

# Regulation of SLD5 gene expression by miR-370 during acute growth of cancer cells

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# SCIENTIFIC REPORTS



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## Regulation of SLD5 gene expression by miR-370 during acute growth of cancer cells

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SLD5 is a member of the GINS complex, essential for DNA replication in eukaryotes. It has been reported that SLD5 is involved in early embryogenesis in the mouse, and cell cycle progression and genome integrity in *Drosophila*. SLD5 may be involved in malignant tumor progression, but its relevance in human cancer has not been determined. Here, we found strong SLD5 expression in both human bladder cancer tissues from patients and cell lines. Knockdown of SLD5 using small interfering RNA resulted in reduction of cell growth both *in vitro* and an *in vivo* xenograft model. Moreover, we found that high levels of SLD5 in bladder cancer cells result from downregulation of microRNA (miR)-370 that otherwise suppresses its expression. High level expression of DNA-methyltransferase (DNMT) 1 and IL-6 were also observed in bladder cancer cells. Knockdown of IL-6 led to downregulation of DNMT1 and SLD5 expression, suggesting that IL-6-induced overexpression of DNMT1 suppresses miR-370, resulting in high SLD5 expression. Our findings could contribute to understanding tumorigenic processes and progression of human bladder cancer, whereby inhibition of SLD5 could represent a novel strategy to prevent tumor growth.

In order to carry out nuclear DNA replication during G1 and S phase, many factors are recruited to the chromosomal origin, one of which is the GINS complex composed of SLD5 and partner of Sld5 (Psf) -1, -2, and -3, identified in eukaryotes in 2003<sup>1</sup>. The GINS complex has been reported to regulate DNA polymerase  $\epsilon$  (DNA poly  $\epsilon$ )<sup>1</sup>. In budding yeast, cyclin-dependent kinases (CDK) activate and load GINS complexes at origins<sup>1,2</sup> to regulate the initiation of DNA replication.

Recently, several studies have shown that each GINS member is associated with malignant progression in several different tumor histotypes, i.e., PSF1 for breast cancer<sup>3</sup>, colon cancer<sup>4</sup>, and lung cancer<sup>5</sup>, PSF2 for cholangiocarcinoma<sup>6</sup>, and PSF3 for colon cancer<sup>7</sup> and non-small cell lung cancer<sup>8</sup>. Although we previously reported on the levels of expression of SLD5 in different cancer cell lines<sup>7</sup>, to the best of our knowledge there have been no reports thus far on its expression and malignant tumor progression. We did report that targeted disruption of the SLD5 gene led to disturbance of epiblast proliferation and resulted in early embryonic lethality<sup>9</sup>. Moreover, we found that SLD5 is involved in protection from DNA damage in mice<sup>10</sup>. It has also been suggested that SLD5 plays a role in maintaining genome integrity in *Drosophila*<sup>11</sup>. Therefore, it may be hypothesized that SLD5 has a specific function in tumorigenesis.

Most bladder cancers are non-invasive transitional cell carcinomas, but recurrence and progression rates can be very high. It has been suggested that malignant progression in bladder cancer is associated with chromosomal abnormalities<sup>12,13</sup> and gene mutations in RB1 and p16<sup>13</sup>, TP53<sup>14</sup>, G1 checkpoint protein<sup>15</sup> and/or cyclin D1<sup>16,17</sup>. Apart from these genes, other molecular targets for suppressing tumor progression are likely to exist in bladder cancer.

MicroRNAs (miRNAs) are endogenous RNAs containing approximately 18–25 nucleotides which regulate gene expression and translation by binding to 3' UTRs of target genes<sup>18,19</sup>. So far, over 1000 miRNAs have been discovered, and it has been proposed that around 60% of genes are regulated by miRNAs<sup>20,21</sup>. It is widely accepted that changes of miRNA expression in tumors compared with normal tissues affect malignant progression and that such miRNAs may be therapeutic targets. In bladder cancer, several studies have suggested that changes of miRNA expression are involved in tumorigenesis<sup>22–24</sup>.

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The aim of the present study was to analyze whether SLD5 expression is associated with tumor growth and if so, how its expression is regulated genetically or epigenetically in tumors, including by miRNAs. Our data suggest that expression of miR-370 is negatively regulated in bladder cancer cells, resulting in upregulation of SLD5 to induce tumor growth. Here, we show how miR-370 is suppressed in association with the functions of IL-6 and DNMT1, and discuss the effectiveness of SLD5 suppression as a potential therapeutic benefit.

## Results

**SLD5 is highly expressed in human bladder cancer tissues.** In the tested human cancer arrays, strong SLD5 expression in bladder cancer was found exclusively in transitional cell carcinoma (Fig. 1A). Because SLD5 is a member of the GINS complex, which regulates DNA replication, we asked whether SLD5 expression is related to cell cycle activity and cellular growth. For this, we stained tissues with anti-SLD5 together with anti-Ki-67 antibodies. As shown in Fig. 1B, most of the SLD5-positive cells also expressed Ki-67, suggesting that SLD5 marks the proliferating cancer cells. Some cells in normal bladder tissue also expressed SLD5, but these were Ki-67 negative, suggesting a different role of SLD5 in normal cells.

To assess the expression level of SLD5 in cancer cells relative to normal cells, we analyzed SLD5 mRNA expression in human bladder cancer cell lines (T24 and KMBC2), in human normal bladder cells (HNBC), and human umbilical vein endothelial cells (HUVECs) using quantitative (q) RT-PCR analysis. Extremely high expression of SLD5 was observed in cancer cells relative to normal cells (Fig. 1C). We also assessed SLD5 protein expression in bladder cancer cells by Western blotting. As observed for mRNA, bladder cancer cells also expressed higher amounts of SLD5 protein than normal cells (HUVEC and HNBC) (Fig. 1D). Moreover, we confirmed that most of the cultured bladder cancer cells expressed SLD5 protein exclusively in their nuclei (Fig. 1E). T24 cells divided faster than HUVECs (Fig. 1F), suggesting that SLD5 expression may correlate with the rate of cell proliferation and possibly acute tumor growth.

**Knockdown of SLD5 reduces the number of cells in S phase and Ki-67-positive cells.** Next, we analyzed whether knocking down SLD5 affects the growth of bladder cancer cells. For this, we silenced SLD5 in bladder cancer cells using SLD5-specific siRNA. We confirmed the knockdown effect using qRT-PCR and Western blotting (Fig. 2A,B) and found that attenuation of SLD5 expression inhibits the growth of both T24 and KMBC2 cancer cells (Fig. 2C). It is possible that silencing SLD5 might affect the expression of other GINS members as well, but we found that levels of these were not altered (Supplementary Fig. S1).

When cells were labelled with EdU to analyze DNA synthesis, EdU-positive cells were significantly reduced on silencing SLD5 (Fig. 2D). Similarly, Ki-67 positivity was reduced by knocking down SLD5 (Fig. 2E). Cell cycle analysis showed that the fraction of cells in S phase was reduced by knocking down SLD5 (Fig. 2F). Therefore, we conclude that SLD5 positively regulates the cell cycle of bladder cancer cells.

**Suppression of tumor growth by injection of SLD5 siRNA *in vivo*.** The above *in vitro* experiments had clearly demonstrated that silencing SLD5 suppresses proliferation of cancer cells. Next, we asked whether silencing of SLD5 effectively inhibits tumor growth *in vivo*. To assess this, T24 cells were inoculated into nude mice and scrambled (control) or SLD5-specific siRNA mixed with atelocollagen was injected into the tumors once they had reached approximately 50 mm<sup>3</sup> in diameter. After injection of SLD5-specific siRNA, tumor growth was clearly suppressed for at least 20 days (Fig. 3A,B).

To confirm reduced expression of SLD5, we performed qRT-PCR analysis using RNA from scrambled or SLD5-specific siRNA-injected tumors 20 days after treatment. We confirmed that the level of SLD5 in the tumors injected with specific siRNA was half that of control tumors (Fig. 3C). Moreover, the fraction of SLD5- and Ki-67-double positive cells was also reduced in siRNA-injected tumors, as confirmed by immunohistochemistry (Fig. 3D).

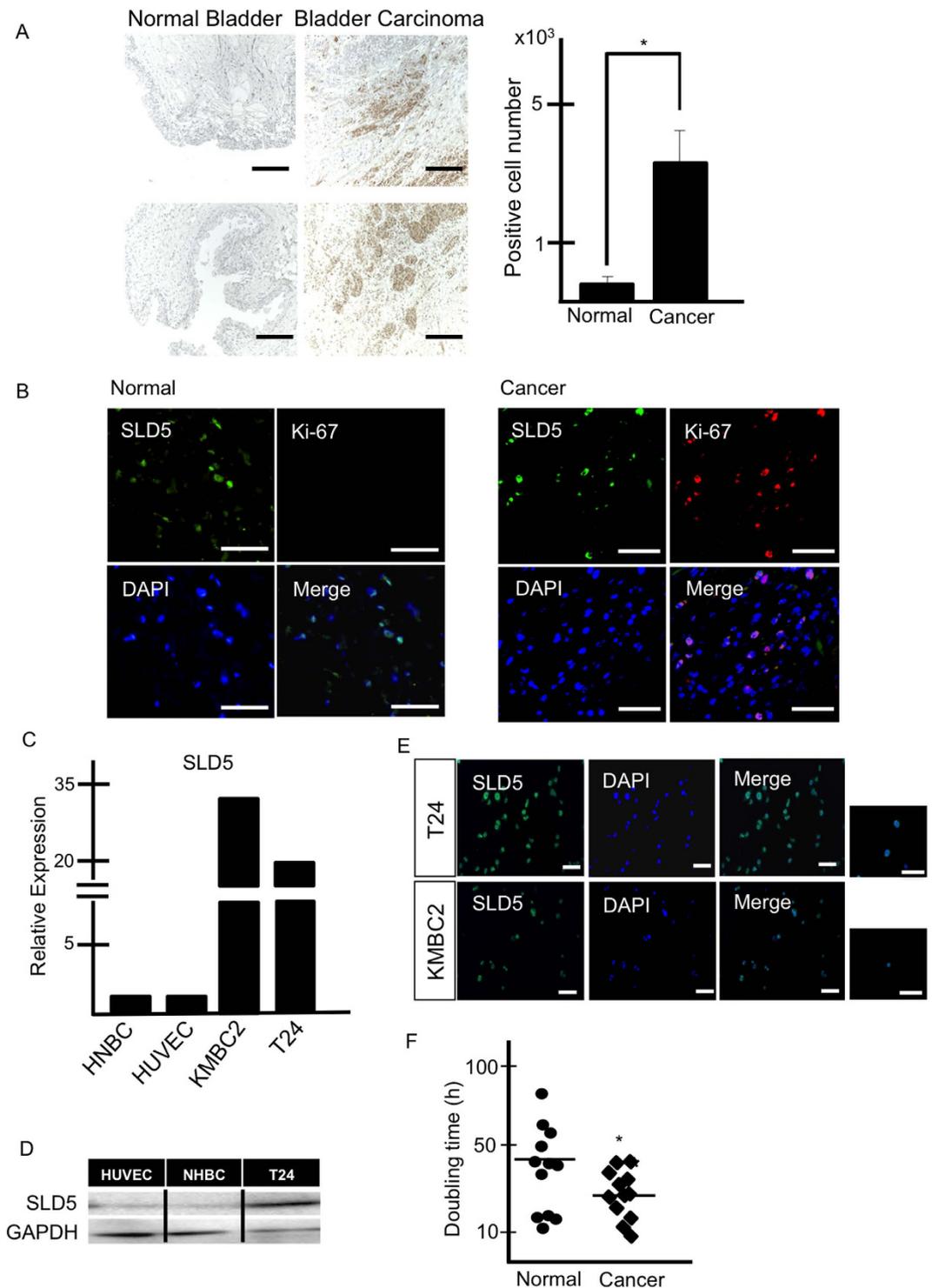
**miR-370 downregulation results in SLD5 upregulation in human bladder cancer cells.** SLD5 expression is lower in normal cells than in bladder cancer cells (Fig. 1). This suggests that miRNAs negatively regulating SLD5 expression are downregulated in cancer cells. To identify candidate miRNAs responsible for this, we sought those which interact with the 3'-UTR region of the SLD5 gene using Targetscan, as in previous reports showing an association with bladder cancers<sup>22–24</sup>. Among several candidates, we found that a miR-370 target sequence is located at the 3'-UTR of the SLD5 gene. We confirmed that miR-370 expression was indeed suppressed in T24 and KMBC2 cells (Fig. 4A,B).

To document direct binding of miR-370 to the 3'-UTR of the SLD5 gene, we performed luciferase reporter assays. T24 cancer cells were transfected with pMIR luciferase vector with an inserted 3'-UTR region of SLD5 containing the miR-370 target sequence. Addition of a miR-370 mimic led to reduction of the luciferase activity of the SLD5 3'-UTR, whereas miR-214 as a negative control had no effect (Fig. 4C). Therefore, we conclude that miR-370 directly binds to the 3'-UTR of the SLD5 gene.

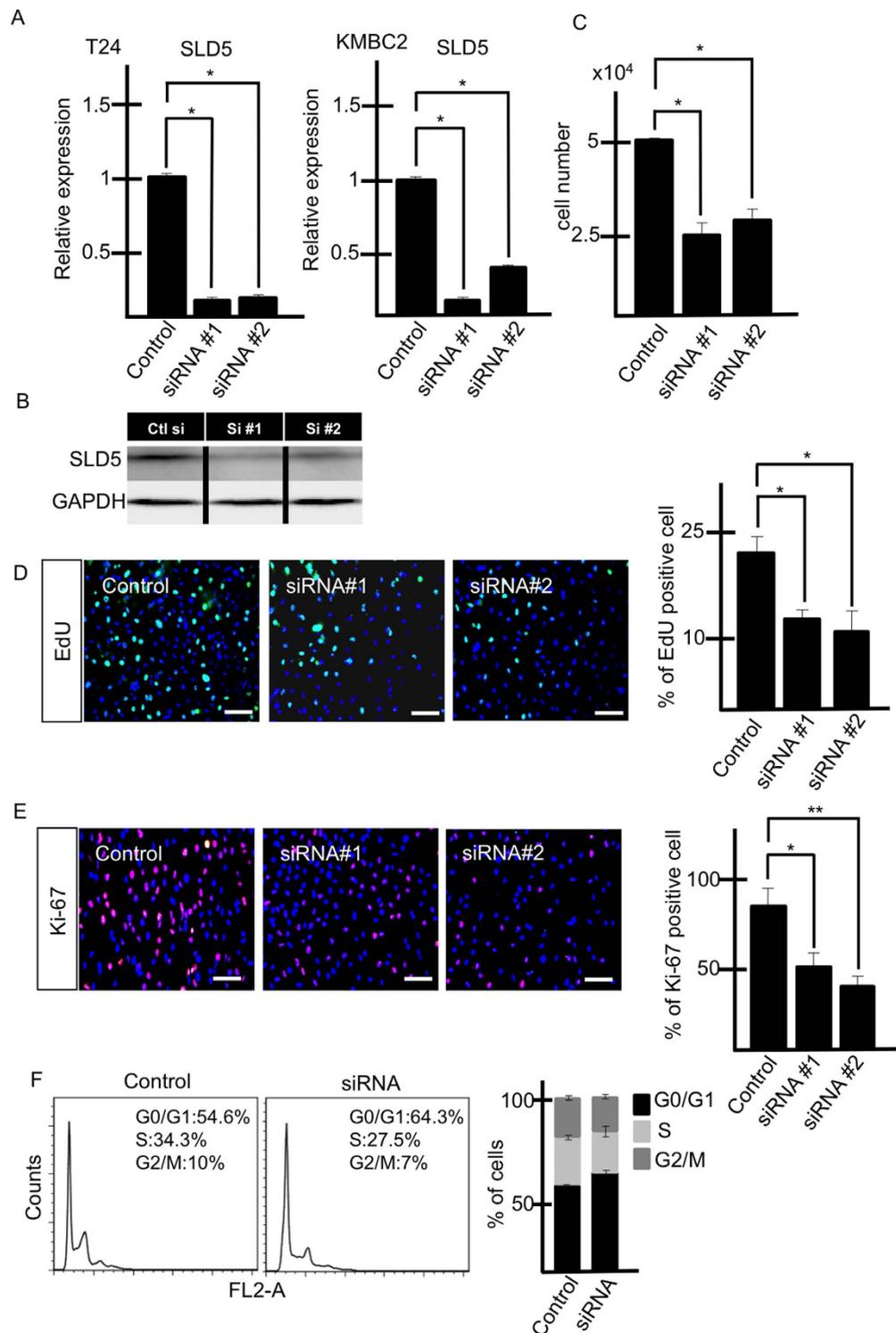
Next, we investigated how miR-370 affects SLD5 expression. We overexpressed it in T24 bladder cancer cells using miRNA mimic transfection methods, and assessed SLD5 mRNA and SLD5 protein levels using qRT-PCR (Fig. 4D) and Western blotting (Fig. 4E), respectively, after 48 hours. The results suggested that miR-370 inhibits mRNA expression in T24 cells, resulting in a reduction of SLD5 protein expression.

Next, we investigated whether miR-370 silencing affects SLD5 expression in normal cells. We knocked down miR-370 in normal cell lines (HUVEC) using miR-370 inhibitor transfection methods and assessed SLD5 mRNA and SLD5 protein levels using qRT-PCR and Western blotting, respectively (Supplementary Fig. 3). The results suggested that inhibition of miR-370 led to increased SLD5 mRNA and protein expression in normal cells.

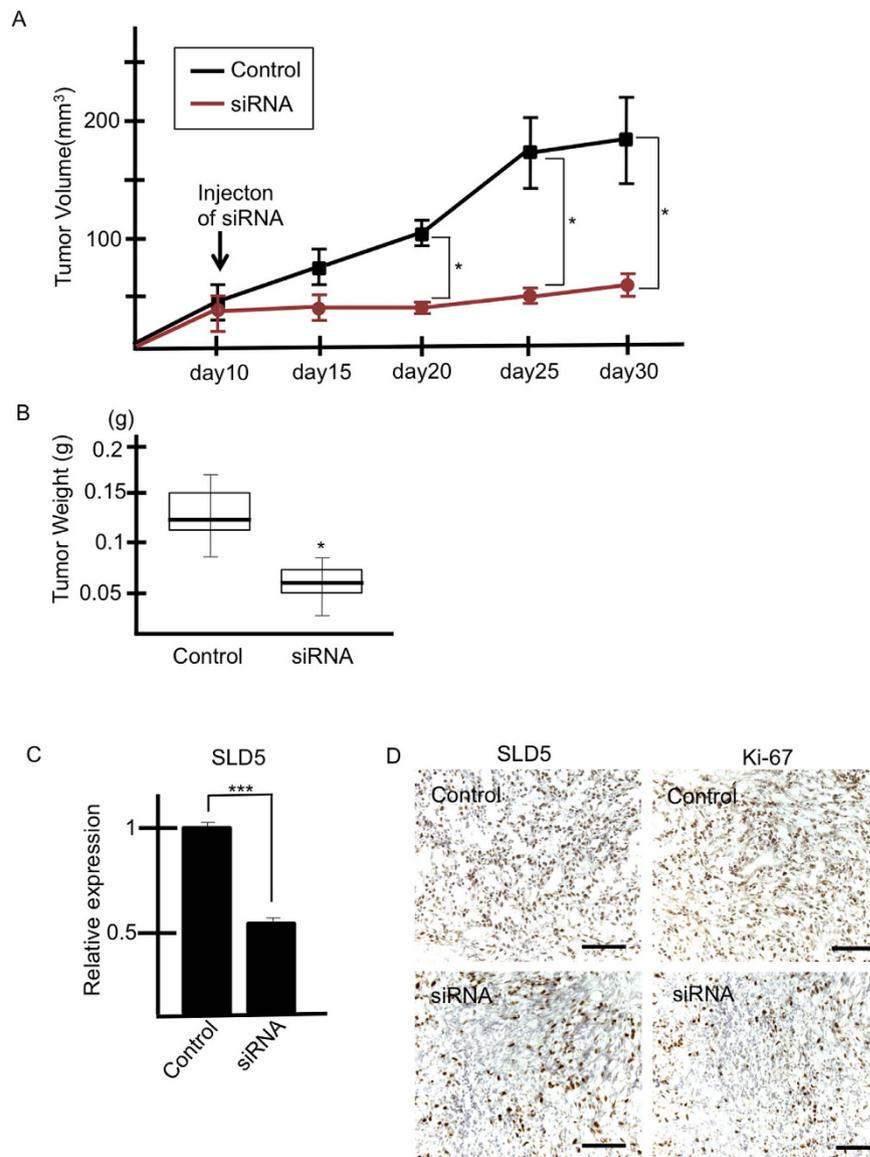
As observed in the siRNA experiments (Fig. 2), miR-370 overexpression also inhibited the proliferation of bladder cancer cells (Fig. 4F). Moreover, we analyzed cell kinetics and how the cell cycle was affected by miR-370.



**Figure 1. Expression of SLD5 in human bladder cancer tissue.** (A) Representative images of normal bladder tissue and bladder cancer tissue stained with anti-SLD5 antibody. Samples were counterstained with hematoxylin. Brown: SLD5, Blue: nuclei. Scale bars: 100  $\mu$ m. Graph shows the quantitative evaluation of SLD5-positive cells in random fields. Data show the mean  $\pm$  SE, \* $p < 0.05$ , (n = 3). (B) Immunofluorescence staining of normal and bladder cancer tissues. SLD5 (green), Ki-67 (red), and DAPI (blue). Scale bar: 100  $\mu$ m. (C) Comparison of mRNA expression levels of *SLD5* in normal and cancer cells. Human normal bladder cells (HNBC), human umbilical vein endothelial cells (HUVEC), and human bladder cancer cells (KMBC2, T24) were analyzed by qRT-PCR. Normal cell expression was set at unity. *GAPDH* was used as an internal control. (D) Comparison of SLD5 protein expression by Western blotting. HUVECs, HNBC and T24 bladder cancer cells (T24) were used. (E) Immunocytochemical analysis of T24 and KMBC2 with anti-SLD5 antibody (green) in cancer cells. Nuclei were stained with DAPI (blue). Scale bars: 100  $\mu$ m. (F) Comparison of doubling time in HUVECs (Normal) and T24 (Cancer). Bars show the mean  $\pm$  SE, \* $p < 0.05$ .



**Figure 2. Relevance of SLD5 expression for cell cycle progression.** Bladder cancer cell lines (T24, KMBC2) were transfected with two types of SLD5-specific siRNA (#1 and #2). (A) Quantitative evaluation of *SLD5* mRNA affected by siRNA. Data are mean  $\pm$  SE, \* $p < 0.05$  ( $n = 3$ ). (B) Western blotting of SLD5 in control scrambled siRNA (Ctl) and *SLD5*-specific siRNA #1 and #2 treated cells. (C) Comparison of cell growth after siRNA transfection (T24). Data are mean  $\pm$  SE, \* $p < 0.05$  ( $n = 3$ ). (D,E) Cell growth (T24) affected by SLD5 silencing. (D) Uptake of EdU (green). Nuclei were stained with Hoechst 33342 (blue). EdU-positive cells were counted and normalized to nuclear numbers, and are shown graphically. Data are mean  $\pm$  SE, \* $p < 0.05$  (3 random fields) Scale bars: 100  $\mu$ m. (E) Ki-67 staining (red). Nuclei were stained by DAPI (blue). Ki-67-positive cells were counted and normalized to the number of all surviving cells. Results are shown as mean  $\pm$  SE, \* $p < 0.05$  \*\* $p < 0.005$  (3 random fields). Scale bars: 100  $\mu$ m. (F) Cell cycle analysis using flow cytometry. After transfection with control or *SLD5*-specific siRNA, cells were analyzed. The right-hand graph depicts the quantitative evaluation by showing mean  $\pm$  SE ( $n = 3$ ).

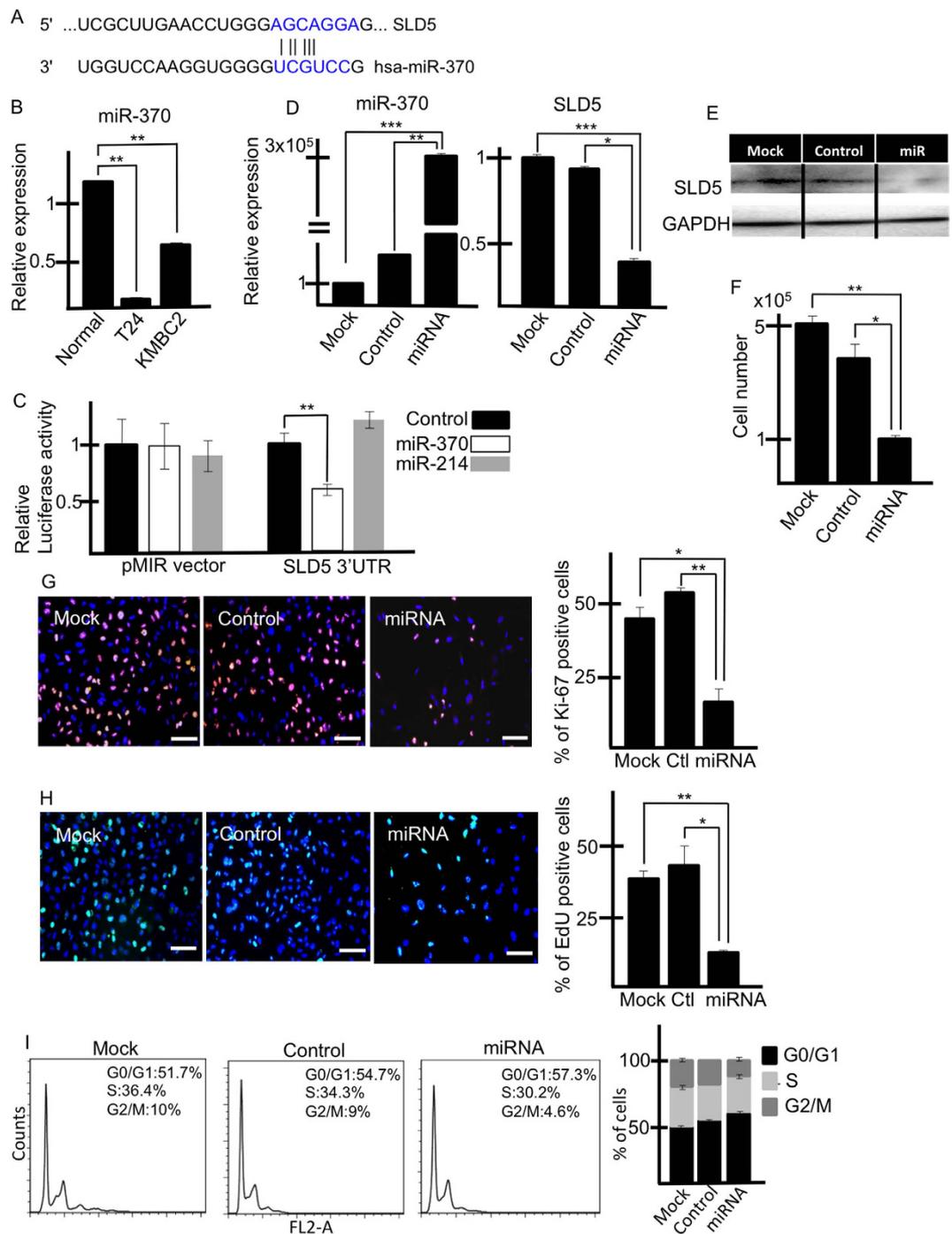


**Figure 3. Silencing of SLD5 attenuates tumor growth.** Bladder cancer cells (T24) were subcutaneously transplanted into nude mice. Control or *SLD5*-specific siRNA was injected on day10 post-inoculation. **(A)** Time course of tumor volume dynamics. Data are mean  $\pm$  SE (n = 3). **(B)** Tumor weight on day 30 post-inoculation. \*p < 0.05 mean  $\pm$  SE. **(C)** qRT-PCR analysis for assessment of *SLD5* mRNA expression in control and *SLD5* siRNA-injected tumors. Data are mean  $\pm$  SE \*\*\*p < 0.0001. **(D)** Xenograft tumors injected with control or *SLD5* siRNA were stained with anti-*SLD5* or Ki-67 antibody (brown). Sections were counterstained with hematoxylin. Scale bars: 100  $\mu$ m.

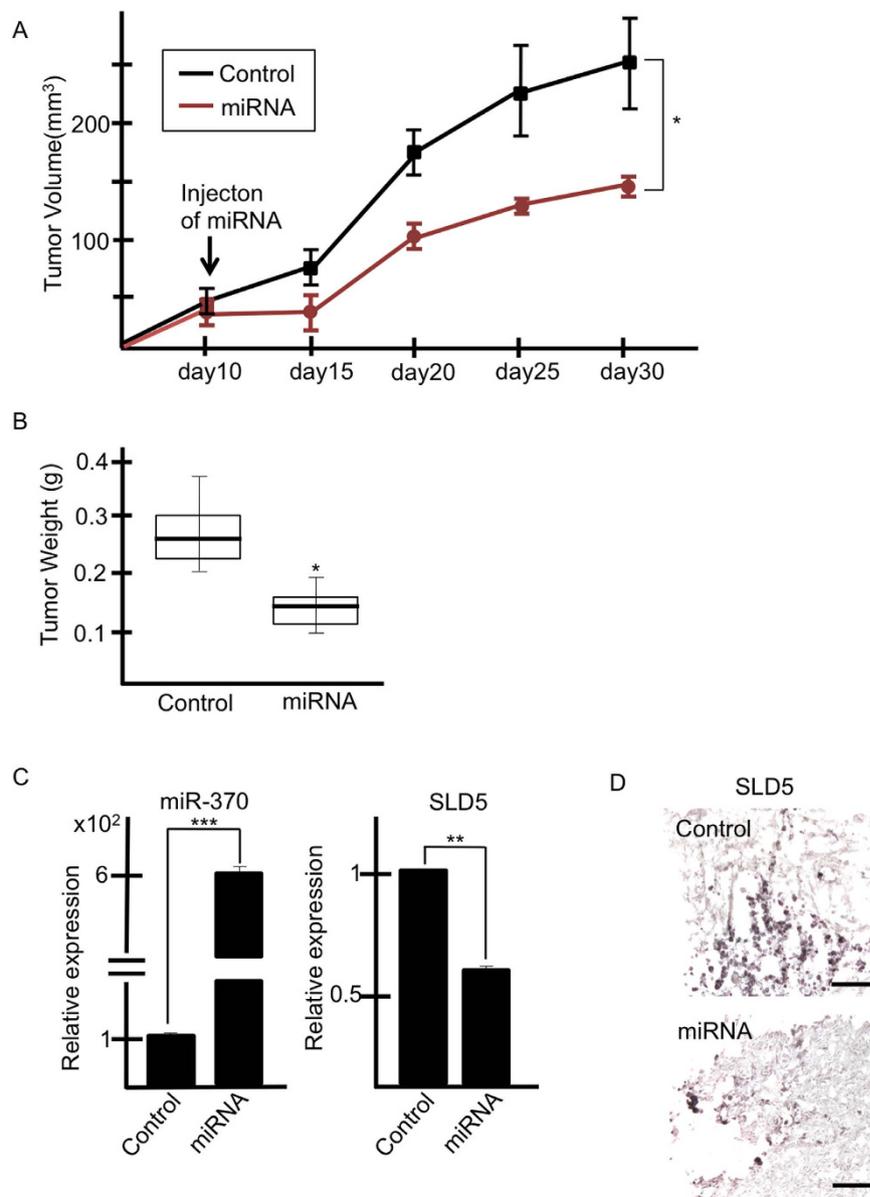
As shown in Fig. 4G,H, the fraction of Ki-67- and EdU-positive cells was reduced in miR-370-transfected cells compared to controls. In terms of the cell cycle, similar to the results from the *SLD5* knockdown study, miR-370 mimic transfection reduced the fraction of cells in S phase relative to control miR transfection (Fig. 4I).

**A miRNA-370 mimic inhibits tumor growth *in vivo*.** *In vitro* experiments had clearly demonstrated that overexpression of miR-370 suppresses proliferation of cancer cells. Next, we investigated whether miR-370 can effectively inhibit tumor growth *in vivo*. To assess this, T24 cells were inoculated into nude mice and control miR or miR-370 mixed with atelocollagen was injected into the tumors once they had reached approximately 50 mm<sup>3</sup> in diameter. After injection of miR-370, tumor growth was clearly suppressed (Fig. 5A,B).

We assessed the level of expression of miR-370 and *SLD5* by qRT-PCR using tumor tissue on day 20 after treatment, and confirmed overexpression of miR-370 and downregulation of *SLD5* in tumors from the miR-370-injected group. Moreover, we confirmed that *SLD5* protein was also suppressed in tumor tissue of miR-370-injected relative to control tumors (Fig. 5C,D).



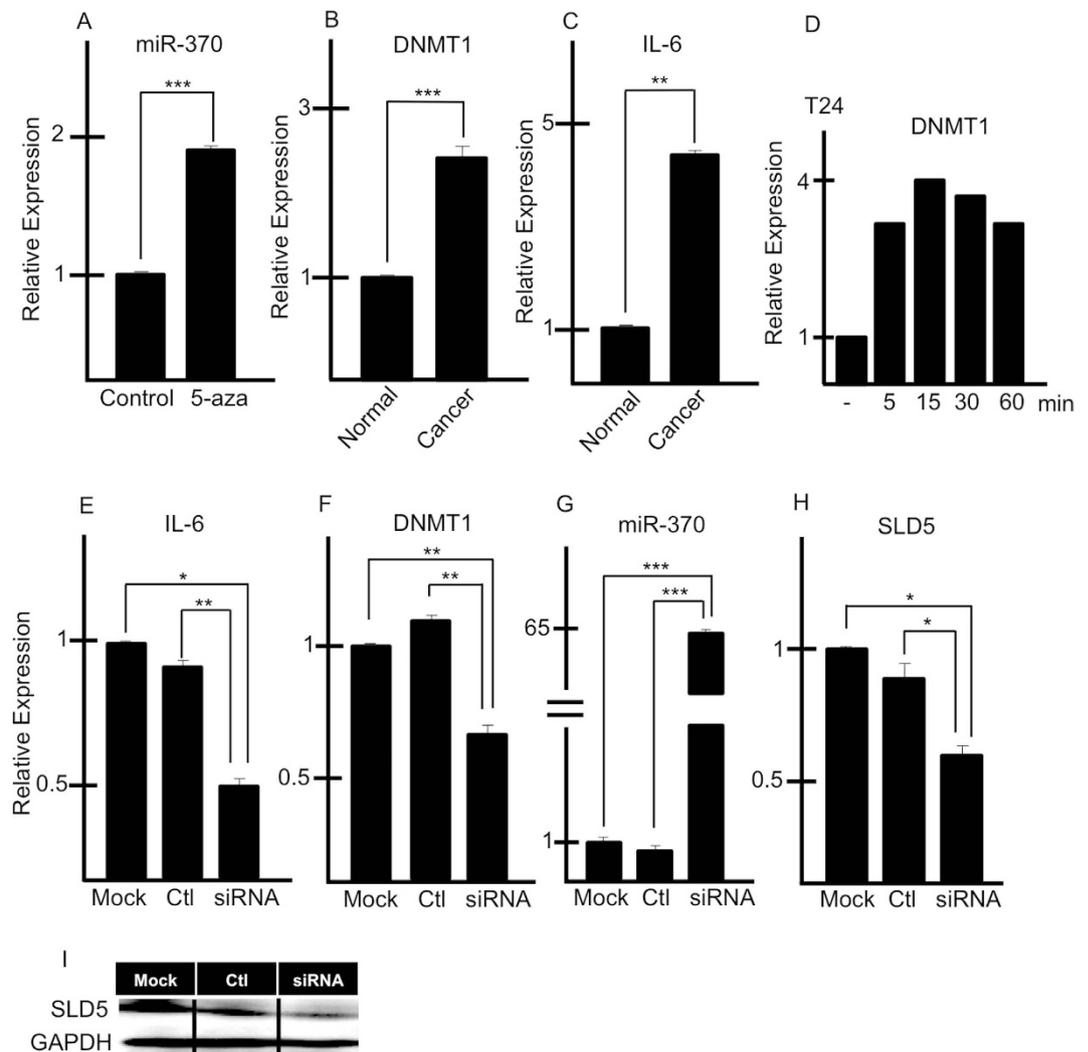
**Figure 4.** *SLD5* mRNA expression is regulated by miR-370. (A) Seed sequence of miR-370 (blue) located in the 3'-UTR of *SLD5*. (B) Expression level of miR-370 in human bladder cancer cells compared to HUVECs (Normal). Data are mean  $\pm$  SE  $^{**}p < 0.001$  ( $n = 3$ ). The normal cell was set at unity. *U6* is an internal control. (C) Luciferase reporter assay. T24 cells were transfected with pMIR luciferase reporter containing 3'-UTR sequences of *SLD5* and miR-370. miR-214 was used as a control. Graph depicts relative luciferase luminescence activity. Control was set at unity. Results are mean  $\pm$  SE,  $^{**}p < 0.005$  ( $n = 3$ ). (D) Attenuation of *SLD5* expression by miR-370 transfection in T24 cells. The level of miR-370 (left) and *SLD5* (right) after transfection of each miR was confirmed by qRT-PCR. Values for mock control cells were set at unity. *U6* or *GAPDH* was used as an internal control. Data are mean  $\pm$  SE  $^{*}p < 0.05$ ,  $^{**}p < 0.001$ ,  $^{***}p < 0.0001$ . (E) Western blotting showing *SLD5* expression in miR-370 mimic-transfected T24 cells. (F) Cell growth (T24) affected by miR-370 mimic transfection. Data are mean  $\pm$  SE  $^{*}p < 0.05$ ,  $^{**}p < 0.001$ . (G) Ki-67 (red) and (H) EdU (green)-positive cells in miRNA-transfected T24 cells. Graphs show quantitative evaluation of Ki-67 or EdU-positive cells. Data are mean  $\pm$  SE,  $^{*}p < 0.05$ ,  $^{**}p < 0.005$  (3 random fields). Scale bars: 100  $\mu$ m. (I) Cell cycle analysis using flow cytometry. Cells were analyzed after transfection with control or miR-370 mimic. The right-hand graph depicts a quantitative evaluation by showing mean  $\pm$  SE ( $n = 3$ ).



**Figure 5. Tumor growth is affected by miR-370.** (A,B) Bladder cancer cells (T24) were subcutaneously transplanted into nude mice. miR-370 mimics or control miR were injected on day10 post-inoculation. (A) Time course of tumor volume dynamics. Data are mean  $\pm$  SE (n = 3). (B) Tumor weight on day 30 post-inoculation. \*p < 0.05 mean  $\pm$  SE (C) qRT-PCR analysis for assessment of miR-370 or *SLD5* mRNA expression in tumors injected with control or miR-370. Data are mean  $\pm$  SE, \*\*p < 0.0005, \*\*\*p < 0.0001 (n = 3). (D) Xenografted tumors injected with control or miR-370 mimics were stained with anti-*SLD5* antibody (brown). Sections were counterstained with hematoxylin. Scale bars: 100  $\mu$ m.

**Regulation of miR-370 in human bladder cancer cells.** It has been reported that the expression of miRNAs is regulated by chromosomal modifications<sup>25</sup>. We analyzed whether miR-370 expression is regulated by DNA methylation by using 5-azacytidine (5-aza), an inhibitor of DNA methylation. After 24 hours of treatment with 5-aza, miR-370 expression was more strongly induced in cancer cells than in control cells (Fig. 6A), suggesting that its expression is suppressed in cancer cells by DNA methylation.

Next, we analyzed DNA methylation-related gene expression by qRT-PCR (Fig. 6B). Of the three DNA methyltransferases (DNMTs) in mammals, i.e., DNMT1, DNMT3a, and DNMT3b, DNMT1 is the most abundant in adult cells<sup>26</sup>. We found that T24 bladder cancer cells express two-fold higher levels of *DNMT1* than normal cells (Fig. 6B). It has been reported that IL-6 expression correlates with DNMT1 expression<sup>27</sup>. Accordingly, we found that *IL-6* expression is also upregulated in bladder cancer cells (Fig. 6C). When T24 cells were stimulated with IL-6, *DNMT1* expression was induced (Fig. 6D), suggesting that an autocrine loop of IL-6 in T24 cells upregulates *DNMT1*. Consistent with this, knocking down *IL-6* using siRNA resulted in reduced *DNMT1* expression (Fig. 6E,F) and concurrent upregulation of miR-370 (Fig. 6G) and reduction of *SLD5* (Fig. 6H). We confirmed that *SLD5* expression was suppressed not only at the RNA but also at the protein level (Fig. 6I). Therefore, we



**Figure 6. Regulation of miR-370 expression in tumors.** (A) miR-370 expression following 5-azacitidine treatment of T24 cells. Results are mean  $\pm$  SE \*\*\* $p$  < 0.0005 ( $n$  = 3). (B,C) qRT-PCR analysis of *DNMT1* (B) and *IL-6* (C) mRNA expression in normal bladder cells (Normal) and a cancer cell line (T24; Cancer). Data are mean  $\pm$  SE, \*\* $p$  < 0.0005, \*\*\* $p$  < 0.00001 ( $n$  = 3). (D) qRT-PCR analysis of *DNMT1* mRNA expression in T24 cells after treatment with IL-6. (E–H) qRT-PCR analysis of *IL-6* (E), *DNMT1* (F), miR-370 (G) and *SLD5* (H) expression in T24 cells 48 hours after transfection of *IL-6*-specific siRNA or control siRNA (ctl). Data are mean  $\pm$  SE, \* $p$  < 0.05, \*\* $p$  < 0.005, \*\*\* $p$  < 0.00001 ( $n$  = 3). (I) Western blotting for detection of *SLD5* in cells as described in (E).

conclude that in bladder cancer cells, up-regulation of IL-6 induces *SLD5* expression to promote acute proliferation of cancer cells.

## Discussion

In this study, we document pivotal roles of *SLD5* in acute proliferation of human bladder cancer cells. In normal cells, *SLD5* expression is blocked by miR-370 expression. However, in cancer cells, IL-6 induces *DNMT1* to inhibit miR-370 expression. Because miR-370 suppresses *SLD5*, in cancer cells *SLD5* expression is restored and cells can proliferate. *SLD5* is a member of the GINS complex, acting on DNA replication<sup>1,18</sup>. We previously reported that *SLD5* is essential for early embryogenesis and that *SLD5*-knockout mice were embryonic lethal because of impaired proliferation of cells in the inner cell mass<sup>7</sup>. In terms of *SLD5* expression in tumors, a few papers have indicated its overexpression, including in breast cancer<sup>3</sup> and colorectal cancer<sup>28</sup>. Here, we report the overexpression of *SLD5* in bladder cancer cells. We analyzed the expression of *SLD5* and its relationship to prognosis of bladder cancer patients ( $n$  = 165) by using public database Prognoscan (<http://www.abren.net/Prognoscan/>). Overall survival rate in that research indicated that *SLD5* expression was significantly associated with poor survival rate ( $p$  = 0.013). Therefore, our basic research contains clinical implication.

In S phase, cell division cycle (cdc) 45, mini chromosome maintenance protein (MCM) 2–7 and GINS complex cooperate as a replicative helicase CMG complex<sup>29</sup>. Especially, GINS is essential for the maintenance of interaction cdc45 and MCM2-7. Because GINS contributes to activity of CMG complex, increasing expression

level of SLD5, a member of GINS complex might leads to aberrant proliferation in cancer cells. Moreover, tumor suppressor gene, Rb1, works as an S phase transition inhibitor is often mutated in bladder cancer patients<sup>30</sup>. Because of this reason, it is possible that Rb1 mutation also leads to SLD5 overexpression in bladder cancer cells.

Regarding other members of the GINS complex, PSF1 was also overexpressed along with SLD5 in bladder cancer cells, but PSF2 and PSF3 levels were similar to those in normal cells such as HUVECs (data not shown). This expression profile in bladder cancer cells is similar to that reported in breast cancer cells<sup>3</sup>. As it has been suggested that psf2 and psf3 act as multicopy suppressors of sld5 and psf1 in budding yeast<sup>1</sup>, our results suggest that SLD5 is critical for proliferation of cancer cells, and therefore that PSF2 and PSF3 overexpression in cancer cells may not be required.

According to our results, expression of SLD5 is significantly upregulated both in human bladder cancer tissue samples as well as in cell lines. Moreover, decreased cell numbers, reduced fractions of cells in S phase, and suppressed tumor growth were observed following silencing of *SLD5*. Furthermore, we identified miR-370 as a negative regulator of *SLD5* gene expression. Many studies showing associations of miR with tumorigenesis have been reported in bladder cancer. In bladder cancer cells, miR-101, -203, and -204<sup>22–24</sup> are downregulated, resulting in induction of aberrant increases of proliferative capacity or cell migration. Although several lines of evidence indicate downregulation of miR-370 in cancer cells<sup>23,31,32</sup>, the relationship between miR-370 and SLD5 has not been reported previously. Based on our results, we hypothesize that miR-370 acts as a negative regulator of SLD5 expression in normal states. In contrast, in bladder cancer cells, miR-370 is downregulated, and SLD5 expression levels are increased to induce proliferation of cancer cells. In Fig. 4, overexpression of miR-370 might affect not only cancer cells but other cells consisting cancer microenvironment. Because endothelial cells, blood cells (data not shown) and fibroblastic cells<sup>33</sup> express GINS complex gene, reduced SLD5 expression by the injection of miR-370 caused attenuation of cellular activity of those cells and cancer cell proliferation and survival could not be supported by such stromal cells.

We found that attenuated miR-370 expression in cancer cells is partly mediated by methylation of the miR-370 promoter, because treatment with 5-aza-2-deoxycytidine (5-aza), a specific inhibitor of DNA methylation<sup>34</sup>, induced the expression of this miRNA. Therefore, bladder cancer cells may have intrinsic signaling pathways for methylating the miR-370 promoter region. So far, many studies have focused on IL-6 as a player in malignant tumor progression by facilitating epithelial-mesenchymal transition (EMT), inflammation, and angiogenesis. Moreover, it has been reported that IL-6 is involved in DNMT1 expression<sup>27,35,36</sup>. Here, we also found that IL-6 and DNMT1 expression were correlated in bladder cancer. It is possible that upregulated DNMT1 leads to hypermethylation of CpG islands in the promoter regions of many genes in bladder cancer cells<sup>37</sup>. In addition to DNMT1 expression, we found that the endogenous demethylation factor, ten-eleven translocation family methylcytosine hydroxylase (TET1)<sup>38</sup> was markedly lower in cancer cells (Supplementary Fig. S2). These results suggest that hypermethylation caused by IL-6 induces suppression of many genes including miR-370, contributing to the upregulation of SLD5. It has also been suggested that miR-370 is normally imprinted and activated only on the maternal allele<sup>35</sup>, so it may be predicted that miR-370 is downregulated even when CpG islands are only slightly methylated.

In summary, to the best of our knowledge, there are no published studies focused on the behavior of SLD5 in human cancer cells. The present study revealed a relationship between SLD5 and miR-370 in human bladder cancer. Although we focused on miR-370, it is possible that there are multiple additional miRNAs that are relevant for SLD5 expression *in vivo*. It has been reported that not only SLD5 but also other GINS members have the potential to regulate cancer proliferation and that these genes might be cancer therapeutic targets. Further studies on the regulation of SLD5 expression are required to develop this moiety as a drug target.

## Methods

**Cell lines and samples.** Human bladder cancer cell lines T24 and KMBC2 (JCRB, Osaka, Japan) were maintained in MEM medium (Sigma, St. Louis, MO) and Ham's F12 medium (GIBCO, Rockville, MD), respectively, with 10% fetal bovine serum (FBS; Sigma, St. Louis, MO) and penicillin/streptomycin (GIBCO, Rockville, MD). Human umbilical vein endothelial cells (HUVECs KURABO, Osaka, Japan) were maintained in HuMedia EG2 (KURABO, Osaka, Japan). For qRT-PCR analysis, normal human bladder RNA sample was purchased from US Biomax (Rockville, MD).

**Immunohistochemistry.** Human carcinoma tissue array specimens and bladder cancer tissue specimens including normal human bladder were purchased from Biochain (Beijing, China) and US Biomax (Rockville, MD), respectively. For immunohistochemical analysis, after deparaffinization, sections were activated with citrate buffer solution. Blocking with skimmed milk in PBST (0.1% TritonX-100 in PBS) for 1 h. Anti-*SLD5* antibody (1:200 Iwaki, Japan) was used as the primary antibody with biotin-conjugated goat anti-rat IgG (1:200, Jackson ImmunoResearch, West Grove, PA) as the secondary antibody. Avidin-biotin complex (ABC) kits (Vector Laboratories, Burlingame, CA) and 3,3'-diaminobenzidine (DAB; Dojindo, Japan) and nickel chloride (NiCl<sub>2</sub>, Wako, Japan) were used for visualization. For staining of nuclear DNA, hematoxylin (Wako) was applied in counterstaining.

For the fluorescent immunohistochemical analysis, rat anti-*SLD5* antibody (1:200; Iwaki), rabbit anti-*SLD5* (1:200 Abcam, Cambridge, UK), rat anti-CD44-PE (1:100 eBioscience), and mouse anti-Ki-67 (1:400 Dako, Carpinteria, CA) were used as primary antibodies and donkey-anti-rat IgG Alexa Fluor 488 (1:200, Jackson ImmunoResearch) and goat anti-mouse IgG Alexa Fluor 546 (1:200, Jackson ImmunoResearch) as secondary antibodies. Nuclear DNA was counterstained with Hoechst 33342 (1:3000, Sigma) or DAPI (1:3000 Dojindo). Stained samples were assessed under a microscope (CTR5500, Leica, Wetzlar, Germany). Images were analyzed using Image J software (NIH).

**Immunocytochemistry.** Cells were fixed with 4% paraformaldehyde for 10 min and blocked with skimmed milk in PBST (0.1% Triton X-100 in PBS) for 1 h. Cells were then incubated with rabbit anti-SLD5 (1:200) or mouse anti-Ki-67 (1:200) overnight at 4 °C, washed with PBS, and then incubated with donkey anti-rabbit Alexa Fluor 488 (1:200, Jackson ImmunoResearch) or goat anti-mouse Alexa Fluor 546 (1:200, Jackson ImmunoResearch) for 1 h at room temperature. Nuclear DNA was counterstained with DAPI (1:3000 Dojindo). Stained samples were assessed under a microscope and images were analyzed using ImageJ software.

**Quantitative Reverse Transcription-PCR.** Total RNA was isolated from cells or FFPE sections using the RNeasy plus mini kit (Qiagen, Chatsworth, CA) and the Nucleospin total RNA FFPE (MACHEREY-NAGEL, Germany), respectively, according to the manufacturer's protocol. Total RNAs were reverse transcribed using the PrimeScript RT reagent kit (Takara bio, Shiga, Japan). Quantitative RT-PCR (qRT-PCR) was performed using SYBR Premix Ex Taq II (Takara) on Mx3000P (Stratagene, La Jolla, CA) as previously described<sup>39</sup>. Baseline and threshold were adjusted according to the manufacturer's protocol. Levels of the specific amplified genes were normalized to the level of GAPDH. All primers are listed in Supplementary Table 1 and were purchased from GeneDesign, Inc (Osaka, Japan).

For detection of miRNA expression, total and small RNAs were extracted using NucleoSpin miRNA kits (Takara clontech) followed by small RNA reverse transcription using Mir-X miRNA First-Strand Synthesis Kits (Takara clontech). The qRT-PCR reaction was performed according to the manufacturer's protocol using specific primers (Supplementary Table 1). The  $2^{-\Delta\Delta Ct}$  method was used to calculate results and amplified genes were normalized to the level of U6.

**RNA interference.** SLD5 and IL-6 expression in human bladder cancer cell lines was transiently knocked down with small interfering RNA (siRNA) (Invitrogen, Carlsbad, CA or Thermofisher Scientific Inc, Massachusetts, respectively). Lipofectamine 2000 (Invitrogen) was used for the transfection of siRNA, following the manufacturer's instructions. We used two different siRNA oligonucleotides for SLD5 and scrambled siRNA (Invitrogen). The Pre-hsa-miR-370 mimic (GeneDesign, Osaka, Japan) was transfected into T24 bladder cancer cells following the manufacturer's instructions.

**Western blotting.** Total cells were lysed using radioimmunoprecipitation assay (RIPA) lysis buffer. Normal human bladder whole cell lysate was purchased from Abcam. Proteins were heated for 3 min at 95 °C, loaded onto SDS-polyacrylamide gels and then transferred onto polyvinylidene difluoride membranes (Millipore, Jaffery, NH). Blocking was with 3% BSA prior to staining with anti-SLD5 antibody (1:500 Abcam) or anti-GAPDH (1:3000; Millipore) as primary antibodies overnight at 4 °C. Blots were developed with anti-rabbit HRP and anti-mouse HRP antibodies (Jackson ImmunoResearch) using enhanced chemiluminescence (Pierce Biotechnology Inc, Rockford). Images were analyzed with ImageJ software.

**Measurement of doubling time.** To calculate cell growth rate, bladder cancer cell (T24) and normal cells (HUVEC) were cultured for 6, 12, 24 or 48 hours. Cells were counted and numbers were calculated using the following formula:  $\text{duration} \times \log 2 / \log (\text{final cell number}) - \log (\text{initial cell number})$ .

**Flow cytometry and EdU incorporation assay.** To analyze DNA content, cells were fixed in 70% ethanol and suspended in propidium iodide (PI)/RNase/Triton X-100 staining solution, and then analyzed by FACS caliber (BD Biosciences). Cells were plated into 24-well plates and transfected with siRNA and miRNA. After transfection, EdU incorporated into cells was visualized using Click-iT EdU Alexa Fluor 488 imaging kits (Invitrogen, Carlsbad, CA, USA), according to the manufacturer's instructions.

**Analysis of xenografted tumor with siRNA and miRNA *in vivo* transfection.**  $2 \times 10^6$  T24 cells suspended in 100  $\mu$ l of phosphate-buffered saline (PBS) were subcutaneously injected into the dorsal surfaces of 6-week-old female KSN/Slc nude mice (Slc, Shizuoka, Japan). Two weeks after transplantation, the developed tumors were treated with control scrambled RNA, SLD5 siRNA, or miR-370 mimic suspended in atelocollagen (Atelogene, KOKEN, Tokyo, Japan) to facilitate siRNA introduction into tumors, according to the manufacturer's instructions. Twenty days after atelocollagen injection, xenograft tumor volumes and weights were measured, RNA was isolated from tumors for the analysis of SLD5 or miR-370 expression by qRT-PCR, and tumor tissues were fixed in 4% paraformaldehyde for immunohistochemistry analysis. Tumor volumes were calculated using the following formula:  $(\text{major axis}) \times (\text{minor axis})^2 \times 0.5$ . All animal experiments were approved by the Animal Research Committee of Osaka University. All experiments were carried out under the guidelines of Osaka University Committee for animal and recombinant DNA experiments.

**Luciferase reporter assay.** The 3' UTR region of SLD5, containing miR-370 target sites, was amplified using a standard PCR protocol with the following linker primers: 5'-ATT ACT ACT AGT GCA TAA ACA GCC AGG CAT GGT GAC-3' (forward), 5'-AAC CAT AAG CTT TAG TAG AGA TGG GTT TAG TAG AG-3' (reverse). cDNA was inserted into SpeI and HindII sites of pMIR-REPORT Luciferase vectors (Applied Biosystems, Carlsbad). T24 bladder cancer cells were plated in 96-well plates and transfected with or without miR-370 mimic and luciferase vector. After transfection, luciferase reporter activity in the cells was measured using the Dual-Glo Luciferase Assay System (Promega, Madison, WI). We used the pRL-TK vector (Promega) for normalized values of reporter activity, and a miR-214 mimic was used as a negative control. All transfections were performed using Lipofectamine 2000 (Invitrogen) according to the manufacturer's instructions.

**DNA demethylation.** T24 cells were treated with 1  $\mu$ M 5-aza-2-deoxycytidine (Sigma Aldrich). After 24 h, miRNA was isolated and cDNA synthesized for qRT-PCR analysis of miR-370. Levels of the specific amplified genes were normalized to the level of U6.

**hIL-6 treatment in a normal cell line (HUVEC) and a cancer cell line (T24).** Normal and cancer cell lines were treated with 100 ng/ml human recombinant IL-6 (Biolegend, San Diego, CA). RNA from recovered cells after treatment was isolated for qRT-PCR analysis of DNMT1. Levels of the specific amplified genes were normalized to the level of GAPDH.

**Statistical analysis.** Results are expressed as the mean  $\pm$  standard error (SE). Student's t-test was used for statistical analysis. Differences were considered statistically significant if the p-value was less than 0.05.

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## Author Contributions

K.Y. performed most of the experiments and wrote the article; H.N., T.W., H.Y., F.M., T.I. and H.K. supported experiments and discussed the project; N.T. conceived the project and wrote the article.

## Additional Information

**Supplementary information** accompanies this paper at <http://www.nature.com/srep>

**Competing financial interests:** N.T. is an employee of GeneStem Co., Ltd. which produces anti-SLD5 antibody on a commercial basis.

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