Knockdown of SLC39A4 Expression Inhibits the Proliferation and Motility of Gallbladder Cancer Cells and Tumor Formation in Nude Mice

Min Li 1,*, Kun Fan 1,*, Bohao Zheng 1,*, David Zekria 2, Tao Suo 1, Han Liu 1, Sheng Shen 1, Houbao Liu 1, Xiaoling Ni 1
1Department of General Surgery, Zhongshan Hospital, Fudan University, Shanghai, 200032, People’s Republic of China; 2Department of Radiology, Leicester Royal Infirmary, Leicester, LEI 5WW, UK
*These authors contributed equally to this work

Purpose: Gallbladder cancer (GBC) is a common malignancy of the biliary tract and is characterized by rapid progression and early metastasis. Elucidating the molecular mechanisms of GBC could help to develop better treatment strategies.

Materials and Methods: Human GBC cell lines (GBC-SD and NOZ) were applied to determine the capacity of the proliferation and migration of cells using the MTT assay, colony formation, wound-healing assay as well as the Transwell™ assay. A nude xenograft was used to evaluate tumor growth in vivo.

Results: Using two types of GBC cell lines, we found that absence of solute carrier family (SLC) 39A4 (which encodes the zinc transporter ZRT/IRT-like protein [ZIP]4), could suppress the proliferation and migration of cells. Additionally, absence of ZIP4 could impair growth of xenografts in nude mice. While, over-expression of SLC39A4 could promote the GBC cell proliferation and migration, and inhibit apoptosis. We revealed that SLC39A4 might affect GBC progression by modulating the signaling pathways responsible for the survival, energy supply and metastasis of cells, and indicated that SLC39A4 could serve as a novel therapeutic target for GBC.

Conclusion: SLC39A4 promoted the viability and motility of GBC cells, and tumor formation in nude mice. We demonstrated an oncogenic potential for SLC39A4.

Keywords: gallbladder cancer, SLC39A4 (ZIP4), proliferation, migration, potential therapeutic target

Introduction

Gallbladder cancer (GBC) is aggressive, is more common in females, and is characterized by rapid progression and early metastasis. Treatment options are surgery, chemotherapy and radiotherapy. 1,2 However, GBC is usually diagnosed late due to a lack of early signs and clinical symptoms, which limits therapy choices and undermines a better prognosis. Thus, identification of specific diagnostic markers and elucidation of the underlying molecular mechanisms of GBC are very important. Indeed, the molecular pathology of GBC is poorly understood despite extensive research efforts. 3,4

Zinc is a vital element in the human body. It is not only a catalytic cofactor for several enzymes, it also has key roles in the signal transduction involved in cell differentiation, tissue development, and metabolism. 5,6 Zinc deficiency is associated with several diseases, 7–10 but zinc levels in most tumor cells are increased due to abnormal overexpression of zinc importers, which allow them to survive. 11
Solute carrier family (SLC) 39 (also known as ZRT/IRT-like protein [ZIP]) is responsible for transferring zinc from the extracellular space and organelles into the cytosol. Among them, ZIP4 (which is encoded by SLC39A4) is the major transporter for zinc uptake. Its aberrant expression has been found in different types of cancers. In hepatocellular carcinoma (HCC), SLC39A4 suppresses the apoptosis and promotes the migration of cells. In addition, SLC39A4 mediates drug resistance in non-small-cell lung cancer (NSCLC). Moreover, it serves as a prognostic marker in multiple cancer types. Nevertheless, the role of SLC39A4 in GC is not clear. We wished to assess the impact of SLC39A4 on the proliferation and migration of GBC cells. We also wished to clarify the changes of signaling pathways in response to SLC39A4 deficiency.

**Materials and Methods**

**Cell Culture and Construction of a Stable Cell Line**

Two human GBC cell lines, GBC-SD and NOZ, were obtained from the Japanese Collection of Research Bioresources Cell Bank (Tokyo, Japan). Both cell types were cultured in RPMI-1640 medium supplemented with 10% fetal bovine serum (FBS) at 37°C in an atmosphere of 5% CO₂. HEK293T cells were maintained in Dulbecco’s modified Eagle’s medium supplemented with 10% FBS at 37°C in an atmosphere of 5% CO₂.

GBC-SD cells and NOZ cells were infected with lentivirus encoding SLC39A4 short hairpin (sh) RNA or control lentivirus co-expressed with green fluorescent protein (GFP) according to manufacturer instructions. Briefly, HEK293T cells were used for lentivirus production. Stable cell lines expressing shSLC39A4 or control shRNA were sorted against GFP fluorescence 48 h after infection using fluorescence activated cell sorting and maintained in growth medium supplemented with puromycin (2 µg/mL) for one week.

**Cell Proliferation**

Cells were plated in 96-well plates at 2×10³/well and cultured in complete medium for 5 days. 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; catalog number, JT343; Geneview) was added and incubation allowed for 30 min. Then, the absorbance of the formazan solution was read using a spectrophotometer at 490 nm. All experiments were carried out in triplicate.

**Colony Formation**

Cells were seeded in six-well plates at 1×10³/well and then incubated for 13 days (GBC-SD) or 8 days (NOZ). Incubation was followed by staining with 0.1% Crystal Violet for 20 min at room temperature. Colonies containing ≥50 cells were counted manually under a microscope. Each assay was undertaken in triplicate.

**Wound Healing**

Cells were plated in 96-well plates and cultured overnight. Wounds were made in confluent monolayer cells using the 96 Wounding Replicator (V&P Scientific, San Diego, CA, USA). Cells were cultured in medium supplemented with 0.5% FBS. Wound healing was detected at 0 h and 48 h within scraped lines. Representative fields at different time points were photographed (×100 magnification). Migration area was analyzed by Celigo™ (Nexcelom, Lawrence, MA, USA).

**Transwell™ Migration**

Cells were plated in the upper chamber of the apparatus with serum-free medium in the Transwell with inserts of pore size 8-µm (3422; Millipore, Billerica, MA, USA). The lower chambers were filled with complete culture medium. After incubation for 24 h, cells on the upper-membrane surface were removed by cotton tips. Then, membranes were washed with phosphate-buffered saline (PBS), fixed with 4% paraformaldehyde, washed twice with PBS, and stained with 0.2% Crystal Violet. The migrated cells from nine fields were counted under a light microscope (×200 magnification). All experiments were done in triplicate.

**Distribution of the Cell Cycle**

Cells were treated as indicated. Approximately 48 h later, cells were harvested, washed with ice-cold PBS, and fixed in 70% ethanol overnight at −20°C. Subsequently, fixed cells were washed thrice with PBS and stained with propidium iodide (PI) solution (50 µg/mL PI and 100 µg/mL RNase A in PBS) in the dark for 30 min. DNA content was measured on a FACSscan flow cytometer (BD Biosciences, San Jose, CA, USA) with FlowJo 7.6.1 (Tree Star, Ashland, OR, USA). Experiments were repeated thrice.

**Apoptosis**

Pellets of GBC-SD cells and NOZ cells were washed with ice-cold PBS and resuspended in 200 µL of 1× binding buffer. Subsequently, cells were incubated with 10 µL of Annexin V-allophycocyanin (Beyotime Institute of...
Technology, Beijing, China) for 15 min in the dark at room temperature. Samples were detected using a BD Accuri™ C6 Plus flow cytometer (BD Biosciences). Data were quantified and analyzed using a Guava easyCyte™ flow cytometer (EMD Millipore, Burlington, MA, USA).

**Xenograft Model in Nude Mice**

Animal studies were conducted in compliance with the regulations on management of animal welfare set by the Chinese Association for Laboratory Animal Sciences (Beijing, China). The protocol for animal experiments was approved by the Ethics Committee of Zhongshan Hospital (Shanghai, China).

Briefly, $4 \times 10^6$ stably transfected NOZ cells were suspended in 200 μL of PBS and injected (s.c.) into female BALB/c nude mice (4 weeks). Nude mice were sacrificed after 18 days. Tumor masses were removed surgically and weighed. Tumor volume was calculated as: tumor volume = (length × width$^2$) × 0.5.

**RNA Extraction and Real-Time Reverse Transcription-Quantitative Polymerase Chain Reaction (RT-qPCR)**

RNA was extracted using TRIzol™ Reagent (Invitrogen, Carlsbad, CA, USA) and used for complementary (c) DNA synthesis with PrimeScript™ RT Reagent Kit with gDNA Eraser (TaKaRa Biotechnology, Shiga, Japan). RT-qPCR was conducted with SYBR™ Green PCR Master Mix (Applied Biosystems, Foster City, CA, USA) using the ABI Prism 7500 Real-time PCR System (Applied Biosystems).

The following primer sequences (forward and reverse, respectively) were used: ZIP4, 5'-CGGGCAGATGTGAAAGTACG-3' and 5'-ACAGCAGCAGGACGCT-3'; glyceraldehyde 3-phosphate dehydrogenase (GAPDH), 5'-TACTTCACACGAGACACA-3' and 5'-CACCC TGTGCTGATGCCAAA-3'; BMP4, 5'-CCTGTTGACCTGCA-3' and 5'-GATTTGCTGCACTAC-3'; SYK, 5'-CTGTCCTTCTGCACTAC-3' and 5'-GATTTGCTGCACTAC-3'; SGK1, 5'-GAAGCAGAGGAGGATGCTCTC-3' and 5'-GGTGGAGACCATGCAGCTGAGC-3'; CDK4, 5'-AGTTTGAGAAGGATGCTCTC-3' and 5'-GATTTGCTGCACTAC-3'; EGFR, 5'-AGGAGCAGAGGTACCAAAGCTC-3' and 5'-ATGAGGACATAACAGGACCACC-3'; MET, 5'-TATGTTGACCTGCACTAC-3' and 5'-ACGCTTCAAGATGTGAAAGTACG-3'. Analyses of melting curves were done to confirm specificity. The $2^{-\Delta\Delta C_t}$ method was used for quantification. Each sample was run in triplicate, and GAPDH was used as an internal control.

**Western Blotting**

Cells were washed with cold PBS. Proteins were extracted with lysis buffer supplemented with a proteinase inhibitor. Cell lysates were centrifuged at 12,000 $\times$ g for 10 min at room temperature. Supernatants were collected and protein concentrations were quantified using a bicinchoninic acid kit (P0010S; Beyotime Institute of Technology). Then, proteins (35 mg) with loading buffer were resolved by sodium dodecyl sulfate–polyacrylamide gel electrophoresis. Western blots were developed using anti-ZIP4 (orb395779; Biorbyt, Cambridge, UK), anti-GAPDH (sc-32,233; Santa Cruz Biotechnology, Santa Cruz, CA, USA), anti-GJA1 (ab51067; Abcam, Cambridge, UK), anti-SGK1 (12,103; Cell Signaling Technology, Danvers, MA, USA), anti-MET (ab51067; Abcam), anti-VEGFC (ab83905; Abcam), mouse anti-rabbit IgG (7074; Cell Signaling Technology) and rabbit anti-mouse IgG (7076; Cell Signaling Technology).

**Statistical Analysis**

Each experiment was carried out at least three times independently. Data are the mean ± SD. Comparisons between two groups were carried out using two-tailed unpaired $t$-tests. $P < 0.05$ was considered significant.

**Results**

**Establishment of SLC39A4 Stably Knocked-Down GBC Cells**

Studies have uncovered a tumor-promoting role of ZIP4 (SLC39A4) in various cancer types. $^{14,15,17,18}$ We wished to explore its potential relationship to GBC. First, we knocked-down SLC39A4 expression in a GBC cell line (GBC-SD) with lentivirus expressing three SLC39A4-specific shRNA sequences. shSLC39A4-2 had relatively higher efficiency in downregulating its expression of mRNA and protein (Figure 1A and B). Thus, lentivirus harboring shSLC39A4-2 was chosen to infect another GBC cell line: NOZ. Real-time RT-qPCR and Western blotting showed that expression of ZIP4 (SLC39A4) had been inhibited significantly (Figure 1C and D).
Knockdown of SLC39A4 Expression Suppressed Cell Proliferation and Colony Formation

ZIP4 has been shown to repress apoptosis and promote the cell cycle in pancreatic cancer cells and HCC cells.\textsuperscript{13,17} We wondered if this was also the case in GBC cells. Cell proliferation was inhibited significantly by silencing of SLC39A4 expression in NOZ cells (Figure 2A) and in GBC-SD cells (Figure 2B). The ability to form colonies was also decreased dramatically in SLC39A4 knocked-down cells (Figure 2C and D), which suggested that ZIP4 was required for the proliferation of GBC cells.

Knockdown of SLC39A4 Expression Impeded Cell Migration

It has been reported that ZIP4 is overexpressed in NSCLC, and is associated with enhanced migration of cells.\textsuperscript{14} In HCC, ZIP4 expression is also activated and cell migration increased.\textsuperscript{17} Next, we sought to discover what would happen after SLC39A4 expression was silenced. In the wound-healing

\begin{figure}
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\caption{Establishment of SLC39A4 stably knocked-down cell lines in GBC-SD cells and NOZ cells. (A and B) Knockdown efficiency of three shSLC39A4s was evaluated in GBC-SD cells by real-time RT-qPCR (A) and Western blotting (B) and shSLC39A4-2 was selected to carry out subsequent experiments. (C and D) Confirmation of SLC39A4 expression in NOZ shSLC39A4-2 (NOZ/shSLC39A4-2) and control (NOZ/shCtrl) cells by real-time RT-qPCR (C) and Western blotting (D). The results of real-time RT-qPCR are the mean ± SD (unpaired t-test). \(*p < 0.05, **p < 0.01, ***p < 0.001,\) compared with NOZ/shCtrl or GBC-SD/shCtrl.}
\end{figure}
assay, NOZ/shSLC39A4 cells and GBC-SD/shSLC39A4 cells migrated much more slowly than control cells (Figure 3A and B). Also, in the Transwell migration assay, silencing of SLC39A4 expression inhibited transmigration of NOZ cells considerably. An identical result was obtained in GBC-SD/shSLC39A4 cells (Figure 3C and D). Taken together, these data indicated that ZIP4 facilitated migration of GBC cells.

Knockdown of SLC39A4 Expression Induced the Apoptosis and Cycle Arrest of Cells

To reveal the mechanisms underlying repression of cell proliferation, we analyzed the cell-cycle distribution of GBC cells after knockdown of SLC39A4 expression. After knockdown of SLC39A4 expression, the number of apoptotic GBC-SD cells and NOZ cells increased (Figure 4A–D). We observed accumulation of GBC-SD/shSLC39A4-2 cells in the G0/G1 phase compared with that in control cells, and an obvious reduction in the number of GBC-SD/shSLC39A4-2 cells in the S phase (Figure 4E and F). Knockdown of shSLC39A4 expression also led to increased proportions of NOZ cells in the G0/G1 phase and decreased proportions in the S phase (Figure 4G and H), suggesting that cell-cycle arrest and apoptosis contributed considerably to inhibition of cell proliferation.

Overexpression of SLC39A4 Promoted the Proliferation, Cycle Progression, and Migration of Cells

To exclude the off-target effects of shRNA knockdown, we overexpressed SLC39A4 in GBC-SD cells and NOZ

![Figure 2 Knockdown of SLC39A4 expression suppressed cell proliferation and colony formation. (A and B) MTT assays were undertaken to evaluate proliferation of NOZ/shSLC39A4-2 cells (A) and GBC-SD/shSLC39A4-2 cells (B). (C and D) Colony-formation assays of NOZ/shSLC39A4-2 cells (C) and GBC-SD/shSLC39A4-2 cells (D). Images and quantitation are shown. Data are the mean ± SD (unpaired t-test) *p<0.05, **p<0.01, ***p < 0.001, compared with NOZ/shCtrl or GBC-SD/shCtrl.](image-url)
cells to observe the growth and migration of cells. SLC39A4 was efficiently overexpressed in GBC-SD and NOZ cells (Figure 5A). As expected, GBC-SD cells and NOZ cells transfected with exogenous SLC39A4 exhibited enhanced viabilities (Figure 5B). The proportion of GBC-SD cells in the G0/G1 phase showed an obvious decline, but an increase in the proportion of GBC-SD cells in the S and G2/M phases after SLC39A4 overexpression was documented (Figure 5C and D). In SLC39A4-overexpressed NOZ cells, the number of G0/G1-phase cells also showed a decline but their number in the G2/M phase increased (Figure 5E and F). These results indicated that the division activity of GBC-SD cells was motivated considerably by increased expression of SLC39A4. Besides, more GBC-SD cells and NOZ cells with SLC39A4 overexpression transmigrated than cells with the control vector (Figure 5G–J).

**Knockdown of SLC39A4 Expression Repressed Tumor Formation in Nude Mice**

SLC39A4 expression was related positively to the proliferation and colony formation of GBC cells in vitro. Hence, we proceeded to discover if SLC39A4 expression could affect tumor formation in vivo. After subcutaneous injection of control cells or NOZ/shSLC39A4 cells, xenografts were allowed to grow freely up to 18 days. As expected, tumors derived from NOZ/shSLC39A4 cells were restrained obviously compared with those in control groups (Figure 6A and B), and the average weight of tumors was also lighter (Figure 6C). In accordance with these findings, expression of c-Met and Ki-67 was downregulated in tumor tissues of mice with knockdown of SLC39A4 expression (Figure 6D). These results suggested that SLC39A4 might serve to potentiate tumorigenesis.

**Downregulation of SLC39A4 Expression Influenced the Signaling Pathways Involved in Tumor Progression**

Based on the results detailed above, we tried to identify the signaling pathways that were affected by SLC39A4 insufficiency. As expected, expression of genes that regulated cell proliferation, such as EGFR, CDK4, and c-MET, was decreased (Figure 7A and B).\(^\text{19,20}\) Besides, expression of genes that promote metastasis and vascular formation, such as GJA1 and SYK, was repressed (Figure 7A and B).\(^\text{21–23}\) In particular, expression of VEGFC, a positive regulator of lymph-node metastasis and lymphangiogenesis,\(^\text{24,25}\) was downregulated in response to SLC39A4 absence (Figure 7B). Inflammation and modulation of the immune microenvironment have key roles in influencing tumor progression.\(^\text{26,27}\) RUNX3 aids in the control of immunity and inflammation.\(^\text{28}\) After silencing of...
Knockdown of SLC39A4 expression induced apoptosis and cell-cycle arrest. (A–D) Apoptosis analyses were done in GBC-SD cells and NOZ cells using FACS (A and C), and apoptotic percentages were calculated by normalizing values to the total number of cells (B and D). Data are the mean ± SD (unpaired t-test). **p<0.01, compared with NOZ/shCtrl or GBC-SD/shCtrl. (E–H) Cell-cycle distributions were analyzed by FACS (E and G) and quantitation are shown in (F and H), after normalization to the total cell number. Data are the mean ± SD (unpaired t-test). *p < 0.05, compared with control cells.
Figure 5 Overexpression of SLC39A4 promoted the growth and migration of cells. (A) Real-time RT-qPCR was done to confirm expression of exogenous SCL39A4 in GBC-SD cells and NOZ cells. mRNA expression was normalized to that of GAPDH. (B) MTT assays were undertaken to evaluate cell proliferation. (C–F) Cycle distributions were analyzed in GBC-SD cells and NOZ cells overexpressing SLC39A4 by FACS (C and E) and data are presented in (D and F) normalized to the total cell number. (G–J) Transwell-migration assays were conducted to observe the transmigration capability of NOZ/SLC39A4 cells and GBC-SD/SLC39A4 cells. (G and I) Representative images (magnification, ×200) and (H and J) quantitation are shown. Data are the mean ± SD (unpaired t-test). *p < 0.05, **p < 0.01, ***p < 0.001, compared with control cells.
SLC39A4 expression, RUNX3 expression was reduced (Figure 7A). Also, RUNX3 has been found to correlate with anchorage-independent growth in pancreatic cancer cells. Moreover, expression of BMP4 and SGK1 was downregulated (Figure 7A and B), both of which have been found to favor tumor-cell survival and whose inhibition could lead to
apoptosis. All the above data illustrated that ZIP4 deficiency could affect the different signaling pathways involved in tumor progression.

**Discussion**

The gallbladder stores bile and is located under the liver. GBC is the most common malignancy of the biliary tract, and often occurs in women. There are limited treatment options for GBC because it is often detected late and specific targets for therapy have not been identified. Therefore, obtaining specific diagnostic markers for GBC is very important to better understand and treat this disease. However, little is known about the mechanisms of GBC.

Zinc is an essential mineral for life. It is indispensable for most enzymes to carry out catalytic activities and for nucleic acids to be synthesized. Zinc inadequacy leads to growth retardation, impaired immune function, and delayed healing of wounds. Conversely, excess zinc can cause disorders. For instance, a large intake of zinc results in low copper status and reductions in the levels of high-density lipoproteins. Moreover, the zinc level has been found to be increased in some tumor cells. As a result, zinc concentration must be controlled tightly.

In humans, intracellular zinc homeostasis is regulated by two major zinc transporter families: the SLC30 (ZnT) family and SLC39 (ZIP) family. The latter is responsible for zinc influx and could play a crucial part in malignancies in humans. For instance, SLC39A6 promotes the proliferation and invasion, and inhibits apoptosis, in esophageal squamous cell carcinoma (ESCC) cells. Suppression of SLC39A7 can abrogate survival of colorectal cancer cells. Also, in metastatic breast cancer, SLC39A10 expression is correlated positively with lymph-node metastasis. Expression of SLC39A4 (which encodes ZIP4) has been reported to be correlated positively with progression of pancreatic cancer. Also, activated ZIP4 inhibits apoptosis of HCC cells and enhances their cell cycle and migration. In NSCLC, SLC39A4 expression has been shown to be associated with strengthened cell migration, cisplatin resistance, and poor survival. SLC39A4 could even serve as a prognostic marker in ESCC. All the observations mentioned above indicate that SLC39A4 could be used as a therapeutic target for human cancers. We wished to determine the effects of SLC39A4 on proliferation of GBC cells in vitro and in vivo, as well as the molecular signaling mechanisms involved. We discovered that down-regulation of SLC39A4 expression could repress the growth and migration of cells significantly, as well as tumor formation in nude mice. Taken together, these results implied an oncogenic character for SLC39A4 in GBC cells.

Gene expression analysis showed that SLC39A4 regulated the expression of VEGFC and SYK. The latter is downstream of SRC kinases and VEGFC is the ligand of VEGFR3. SYK is involved in vascular development, and VEGFC has been shown to promote lymphangiogenesis and lymph-node metastasis. These phenomena are consistent with the effects of SLC39A10 mentioned above. c-MET and CDK4 are important oncogenes which regulate GBC cell survival, metastasis and cell

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**Figure 7** Downregulation of SLC39A4 expression influenced the malignant-development potential of GBC-SD cells. (A) mRNA expression of target genes in GBC-SD/shSLC39A4-2 cells by real-time RT-qPCR. Results are representative of three independent experiments, and are shown as the mean ± SD (unpaired t-test). *p < 0.05, ***p < 0.001, compared with GBC-SD/shCtrl. (B) Protein expression of indicated target genes in GBC-SD/shSLC39A4-2 cells by Western blotting.
Combination of CDK4/6 and c-MET inhibitor have been applied for the treatment of glioblastoma.\(^4^\)\(^-^\)\(^2^\) Here, we showed that CDK4 and c-MET were regulated by SLC39A4. Both mRNA and protein abundance of CDK4 and c-MET was reduced in GBC cells with silenced SLC39A4. Thus, we proposed that SLC39A4 knockout suppressed GBC growth and migration partly through downregulation of CDK4 and c-MET.

Owing to inhibition of ZIP4 expression, zinc concentrations may change in the cytosol of tumor cells. We found that various aspects of the behaviors and signaling pathways of GBC cells were influenced if SLC39A4 expression was downregulated. Considering zinc as a vital element for numerous enzymes to function normally, we wondered if these outcomes were caused by changes in the zinc concentration, or if ZIP4 participated in these pathways directly. Additional investigations are needed to clarify the exact roles of ZIP4 in GBC.

**Conclusions**

We demonstrated an oncogenic potential for SLC39A4. Suppression of SLC39A4 expression in GBC cells weakened the viability and motility of GBC cells, and tumor formation in nude mice. Conversely, high expression of SLC39A4 could promote the growth and migration of tumor cells. Based on these findings, we speculate that SLC39A4 is very likely to be a novel target for developing new strategies for treating GBC. Importantly, this study may help us gain some novel insights of the molecular events triggering GBC progression.

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**Disclosure**

The authors report no conflicts of interest in this work.

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