AQP5-1364A/C polymorphism and the AQP5 expression influence sepsis survival and immune cell migration: a prospective laboratory and patient study

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Abstract

Background: The C-allele of the aquaporin (AQP5) -1364A/C polymorphism is associated with decreased AQP5 expression but increased 30-day survival in patients with severe sepsis. AQP5 expression might affect survival via an impact on cell migration. Consequently, we tested the hypothesis that (1) Aqp5 knockout (KO) compared to wild type (WT) mice show an increased survival following lipopolysaccharide (LPS) administration, and that (2) AQP5 expression and the AQP5 -1364A/C polymorphism alters immune cell migration.

Methods: We investigated Aqp5-KO and wild type mice after intraperitoneal injection of either E.coli lipopolysaccharide (LPS, serotype O127:B8, 20 mg/kg) or saline. Furthermore, neutrophils of volunteers with the AA-AQP5 or AC/CC-AQP5 genotype were incubated with 10^-8 M Chemotactic peptide (fMLP) and their migration was assessed by a filter migration assay. Additionally, AQP5 expression after fMLP incubation was analyzed by RT-PCR and Western blot. Moreover, migration of AQP5 overexpressing Jurkat cells was studied after SDF-1α-stimulation. We used exact Wilcoxon–Mann–Whitney tests; exact Wilcoxon signed-rank tests and the Kaplan–Meier estimator for statistical analysis.

Results: Fifty-six percent of Aqp5-KO but only 22% of WT mice survived following LPS-injection. WT mice showed increased neutrophil migration into peritoneum and lung compared to Aqp5-KO mice. Target-oriented migration of neutrophils was seen after 0.5 h in AA-genotype cells but only after 1.5 h in AC/CC-genotype cells, with a threefold lower migrating cell count. AQP5 overexpressing Jurkat cells showed a 2.4 times stronger migration compared to native Jurkat cells.

Conclusion: The AQP5 genotype may influence survival following LPS by altering neutrophil cell migration.

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Background

Although wide variability exists regarding the outcome from severe sepsis, some of this variability may be caused by genetic variability, as shown in twin and adoption studies [1–4]. A potential candidate gene for investigation is the gene encoding aquaporin (AQP) 5 [5], which mediates key mechanisms of inflammation that prevail in sepsis, including cell migration and proliferation [6], activity of the renin–angiotensin–aldosterone system (RAAS) [7], and the transport of water across biological membranes [8]. Previously, we described a functional, and common (70% AA, 23% AC, 7% CC) single nucleotide (−1364A/C, rs3759129) polymorphism in the AQP5 gene promoter [7]. Substitution of C for A at position—1364 was associated with a decreased AQP5 messenger RNA and AQP5 protein expression.

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expression. It was also a strong, independent prognostic factor for decreased 30-day mortality in patients with severe sepsis [9]. The estimated hazard ratio of nearly 4 for the AC/CC-genotypes compared with the homozygous AA genotype not only suggests that the C-allele of the AQP5-1364A/C polymorphism may have important effects on the AQP5 expression in severe sepsis, but also that there is a potential relevance of the AQP5 expression in severe sepsis. However, the underlying molecular and pathophysiological alterations linking a decreased AQP5 expression in C allele carriers to 30-day mortality is unknown.

In the search for these mechanisms, our previous research [7] suggested an impact of a decreased AQP5 expression in C-allele carriers on sepsis survival via its influence on the renin–angiotensin–aldosterone system, which however could not be confirmed [9]. Therefore, our research focused on further possible mechanisms. Since AQP5 expression also mediates cell migration, which is yet another inflammatory key mechanism and involves transient formation of membrane protrusions (lamellipodia and membrane ruffles) at the migrating cell's leading edge [6], we speculated that the AQP5 expression might influence sepsis survival due to its influence on immune cell migration. During cell migration aquaporins facilitate water influx through the cell membrane and cause a local hydrostatic pressure causing cell membrane expansion [6]. This expansion is stabilized by actin polymerization and mediates migration [6]. In addition it was previously demonstrated that Aqp5-KO mice show increased bacterial load after infection but decreased NFκB expression and mucus secretion compared to WT mice [10]. To this end, a decreased AQP5 expression may decrease immune cell migration and may subsequently alter the inflammatory chain during sepsis. The migration of neutrophils is essential in immune response during sepsis [11] and its regulation is critical for sepsis outcome as sufficient neutrophil migration is critical for pathogen clearance but excessive neutrophil migration is associated tissue damage and subsequent organ dysfunction [12]. In addition regulation of AQP5 expression in neutrophils has not been studied yet and was therefore addressed with this study.

Accordingly, we prospectively tested the following hypotheses:

1. The survival rate in Aqp5-KO mice is different than in wild type mice after lipopolysaccharide induced inflammation.
2. The AQP5 expression and the AQP5-1364A/C genotype is associated with impaired cell migration.

### Methods

#### Mice

Experiments were conducted in accordance with German and local governmental regulations, complying with both the European Community and the American Physiological Society and the Guide for the Care and Use of Laboratory Animals (National Institute of Health publication 85-23, revised 1996) guidelines. Aqp5 (−/−) [knockout (KO)] and wild type recombinant inbred from 129SvJ/Black Swiss mice were provided by Prof. Anil Menon, Cincinnati, and generated as previously described [13].

#### Genotyping of mice

Mice were genotyped with mouse DNA extracted from an ear biopsy as described previously [14].

#### Animal preparation

We investigated Aqp5-KO (−/−) and wild type (+/+) mice after an intraperitoneal (i.p.) injection of either *Escherichia coli* lipopolysaccharide (LPS, 20 mg/kg in saline [15], serotype O127:B8, Sigma-Aldrich, Taufkirchen, Germany) or an equivalent amount of pyrogen-free saline. After 4.5 h and 24 h post LPS/saline injection 100 µl retro orbital blood was collected. The serum layer was removed after clotting and centrifugation and stored at −80 °C. IL-10 and TNF-α concentrations were determined using Legend Max Elisa Plates (BioLegend, San Diego, CA, USA). The survival and the state of health (score sheet) of the mice were observed every 6 h. In the score sheet points (zero for no abnormality and 20 for worst case) were given for body weight, general condition, behavior, clinical findings and secretions. Due to these points the disease score was calculated (Additional file 1).

Eight Aqp5-KO (−/−) and eight wild type (+/+) mice were killed 3 h after LPS or saline injection by cervical dislocation to study neutrophil migration. Peritoneal lavage and lungs were harvested and right lungs were snap frozen whereas left lungs were stored in formalin and paraffin-embedded (Paraplast Xtra-plus, Sigma-Aldrich, Munich, Germany).

#### MPO activity assay

Myeloperoxidase (MPO) activity was measured as a marker for neutrophil infiltration. The right lungs were homogenized in buffer containing 10 mM NEM (Sigma-Aldrich, Taufkirchen, Germany) and then solubilized in buffer containing 0,5% HTA-Br. MPO activity was determined by the use of Myeloperoxidase activity assay kit (abcam, Cambridge, UK). In addition protein amount was determined using RotiQuant universal (Roth, Karlsruhe, Germany).
Flow cytometric analysis of peritoneal lavage
For flow cytometry (FACS) cell suspensions were collected by a lavage of the peritoneum with 5 ml PBS containing 5 mM EDTA. For FACS analysis, cells were stained with the appropriate antibodies (APC antimouse Ly6g/Ly6C (Gr-1) clone RB6-8C5, BioLegend, San Diego, CA, USA; Antimouse CD11c FITC, clone N418, eBiolscience, Frankfurt, Germany), after blocking FcRs with anti-CD16/32. Cells were analyzed on a BD FACSCanto II (Heidelberg, Germany). Data were analyzed with Flow-Software 2.

Neutrophils staining in lung tissue with naphthol AS-D chloroacetate (NASDCL)-esterase
Naphthol AS-D chloroacetate (NASDCL)-esterase was used to detect neutrophils on tissue sections. Paraffin left lung sections (1–2 μm thick) were mounted on slides and de-paraffinized. Staining with naphthol AS-D chloroacetate staining solution (containing naphthol-AS-D-chloracetat in dimethyl sulfoxide dimethyl, pararosaniline and sodium nitrite) was performed for 30 min. Finally, the sections were counterstained with Mayer’s hematoxylin.

Patients
The study was reviewed and approved by the Medical Faculty’s Ethics Committee of the Duisburg-Essen University. The samples used in this study have been used for several other studies before [3, 4, 9] thus no additional material had to be collected. Patients with sepsis were considered eligible when they fulfilled the 2001 American College of Chest Physicians/Society of Critical Care Medicine Consensus conference guidelines [16]. Patients with sepsis were enrolled on the day when the sepsis was first diagnosed and arterial and venous blood was sampled. Afterwards, blood tests, genotyping, cytokine determination, microbiology cultures, and RNA extraction were performed. Informed consent for all the patients was obtained from the patient’s guardian.

Genotyping of patients and healthy volunteers
Genotyping was performed as previously described [7].

RNA isolation and cytokine concentration measurements
RNA and cytokine extraction and measurement was performed as described [4].

Immunocytochemistry
Neutrophils were isolated from 30 ml of venous blood obtained from four healthy donors after ethics committee approval (ethics committee of the Essen University Hospital, Essen, Germany) and a written informed consent. Immunostaining was performed with an AQP5 antibody [AQP5 G-19; sc-9890] as described previously [17].

Stable transfection of jurkat T-cells
Stable transfection with Jurkat T-cells was performed using an AQP5_pReceiver EX-T1015-M09 vector (Human Full ORF-Clone, NM-001651, pReceiver-M09) or a pReceiverM09 (GeneCopeia, Rockville, MD, USA). The vector was linearized with Scal (New England Biolabs, Ipswich, MA, USA) and transfected into Jurkat cells by electroporation using the Gene Pulser Xcell Electroporation System (BioRad, Munich, Germany). The cells were maintained in RPMI plus 10% FKS, plus Pen/Strep plus 1.1 mg/ml gentamicin disulfate. Positive clones were selected using limiting dilution [18].

Western blot
For protein extraction, stable transfected Jurkat cells or Chemotactic peptide (N-Formyl-l-methionyl-l-leucyl-l-phenylalanine, fMLP) stimulated HL-60 (10⁻⁸M fMLP for 6 h) promyelocyte cells were lysed with RIPA buffer and proteins were extracted by shaking at 4 °C. Western Blot was performed as described [19].

Real-time PCR
RNA from blood cells of septic patients was extracted as described above and 0.5 μg RNA was used to synthesize cDNA. In addition RNA from fMLP stimulated HL-60 cells (10⁻⁸M for 6 h) was extracted and 1 μg RNA was used for cDNA synthesis. Primers for AQP5 and the housekeeping gene actin were used and the real-time PCR was performed as described [19, 20].

Neutrophils migration assay
Blood samples (30 ml) from 8 AQP5-genotyped healthy volunteers (EDTA-Monovetten Sarstedt, Nümbrecht, Germany) were centrifuged at 500g for 30 min using 30 ml Polymorphprep (Fresenius Kabi, Oslo, Norway). Neutrophils were collected. Their purity was determined using “Scil Vet ABC” cell counter (Scil, Viernheim, Germany) and was above 90%. 5 * 10⁵ cells were deposited in 200 μl RPMI into the upper compartment of a filter migration assay system containing a polycarbonate membrane filter (5 μm pore size, BD, Heidelberg, Germany). The lower compartment contained 10⁻⁸ M fMLP (Sigma-Aldrich, Taufkirchen, Germany) in 500 μl RPMI or control media. The cells were incubated at 37 °C with 5% CO₂ in air for 0.5 and 1.5 h in duplicate for each sample as described before [21]. The migrated cells were counted after 0.5 and 1.5 h using the MUSE Count & Viability Kit (Merck Millipore, Darmstadt, Germany). On the one hand total migrated cells (all cells which migrated to RPMI containing 10⁻⁸ M fMLP) and on the other hand target oriented migrated cells (difference between cells migrated to RPMI containing 10⁻⁸ M fMLP and spontaneously
migrated cells to RPMI alone) were counted and calculated.

For the migration assay with Jurkat T-cells, stable transfected cells were used. Polycarbonate membrane filters were coated with 6.3 μg fibronectin and 5 * 10^5 cells were deposited in 200 μl RPMI into the upper compartment. The lower compartment contained 100 ng/ml SDF-1α (PROSPEC, East Brunswick, NJ, USA) in 500 μl RPMI or control media. Cells were counted using the MUSE Count & Viability Kit after 4.5 h and again after 24 h.

Statistical analysis
Continuous variables are summarized as boxplots. For the comparison of continuous variables of two independent groups, we used exact Wilcoxon–Mann–Whitney tests; for two dependent groups exact Wilcoxon signed-rank tests. The time-to-event survival variable was summarized as the Kaplan–Meier estimator. The reported p-values are two-sided and within each panel adjusted for multiplicity by Bonferroni’s correction. A p value of <0.05 was regarded as statistically significant. All statistical analyses were done using GraphPad Prism 6 (La Jolla, CA, USA) or IBM SPSS Statistics 21.

Results
Survival of Aqp5-KO and wild type mice after intraperitoneal LPS injection
Seven-day survival following LPS injection differed between Aqp5-KO and WT mice. Fifty-six percent (10/18) of the Aqp5-KO mice but only 22% (4/18) of the WT mice survived for seven days (Fig. 1a). Overall, WT mice had a higher risk of death (fully adjusted hazard ratio (HR):2.48, 95% CI 1.04–5.90) compared to KO mice. After 96 h, WT mice showed a significantly increased disease score compared to KO mice (Fig. 1b). To clarify if different cytokine expressions might be responsible for differences in survival in wild type and KO mice, we examined serum concentrations of tumor necrosis factor alpha (TNF-α), as one of the early pro-inflammatory cytokines [22], and of interleukin-10 (IL-10), as an important anti-inflammatory cytokine in sepsis [23]. We did not observe evidence for differences in cytokine concentrations of IL-10 and TNF-α in the blood of the Aqp5-KO and WT mice (Fig. 1c, d).

Migration of neutrophils in mice tissues
First we analyzed total cell count in the peritoneal lavage of Aqp5-KO and WT mice. Aqp5-KO mice had significantly deceased cell count in the lavage compared to WT mice (Fig. 2a). Furthermore we analyzed the percentage of CD11c Ly6G/C positive cells in the peritoneum of Aqp5-Ko and WT mice at 3 h after i. p. LPS administration compared to saline injected control mice with flow cytometry (Fig. 2b). In WT mice percentage of CD11c Ly6G/C positive cells significantly increased after 3 h LPS compared to saline (Fig. 2c), whereas no significant increase was detected in Aqp5-KO mice 3 h after LPS compared to saline injection. Hence, we found a significantly higher amount of CD11c Ly6G/C positive cells in the peritoneum, of WT mice compared to Aqp5-KO mice (Fig. 2d) after LPS administration.

In the lungs of WT mice a higher number of neutrophils were detected compared to Aqp5-KO mice after NASDCL-Esterase staining on fixed lung samples (Fig. 2e). In line with this MPO activity increased in lungs of WT mice after 3 h LPS in comparison with saline injection, but not in Aqp5-KO mice (Fig. 2f). Therefore 3 h after LPS administration WT mice showed significantly higher MPO activity compared to Aqp5-KO mice (Fig. 2f) in lungs.

AQP5-dependent cell migration of a human cell line
We used the Jurkat cell line for several reasons: (1) Jurkat cells can be transfected by routine protocols [28], (2) migration assay for Jurkat cells is well established using the chemoattractant SDF-1α [29], and (3) we could address AQP5’s impact on granulocytes’ (neutrophils) and lymphocytes’ (Jurkat) migration. As shown in Fig. 3a and validated by Western blot, Jurkat cells could be successfully stable transfected with an AQP5 overexpression vector and with a control vector. AQP5 transfected Jurkat cells showed a 2.4-fold increase in total cell migration compared to cells transfected with the control vector (Fig. 3b).

Genotype-dependent AQP5 expression and migration of human neutrophils
Neutrophils from healthy donors with AA genotype showed a considerable amount of AQP5 protein while no clear AQP5 protein expression could be detected in CC genotype carriers (Fig. 3c). Neutrophils of AA-genotype carriers already showed target-oriented migration after 0.5 h, while neutrophils from AC/CC-genotypes did not (Fig. 4a, b). After 1.5 h, neutrophils from AC/CC-genotype carriers also showed target-oriented migration, but the amount of migrating cells was threefold lower than in AA-genotype carriers (Fig. 4c, d). In addition, incubation of the neutrophil cell line HL-60 with the chemotactic peptide fMLP increased the AQP5 mRNA expression (Fig. 5a). In line with that an increase in AQP5 protein expression was also evident by Western blot (Fig. 5b).
To conclude, we examined if AQP5 expression in septic patients differs between A and C-allele carriers as we showed it in neutrophils of healthy donors. The AQP5 mRNA expression was greater in the blood of septic patients carrying the AQP5 AA genotype compared to those carrying the AC genotype (Fig. 5c). In addition, TNF-α concentration in the blood of septic patients was determined depending on AQP5 genotype. Again as in mice (Fig. 1d), there was no evidence for AQP5-genotype dependent differences in TNF-α concentration (Fig. 5d).

**Discussion**

This study suggests that AQP5 might be an important protein in LPS induced systemic inflammation and identifies the influence of AQP5 expression on immune cell migration as a potential mechanism by which the AQP5 expression affects survival in sepsis. Furthermore, we showed that the C-allele, associated with decreased AQP5 expression and increased survival from severe sepsis, is associated with a decreased neutrophil cell migration in vitro. Thus, the link between the AQP5-1364A/C polymorphism and sepsis survival could be due to AQP5 expression and its impact on neutrophil cell migration. Less cell migration can cause less tissue damage caused
by infiltrating neutrophils and pro-inflammatory mediators [12, 24]. Another possible mechanism which was not examined in this study is the potential impact of AQP5 expression on neutrophil maturation. If AQP5 influences maturation the impact on cell migration would only be indirectly but it would still have the same impact on course of sepsis.

Despite great efforts in research and clinical treatment made in the last 30 years, the sepsis mortality remains unabatedly high [25]. Accordingly, new approaches are urgently needed. Our previous association study suggested that AQP5 seems to be a key protein in severe sepsis. Substitution of C for A at position −1364 was associated with decreased AQP5 protein expression and was a strong and independent prognostic factor for decreased 30-day mortality in patients with severe sepsis [9]. Therefore this study is in line with our previous study, which also showed a twofold decreased AQP5 mRNA and protein expression in C-allele carriers compared to A-allele carriers [7]. Our investigation of Aqp5-KO mice in comparison with WT mice now confirms our hypothesis that the AQP5 expression itself has a particularly strong influence on the mortality of sepsis and that this influence is associated with decreased neutrophil migration.

Findings from Vassiliou et al. support our results [19, 26] that aquaporins seem to be important target proteins

![Fig. 2](image-url)
in sepsis, as they identified AQP1 as a gene that is up-regulated in LPS induced inflammation. Importantly, an AQP1 upregulation not only took place in lung tissue [19] but also in leukocytes. Vassiliou et al. showed that the AQP1 expression induced by LPS was associated with increased cell membrane permeability [26]. In our study, we examined cell migration and showed that increased AQP5 expression is accompanied by increased neutrophil migration. The precise mechanism of AQP5 induced immune cell migration has not been clarified yet. Recently the mechanism of AQP9 induced neutrophil migration could be elucidated [27] and a similar mechanism was described for AQP1 [6]. Basing on these findings we can speculate on the mechanism of AQP5 induced cell migration. AQP5 could be phosphorylated by fMLP pathway and then translocated to the plasma membrane. Furthermore AQP5 expression could be increased by fMLP pathway, possibly via the transcription factor nuclear factor of activated T cells 1 (NFAT1) [28]. Increased AQP5 expression and localization of AQP5 to the plasma membrane leads to an influx of water and creation of a lower concentration of actin monomers which directs a flow of actin monomers to the site. Actin monomers polymerize and form a stable lamellipodium which leads to cell migration [6].

In this study, we also showed that AQP5 overexpression in the Jurkat lymphocyte cell line enhances SDF-1α induced migration. Interestingly, although SDF-1α is found ubiquitously throughout the body, lymphocytes from septic patients show increased migration towards SDF-1α [30].

We also demonstrate that fMLP induces AQP5-dependent neutrophil cell migration through a membrane. The process seems to amplify itself as we showed that fMLP increases AQP5 expression in the promyelocyte cell line HL-60. Neutrophil cell migration is an important mechanism occurring in sepsis [31, 32]. On the one hand, a decrease in neutrophil cell migration to the infected organs is associated with a bad outcome of sepsis, due to incomplete bacteria eradication [33]. On the other hand, neutrophils release large amounts of proteases and reactive oxygen species, which not only
kill bacteria but also damage host tissues [34]. Therefore, an overwhelming neutrophil cell migration to the tissues can cause multiple organ failure, which leads to death [34].

Limitations of our study should be mentioned. To our surprise, we could not demonstrate in our mouse model of LPS induced inflammation that Aqp5-KO is associated with a decreased cytokine concentration. The lack of a genotype dependent effect in cytokine concentrations could be due to the wide variability in cytokine production, which is acknowledged both in mice [35] and humans [36, 37], and cytokine secretion in itself as well as the response to corticoids is influenced by genetic variations [3, 4]. In contrast to humans mice are inbreed littermates with no genetic variation and cytokine differences should have been detected. Other studies could detect differences in TNF-α expression between KO and wild type mice of other genes [38]. Therefore AQP5 might not influence TNF-α pathway and other inflammatory proteins might be responsible for the differences occurring in Aqp5-KO and WT animals.

**Fig. 4** Migration assay of human primary neutrophils. Migration assays of human primary neutrophils unstimulated or fMLP (target) stimulated using cell culture inserts. Migrated cells were counted after 0.5 and 1.5 h (n = 4). Neutrophils from AA-genotype carriers showed target-oriented migration already after 0.5 h (a p = 0.029; c p = 0.056). Neutrophils from AC/CC-genotypes did not demonstrate target-oriented migration until 1.5 h, and, when compared to AA-genotype carriers, the number of migrated cells was threefold less than the AC/CC-genotype carriers (b p = 0.029; d p = 0.029).
Conclusion
In summary, the AQP5 genotype and AQP5 protein expression seem to alter neutrophil cell migration and may influence survival in sepsis by altering neutrophil cell migration. Therefore AQP5 might be a key protein in inflammation and depict a novel target for developing sepsis therapeutics.

Additional files

**Additional file 1.** Score sheet for monitoring animal health.

**Additional file 2.** Raw data from the figures are available as an excel file.

Abbreviations
AQP: aquaporin; KO: knockout; WT: wild type; LPS: lipopolysaccharide; TNF: tumor necrosis factor; fMLP: chemotactic peptide; IL: interleukin.

**Authors’ contributions**
KR carried out the molecular genetics and animal studies, analyzed the data and drafted the manuscript. MU and LB analyzed the data and helped to draft the manuscript. AB carried out the immunostaining. AM designed and provided the AQP5-KO mice. STS helped to draft the manuscript and participated in study design. AS performed the statistical analysis. ZB carried out the molecular biological analyses. WS, JP and MA designed the study, analyzed the data and helped to draft the manuscript. All authors read and approved the final manuscript.

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**Fig. 5** AQP5 expression in HL-60 promyelocytes after fMLP administration and in whole blood of septic patients. **a** AQP5 mRNA expression in HL-60 cells stimulated with $10^{-8}$ M fMLP for 6 h. FFMLP significantly increased AQP5 expression (n = 5; p = 0.03). **b** AQP5 protein expression in fMLP stimulated HL-60 cells for 6 h. AQP5 protein amount is increased in fMLP ($10^{-8}$ M)-stimulated cells (lane 2) compared to control (lane 1). **c** AQP5 RNA expression measured with RT-PCR in whole blood of septic patients (n = 14, 11); AA-genotype carriers showed an increased AQP5 mRNA expression compared to AC-genotype carriers (p = 0.025). No RNA was available from CC genotype carriers. **d** TNF-alpha serum concentrations of septic patients. No differences could be detected between genotypes (p $> 0.05$); Data derive from serum of septic patients (n = 65, 33, 3 a-d); data are shown as boxplots (min to max).
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Competing interests
The authors declare that they have no competing interests.

Consent for publication
Not applicable because the manuscript does not contain any individual person's data.

Availability of data and material
Raw data can are available as Additional file 2.

Ethics approval and consent to participate
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