The protein kinase Cmk2 negatively regulates the calcium/calcineurin signalling pathway and expression of calcium pump genes PMR1 and PMC1 in budding yeast

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Abstract
Through a genome-wide screen we have identified calcium-tolerant deletion mutants for five genes in the budding yeast Saccharomyces cerevisiae. In addition to CNB1 and RCN1 that are known to play a role in the calcium signalling pathway, the protein kinase gene CMK2, the sphingolipid homeostasis-related gene ORM2 and the gene SIF2 encoding the WD40 repeat-containing subunit of Set3C histone deacetylase complex are involved in the calcium sensitivity of yeast cells to extracellular calcium. Cmk2 and the transcription factor Crz1 have opposite functions in the response of yeast cells to calcium stress. Deletion of CMK2 elevates the level of calcium/calcineurin signalling and increases the expression level of PMR1 and PMC1, which is dependent on Crz1. Effects of Cmk2 on calcium sensitivity and calcium/calcineurin signalling are dependent on its kinase activity. Therefore, Cmk2 is a negative feedback controller of the calcium/calcineurin signalling pathway. Furthermore, the cmk2 crz1 double deletion mutant is more resistant than the crz1 deletion mutant, suggesting that Cmk2 has an additional Crz1-independent role in promoting calcium tolerance.

Keywords: Saccharomyces cerevisiae, Cmk2, Pmr1, Calcineurin, Crz1, Calcium signalling

Background
Calcium ions regulate many cellular processes in both prokaryotes and eukaryotes [1–5]. Regulation of intracellular calcium homeostasis and the calcium/calcineurin signalling pathway are highly conserved in eukaryotic cells [6, 7]. In the budding yeast Saccharomyces cerevisiae, cytosolic calcium homeostasis is regulated by Ca2+ transporters and sequestrators in the plasma and organellar membranes. Transient increases in cytosolic Ca2+ activate the calcium/calcineurin signalling pathway. Sustained Ca2+ accumulation in the cytosol is prevented by a Ca2+ sequestration system composed of the calcium pump Pmc1 and the Ca2+/H+ exchanger Vcx1 in the vacuolar membrane as well as the calcium pump Pmr1 and the Ca2+/H+ exchanger Gdt1 in the ER/Golgi secretory pathway [8–10]. In our recent studies, we have demonstrated that Rch1 is a novel negative regulator of calcium uptake in the plasma membrane in the budding yeast and the human yeast pathogen Candida albicans [11–15].

As a eukaryotic model organism, S. cerevisiae has been proven to be a valuable genetic tool to elucidate mechanisms regulating calcium homeostasis and calcium signalling in eukaryotic cells as well as to characterize gene-drug and pathway–drug interactions [16–20]. In our previous study, from all nonessential genes in the genome we have identified 120 genes that are involved in the tolerance of the budding yeast to high levels of extracellular calcium [21]. To explore genes involved in the sensitivity of yeast cells to extracellular calcium, we screened the same diploid deletion mutant set for calcium-tolerant gene deletion mutants. In this study, we have identified a total of five genes whose deletion leads to the tolerance of S. cerevisiae cells to high levels of extracellular calcium. Furthermore, we show that the protein kinase Cmk2 is a negative feedback controller of the calcium/calcineurin signalling pathway.
Methods
Strains and reagents
The S. cerevisiae haploid wild type BY4741, the diploid wild type BY4743 strain and their isogenic deletion mutant sets used in this study were purchased from Invitrogen Inc. [21–23] (Table 1) Yeast cells were grown at 30°C in YPD medium (1% yeast extract, 2% peptone, 2% glucose) or SD medium (0.67% yeast nitrogen base without amino acids, 2% glucose, and auxotrophic amino acids as needed). Chemicals were purchased from Sangon Biotech (Shanghai, China), except that cyclosporine A (CsA) and O-nitrophenyl-β-D-galactopyranoside (ONPG) were purchased from Sigma (Beijing, China) and the antibiotic nourseothricin was purchased from Golden Biotechnology Inc. (Missouri, USA).

Genome-wide screen for calcium-tolerant mutations
For primary screens for calcium-tolerant strains, the collection of homozygous diploid deletion mutants of nonessential 4757 genes was replicated onto YPD plates with 0.4 M CaCl2 but not on YPD plates without supplemented CaCl2, surrounding mutants on YPD plates containing 0.4 M CaCl2 but not on YPD plates without supplemented CaCl2, was identified as a potential calcium-tolerant mutant. Primary screen was repeated two times. Tolerant mutants were subjected to a secondary screen by a serial dilution assay method as described [22, 24]. The suppressive effects of cyclosporine A on the phenotypes of calcium-tolerant mutants were examined by supplementing 50 µg/ml CsA into YPD plates with or without CaCl2 [25].

Construction of double-gene deletion mutants
To study the interaction between CMK2 and genes encoding components of the calcium/calcineurin signalling pathway in calcium sensitivity, we created their double-gene deletion mutants in the BY4741 genetic background. First, kanMX4 markers in single-gene deletion mutants for CMK2 and VCX1 (cmk2::kanMX4 and vcx1::kanMX4) were replaced by the natMX4 in the plasmid p4339 to generate cmk2::natMX4 and vcx1::natMX4 strains as described previously [26]. The cmk2::natMX4 cassette was PCR amplified from the genomic DNA of the cmk2::natMX4 strain with a pair of primer CMK2-F (5’-GATATTTGTGCAAGATCGACGGAGC-3’) and CMK2-R (5’-ATGCCATGAA GCTGCTCCTG 3’), which locate 100-bp upstream and downstream, respectively, of the open reading frame (ORF) of the CMK2 gene, and used for replacing the CMK2 ORFs in kanMX4 mutants for target genes to generate their double-gene deletion mutants. The deletion of CMK2 gene in these double-gene deletion mutants was confirmed by PCR amplification with detection primer pair CMK2-DF (5’-GAGGCTTATC TTAGAACC GGAGC-3’) and CMK2-DR (5’-AGAACCGGTT AGGC AACTAC 3’), which flank the CMK2-F and CMK2-R. Similarly, the double-gene deletion mutant between CNB1 and VCX1 was constructed (Table 1).

Galactosidase activity assay
To measure the calcineurin dependent response element (CDRE)-driven β-galactosidase activity in the wild type and the cmk2/cm2 mutant, we integrated the Stul-pre-cut plasmid DNA containing the 4 × CDRE-lacZ reporter into the AUR1 locus of these strains as described [12, 27]. The PMCI-lacZ reporter and PMR1-lacZ reporter were described in our previous study [28]. The β-galactosidase activity was determined using the substrate ONPG as described [29–31]. Data are mean ± SD from six independent experiments.

DNA manipulation
To clone the full-length gene CMK2 into the centromeric vector pHAC111 [26], a DNA fragment containing the 776-bp promoter, the 1344-bp open reading frame (ORF) and the 341-bp terminator region of CMK2 was amplified with a pair of primers ScCMK2-clonF (5’-gtagttgttg ctgctgca gGACAAGCT AATAGCGCAA CGC-3’) and ScCMK2-clonR (5’-aagacgca gcagtgaat cTGACACTGA CGTTAGCGAT GACT-3’).
sequence from pHAC111), and cloned between PstI and EcoRI sites in pHAC111 through homologous recombination, which yielded pHAC111-CMK2.

Next, we amplified the DNA fragment CMK2-UP containing the 842-bp promoter and the 225-bp CMK2 ORF region upstream of the AAG codon (lysine) with primers ScCMK2-mutant-F1 (5′ CCTTTTCTAGA GCGT TTTCTG TG 3′) and ScCMK2-mutant-R1 (5′ CAAT AAGATG GCTATAGCAA CATCTTCATT TGTTG 3′; underlined is the mutated codon of AAG). Similarly, the DNA fragment CMK2-DOWN containing the 1115-bp CMK2 ORF region downstream of the AAG codon and the 414-bp terminator was PCR amplified with primers ScCMK2-mutant-F2 (5′ GAAGATGTTG CTATAGC- CAT CTATGGAG AAGGCAATTGC 3′; underlined is the mutated codon of AAG) and ScCMK2-mutant-R2 (5′ CTCAGAGAAG AACCACGGTG 3′). The two fragments CMK2-UP and CMK2-DOWN were fused together by PCR with primers ScCMK2-clonF and ScCMK2-clonR, and the fused product was cloned between PstI and EcoRI sites in pHAC111 through homologous recombination, yielding pHAC111-CMK2M, expressing a point (K76A) mutant form of Cmk2 under homologous recombination, yielding pHAC111-CMK2M, expressing a point (K76A) mutant form of Cmk2 under its own promoter. All constructs were confirmed by DNA sequencing.

**Statistical analysis**
Data are presented as means ± SD. Significant differences were analyzed by GraphPad Prism version 4.00 (USA). P values of <0.05 were considered to be significant.

**Results**

**Screen for calcium-tolerant gene deletion mutants**
In our previous study [21], through a genome-wide approach we have identified 120 genes that are involved in the tolerance of *S. cerevisiae* cells to high levels of extracellular calcium. To explore genes involved in the sensitivity of yeast cells to extracellular calcium, we screened the same deletion library for calcium-tolerant gene deletion mutants. As a result, we revealed that deletion of *CNB1, RCNI, CMK2, ORM2* or *SIF2* caused yeast cells to be tolerant to 0.4 M CaCl₂ and these phenotypes were more dramatic in the presence of 0.6 M CaCl₂ (Fig. 1). The calcium tolerant-phenotypes of yeast cells lacking *RCNI*, *ORM2* or *SIF2* on YPD plate containing 0.4 M CaCl₂ could be slightly suppressed by the addition of the cyclosporin A (CsA), the specific inhibitor of calcineurin (Fig. 1).

*RCNI* encodes a protein involved in calcineurin regulation during calcium signalling and has similarity to human DSCR1 that is found in the Down Syndrome candidate region [32]. *CNB1* encodes a regulatory subunit of calcineurin, a Ca²⁺/calmodulin-regulated type 2B protein phosphatase, which dephosphorylates the transcription factor Crz1, and promotes its nuclear localization [9]. Our observation that deletion of *CNB1* leads to a calcium-tolerant phenotype in the wild type BY4743 background is consistent with previous studies [33, 34], showing that inactivation of calcineurin including deletion of *CNB1* restores the calcium-tolerance of yeast cells lacking *PMCI*, since calcineurin decreases calcium tolerance of *pmcl* cells by inhibiting the function of the vacuolar H⁺/Ca²⁺ exchanger Vcx1. Therefore, it is not surprising to note that the difference in calcium sensitivity between the wild type and the *cnb1/cnb1* mutant is abolished by CsA, since it suppresses the function of calcineurin in the wild type (Fig. 1). However, interestingly, yeast cells lacking *CMK2* were sensitive to calcium stress in the presence of CsA as compared to the wild type (Fig. 1). This suggests that the calcium tolerance due to deletion of *CMK2* requires the presence of a functional calcineurin.

**Calcium tolerance of cmk2 cells requires the calcium/calcineurin signalling in response to high levels of extracellular calcium**

As homologs of mammalian Cam Kinase II, Cmk2 and its paralog Cmk1 are calmodulin-dependent protein kinases (CaMK) that play a role in stress response in *S. cerevisiae* [35, 36]. The yeast CaMK, encoded by both *CMK1* and *CMK2*, and the calcineurin act independently in promoting survival of pheromone-induced growth arrest [35]. However, deletion of *CMK1* did not lead to a calcium-tolerant phenotype of the wild type BY4743 cells as deletion of *CMK2*, neither did affect the calcium-tolerant phenotype of yeast cells lacking *CMK2* (Fig. 2a). This indicates only Cmk2, but not Cmk1, is involved in the response of yeast cells to high levels of external calcium. To examine if the calcium-tolerant phenotype of yeast cells lacking *CMK2* is related to the calcium/calcineurin signalling pathway, we constructed double-gene deletion mutants between *CMK2* and genes encoding various components of the calcium/calcineurin signalling pathway.

Although deletion of *CNB1* causes yeast cells to be calcium-tolerant in the haploid BY4741 background as in the diploid BY4743 background, further deletion of *CNB1* led to a calcium-sensitive phenotype for yeast cells lacking *CMK2* (Fig. 2b). In contrast, deletion of *VCX1* did not have an impact on the calcium sensitivity of the wild type BY4741 cells harboring a functional calcineurin, and neither did on the calcium-tolerant phenotype for yeast cells lacking *CMK2* (Fig. 2b and c). Like in yeast cells lacking *CMK2*, further deletion of *CNB1* also led to a calcium-sensitive phenotype for yeast cells lacking *VCX1* (Fig. 2c). As expected [10], deletion of *CRZ1* alone led to a hypersensitivity of
BY4741 cells to calcium stress (Fig. 2b). Yeast cells lacking both CRZ1 and CMK2 showed a hypersensitive phenotype on YPD plate containing 0.4 M CaCl₂ with a similar degree to cells lacking CRZ1 alone. In contrast, yeast cells lacking both CRZ1 and CMK2 showed a sensitive phenotype on YPD plate containing 0.2 M CaCl₂ in a less degree than cells lacking CRZ1 alone (Fig. 2d). However, all these phenotypes could be suppressed by the specific inhibitor of calcineurin, CsA (Fig. 2d). This indicates that Cmk2 and Crz1 have independent and opposite functions in the calcium sensitivity of yeast cells. Taken together, our data suggest that the calcium tolerance due to deletion of CMK2 is dependent on Crz1 under high concentrations of calcium, but is independent of Crz1 under lower concentrations of calcium.
Deletion of CMK2 leads to increased expression of PMR1 and PMC1 due to activation of the calcium/calcineurin signalling

Next, we examined if deletion of CMK2 would activate the calcium/calcineurin signalling. We examined the calcium-dependent response element (CDRE)-lacZ activity in the wild type and the cmk2 mutant. As compared to the wild type, the CDRE-lacZ activity in the cmk2 mutant was significantly increased in the absence or presence of 0.2 M CaCl₂ (Fig. 3a). This suggests that deletion of CMK2 elevates the activation levels of the calcium/calcineurin signaling independent of calcium stress.

Activation of calcium/calcineurin signalling induces the expression of PMR1 and PMC1, but not VCX1 [21, 34, 37]. Next, we examined the expression of PMR1 and PMC1 using their lacZ reporters in the wild type, the crz1 mutant and the cmk2 mutant. The lacZ activities of PMR1 and PMC1 were induced in the wild type in response to 0.2 M CaCl₂ (Fig. 3b and c). This is consistent with the previous observation [34]. In the absence of 0.2 M CaCl₂, there was no significant difference in the lacZ activities of PMR1 or PMC1 between the wild type and the crz1 mutant, but there was a significant increase in the lacZ activities of PMR1 and PMC1 in the cmk2 mutant as compared to the wild type (Fig. 3b and c).

In the presence of 0.2 M CaCl₂, there was a significant decrease in the lacZ activities of both PMR1 or PMC1 in the crz1 mutant as compared to the wild type (Fig. 3b and c). In contrast, there was a significant increase in the lacZ activity of both PMR1 and PMC1 in the cmk2 mutant as compared to the wild type (Fig. 3b and c). Therefore, deletion of CMK2 increase the expression level of both PMR1 and PMC1.

Enhanced CDRE-lacZ activity due to deletion of CMK2 is dependent on Crz1

As we described above, expression of both CDRE-lacZ and PMR1-lacZ reporters was increased in the absence of added calcium in the cmk2 mutant as compared to the wild type (Fig. 3a and b). To examine if this effect is due to enhanced Crz1 activity, we measured the activity of CDRE-lacZ in the crz1, the cmk2 and the crz1/cmk2 mutants. As expected, in the absence or presence of 0.2 M CaCl₂, the activity of CDRE-lacZ was almost abolished in both the crz1 mutant and the crz1/cmk2 mutant, albeit increased in the cmk2 mutant (Fig. 3d). In addition, the activity of CDRE-lacZ was significantly lower in either the crz1 mutant or the crz1/cmk2 mutant than the wild type strain in the absence of added calcium, although no significant difference in the activity of CDRE-lacZ was observed between the crz1 mutant and the crz1/cmk2 mutant in the absence or presence of 0.2 M CaCl₂ (Fig. 3d). Taken together, these data suggest that the enhanced CDRE-lacZ activity in the cmk2 mutant was dependent on Crz1 in the absence or presence of calcium stress and that a basal level of Crz1 activity is present in yeast cells growing in YPD medium.

Effect of Cmk2 on the CDRE-lacZ activity is dependent on its catalytic activity

We examined if the catalytic activity of the protein kinase Cmk2 is responsible for its suppressive effect on the
CDRE-lacZ activity. The critical lysine residue in ScCmk2 was identified from the amino acid alignment between *S. cerevisiae* ScCmk2, *Schizosaccharomyces pombe* SpCmk2 [38] and *S. cerevisiae* ScRck2 [26, 39] (Additional file 1: Figure S1). We first constructed the plasmids pHAC111-CMK2 and pHAC111-CMK2M, expressing the wild type Cmk2 and its kinase-dead point mutant (K76A) variant of Cmk2. Introduction of pHAC111-CMK2, but not pHAC111-CMK2M, partially suppressed the calcium-tolerant phenotype of the cmk2 mutant (Fig. 4a). This suggests that the catalytic activity of Cmk2 is required for the function of Cmk2 in the sensitivity of yeast cells to high levels of extracellular calcium stress.

Next, we introduced the vector pHAC111 to the wild type BY4741 with the CDRE-lacZ reporter and the cmk2 mutant with the CDRE-lacZ reporter, respectively. In addition, pHAC111-CMK2 and pHAC111-CMK2M were introduced into the cmk2 mutant strain with the CDRE-lacZ reporter, respectively. As expected, in the absence or presence of 0.2 M CaCl₂, the cmk2 mutant carrying the pHAC111 vector showed a higher CDRE-lacZ activity than the wild type carrying pHAC111 (Fig. 4b). Introduction of pHAC11-CMK2, but not pHAC11-CMK2M, suppressed the enhanced effect of cmk2 deletion on the CDRE-lacZ activity in the absence or presence of 0.2 M CaCl₂ (Fig. 4b). This indicates that the negative effect of Cmk2 on the CDRE-lacZ activity is dependent on its kinase activity.

![Fig. 4](image-url)

Fig. 4 Effects of Cmk2 on both calcium sensitivity and CDRE-lacZ activity are dependent on its catalytic activity. (a), cells of the wild-type BY4741 and the cmk2 mutant carrying the pHAC111 vector, pHAC111-CMK2 or pHAC111-CMK2M were grown at 30 °C in liquid SD-LEU medium overnight, serially diluted by 10 times and spotted on YPD plates with or without 0.6 M CaCl₂, respectively. Plates were incubated for 2 to 3 days at 30 °C. (b), Galactosidase activities of 4xCDRE-lacZ in the wild-type (WT) BY4741 and the cmk2 mutant, carrying the pHAC111 vector, pHAC111-CMK2 or pHAC111-CMK2M as indicated. They were grown at 30 °C in liquid SD-LEU medium overnight before they were inoculated to, and grown in, YPD medium to log phase. Cells were further grown in the absence or presence of 0.2 M CaCl₂ for 2 h before they were collected for protein extraction and lacZ activity assay. Symbols # and * show statistically significant differences (P < 0.05) between the wild type and each of other strains in the absence or presence of 0.2 M CaCl₂, respectively. Values were means of six independent experiments.
Discussion
Through a chemical genetic screen, we have identified five genes, *CNB1, RCN1, ORM2, SIF2* and *CMK2*, that are involved in the sensitivity of *S. cerevisiae* cells to high levels of extracellular calcium. Two genes, both *CNB1* and *RCN1*, encode regulators of calcineurin. Calcineurin not only induces the expression of *PMC1* and *PMR1* through the action of the transcription factor Crz1, but also inhibits in a Crz1-independent fashion the activity of Vcx1, the high capacity, low affinity vacuolar Ca\(^{2+}\) exchanger [37, 40]. The calcium-tolerance of yeast cells lacking *CNB1* is likely due to the release of the inhibitory effect of calcineurin on Vcx1 activity, which is supported by our observation that further deletion of *VCX1* reversed the phenotype of cells lacking *CNB1* to be hypersensitive to calcium stress with a similar degree to that of cells lacking *VCX1* in the presence of CsA (Fig. 2c).

*ORM2* encodes a regulator of the sphingolipid homeostasis. Our finding is consistent with previous observations that sphingolipids play roles in calcium homeostasis, protein trafficking/exocytosis, longevity and cellular aging, nutrient uptake, and the interaction of sphingolipids and antifungal drugs in *S. cerevisiae* [41]. In mammalian cells, sphingosine 1-phosphate is a signalling lipid known to be involved in calcium homeostasis [42–44]. *SIF2* encodes the WD40 repeat-containing subunit of Set3C histone deacetylase (HDAC) complex [45]. The Set3C binds histone H3 dimethylated at lysine 4 to mediate deacetylation of histones in 5′-transcribed regions [46]. Interestingly, the chaperone mammalian relative of DnaJ (Mrj) decreases the occupancy of nuclear factor of activated T cells (NFAT), the mammalian functional counterpart of yeast Crz1, on the tumor necrosis factor-alpha promoter in cardiomyocytes in an HDAC-dependent manner [47]. A similar role might exist for Set3C on the regulation of Crz1 in gene transcription in response to calcium stress.

*CMK2* encodes a calmodulin-dependent protein kinase, and expression of *CMK2* is induced by the calcium/calcineurin signalling pathway [48, 49]. Yeast cells lacking *CMK2* are tolerant to high levels of external calcium, while cells lacking the *CRZ1* are calcium-sensitive. Therefore, Cmk2 and Crz1 have opposite functions in the calcium sensitivity of yeast cells. In addition, we show here that deletion of *CMK2* elevates the calcium/calcineurin signalling and increases the expression level of *PMR1* and *PMC1* in response to calcium stress (Fig. 3). The effects of Cmk2 on calcium sensitivity and calcium/calcineurin signalling are dependent on its catalytic activity (Fig. 4). Taken together, these data suggest

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Fig. 5 Mode of Cmk2 function in the feedback regulation of calcium/calcineurin signaling pathway. In response to high levels of extracellular calcium, Ca\(^{2+}\) enters the cell through unknown pathways, and both calcineurin and Crz1 are activated to induce expression of target genes including *CMK2, RCN1, PMC1, PMR1, RCN2, CRZ1*, etc (Induced Expression). Arrows indicate activation, and T bars represent inhibition. A dashed line means that more data is needed to confirm.
that Cmk2 functions as a negative feedback controller for the calcium/calcineurin signalling pathway, which is mediated by Crz1 through calcineurin in response to high levels of extracellular calcium stress (Fig. 5). How Cmk2 regulates the calcium signalling pathway remains to be determined. A plausible possibility would be that Cmk2 directly phosphorylates and negatively regulates calcineurin or Crz1 (Fig. 5). In contrast, Rcn1 is a feedback controller of calcineurin with dual effects [48, 50]. Another possibility is that Cmk2 indirectly affects Set3C, which in turn regulates the activity of Crz1.

Interestingly, we have observed that the cmk2 crz1 double deletion mutant is more tolerant to 0.2 M CaCl₂, albeit not to a higher level of calcium (0.4 M CaCl₂), than the crz1 deletion mutant (Fig. 2d). This indicates that Cmk2 has an additional function in calcium tolerance, which is independent of Crz1. However, supplementation of CsA in the medium suppresses the calcium sensitivity of both the crz1 and cmk2/crz1 mutants at both 0.2 M CaCl₂ and 0.4 M CaCl₂ (Fig. 2d). This indicates that the additional Crz1-independent function of Cmk2 is dependent on calcineurin function (Fig. 5). This additional function could be mediated by the Vcx1, whose activity can be inhibited by calcineurin [34]. Nevertheless, the direct phosphorylation and thereby regulation of Vcx1 activity by Cmk2 could not be excluded. This might also explain what we have observed that higher promoter activities of CDRE-lacZ and PMR1-lacZ in the cmk2 mutant than those in the wild type strain under normal growth conditions (Fig. 3), when the calcium/calcineurin signaling pathway is not activated.

Additional file

Additional file 1: Figure S1. Amino acid sequence comparison between ScCMK2, SpCMK2 and ScRCK2. (PDF 88 kb)

Abbreviations
CaMK: Calmodulin-dependent protein kinase; CDRE: Calcineurin dependent response element; CsA: Cyclosporine A; ER: Endoplasmic reticulum; NFAT: Nuclear factor of activated T cells; PCR: Polymerase chain reaction; YPD: Yeast peptone dextron

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Authors’ contributions
HX, TF and HY carried out experiments and analyzed the data, LJ designed the study and wrote the manuscript and analyzed the data. All authors read and approved the final manuscript.

Authors’ information
N/A.

Ethics approval and consent to participate
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Competing interests
The authors declare that they have no competing interest.

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