Research Article

Molecular Cloning and Bioinformatics Analysis of a New Plasma Membrane Na^+/H^+ Antiporter Gene from the Halophyte *Kosteletzkya virginica*

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A new plasma membrane Na^+/H^+ antiporter gene (named as *KvSOS1*) was cloned from the halophyte *Kosteletzkya virginica* by reverse-transcription-polymerase chain reaction (RT-PCR) and rapid amplification of cDNA ends (RACE) technology, which is a homologue of SOS1 (salt overly sensitive 1). The full-length cDNA is 3850 bp and contains an open reading frame (ORF) encoding a protein of 1147 amino acids with a molecular weight of 127.56 kDa and a theoretical pI of 6.18. Bioinformatics analysis indicated that the deduced protein appears to be a transmembrane protein with 12 transmembrane domains at the N-terminal region and a long hydrophilic tail in cytoplasm at its C-terminal region and shares 72–82% identity at the peptide level with other plant plasma membrane Na^+/H^+ antiporters.

1. Introduction

The salinization of soil has become a widespread environmental problem and an important factor in limiting agricultural productivity worldwide. At present, more than 800 million hectares land in the world is affected by salinity and this amount accounts for more than 6% of the world's total land area [1]. Even worse, the saline soil is still rapidly expanding due to irrigation, improper drainage, entry of seawater in coastal areas, and salt accumulation in arid and semiarid regions [2, 3]. So there is an urgent need to develop salt-tolerant crops which can grow in saline environments to overcome farmland salinization as well as enable agriculture in marginal lands [4–6].

As far as we know, the detrimental effects of salt stress on plant can be summarized into three main aspects. Firstly, saline soil leads to osmotic stress, which makes plants hard to

take up water from the soil. Secondly, salt stress may induce ionic toxicity. The increase of Na^+ and Cl^- concentration in the cytosol can negatively affect enzymes and lipids in the cells. When Na^+ and Cl^- concentrations increase to the toxic threshold, cells tend to die. At last, high soil salt concentration can also induce oxidative stress, which can cause a series of oxidative damage [1]. Fortunately, salt-tolerant plants have evolved some special mechanisms of salt tolerance which are to minimize the accumulation of toxic ions in plant tissue, partition them in the apoplast and vacuole, increase the synthesis of osmotic adjustment substances such as proline and betaine for maintaining tissue water status, and enhance antioxidant capacity to prevent the occurrence of oxidative stress [7]. Based on the above physiological mechanism, researchers have successively cloned many genes related to salt stress in various plants (e.g., genes encoding ion transporters, osmolytes, antioxidant enzymes,

components of calcium signaling, and others) [8]. Among them, Na^+/H^+ antiporter genes have been proved to play an important role in salt tolerance, which are considered as promising genes for breeding salt-tolerant crops via genetic engineering. Under salinity, the great problem faced by plants is to maintain Na^+ homeostasis in the cytosol because low cytosolic Na^+ is crucial for cell metabolism [9–11], and this can be achieved by Na^+/H^+ antiporters located in vacuolar membrane and plasma membrane. The vacuolar Na^+/H^+ antiporters (Na^+/H^+ exchangers, NHXs) can actively transport excessive Na^+ into the vacuole for Na^+ compartmentation, while the plasma membrane-located Na^+/H^+ antiporters are responsible for Na^+ exclusion from the cytosol to the external medium [12, 13]. Since studies on *Arabidopsis* showed that overexpression of either the vacuolar membrane Na^+/H^+ antiporter AtNHX1 or the plasma membrane Na^+/H^+ antiporter AtSOS1 could improve the salt tolerance of transgenic plants [14, 15], more and more studies have focused on the cloning and function of Na^+/H^+ antiporter genes from other plant species [16–23]. In the future, more attention should be paid to develop Na^+/H^+ antiporter genes and other salt tolerance genes from halophytes because of their inherent and excellent salt resistance.

Kosteletzkya virginica (L.), also commonly known as seashore mallow, is a perennial facultative halophytic species in the Malvaceae family, natively distributing in coastal areas containing 0.3 to 2.5% sodium salt (mainly NaCl) from Long Island along the Atlantic coast of the U.S. west to eastern Texas and is also found in coastal areas of Eurasia [24–26]. Because of its economic values and the tolerance to saline soils, this species has been introduced in China and recommended as a potential cash crop for alternative saline agriculture [10, 27]. Cloning some crucial salt stress response genes from such halophyte sources and investigating their characterizations and functions should be valuable for further understanding the molecular mechanism of plant salt tolerance and also helpful for breeding salt-tolerant crops. So this work aimed to isolate a new plasma membrane Na^+/H^+ antiporter gene from *Kosteletzkya virginica* and investigate its characterizations, which might not only help to understand the salt tolerance mechanism but also provide valuable genes related to salt tolerance for molecular breeding of salt-tolerant crops.

2. Materials and Methods

2.1. Plant Materials and Growth Conditions. The seeds of *Kosteletzkya virginica* were collected from Yellow River Delta, Shandong Province, China. The seeds were soaked in concentrated sulfuric acid for 20 min to remove the hard shell and then thoroughly rinsed with deionized water. Subsequently, the processed seeds were sown in plastic flowerpots (with drain holes in bottom) containing washed sand and grown in the artificial climatic chambers (Huier, China), which was controlled under 28/25°C (day/night) with a daily photoperiod of 14 h and relative air humidity of 65%. Seedlings were sufficiently watered with 1/2 Hoagland nutrient solution every 3 days. Salt treatments were conducted by adding

NaCl to 1/2 Hoagland nutrient solution. For the isolation of Na^+/H^+ antiporter gene, 3-week-old seedlings were treated by 200 mM NaCl for 24 h, and their roots were carefully removed from sands, washed with deionized water, and then harvested. The samples were rapidly frozen in liquid nitrogen and stored at -80°C for the next experiments.

2.2. Cloning of *KvSOS1* cDNA by RT-PCR and RACE. Total RNA was extracted from the above mentioned roots using RNAiso Plus (TaKaRa, Japan) according to the manufacturer's instruction. Quality and quantity of total RNA were measured by using a NanoDrop-2000c spectrophotometer (Thermo Fisher Scientific, USA). The first-strand cDNA was synthesized according to the instruction of TransScript All-in-One First-Strand cDNA Synthesis SuperMix for PCR (Transgen, China). Based on sequence alignments of the conserved regions of reported SOS1 genes from various plant sources (*Theobroma cacao*, EOY01238.1; *Populus trichocarpa*, XP_002315837.2; *Ricinus communis*, XP_002521897.1; *Bruguiera gymnorhiza*, ADK91080.1), a set of degenerate primers (DP-F, DP-R; sequences given in Table 1) were designed and used for the amplification of core fragment of SOS1 from *Kosteletzkya virginica*. The first strand cDNA was used as the template for PCR amplification under the following conditions: 95°C for 5 min, 35 cycles of 95°C for 40 s, 56°C for 30 s, 72°C for 2 min, and 72°C for 10 min. The amplified fragment was ligated into the pGEM-T easy vector (Promega, USA) and sequenced. After the fragment was confirmed to be part of *KvSOS1* gene by NCBI blast, the 5' and 3' ends of the full-length cDNA were further amplified according to the instruction of SMART RACE cDNA Amplification Kit (Clontech, USA). Gene specific primers and nested primers were designed according to the core cDNA sequence. They are as follows: 5'-GSP, 5'-NGSP, 3'-GSP, and 3'-NGSP, as shown in Table 1. The nested PCR was performed in 5' and 3' RACE. The PCR products were separated by 1% agarose gel electrophoresis. DNA from the target band was excised from the gels and purified, then ligated into the pGEM-T easy vector (Promega, USA) and sequenced. Finally, the above obtained sequences were spliced and assembled into the full-length cDNA, which was designated as *KvSOS1*.

3. Results and Discussion

3.1. Cloning and Characterization of the *KvSOS1* cDNA. The full-length cDNA of *KvSOS1* (GenBank accession: KJ577576) was obtained by RT-PCR and RACE methods (specified in Materials and Methods). As shown in Supplementary Figure 1 in Supplementary Material available online at <http://dx.doi.org/10.1155/2014/141675>, the full-length cDNA is 3850 bp, consisting of 5'-untranslated region of 93 bp, an uninterrupted open reading frame (ORF) of 3444 bp, and 3'-untranslated region of 313 bp. The predicted ORF of *KvSOS1* encodes a protein of 1147 amino acids with a molecular weight of 127.56 kDa and a theoretical pI of 6.18.

3.2. Bioinformatics Analysis of *KvSOS1*. Conserved domain analysis using CDD of NCBI revealed that the putative

TABLE 1: The primers used in this study.

Primer	Sequence (5'-3')
DP-F	5'-GG(A/G)GAATCCTT(A/G)ATGAA(C/T)GATGGGAC-3'
DP-R	5'-C(T/C)A(G/A/T)AGC(G/A)CTTTCCTGCCA(C/T)AG-3'
5'-GSP	5'-GCTATCCCCAAAAGCAATTCCAACCGC-3'
5'-NGSP	5'-CCAAGTGAGACTTTGGCCAGAAATT-3'
3'-GSP	5'-GTGCATCCAACCTTTAGTCATGGGAG-3'
3'-NGSP	5'-GTGACAGAATACTTTCAGTACTAAGGTC-3'

FIGURE 1: Analysis of conserved domains in KvSOS1.

protein belongs to the sodium/hydrogen exchanger family (Figure 1), which generally contains 10–12 transmembrane regions at the amino-terminus and a large cytoplasmic region at the carboxyl terminus. Hydropathy plot analysis using the TMpred program further indicated that the obtained KvSOS1 encodes a predicted transmembrane protein. The N-terminal region includes 12 predicted transmembrane domains, while its C-terminal region has a long hydrophilic tail in cytoplasm (Figure 2). This is consistent with previously reported for plant plasma membrane NHAs [28–30].

Multiple sequence alignments demonstrated that the deduced amino acid sequence of KvSOS1 is 82%, 74%, 73%, 73%, and 72% identical to those homologues from *Theobroma cacao*, *Populus trichocarpa*, *Citrus sinensis*, *Ricinus communis*, and *Populus euphratica*, respectively, which are all plasma-type Na⁺/H⁺ exchangers. The highest degree of sequence similarity, especially, locates in the transmembrane regions, where it reaches almost 88% between KvSOS1 and TcSOS1 (Figure 3).

Phylogenetic analysis of some Na⁺/H⁺ antiporters from various plants showed that KvSOS1 formed a cluster with other plant plasma membrane SOS1 homologues and is most closely related to the *Theobroma cacao* homologue (GenBank accession EOY01238.1), which was different from the cluster of plant vacuolar NHX1 homologues (Figures 4 and 5). All these results implied that the obtained KvSOS1 is a plasma membrane type Na⁺/H⁺ antiporter gene.

4. Conclusion

Kosteletzkya virginica has been proved to be a promising halophyte and has been introduced in China and recommended as a potential cash crop for alternative saline agriculture. In addition, some crucial salt stress response genes can be cloned from it for molecular breeding of salt-tolerant crops. In our

FIGURE 2: Plot of hydrophobic and hydrophilic areas in KvSOS1.

study, the full-length cDNA of KvSOS1 was isolated from *Kosteletzkya virginica*. Bioinformatic analysis predicted that it encodes a putative plasma membrane Na⁺/H⁺ antiporter, which has the typical characteristics of other homologous genes. In order to investigate its characterization and function in salt tolerance, the further study would focus on the expression pattern and genetic transformation of KvSOS1, which might provide guidance for molecular breeding of salt-tolerant crops.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

FIGURE 3: Alignment of KvSOS1 with other SOS1 homologues derived from *Theobroma cacao* (TcSOS1, EOY01238.1), *Populus trichocarpa* (PtSOS1, XP_002315837.2), *Ricinus communis* (RcSOS1, XP_002521897.1), *Bruguiera gymnorrhiza* (BgSOS1, ADK91080.1), *Cucumis sativus* (CsSOS1, XP_004150155.1), *Vitis vinifera* (VvSOS1, NP_001268140.1), and *Zygodphyllum xanthoxylum* (ZxSOS1, ACZ57357.1). Identical peptides highlighted in black.

FIGURE 4: Phylogeny of KvSOS1 and other Na^+/H^+ antiporter proteins derived from *Populus trichocarpa* (PtSOS1, XP_002315837.2), *Populus euphratica* (PeSOS1, ABF60872.1), *Ricinus communis* (RcSOS1, XP_002521897.1), *Theobroma cacao* (TcSOS1, EOY01238.1), *Citrus sinensis* (CsSOS1, XP_006492282.1), *Glycine max* (GmSOS1, AFD64746.1), *Lolium perenne* (LpSOS1, AAY42598.1), *Solanum lycopersicum* (SISOS1, NP_001234698.1), *Limonium gmelinii* (LgSOS1, ACF05808.1), *Arabidopsis thaliana* (AtSOS1, AF256224.1), *Theobroma cacao* (TcNHX1, XP_007030791.1), *Populus euphratica* (PeNHX1, ACZ05630.1), *Solanum torvum* (StNHX1, AEN04067.1), *Triticum aestivum* (TaNHX1, AAS17949.1), *Arabidopsis thaliana* (AtNHX1, NP_198067.1), *Halostachys caspica* (HcNHX1, ADK62565.1), *Zoysia japonica* (ZjNHX1, ABY19311.2), and *Aeluropus littoralis* (AINHX1, AAV80466.1).

1 M E E L K E N L Y V L P S R I L Q E I S S A S S E P V D A V
 1 ATGAGGAACTGAAGGAGAATCTGTACGTCTTACCCTCGCGAATTTTGAAGAGATTAGCTCTGCTTCATCGGAGCCCGTTGATGCGGTT
 31 I F V G I S L A L G I A C R H I F R G T K V P Y T V A L L I
 91 ATCTTCGTTGGGATTCTCTGGCATTAGGAATTGCTTGCCGACATATTTTCCGCGGTACCAAAGTTCCTTACACTGTCGCTTTGCTTATC
 61 I G I G L G S I E Y G T S H K L G R I G D G I R I W N N I D
 181 ATTGGCATCGGTCTCGGCTCTATTGAATATGGTACAAGTCATAAATTAGGAAGGATTGGAGATGGTATTGCTATTGGAACAACATTGAT
 91 P D L L L A V F L P A L L F E S A F S M E V H Q I K R C M A
 271 CCTGACCTTCTATTAGCTGTTTCTACCTGCTTTACTTTTTGAGAGTGCATTTTCTATGGAAGTGCACCAGATAAAGAGGTGTATGGCA
 121 Q M V L L A G P G V L I S T F C L G S A L K L T F P Y K W D
 361 CAAATGGTTCTACTTGCTGGTCTGGAGTTCTTATTTGACCTTCTGCCTTGGATCTGCTCTGAAGCTCACTTTTCCATACAAGTGGGAC
 151 W K T S L L L G G L L S A T D P V A V V A L L K D L G A S K
 451 TGGAAACATCGCTGTTGCTTGGGGGACTTCTTAGTGCCACTGATCCTGTTGCTGTTGTGGCATTATTGAAGGATCTTGGTGCTAGCAAA
 181 K L S T I I E G E S L M N D G T A I V V Y Q L F Y K M V L G
 541 AAAGTGAAGCAGCATAATTGAAGCGAATCCTTGATGAATGATGGGACAGCAATTGTGGTCTATCAGTTATTCTATAAAATGGTACTTGGAA
 211 K S F S W D A I I E F L A K V S L G A V A V G I A F G I A S
 631 AAGAGCTTTAGTTGGGATGCCATTATTGAATTTCTGGCCAAAGTCTCACTTGGAGCCGTGGCGGTTGGAATTGCTTTTGGGATAGCATCA
 241 V L W L G F I F N D T V I E I T L T L A V S Y I V Y F T A Q
 721 GTTTTGTGGCTCGGATTTATTTTCAATGATACAGTGATTGAGATTACATTGACACTTGCTGTGAGCTACATTGTTTACTTCACTGCTCAA
 271 E G I E V S G V L A V M T L G M F Y A A V A K T A F K G D S
 811 GAAGTATTGAAGTTTCCGGTGTGTTGGCAGTGATGACATTAGGAATGTTTATGCGGCTGTTGCAAAGACAGCCTTAAGGGTGATAGC
 301 Q Q S L H H F W E M V A Y I A N T L I F I L S G V V I A E G
 901 CAGCAGAGCTTGACCACTTTTGGGAAATGGTTGCCTATATTGCAAAATACATTAATTTTTCATCCTGAGTGGGGTGTATAGCTGAGGGC
 331 I L G D D K I F H N N E H S W G Y L I L L Y I F I Q V S R C
 991 ATTCTTGCGCATGATAAGATATTTTCAACAATGAACATTCTTGGGCTATCTGATTCTTTGTACATCTTTATCCAAGTATCACGCTGC
 361 I V V G A L Y P F L R Y L G Y G L D L K E A V I L I W S G L
 1081 ATTGTTGTTGGAGCATTATATCCATTTTACGATATCTTGATATGGTTTGGATTAAAGGAAGCCGTCATCCTAATATGGTCAGGGCTT
 391 R G A V A L S L S L S V K R S SD G S Q Y I S S E T G S L F
 1171 CGAGGGGCTGTTGCATTATCACTTTCTCTATCTGTAAAGCGTCCAGTGACGGCTCACAATATATCAGTTCTGAAACAGGAAGCCTGTTT
 421 I F F T G G I V F L T L I V N G S T T Q F V L H F L D M D K
 1261 ATTTTCTTCACTGGTGAATTGTATTCTTGACACTTATTGTGAACGGATCAACTACACAGTTTCGTTTACATTTTCTGGATATGGATAAA
 451 L S A A K K R I L D Y T K Y E M L N K A F E A F E D L G D D
 1351 CTATCAGCAGCCAAAGAGCGTATTCTGGAATACACAAAGTATGAAATGTTGAACAAAGCATTGAGGCTTTTGAAGACCTTGGAGATGAT
 481 E E L G P A D W P T V K R Y I A S L N D L E G D P V H P H T
 1441 GAGGAACTTGGACCTGCTGATTGGCCACGGTAAAGAGATACATTGCAAGCTTAAACGATTGGAGGGGACCTGTGCATCCTCACACT
 511 E S E A D N N L D P S N L K D I R V R L L N G V Q S S Y W G
 1531 GAATCTGAAGCTGATAAATCTGGACCCCTCAAATTTGAAAGATATACGAGTACGGCTTTTAAATGGTGTCCAGTCATCACTGAGGGA
 541 M L D E G R I S Q T T A N L L M Q S V D E A I DV A S H E P
 1621 ATGCTTGATGAAGGAGAATTTCAACAACTACAGCAAATCTATTGATGCAATCTGTAGATGAAGCTATTGATGTGGCATCTCATGAACCT
 571 L C D W K G L K S N V H F P N Y Y K F L Q S S M F P Q K L I
 1711 TTATGTGATTGGAAGGGCTTAAATCTAATGTTTATTTCCCAAATTATTACAAGTTTCTTCACTCAAGTATGTTCCCTCAAAAAGTATTGAT
 601 T Y F T V E R L E N A C C V C A A F L R A H R I A R R Q L H
 1801 ACGTATTTCACTGTGGAAGGCTGGAAAATGCATGCTGTGTTTGTGCTGCATTTCTTCGAGCTCATAGAATTGCACGACGGCAGCTTCAT
 631 E F I G D S V V A S T V I S E S E A E G E E A R K F L E D V
 1891 GAGTTTATAGGTGACAGTGTGTTGCTTCTACTGTAATTTCTGAAAGTGAGGCTGAAGGAGAAGAGGCAAGGAAGTTTGGGAAGATGTC
 661 R I T F P Q V L R V V K T R Q V T Y S V L N H L I E Y L Q N
 1981 CGTATAACTTTTCCGAGGTTTTCGCTGTTGTTAAGACAAGACAAGTTACCTACTCGGTATTGAACCATCTGATTGAGTATTTACAAAAC
 691 L E K V G L L E E K E M L H L H D A V Q T D L K K L L R N P
 2071 CTCGAGAAGGTTGGGTTACTGGAAGAAAAGAAATGCTTCATCTTCATGATGCTGCTCCAGACTGACTGAAGAAGCTTTTAAAGGAATCCT
 721 P L V K I P K I N D L I S A H P L L G A L P S T I C E K L L
 2161 CCCTTGGTAAAGATTCCGAAGATAAATGATCTAATAAGTGCCCATCCTTTGCTAGGCGCCCTTCTTCTACTATCTGCGAGAAACTATTA

(a)

FIGURE 5: Continued.

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751 G Y T K E K M K T R G M T L Y K E G S K S N G I W L V S N G
2251 GGTATACAAAAGAAAAATGAAACTCGTGGTATGACACTTTACAAAGAGGGCTCTAAATCAAATGGTATTGGCTAGTTTCAAACGGT
781 V V K W T S R S I R N K H S V H P T F S H G S T L G L Y E V
2341 GTTGTCAGTGGACGAGTAGGAGCATAAGAAACAAGCATTCACTGTCATCCAACCTTTAGTCATGGGAGTACGTTGGGCTGTACGAAGTA
811 L V G K P Y I C D M V T D S V V L C I F I E S D R I L S V L
2431 TTGGTTGGAACCATACATCTGCGACATGGTCACAGATTCCGTGGTTCTGTATTTTATTGAGAGTGACAGAATACTTTCAGTACTA
841 R S D P D I E D F L W R E S A L V L A K L L V P Q I F E K M
2521 AGGTCGGATCCTGACATAGAAGATTTCTCTGGCGGAAAGTGCTCTTGTCTCGCCAACTCTTGGTTCCTCAAATATTTGAGAAAATG
871 A L H D L R A L V A E R S S M K T Y I A G E T I E V S H Q L
2611 GCACTGCACGATTAAAGAGCTCTGTAGCAGAAAGTCGTCGATGAAGACATACATTGCAGGGGAAACAATAGAAGTGTCACACCAATTG
901 I G F L L E G F A K P L L A Q E E L I T S P A V L L P S Q G
2701 ATTGGCTTCTTGTGGAAGGGTTCGCGAAACCTTTACTTGTCAAGAAGAACTCATCACATACCAGCAGTTCTTTTGCCTTCACAAGGG
931 N Q S F L Y A D K S G S A T T S F S H Q R S G Y Q L E T R G
2791 AATCAAAGTTTCTTATATGCAGATAAATCAGGTTCTGCAACAACAGCTTTTCTCATCAGCGATCTGGGTATCAACTTGAGACAAGAGGA
961 S I I Y Q V E T R A R A I I F D I A T L E A N R V L R R N S
2881 AGCATAATATATCAAGTTGAGACTAGAGCAAGAGCAATTATTTTGATATTGCAACACTTGAAGCCAATAGAGTTTTCGGGAGAAATTC
991 S S F T H S H K S L I R E H G G L M S W P E N F F S G R Q H
2971 TCCTCATTTACCCATTACACAAAAGTTTAATTAGAGAACATGGGGGTCTTATGAGTTGGCCTGAAAACCTTCTCAGCGGAAGACAACAT
1021 T Q N H E E S D Q Q V N S L S A R A M Q L S I F G S T V D L
3061 ACACAAAATCATGAAGAAAGTGATCAACAAGTAAACAGCTTATCTGCAAGAGCCATGCAGTTAAGCATCTTTGGCAGCACGGTTGATTG
1051 P R R S R S L S R M H Q S K P A Y N R S Y D R I L S F P G H
3151 CCACGGCGCAGCCGAGTTTATCAAGGATGCATCAATCTAAACCAGCATACAACCGGTCATATGATAGAATTCTTTCATTCCCTGGACAT
1081 P L V S G R S E G S V T M R K N L E E G R K I T R P L P P A
3241 CCACTGGTTTCTGGCAGATCAGAAGGATCTGTTACAATGAGGAAGAATCTCGAAGAAGGGAGAAAGATAACAAGACCGTTACCCCCGCA
1111 Q A K N T D S K E G H G N D E S D E D E I L V R I D S P S G
3331 CAAGCGAAGAACACGGACTCGAAAGAGGGCCACGGAAATGATGAAAGTGATGAAGATGAAATCCTGGTGAGGATCGATTGCGCTAGCGGA
1141 L S F N Q A S *
3421 CTATCATTTCAACCAAGCTTCTTGA

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(b)

FIGURE 5: The ORF nucleotide sequence and the deduced peptide sequence of *KvSOS1*. Start codon and termination codon are highlighted in red.

Authors' Contribution

Hongyan Wang, Xiaoli Tang and Chuyang Shao should be considered co-first authors.

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