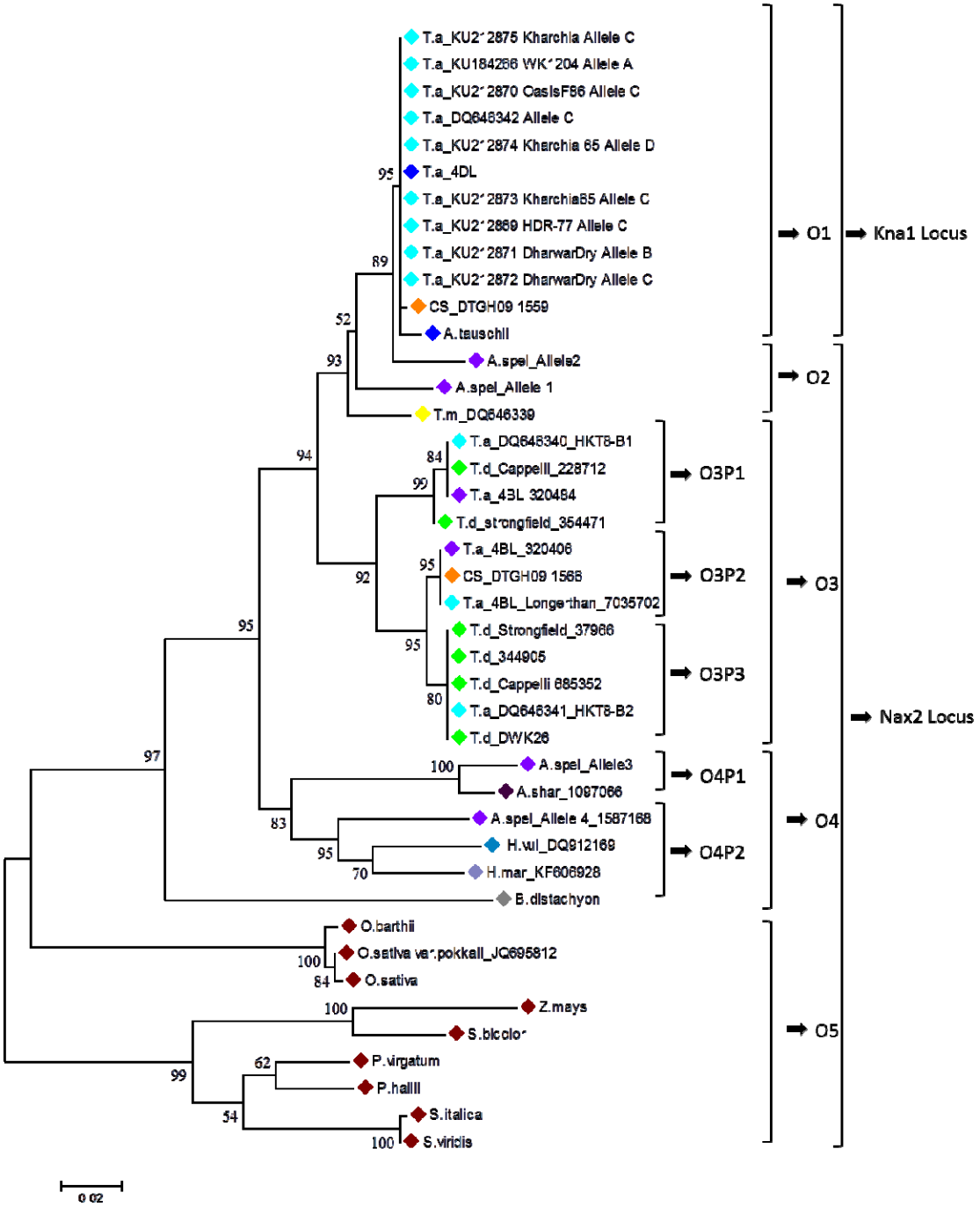

Figure 5. Reconstructed phylogenetic tree showing the orthologous and paralogous comparison of HKT1;5 of *Kna1* and *Nax2* homoeologs relevancy of the gene in different species. O=Ortholog, P=Paralog. Phylogeny based on Maximum Likelihood method with Jukes Cantor (JC) model and 1000 bootstrap resampling.

Comparative analysis between *T. aestivum* and *A. thaliana*

Protein modeling and comparative proteomics indicated structural conservation of protein sequence between *A. thaliana* and *T. aestivum* (Fig. 6a). Further alignment between *T. aestivum HKT1;5-D* and *A. thaliana HKT1* revealed similarity of 54.9 % and 48.4%, respectively, in exon and protein sequences.

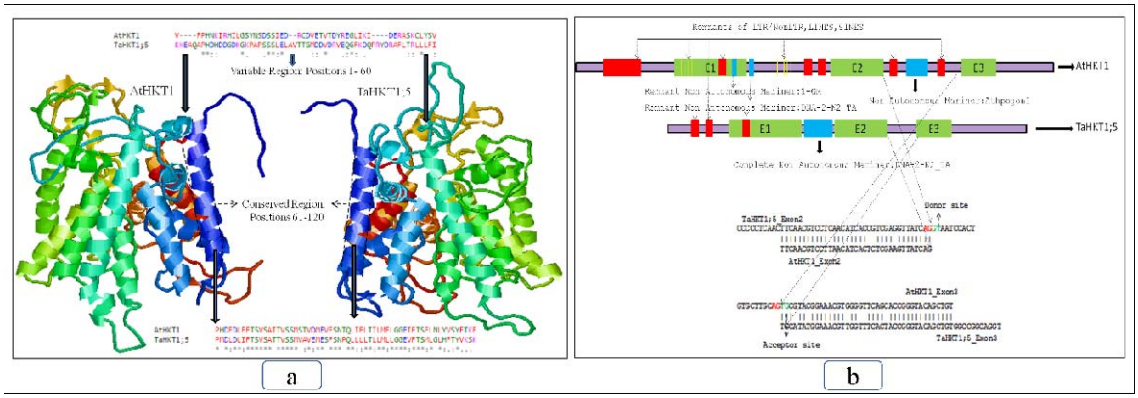

Figure 6. a) Protein homology modelling, tertiary structure, and structural differences in comparison with primary structure of *HKT1* protein between *A. thaliana* and *T. aestivum*. b) Structural organization and insertions transposons/fragments of transposon in *HKT1* gene from A. *thaliana* and *HKT1;5* from *T. aestivum*.

Nucleotide level global alignment with LAGAN revealed the presence of higher conservation in coding or exonic part of *HKT1* sequences between wheat and *A. thaliana*. The comparison of transposable elements inside the gene showed that *A. thaliana HKT1* gene has higher number of transposons or remnants of transposons inside the gene rather than bread wheat. The conservation of CDS region, donor and acceptor splicing sites proved the similarity of the structural organization of this gene between *A. thaliana* and *T. aestivum*. Introns are enriched with the presence of more remnants of different transposon; however, a complete non-autonomous transposon called Mariner in intron1 was highly comparable between these two-diverse species (Fig. 6b). Physico-chemical properties of the protein comparison between *A. thaliana* and *T. aestivum* revealed a higher similarity of this orthologous gene from these two-diverse species (Supplementary Table 5). The structural organization of the corresponding putative proteins sequence also revealed higher similarity. The instability index (II) indicates that *HKT1* protein from *T. aestivum* and *A. thaliana* is stable.

Salinity stress induced phenotypic evaluation in selected allele designated wheat lines

Initial germination experiments and traits analysis with salinity tolerant reference lines and sensitive lines shown in supplementary table. 6. Additional germination related phenotypic analysis was carried out for the bread wheat cultivars as shown in table 4. Statistical results concerning wheat cultivars to salinity stress is listed in table 4. Detailed description of the analysis in cultivars HDR-77 (Allele ‘C’), C306 (Allele ‘C’), Oasis F86 (Allele ‘C’) and WK1204 (Allele ‘A’) is given here. The germination test among these four cultivars showed HDR-77 to be early germinating than other three cultivars. The cultivar WK1204 with ‘A’ allele was later germinating. Final Germination Percentage (FGP) (Fig.7a) was higher for the varieties HDR-77 and C306 at 200 mM than WK1024, while Germination Efficiency Percentage

(GEP) (Fig.7b) was higher in Oasis F86. Fresh Weight (FW) was increased upon increasing the salinity concentration at 200 mM level than control in all varieties, comparatively FW was higher in C306 and HDR-77 than WK1204 (Fig. 7c).

Table 4. Means of traits under different treatments of salt including control and means of phenotypes for each genotype responding to treatment. Letters not sharing similarity are statistically significant to LSD in each rows at P=0.05 level. FGP= Final Germination Percentage, GE= Germination Energy, SVI= Seed Vigor Index, CL= Coleoptile Length, RL= Radicle Length, NoR= Number of radicles, ColL- Leaf Length from Coleoptile, FW= Fresh Weight, DW= Dry Weight, RWC= Relative Water Content, MT= Mean Germination Time, CVt= Coefficient Variation of Germination Time, MR=Mean Germination Rate, U= Uncertainty of the Germination Process, Z=Synchrony. Statistical significance. p<1, *P<0.05, **p<0.01, ***P<0.001, Ns- Not significant.

			Treatment				
Traits	**Control**	**50 mM**	**100 mM**	**150 mM**	**200 mM**	**LSD at P=0.05**	**F test significance**
FGP	96.11a	88.88ab	88.33ab	86.66b	83.88b	7.8	0.03957 *
GE	13.33a	11.11a	10.55a	8.88a	5.55a	8.68	0.47182 Ns
SVI	998.66a	891.99a	635.48b	396.34c	255.15d	116.95	< 2e-16 ***
CL	44.68a	43.72ab	38.61bc	37.68c	36.95c	5.72	0.0205 *
RL	103.60a	100.14a	72.20b	44.45c	29.67d	12.36	< 2e-16 ***
NoR	3.98a	3.98a	3.79ab	3.75ab	3.59b	0.26	9.31e-07 ***
ColL	104.49a	99.40a	82.81b	70.86c	42.47d	8.94	4.39e-16 ***
FW	1.47a	1.42a	1.34a	1.33a	1.08b	0.15	8.83e-05 ***
DW	0.50a	0.49a	0.49a	0.43b	0.42b	0.04	0.000918 ***
RWC	68.93a	66.27ab	63.00bc	61.91bc	59.10c	5.38	0.00613 **
MT	3.24a	3.02a	2.68b	2.67b	2.65b	0.25	2.51e-05 ***
CVt	7.71a	7.11a	7.11a	6.41a	6.39a	1.66	0.4682 Ns
MR	0.38a	0.38a	0.38a	0.34b	0.31b	0.03	0.00052 ***
U	1.70a	1.62a	1.58a	1.53a	1.52a	0.32	0.824 Ns
Z	29.97a	25.50ab	22.77ab	19.20b	15.94b	10.31	10.0772
			Genotypes				
Traits	**C306**	**HDR-77**	**WK1204**	**Oasis F86**	**LSD at P=0.05**	**F test Significance**	
FGP	96.88a	90.22ab	84b	84b	6.98	0.00108 **	
GE	18.22a	9.77b	8.00b	3.55b	7.76	0.00393 **	
SVI	820.48a	717.02a	520.01b	484.58b	104.6	2.47e-08 ***	
CL	49.14a	44.55a	34.06b	33.55b	5.11	3.15e-08 ***	
RL	84.52a	79.01a	60.44b	56.08b	11.05	2.75e-06 ***	
NoR	4.23a	3.89b	3.67bc	3.50c	0.23	9.31e-07 ***	
ColL	94.15a	90.73a	77.30b	65.85c	7.99	4.99e-09 ***	
FW	1.44a	1.41a	1.36a	1.10b	0.14	2.49e-05 ***	
DW	0.55a	0.50b	0.48b	0.34c	0.04	1.73e-13 ***	
RWC	68.24a	65.47ab	62.08bc	59.59c	4.81	0.00416 **	
MT	3.57a	2.78b	2.64bc	2.43c	0.22	6.04e-13 ***	
CVt	10.12a	7.52b	5.08c	5.07c	1.49	5.95e-09 ***	
MR	0.41a	0.38ab	0.36b	0.28c	0.03	4.21e-10 ***	
U	1.93a	1.75a	1.38b	1.29b	0.29	8.45e-05 ***	
Z	36.33a	30.44a	16.24b	7.70b	9.22	1.76e-07 ***	

All the varieties exhibited higher dry weight at 200 mM than control, notably HDR-77 exhibited higher Dry Weight (DW) than the rest of the varieties (Fig. 7d). Relative Water Content (RWC) was higher for C306 than rest of the cultivars at 200 mM (7e). It suggested that the salt sensitive variety Oasis F86 was able to retain and absorb the water efficiently than rest of the cultivar in the absence of salt stress. However, the salinity tolerant cultivar C306 exhibited with more RWC than the rest of the cultivars including Oasis F86. Radicle Length (RL) (Fig. 7 f, h) varied or decreased corresponding to increased salinity concentration, proving the inversely proportional relationship between these two factors. Number of radicles (NoR) (Fig. 7 g, i) in C306 increased at 150 mM and 200 mM than control, while it decreased in salt tolerant cultivar HDR-77 in 200 mM than control. Effect of salinity on coleoptile and radicle size of the seedlings of C306 and HDR-77 is shown in Fig. 7 j and Fig. 7 k respectively.

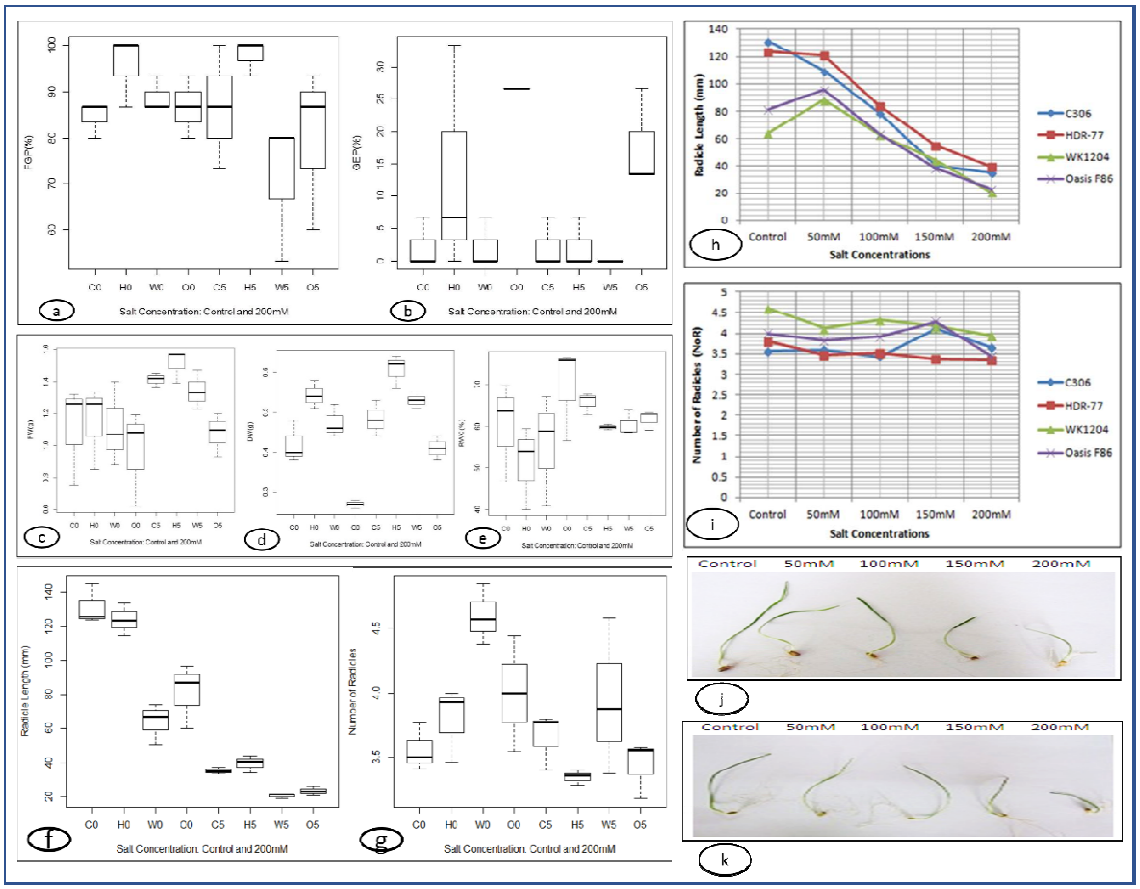

Figure 7. a) Box plot showing the distribution of variation for RL (Radicle Length) among the cultivars between control and high stress (200 mM). b) Box plot showing the distribution of variation for Number of Radicles (NoR) among the cultivars between Control and high salt stress (200 mM). Legend: C0-C306 Control, H0-HDR-77 control, W0-WK1204 control, O0-Oasis F86 control; C5-C306 200 mM, H5-HDR-77 200 mM, W5-WK1204 200 mM, O5-Oasis F86 200 mM. c) Effect of salinity on Number of Radicles

d) Effect of salinity on Radicle Length. e) Reduction of coleoptile and radical size upon increasing salt stress in bread wheat HDR-77.

Our study revealed that the cultivars C306 (Allele ‘C’) and HDR-77 (Allele ‘C’) are more tolerant to salt than WK1204 (Allele ‘A’: g.[830T>C;832T>C;2106;T>C]) and Oasis F86 (Allele ‘C’). Relative Water Content was higher for saline sensitive cultivar and lower for HDR-77, indicates even saline sensitive cultivars able to accumulate or assimilate the water more than the amount of water needed by tolerant cultivars. Mean Germination Time of both sensitive cultivars were increased due to less Mean Germination Rate and these trends are in contrast with tolerant cultivars under stress conditions. Thus, the cultivars WK1204 and Oasis F86 are more susceptible to salt stress than HDR-77 and C306. These analyses revealed that the cultivar HDR-77 is far most tolerant to salinity than all three cultivars, C306 is in the second most tolerant cultivar. The allele ‘C’, was common to both salinity tolerant and susceptible cultivars such as Kharchia and OasisF86, while only one genotype WK1204 had allele ‘A’, which is saline sensitive cultivar as revealed with our analysis. Dharwar Dry is a salinity tolerant cultivar possessed both ‘B’ (*T. aestivum* cultivar Dharwar Dry) and ‘C’ alleles (*T. aestivum* cultivar Dharwar Dry). Similarly, the salinity tolerant cultivar Kharchia 65 possessed both ‘C’ and ‘D’ alleles. It is intriguing that cultivars showed sensitiveness had allele ‘A’ and some of them with allele ‘C’, while the combination of alleles ‘B’ (g.[830T>C;832C>T;2106;T>C]) or ‘D’ (g.[830C>T;832T>C;2106;C>T]) present in *T. aestivum* cultivar Kharchia 65), which is known saline tolerant cultivar.

Greenhouse based germination related phenotypic analysis was carried out for the bread wheat cultivars used for allele mining in this study. Statistical results concerning wheat cultivars to salinity stress is listed in table 5. Mostly saline sensitive bread wheat cultivars OasisF86 (Allele ‘C’), WK1204 (Allele ‘A’) and durum wheat cultivar DWK26 was observed with earlier booting or heading stage in short period (50 DAS) of the time since the salinity exposure. However, salinity stress didn’t induce early (50 DAS) booting and heading in salinity tolerant variety bread wheat variety Kharchia (Allele ‘C’), drought tolerant and heat tolerant varieties HDR-77 (Allele ‘C’) and C306 (Allele ‘C’) (Table 6). The greenhouse-based experiment also revealed that the saline sensitive cultivars tend to have an early maturation and short life cycle revealed through the measurement of early booting and flowering stages (Supplementary table 7). In addition, the salinity tolerance level in salinity tolerant, drought and heat tolerant cultivars were exhibited with higher salinity tolerance as well, while the known sensitive cultivars showed lesser salt tolerance as expected by exhibiting the restricted phenotypic characteristics unlike the tolerant ones.

Table 5. Means of traits under different treatments of salt including control and means of phenotypes for each genotype responding to treatment. Letters not sharing similarity are statistically significant to LSD in each rows at P=0.05 level. PH=Plant Height, NoT=Number of Tillers, NoET=Number of Effective Tillers, FLL=Flag Leaf Length, TS= Total Number of Spikes, SL= Spike Length. Statistical significance: .p<1, *P<0.05, **p<0.01, ***P<0.001, Ns- Not significant.

				Treatment			
Traits	**Control**	**50 mM**	**100 mM**	**150 mM**	**200 mM**	**LSD at P=0.05**	**F test significance**
PH	59.44a	59.27a	57.72a	56.16a	47.88b	7.08	3.20e-05 ***
NoT	5.05a	4.72a	4.66a	4.50a	4.38a	1.15	0.52 Ns
NoET	4.11a	3.38ab	3.38ab	3.00b	2.72b	0.86	0.000274 ***
FLL	55.19a	50.86ab	43.72ab	42.58b	41.94b	12.57	0 0.00907 **
TS	4.11a	3.38ab	3.38ab	3.00b	2.72b	0.86	0.000274 ***
SL	25.66a	23.22ab	21.08ab	19.16bc	15.86c	4.88	1.58e-06 ***

				Genotypes				
Traits	**C306**	**HDR-77**	**Kharchia**	**OasisF86**	**WK1204**	**DWK26**	**LSD at P=0.05**	**F test Significance**
PH	62.76a	61.56a	60.93a	51.70b	50.80b	48.83b	8.13	5.11e-08 ***
NoT	6.60 a	5.00 b	4.86b	4.53bc	3.73bc	3.26c	1.33	9.8e-10 ***
NoET	3.80a	3.80a	3.40ab	3.33ab	2.86ab	2.73b	0.99	0.004342 **
FLL	63.80a	62.90a	59.90a	35.06b	33.21b	26.30b	14.44	3.08e-15 ***
TS	3.80a	3.80a	3.40ab	3.33ab	2.86ab	2.73b	0.99	0.004342 **
SL	25.43a	22.63ab	22.53ab	21.36ab	19.66bc	14.36c	5.6	2.74e-06 ***

Statistical result from greenhouse experiment revealed that bread wheat variety C306 (Allele 'C') is more saline tolerant (PH: 62.76a, LSD at P=0.05, F test significance 5.11e-08, P<0.001) (Table 5) than other bread wheat varieties. Oasis F86 (Allele 'C') is saline sensitive cultivar and comparable with WK1204 (Allele 'A') and durum for salt sensitiveness (Fig. 7). The overall experiment resulted the salinity sensitiveness and tolerance in varieties through the comparison of phenotypes with alleles.

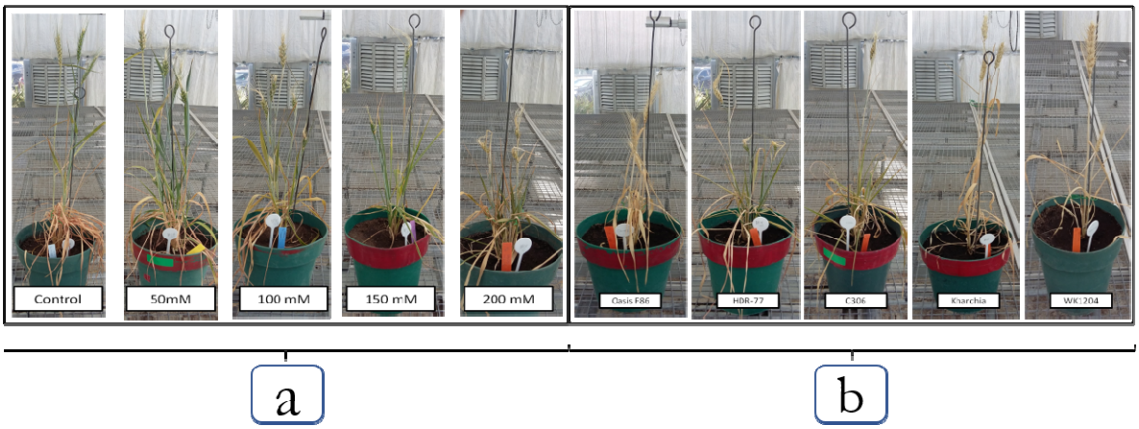

Figure 8. a) Salinity induced growth retardation of the cultivar HDR-77. b) Phenotypic effect of high salinity stress with 200mM on bread wheat cultivars.

Our greenhouse experiment demonstrates that the varieties C306 and HDR-77 even shown slightly better salt tolerance over known salinity tolerant reference line Kharchia. However, wheat variety like Munal#1 evinced slightly better tolerance with germination related salinity stress than varieties WK1204, OasisF86 and Dharwar Dry. Germination related study shewed that the variety WK1204 is slightly better salinity tolerant than the variety OasisF86, while the green house experiment is contrary, as OasisF86 exhibits slightly better phenotypic features with salinity stress than the variety WK1204. Perhaps, a known saline sensitive variety OasisF86 is very sensitive during germination with salinity stress. Nonetheless, the varieties C306 and HDR-77 had better tolerance with germination related salt stress and with greenhouse study (Supplementary Table 8). A comparative germination related salt stress experiment derived data is shown in (Supplementary Table 9).

The most common allele 'C' was present both in saline tolerant and sensitive cultivars, while Allele 'A', 'B' and 'D' were rare and observed with both tolerant and sensitiveness depends upon the cultivars. The current study has deciphered the most redundant allele in both tolerant and sensitive cultivars that is allele 'C', however additional study is needed to understand why this allele is widely present among the studied cultivars. However, some cultivars were observed other alleles or combination of alternative form of gene with heterozygotic nature. Therefore, there is a need of additional studies on various genes linked through metabolic pathways or even transcription factors. Nonetheless, the current study provided the presence of multiple variants or alleles in wheat and progenitor species and the plausible comparison with phenotypes during salinity exposure. Overall, this current study proved the salinity tolerance and sensitiveness in known tolerant, sensitive and unknown lines of bread wheat and compared with *HKT1;5* alleles. However, salinity tolerance is complex physiological and genetic trait, hence further studies such as breeding and pre-breeding strategies are necessary to develop a better tolerant variety with sustainable yield.

DISCUSSION

To better understand the sequence variation of *TaHKT1;5-D* gene in bread wheat, the whole gene 3Kb sequence was amplified and sequenced in bread wheat varieties and wild species. Only three SNP were identified in 3Kb sequence, which indicates high sequence conservation in *TaHKT1;5-D* gene within bread wheat. However, large numbers of inter-specific SNPs were found when gene sequences of bread wheat and wild relatives were aligned. In congruence with previous studies (Huang *et al*., 2008), we also could not amplify *HKT1;5* gene in *T. urartu*, which supports the conclusion that the gene might be absent in *T. urartu*. Thus, none of the variant of *HKT1;5* gene was observed in A genome of wheat (Huang *et al*., 2008). However, *TmHKT1;5-A* from *T. monococcum* was presumed as strong candidate gene for *Nax2* locus, which is homeolog to *Kna1* locus with a single variant located at distal 5AL chromosome in bread wheat (Byrt *et al*., 2007). Being an allogamous species and B genome progenitor, *A. speltoides*

tends to have slightly higher mutations/variant form of gene, thus we could achieve amplification only in some accessions. Hence specific attention must be given in this species to excavate the hidden form of this gene to answer whether it is due to supernumerary nature of B chromosomes with more intraspecific nucleotide variations (Jones et al., 2008). Also, this kind of study would provide better understanding of orthologous gene variations and gene flow of wild relatives and domesticated species. Gene flow from wild to domesticated species through pre-breeding entails the conservation of genetic diversity (Papa *et al*., 2005) and alien transfer of genes with functional relevancy (Jauhar *et al*., 2003).

Phylogenetic analysis among the orthologs of *HKT1;5* from diverse species of poaceae revealed the existence of five groups based on the closeness of each species or chromosomal relevancies. The existence of multiple variants in putative B genome donor *A. speltoides*, would provide a clue for the presence of 3 variants in 4BL chromosome (Huang *et al*, 2008) of bread wheat. However, the *A. speltoides* allele 1 and 2 (from ortholog group II), revealed the similarity with the gene from 4DL chromosome and exhibited the vicinity to *T. monococcum* highlights the homeolog comparisons of *Nax2* locus and *Kna1* locus. Further comparison of ortholog group III and IV, revealed the presence of maximum variants in B genome itself or B genome related sitopsis section species *A. sharonensis.* Further the ditelosomic line experiments revealed that one homeolog is exclusively present on 4DL chromosome and the other one on 4BL of bread wheat, the possibility that this homeolog region from 4BL could have ancestrally translocated to 4DL through homoeologous recombination after AABB+DD polyploidization event, perhaps any mutation at chromosome 5B encoded mutant *Ph1* locus would have not suppressed the homoeologous recombination (Koo et al., 2020, Gaeta et al., 2009, Qi et al., 2007) or *Ph1-inhibiting gene* called *PhI* kind of gene would have rendered homoeologous recombination by inhibiting Ph^1 gene (Jauhar 2003). A recent study with resequencing of diploid progenitors of AADD allotetraploid line revealed the similar trend about the nonreciprocal homeologous exchange between 4A and 5D (Zhang et al., 2020) on the regions of ancient translocations of long arms between chromosomes 4A and 5A in *T. urartu* (Ling et al., 2018).

Several previous genetic analyses also compared the monocots and dicots using different genes and genomic data (Subburaj *et al*., 2016, Sarma *et al*., 2014, Cserhati., 2015). It is noteworthy that the model plant *A. thaliana* and its close relatives from family *Brassicaeae* reportedly represented a vital role in evolution and functional relevancies of plant kingdom (Koornneef and Meinke *et al*., 2010; Hu *et al*., 2011).

Extensive comparative genomics for *HKT* gene between *Arabidopsis* and monocots (Horie *et al*., 2009) and between *A. thaliana* and wheat were also reported (Peng *et al*, 2015). Since *A. thaliana* is a model dicot species, further comparative genomics analysis was done with wheat gene to understand the conservation and variations of the *HKT* gene. Previous report also compared the model dicot species *A.*

thaliana with wheat and barley (Peng *et al*, 2015). Alignment of the proteins *AtHKT1* and *TaHKT1;5-D*, disclosed the identity of numerous amino acids and conservative substitution of amino acids. Secondary structure and physio-chemical properties also revealed the close vicinity between the two proteins. Homology modelling based tertiary structure prediction exhibited a least structural change between these orthologous proteins in accordance with the comparison of primary, secondary, and tertiary structures. The comparison of the presence of negatively charged and positively charged amino acids were closer. The instability index was below 40, stating that the stability of both proteins was not affected. Therefore, there was no maximum structural change or dissimilarity observed between these orthologous proteins.

The remnants of SINE9-OS and TABARE (Retroviral like sequence of plants) were observed in 5' UTR region of the *TaHKT1;5* gene. The exon1 was disrupted with remnants of LTR retrotransposon Copia-1_SSa-I, EnSpm-14_SBi and RTE-11_BF autonomous Non-LTR Retrotransposon. Exon 1 and exon 2 was separated by a complete non-autonomous mariner DNA transposon DNA-2-N2_TA, which occupied almost all intron1. Intron 2 neither produced any blast hits on repeated elements nor showed any transposon signals/signatures. The alignment of the mRNA region (exon 2 and exon 3) from *TaHKT1;5* gene with *A. thaliana HKT1* mRNA, resulted a clear donor and acceptor sites alignment between these two orthologues genes. *A. thaliana HKT1* gene (derived from chromosome 4: Genbank CP002687, Mayer *et al*., 1999) showed higher number of transposon remnants including DNA and Retrotransposons than the gene of *TaHKT1;5*. Therefore, the size of the gene was modulated with 3680 bp and 2087 bp (from start to stop codon) in *A. thaliana* and *T. aestivum*, respectively. Unlike wheat, only one *HKT1* gene was reported in *A. thaliana* (Uozumi *et al*., 2000) and may have been diverged from its ancestor and evolved substantially in *in* Poaceae members of monocots due to the presence of several paralogs (Waters et al., 201). Global alignment between the gene of *AtHKT1* and *TaHKT1;5-D*, revealed higher conservation/similarity in CDS region from both species. However, the introns showed least conservation, for instance remnant of mariner transposon in intron1 of *AtHKT1* showed similarity in *TaHKT1;5-D* with same region. BLASTN analysis showed that the existence of this mariner element in almost all chromosomes of bread wheat at 3120 BLASTN hits with >75% sequence similarity from A, B, D genomes of about 1000 hits per genome. Whereas 740 BLASTN hits in *A. tauchii*, 1068 and 1030 hits in *T. urartu* and *A. speltoides* respectively with >75 % sequence similarity was observed. It appeared that after polyploidization, the D genome reinforced with the multiple movements of this transposon as D genome progenitor species *A. tauschii* exhibited lesser hits for this element than D genome in bread wheat. The BLASTN analysis from the survey sequences (IWGSC-URGI and NCBI wheat) revealed that the entire gene was present in Chromosome 4DL of bread wheat. Chromosome 4BL also found to contain entire length of the gene as reported by Huang *et al*., 2008. *A. speltoides* resulted the presence of multiple variants of the gene in its genome. Any of the A chromosomes did not show reasonable query coverage, although remnants of the gene (region belongs to non-autonomous mariner transposon corresponding to

intron 1) present in almost all chromosomes even from chromosome A. It is important to note that BLASTN analysis revealed the presence of remnants of various parts of the gene (for instance the region of exon 2 and intron 2 of about 20 %) in *T. urartu*, but not the whole gene. However, the presence of the gene was already reported in *T. monococcum* and *A. tauschii* (Byrt *et al*., 2007) and BLASTN hits confirmed the same. The presence of the gene in *T. durum* but absence in *T. urartu*, suggested that during or after polyploidization, it could have inherited from B genome donor rather than A genome donor *T. urartu* (Huang *et al*., 2008).

Salinity tolerance assessment was made during the germination of drought and heat tolerant cultivars such as HDR-77 (Allele 'C': g.[830T>C;832T>C;2106;C>T]) and C306 (Allele 'C') respectively and salt sensitive cultivar Oasis F86 (Allele 'C') along with an unknown cultivar for salinity tolerance namely WK1204. The bread wheat variety HDR-77 was reported to have higher water-binding characteristics and seed membrane stability, which confers increased drought tolerance (Chatterjee and Nagarajan, 2006). The cultivar C306 is a traditionally known drought and heat tolerant variety, which is being extensively used for the cultivation in rain-fed areas. The varieties WK1204 and Oasis F86 are known to yellow rust resistance (Thapa D.B. 2020) and saline sensitiveness (Kazi and Hettel, 1995) respectively. Our study revealed that saline tolerant cultivars were able to elongate their radicle length more than the sensitive cultivars, while the radicle numbers were less in tolerant cultivars than the sensitive cultivars. Saline sensitive cultivars were able to produce a greater number of radicles than the tolerant ones and it is contrast to radicle length, similar study was previously reported (Rahman et al., 2016). Green house study revealed that generally abiotic stress tolerant cultivars are tolerant to salinity as well, even the drought or heat tolerant varieties such as HDR-77 or C306 showed better salinity tolerance and exhibited increased yield related parameters. The prevalent 'C' allele is observed in more lines especially in tolerant lines, while the allele 'A' was specific to WK1204. Our study revealed that varieties Dharwar Dry and WK1204 are saline sensitive and comparable to reference salinity sensitive line Oasis F86, which possess allele 'C'. Few wheat lines observed with, heterozygous forms such as Dharwar Dry with 'B','C' and Kharchia 65 with 'C''D'. We presume that homozygous form of 'A' allele is a rare form of allele only present in WK1204, while at heterozygous form 'B' and 'D' allele were observed in Dharwar Dry and Kharchia 65 lines along with their 'C' allele. Hence, we conclude that salinity tolerance is a complex form of physiological trait and further attention with genetic focus for the validation of other interacting genes and factors of salinity tolerance is needed. Nonetheless, our study on SNPs/allele mining, phenotypic evaluation with possible correlation of alleles in wheat with an evolutionary focus would be an asset to develop a sustainable salinity tolerant wheat variety with marker assisted selection and breeding programs.

CONCLUSION

This study proved the presence of multiple variants of the *HKT1;5* gene in bread wheat and its wild relatives. The paralogous/orthologous variants or major alleles exhibited number of SNPs variation, while minor alleles showed lesser SNPs variations. The alleles comparison with salinity phenotypes provided the evidence of *HKT1;5* alleles behind in salinity tolerance and sensitiveness. SNPs/alleles and ortholog/paralog variants identified from this study, can be extensively used at intra or inter-specific breeding to enhance the salinity tolerance in bread or durum wheat, upon introgression of this gene from the compatible wild relatives. Conservative nature of this gene between monocots and dicots was also elucidated. Allele mining and evolutionary focus would be an asset for pre-breeding strategies for conservation of genetic diversity and genetic resources with an alien transfer of genes to excavate enigmatic properties of wild relatives. Overall, these analyses highlight the presence of multiple allelic variants in wheat as well from their progenitors from different geographical origins.

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AUTHOR CONTRIBUTIONS

Devised and designed the experiments: KT, PM, AKJ. Performed the experiments, data generated, analysed, and wrote the manuscript: KT. Assisted in revisions: PV, DS, RV, AKJ, PM. Critical suggestions for manuscript improvement: PV, HR, GV, PG, AL, CC, EP. Contributed reagents/materials/analysis tools: KT, PM, AKJ, DT, SP.

DECLARATION

The authors have declared no competing interests.

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