INTRODUCTION

Chromosome instability (CIN) plays a role in the multistep oncogenesis of cancer cells. CIN involves the unequal distribution of chromosomes to daughter cells upon mitosis, resulting in the loss or gain of chromosomes and/or chromosome rearrangements during cell division and, ultimately, aneuploidy. Therefore, cancer cells are characterized by aneuploidy and CIN.1 Although CIN can be caused by multiple failures, centromere dysfunction is one of main reasons. Several studies have demonstrated a relationship between CIN and centromere dysfunction in cancer cell lines.2-6 Overexpression of CENP-A, which is a key determinant protein for centromere formation,7 has been found in several human malignancies, including hepatocellular carcinoma,8,9 breast cancer10 and ovarian cancer.11 CENP-A expression predicts poor outcomes in patients with these cancers.3

Centromere proteins are classified into several functional groups as follows: CENP-C, CENP-H/I/K/M, CENP-L/M, CENP-O/P/Q/R/U, CENP-T/W and CENP-S/X.5,12-15 In the present study, among centromere proteins, we focused on CENP-U (CENP-50/U), which forms a complex with other CENP-O class proteins (Cenp-OO/P/Q/R/U).16 CENP-50/U localizes to the centromere throughout the cell cycle and has been described as KLIP1/MLF1IP.17-19
As the centromere is essential for chromosome segregation during cell division, disruption of most centromere proteins in mice can cause embryonic lethality, accompanied by enlarged nuclei containing an increased number of nucleoli, nuclear bridging, chromosome condensation and spindle fiber bundling. For example, Cenpc null mutation results in embryonic lethality at 3.5 d, with a chromosome missegregation phenotype and mitotic arrest.20,21 Cenpd null mice fail to survive beyond 6.5 d post-conception, whereas heterozygous knockout mice are healthy and fertile.22 Therefore, it is difficult to generate centromere protein-deficient mice. As this lethality is caused by a chromosome segregation defect, cultured cells with knockout of centromere proteins should be lethal. However, we demonstrated that chicken DT40 cells with knockout of each CENP-O complex protein were viable,23,24 (Table S1), suggesting that cells with knockout of the CENP-O complex proteins were good models to study CIN. Next, we attempted to generate centromere protein-deficient mice and demonstrated that Cenpr knockout mice were viable,25 (Table S1). Conversely, Cenp50/U-deficient mouse ES cells died after they exhibited abnormal mitotic behavior, unlike the chicken DT40 cells.26 In addition, Cenp50/U-deficient mice died during early embryogenesis (at approximately E7.5),27 (Table S1). Although these results were unexpected, the phenotype of cells with knockout of CENP-O complex proteins was weak compared with cells with other centromere proteins knocked out. Therefore, in this study, we generated Cenp50/U conditional knockout mice and characterized their function related to carcinogenesis.

We have demonstrated previously the essential role of Cenpr, which forms a complex with other Cenp-O proteins (Cenp-O/P/Q/R/U), in the two-stage skin carcinogenesis model. Cenpr functions as a tumor suppressor in early papilloma development, but as an oncogene at the late papilloma stage and in malignant conversion.28 Based on this study, we hypothesized that other Cenp-O complex proteins function in skin carcinogenesis. We therefore generated K14(Keratin 14)CreER-Cenp50/Ufl/fl mice to investigate the function of Cenp50/U in vivo, especially in the process of carcinogenesis.

It has been shown previously that the pathology of the two-stage chemically induced skin carcinogenesis mouse model is almost identical to the development of human skin cancers, and therefore it is an ideal model to study skin cancer initiation and growth.25,26 In the first step of this chemically induced carcinogenesis protocol, mice were treated with a low dose of the mutagen 7,12-dimethylbenz(a)anthracene (DMBA) to start tumor development. This first chemical treatment step led to “tumor initiation.” In the second step, mice were treated continuously with TPA to stimulate epidermal tumor proliferation. This second chemical treatment step influenced “tumor promotion.” During tumor promotion, benign tumors, known as papillomas, are thought to develop from additional mutations caused by TPA chemical treatment. After prolonged treatment (~20 wk), some papillomas will progress into carcinogenic tumors such as squamous cell carcinomas (SCC). The roles of numerous genes and cell-signaling pathways in skin tumor development could be explored with this two-stage skin carcinogenesis model using genetically engineered mouse models.27-33 In this study, we demonstrated the function of Cenp50/U in the epidermis and its role in two-stage skin carcinogenesis. We clarified the effects of Cenp50/U deficiency on mouse normal epidermis and its oncogenic role in papillomagenesis.

2 | MATERIALS AND METHODS

2.1 | Mice

This study was carried out in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the Ministry of Education, Culture, Sports, Science, and Technology of Japan. The protocol was approved by the Committee on the Ethics of Animal Experiments of Chiba Cancer Center (Permit Number: 13-18). All efforts were made to minimize suffering. C57BL/6 strain mice were used as recipients for targeted ES cells and as the background strain in this study. The generation of mice carrying the floxed allele of the Cenp50/U gene (Cenp50/Ufl/fl) is described elsewhere.24 Cenp50/Ufl/fl mice were mated to K14-CreER transgenic mice.38 K14-CreER transgenic mice were obtained from the Jackson Laboratory (JaX 005107). To disrupt Cenp50/U, K14CreER-Cenp50/Ufl/fl and Cenp50/Ufl/fl mice at 8 wk old were injected subcutaneously 5 times with 1.0 mg/mL of tamoxifen (Sigma-Aldrich) that had been prepared following the manufacturer’s instructions by completely dissolving tamoxifen into 1 mL of 100% ethanol at 50°C, adding 9 mL of sunflower oil and mixing this thoroughly by vortexing.

2.2 | Skin carcinogenesis

7,12-Dimethylbenz(a)anthracene (DMBA) was purchased from Sigma-Aldrich, Merck Millipore and TPA was purchased from Calbiochem, Merck Millipore. DMBA is used as a carcinogen and TPA as a promoter. At 8-10 wk of age, the backs of the mice were carefully shaved with electric clippers. At 2 d after shaving, DMBA (25 µg per mouse in 200 µL of acetone) was applied to shaved dorsal back skin. Then, 3 d after the first DMBA treatment, TPA (10 µg per mouse in 200 µL of acetone) was applied. After 4 rounds of this single DMBA and TPA treatment, the mice were treated with TPA twice weekly for 20 wk. The papilloma number and size (mm in diameter) were recorded from 8 wk up to 20 wk, and carcinoma development was monitored up to 34 wk.

2.3 | RT-PCR and quantitative real-time RT-PCR

Total RNA was isolated from normal skins, papillomas, carcinomas and metastatic carcinomas of mice after 40 wk of DMBA/TPA treatment using TRizol (Invitrogen) in accordance with the manufacturer’s protocol. cDNA was generated with the iScript™ Select cDNA Synthesis Kit (Bio-Rad) using 100 ng of DNase-pretreated total RNA. RT-PCR was performed using the Prime Taq...
DNA Polymerase Kit (Genet Bio). The products were subjected to agarose gel electrophoresis. Gapdh was used as a control. qRT-PCR was performed using GeneAce SYBR® qPCR Mix α Low ROX following the manufacturer's protocol (NIPPON GENE). mRNA expression levels were measured using the Applied Biosystems® 7500 system (Life Technologies) and normalized to the levels of Gapdh. The following primers were used to generate Cenp-50/U: Forward Primer; TCT ACG CAG AAG ACG AGC TG and Reverse Primer; GTT GCT GTG ACC TTG GTC CT, Cenp-ε: Forward Primer; CAC AGA AAG GCA CCA TCA AA and Reverse Primer; AGT TGT CTG TTT CCC TCC AA, Cenp-ο: Forward Primer; AGA ACT GCG GCA ACA ACG and Reverse Primer; ACG ACT GTG CAA CTG CCC. Cenp-ρ: Forward Primer; ATC AGG AAA TTG CAA CAT GG and Reverse Primer; TTC GGA AAA ACA TGA GCA AA, Cenp-q: Forward Primer; GGA TTT GCC AGA GAA TGA GG and Reverse Primer; TTC GGA AAA ACA TGA GCA AA, Cenp-r: Forward Primer; GAG GTT GCC AGG TTG TTC TC, and Cenp-s: Forward Primer; ATC AGG AAA TTG CAA CAT GG and Reverse Primer; TTC GGA AAA ACA TGA GCA AA.

2.4 | Cell cycle analysis

Mouse keratinocyte cells were isolated from Cenp-50/Ufl/fl and K14CerER²R, Cenp-50/Ufl/fl mice. Isolation of keratinocyte cells from mice was performed as described previously.²⁹ These were then collected and fixed in 70% ethanol at 4°C for 30 min. The fixed cells were stained with propidium iodide (50 μg/mL) containing 200 μg of RNase A/mL and 1% Triton X-100 at 37°C for 40 min. Flow cytometry analysis was conducted using a JSAN instrument (Japan-made sorter, analyzer) (Bay Bioscience). Approximately 1.0 × 10⁵ cells were scanned to analyze the DNA content. Necrotic cells were excluded, and the percentages of cells in the G₁, S and G₂/M phases of the cell cycle were measured.

2.5 | Immunoblotting

Proteins were extracted from different cells using T-PER Protein Extraction Reagent (Thermo). Protein concentrations were quantified by the Quick Start Bradford Protein Assay (Bio-Rad). Denatured proteins (Mouse normal skins: 60 μg) were then analyzed using 15% e-PAGEs (ATTO). After electrophoresis, these were transferred onto a polyvinylidene difluoride (PVDF) membrane (Merck Millipore). The membrane was blocked with 0.5% skimmed milk or 1% BSA in phosphate-buffered saline solution (pH 7.6) containing 0.1% Tween-20 (PBS/T) and then the SNAPi.d. 2.0 Protein Detection system (Merck Millipore). Primary antibodies were used as follows: anti-Cenp-50/U (anti-rabbit polyclonal),¹⁷ anti-Ac-tin (Sigma-Merck Millipore), HRP-conjugated secondary antibodies were used at a dilution of 1:2000 and developed using the ECL Prime Western Blotting Detection Kit (GE Healthcare). Exposure for chemiluminescent samples or membrane analysis for the blots was performed with a LAS 4000 system (GE Healthcare).

2.6 | Immunofluorescence

Samples were fixed with 4% paraformaldehyde at 4°C overnight. The endogenous peroxidase activity in the specimens was blocked by treatment with 0.3% H₂O₂ and samples were then rinsed with PBS. Sections were incubated with primary antibodies against anti-Ki67 (1:100 16A8, Biolegend) and anti-keratin 14 (1:500 Poly19053, Biolegend) diluted in blocking buffer overnight at 4°C. Secondary antibodies were Alexa Fluor 488-conjugated anti-rat antibody (1:100, Molecular Probes, Invitrogen) and Alexa Fluor 568-conjugated anti-rabbit antibody (1:100, Molecular Probes, Invitrogen). Nuclei were counterstained with Hard Set Mounting Medium with DAPI (Vector). All fluorescence images were obtained on a Leica TCS SPE confocal microscope equipped with a DMI400B (×10/0.40, ×20/0.70 and ×40/1.25 oil immersion objectives).

2.7 | TUNEL staining

Apoptotic scores were obtained by terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay using an in situ Apoptosis Detection Kit (TaKaRa). Samples were digested with proteinase K (20 μg/mL) and labeled with TUNEL reaction mixture for 90 min at 37°C. All images were obtained on a Leica TCS SPE confocal microscope equipped with a DMI400B (×10/0.40, ×20/0.70 and ×40/1.25 oil immersion objectives). Data from all fields and all samples were pooled to obtain the apoptotic index, which was the percentage of TUNEL-positive cells of total cells manually counted in randomly selected fields, and compared between Cenp-50/Ufl/fl and K14CerER²R, Cenp-50/Ufl/fl mice.

2.8 | Statistical analysis

At least 3 replicates were performed for all experiments. The significance of differences was calculated using unpaired one- and two-tailed Student t tests or two-way ANOVA. A P-value < .05 was considered significant and a value < .01 was highly significant. These analyses were performed using GraphPad Prism software (GraphPad).

3 | RESULTS

3.1 | mRNA expression profiles of the Cenp-O complex are altered in skin tumors

First, we examined mRNA expression levels of Cenp-O complex (Cenp-O/P/Q/R/U) genes using DMBA/TPA-induced papillomas (n = 10) and carcinomas (n = 10). As a result, Cenp-50/U and Cenp-ρ expression levels differed between papillomas and carcinomas (Figure 1A,B). Conversely, Cenp-α, Cenp-ρ and Cenp-δ had the
same expression levels in papillomas and carcinomas (Figure 1C-E). These results suggested that Cenp-50/U and Cenp-r are more likely to be involved in skin carcinogenesis than Cenp-o, Cenp-p and Cenp-q.

Cenp-r expression levels in carcinomas were significantly higher than in papillomas (Figure 1B). In contrast, Cenp-50/U expression levels in carcinomas were significantly lower than in papillomas (Figure 1A). In other words, Cenp-50/U and Cenp-r exhibited opposite expression profiles in skin carcinogenesis. Our previous study revealed that Cenp-r functions as a tumor suppressor-type gene in papilloma development and as an oncogene-type gene during the progression stage.23 These results suggested that Cenp-50/U plays a opposing role to Cenp-r in skin carcinogenesis.

3.2 | Generation of Cenp-50/U conditional knockout mice

A previous study demonstrated that Cenp-50/U knockout mice die during early embryogenesis (approximately E7.5).24 To investigate the function of Cenp-50/U in vivo, we generated mice harboring conditional alleles of Cenp-50/U, in which exons 4-6 of the Cenp-50/U gene are flanked by loxP sites (Figure 2A). First, we isolated mice with Cenp-50/U fl/+ alleles and finally generated Cenp-50/U fl/fl alleles by intercrossing (Figure 2B). Next, we crossed the Cenp-50/U fl/fl conditional knockout mice with mice carrying a K14CreER allele,34 which is specifically expressed in the epidermis, to generate K14CreER,Cenp-50/U fl/fl progeny in which Cenp-50/U can be rendered non-functional in the skin upon induction with tamoxifen.
Five subcutaneous injections of tamoxifen into K14CreER-Cenp-50/Ufl/fl mice was sufficient to disrupt the floxed Cenp-50/U locus in the dorsal back skin and tail (Figure 2C). In addition, we examined Cenp-50/U expression levels in K14CreER-Cenp-50/Ufl/fl and Cenp-50/Ufl/fl mice after subcutaneous injection with tamoxifen. As a result, mRNA expression and protein of Cenp-50/U was not detected in K14CreER-Cenp-50/Ufl/fl mouse skins by RT-PCR and western blot analyses with anti-Cenp-50/U antibody, respectively (Figure 2D,E).

3.3 | Cenp-50/U deficiency affects cell proliferation in normal skins

As we successfully generated epidermis-specific Cenp-50/U knockout mice, we investigated the function of Cenp-50/U in the epidermis using these mice. First, we performed histological analysis using skins from K14CreER-Cenp-50/Ufl/fl and Cenp-50/Ufl/fl mice after subcutaneous injection with tamoxifen. Hematoxylin-eosin (HE) staining revealed no significant morphological changes (Figure S1A,B). We then carried out immunohistochemical analysis with the cell proliferation marker Ki67 and TUNEL staining to investigate apoptotic cells in skins. As a result, skins from K14CreER-Cenp-50/Ufl/fl mice had a significantly lower number of Ki67-positive cells than skins from Cenp-50/Ufl/fl mice (Figure 3A,B,E). Conversely, TUNEL-positive cells were detected in K14CreER-Cenp-50/Ufl/fl mouse skins, although they were not detected in Cenp-50/Ufl/fl mouse skins (Figure 3C,D,F). However, only a few TUNEL-positive cells per field were detected in K14CreER-Cenp-50/Ufl/fl mouse skins. To confirm these results, we performed cell cycle analysis. The percentage of cells in the G0/G1 phase was significantly lower in skins of K14CreER-Cenp-50/Ufl/fl mice than in those of Cenp-50/Ufl/fl mice (Figure 3G). In addition, the percentage of cells in the subG1 phase was significantly higher in skins of K14CreER-Cenp-50/Ufl/fl mice than in those of Cenp-50/Ufl/fl mice (Figure 3G). However, we did not observe accumulation of G2/M phase in skins of K14CreER-Cenp-50/Ufl/fl mice, suggesting that cells went through mitosis, even if subtle centromere
**Figure 3** Cenp-50/U-deficient mice have a decrease in cell proliferation in normal skins. A, B, Immunostaining patterns of Ki67 (green) in skins from (A) Cenp-50/U^{fl/fl} and (B) K14Cre^{ER}-Cenp-50/U^{fl/fl} mice. C, D, Representative TUNEL staining pattern of TdT (green) in skins from (C) Cenp-50/U^{fl/fl} and (D) K14Cre^{ER}-Cenp-50/U^{fl/fl} mice. Cells were counterstained with Hoechst (red). E, The number of Ki67-positive cells in skins from Cenp-50/U^{fl/fl} (n = 12) (red bar) and K14Cre^{ER}-Cenp-50/U^{fl/fl} (n = 12) (blue bar) mice (P = 4.93E-8). F, The number of TUNEL-positive cells in skins from Cenp-50/U^{fl/fl} (n = 12) (red bar) and K14Cre^{ER}-Cenp-50/U^{fl/fl} (n = 12) (blue bar) mice (P = 4.93E-8). E, The number of Ki67-positive cells in skins from Cenp-50/U^{fl/fl} (n = 7) (red bar) and K14Cre^{ER}-Cenp-50/U^{fl/fl} (n = 7) (blue bar) mice. G, Cell cycle analysis of mouse normal skins from Cenp-50/U^{fl/fl} (n = 6) (red bar) and K14Cre^{ER}-Cenp-50/U^{fl/fl} (n = 5) (blue bar) mice (G_1/G_2 phase: P = .00424, subG_1 phase: P = .000203). DNA content was measured by propidium iodide (PI) staining. H, Trypan blue exclusion test of cell viability. Cells were isolated from Cenp-50/U^{fl/fl} (n = 6) (red bar) and K14Cre^{ER}-Cenp-50/U^{fl/fl} (n = 5) (blue bar) mice (P = .002013). The P-value was calculated by t test (**P < .01). n.s., not significant. n.d., not detected. Error bars represent the standard deviation (SD).
dysfunctions happened. We also performed the trypan blue exclusion test of cell viability using cells isolated from skins of K14Cre\textsuperscript{ER}-Cenp-50/U\textsuperscript{fl/fl} and Cenp-50/U\textsuperscript{fl/fl} mice. As a result, the percentage of dead cells was significantly higher in skins of K14Cre\textsuperscript{ER}-Cenp-50/U\textsuperscript{fl/fl} mice than in those of Cenp-50/U\textsuperscript{fl/fl} mice (Figure 3H). These results suggested that Cenp-50/U deficiency promotes cell death by apoptosis and reduces cell proliferation in mouse skins.

3.4 | Cenp-50/U deficiency partially inhibits papillomagenesis

Our previous study demonstrated the essential function of Cenp-r in the two-stage skin carcinogenesis model.\textsuperscript{23} In this study, our expression analysis suggested that Cenp-50/U functions might be opposite to those of Cenp-r in skin carcinogenesis (Figure 1). To investigate
the role of Cenp-50/U in skin carcinogenesis, we subjected K14CreER-
Cenp-50/Ufl/fl and Cenp-50/Ufl/fl mice to the DMBA/TPA chemical
carcinogenesis protocol, and papilloma development was monitored
for a period of 20 wk. As a result, K14CreER-Cenp-50/Ufl/fl mice ex-
hibited a significantly lower number of papillomas than Cenp-50/Ufl/fl
mice between 10 and 16 wk after initiation (Figure 4A,E, Table S2).
In addition, we classified these papillomas into 3 categories based on
size (<2, 2-6 and >6 mm). K14CreER-Cenp-50/Ufl/fl mice demonstrated
a significant decrease only in small papillomas < 2 mm in diameter
between 10 and 14 wk after initiation (Figure 4B-D, Table S2). These
results suggested that Cenp-50/U mainly functions in early papilloma
development, but has a weak effect on papilloma growth.

3.5 | Cenp-50/U deficiency increases apoptotic cells
in papillomas

Skin carcinogenesis experiments revealed that Cenp-50/U func-
tions in early papilloma development. Therefore, we investigated
cell proliferation and apoptosis in papillomas of K14CreER-Cenp-50/
Ufl/fl and Cenp-50/Ufl/fl mice. First, we performed histological analy-
sis using early-stage papillomas (<6 mm) from K14CreER-Cenp-50/
Ufl/fl and Cenp-50/Ufl/fl mice. As HE staining revealed no significant
morphological changes between these mice (Figure S1C,D), we car-
ried out immunohistochemical analysis with the cell proliferation
marker Ki67 and TUNEL staining to investigate apoptotic cells in
papillomas. Papillomas of K14CreER-Cenp-50/Ufl/fl mice had a signi-
ficantly higher number of TUNEL-positive cells than those of Cenp-50/
Ufl/fl mice (Figure 5C,D,F). Conversely, no significant difference be-
tween K14CreER-Cenp-50/Ufl/fl and Cenp-50/Ufl/fl mice was observed
by immunohistochemical analysis with the cell proliferation marker
Ki67 (Figure 5A,B,E). Therefore, Cenp-50/U deficiency promotes cell
death by inducing apoptotic cells rather than controlling cell pro-
liferation in papillomas. These finding are consistent with previous
reports, which stated that CENP-50/U-deficient DT40 cells undergo
apoptosis.17 Therefore, Cenp-50/U regulates abnormal mitosis both
in mouse and cell lines.

3.6 | Cenp-50/U is required for early
papilloma development, but has little effect on
malignant conversion

To investigate the effects of Cenp-50/U on malignant conversion,
we monitored carcinoma development up to 34 wk after initiation.
The carcinoma incidence was almost the same in K14CreER-Cenp-50/

FIGURE 5 | Cenp-50/U deficiency
induces cell death in papillomas. A, B,
Immunostaining patterns of Ki67 (green)
and K14 (red) in papillomas from (A) Cenp-
50/Ufl/fl and (B) K14CreER-Cenp-50/Ufl/fl
mice. C, D, Representative TUNEL staining
pattern of TdT (green) in papillomas
from (C) Cenp-50/Ufl/fl and (D) K14CreER-
Cenp-50/Ufl/fl mice (P = .00217). Cells were
counterstained with Hoechst (blue). White
inset boxes indicate the magnified region.
E, The number of Ki67-positive cells in
papillomas from Cenp-50/Ufl/fl (n = 12) (red
bar) and K14CreER-Cenp-50/Ufl/fl (n = 12)
(blue bar) mice (P = .144). F, The number
of TUNEL-positive cells in papillomas
from Cenp-50/Ufl/fl (n = 8) (red bar) and
K14CreER-Cenp-50/Ufl/fl (n = 8) (blue bar)
mice. The P-value was calculated by t test
(**P < .01). n.s., not significant. Error bars
represent the standard deviation (SD).
U^fl/fl (58.8%) and Cenp-50/U^fl/fl (59.1%) mice at 34 wk after initiation (Figure 6A). In addition, malignant conversion ratio per papilloma was almost the same in K14Cre^{ER}-Cenp-50/U^fl/fl (13.4%) and Cenp-50/U^fl/fl (16.2%) mice at 34 wk after initiation (Figure 6B). We then performed histological analysis using carcinomas from K14Cre^{ER}-Cenp-50/U^fl/fl and Cenp-50/U^fl/fl mice. HE staining revealed no significant morphological changes between these mice (Fig. S1E,F). These results suggested that Cenp-50/U has almost no effect on malignant conversion.

In this study, we found a decrease in early-stage papillomas in Cenp-50/U-deficient mice due to the induction of cell death, but there was little effect on malignant conversion. Taken together, Cenp-50/U mainly functions in papilloma development, and does not play a role in papilloma growth or malignant conversion (Figure 7).

4 | DISCUSSION

In this study, we specifically knocked out Cenp-50/U in the epidermis, and clarified its function in two-stage skin carcinogenesis. Cenp-50/U deficiency promoted cell death in normal mouse skins. In addition, our skin carcinogenesis experiments revealed that Cenp-50/U deficiency downregulates early papillomagenesis by inducing cell death.

Centromere proteins play a fundamental role in accurate chromosome segregation during mitosis and meiosis. Chromosome segregation errors cause genetic diseases, including some cancers. In addition, several centromere proteins play an important role in embryonic development.4,20-22,35 Therefore, it was difficult to generate centromere protein-deficient mice and perform functional analysis in vivo. As Cenp-50/U-deficient mice also died during early embryogenesis (approximately E7.5),24 its functions were poorly understood. Therefore, we generated Cenp-50/U conditional knockout mice and analyzed its function in vivo.

Cenp-50/U is a centromere protein, but unlike other centromere proteins including CENP-A or CENP-C, CENP-50/U-deficient DT40 cells are viable, suggesting that CENP-50/U-deficient DT40 cells have undergone mitosis, even if mitotic abnormalities happen. However, some populations of CENP-50/U-deficient DT40 cells cause cell death through apoptosis, which is similar to that observed in K14Cre^{ER}-Cenp-50/U^fl/fl mouse skins and papillomas. These observations suggested that Cenp-50/U is critical for cell division to ensure accurate chromosome segregation. By contrast, Cenp-50/U is dispensable in malignant conversion. In this stage, cells might tolerate abnormal mitotic cells like DT40 cells. The essentiality of
CENP-50/U varies between cell types and it is necessary early in papilloma growth.

Recent studies have reported aberrant expression of CENP-50/U in many types of cancers, including breast cancer, prostate cancer, colorectal carcinoma, ovarian cancer, glioblastoma, erythroleukemia, and lymphoma. These might be related to variety of CENP-50/U effects in various cell types and also suggested that CENP-50/U is a molecular marker for cancer therapy. We demonstrated that Cenp-50/U is highly expressed in papillomas and is involved in the early papillomagenesis stage, and this positive correlation suggested that Cenp-50/U can be a molecular marker in carcinogenesis.

Our skin carcinogenesis experiments using K14CreER.Ctenp-50/1fl/fl mice demonstrated that Cenp-50/U deficiency reduces early-stage papillomas. However, Cenp-50/U deficiency has little effect on malignant conversion. Consistent with this, Cenp-50/U is highly expressed in papilloma, but expression was decreased in malignant conversion. This finding suggests that Cenp-50/U functions as an oncogene in papilloma development, but loses this function at the late papilloma stage and in malignant conversion. We previously reported that Cenp-r deficiency increases early-stage papillomas and reduces the conversion of benign papillomas into malignant tumors. These results suggest that Cenp-50/U functions as an oncogene, whereas Cenp-r functions as a tumor suppressor in early papilloma development. Furthermore, based on mRNA expression analysis, Cenp-50/U and Cenp-r exhibit opposite expression profiles in papillomas and carcinomas. Although Cenp-50/U and Cenp-r form the Cenp-O complex, these genes have different functions in the two-stage skin carcinogenesis model. Our previous study demonstrated that Cenp-R localized to the centromere downstream of other CENP-O complex proteins, which can assemble a functional form in the absence of CENP-R, suggesting that CENP-R plays a distinct role compared with other CENP-O complex proteins. Skin carcinogenesis experiments using mice lacking multiple CENP-O complex proteins to elucidate the molecular mechanisms need to be carried out in detail in the future.

In conclusion, we demonstrated that Cenp-50/U plays a major role in early papilloma development and is involved in cell death, including apoptosis. In addition, Cenp-R and Cenp-50/U in the Cenp-O complex have different functions in the two-stage carcinogenesis model. Cenp-50/U may function either alone or in association with other centromere proteins in carcinogenesis, potentially leading to a novel therapeutic strategy.

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CONFLICT OF INTEREST
The authors have no conflict of interest.

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SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section.

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