INTRODUCTION

The epidermal growth factor receptor (EGFR) family, including ERBB1 (EGFR), ERBB2, ERBB3, and ERBB4, is a tyrosine kinase receptor that plays an essential role in development and disease. The EGFR family is abnormally expressed and is associated with metastasis in colorectal cancers. Unlike EGFR and ERBB2 as targets for disease control and cancer therapy, ERBB3 has been underestimated because of its inherently inactive kinase domain. However, ERBB3 is often overexpressed in colorectal cancers and associated with EGFR and ERBB2 expression. Upregulation of ERBB3 can induce resistance of therapeutic inhibitors designed to counteract EGFR and ERBB2 signaling.

In early biochemical studies, the ERBB3 intracellular domain has not been associated with ATP and has no autophosphorylation capacity. Therefore, it was thought that it could be activated only by heterodimerization with other ERBB receptors. Once neuregulins (NRGs) bind to ATP and have catalytic activity in vitro. However, the biological function of ERBB3 kinase activity remains elusive in vivo. Here we have identified the physiological function of inactivated ERBB3 kinase activity by creating Erbb3-K740M knockin mice in which ATP cannot bind to ERBB3. Unlike Erbb3 knockout mice, kinase-inactive Erbb3K740M homozygous mice were born in Mendelian ratios and showed normal development. After dextran sulfate sodium-induced colitis, the kinase-inactive Erbb3 mutant mice showed normal recovery. However, the outgrowth of ileal organoids by neuregulin-1 treatment was more attenuated in Erbb3 mutant mice than in WT mice. Moreover, in combination with the ApcMin mouse, the proportion of polyps less than 1 mm in diameter in mutant mice was higher than in control mice and an increase in the number of apoptotic cells was observed in polyps from mutant mice compared with polyps from control mice. Taken together, the ERBB3 kinase activity contributes to the outgrowth of ileal organoids and intestinal tumorigenesis, and the development of ERBB3 kinase inhibitors, including epidermal growth factor receptor family members, can be a potential way to target colorectal cancer.

KEYWORDS
ApcMin mouse, ERBB3, intestinal organoid, intestinal polyp, tyrosine kinase activity
bind to ERBB3, ERBB2 can phosphorylate the ERBB3 C-terminal tyrosine residues by transphosphorylation, but the mechanism by which kinase-inactive ERBB3 phosphorylates ERBB2 remains unknown.\textsuperscript{12,13} Recently, the purified ERBB3 intracellular domain has been identified as binding to ATP with an affinity similar to that of other active kinases and as having specific autophosphorylation activity.\textsuperscript{14,15} In addition, ERBB3 can form homodimerization and heterodimerization in cell membranes.\textsuperscript{15} Moreover, kinase-inactive human ERBB3(K723M) shows low levels of phosphorylation, indicating that the total ERBB3 phosphorylation includes ERBB3 autophosphorylation.\textsuperscript{14,15} Although this activity of ERBB3 kinase is weaker than that of EGFR, it can phosphorylate other ERBBs and the ligand-induced ERBB3 kinase activity is elevated by the ERBB2-ERBB3 heterodimer.\textsuperscript{15}

Regardless of the kinase activity, the genetic ablation in mice reveals the biological function of ERBB3 under normal physiological and pathological conditions. Homozygous \textit{Erbb3} knockout mice die because of an abnormality of the nervous system during pregnancy,\textsuperscript{16} and the additional defects in development and poor differentiation of several organs have been found in \textit{Erbb3} knockout embryos.\textsuperscript{17} Mammary gland-specific \textit{Erbb3} knockout mouse show that ERBB3 is essential for the formation of ducts during pregnancy and lactation by regulating the survival and differentiation of breast epithelial cells and is also necessary for promoting breast tumor formation.\textsuperscript{18-21} The liver-specific removal of \textit{Erbb3} retards tumor formation in a hepatocarcinogenesis mouse model and diminishes liver fibrosis.\textsuperscript{22,23} Intestine-specific \textit{Erbb3} ablation results in more damage of colonic epithelial cells and slow recovery of epithelial cells in a model of colitis induced by dextran sulfate sodium (DSS).\textsuperscript{24,25} Moreover, in an \textit{Apc\textsuperscript{Min}} intestinal tumor mouse model, intestine-specific \textit{Erbb3} knockout mice have less cellular proliferation within polyps in the intestine than do control mice.\textsuperscript{24}

Previous studies have reported the in vivo function of impaired kinase activity in EGFR family members. Homozygous mice \textit{Egr\textsuperscript{W502}} with V743G substitution are alive and reproducible but have defects in the eyes and skin\textsuperscript{26} and a feeding disorder.\textsuperscript{27} In addition, homozygous \textit{Egr\textsuperscript{W502}} mice with D833G substitution in the DFG motif of the kinase domain die during pregnancy.\textsuperscript{28} The homozygous K750M kinase-dead \textit{Erbb2} mice die at E10.5 because of defects in cardiac trabeculation and in the neural system.\textsuperscript{29} However, the physiological function of ERBB3 kinase activity remains an open question in vivo. In this study, we generated \textit{Erbb3-K740M} with K740M substitution knockin mice in which ATP could not bind to ERBB3, and sought to unveil the function of ERBB3 kinase activity in embryonic development, DSS-induced colitis, ileal organoid formation, and tumorigenesis in the \textit{Apc\textsuperscript{Min}} mouse model.

2 | MATERIALS AND METHODS

2.1 | Generation of \textit{Erbb3} knockin mice with mutation of the ATP-binding site

Detailed information on the generation of knockin mice using the CRISPR/Cas9 tool is described in Data S1. We undertook PCR with DNA lysates from live pups’ toes with \textit{Erbb3-Ex19S1}, \textit{5’-CTCCCTCCTTAACTCTTT-3’}; \textit{Erbb3-Ex19A51}, and \textit{5’-TACAGATGTCCTTTTCTT-3’} primers. The PCR products were subcloned in a pGEM-T Easy Vector System (Promega), and we verified the knockin allele of \textit{Erbb3} (\textit{Erbb3-K740M}, the kinase-inactive mutation of \textit{Erbb3}) by sequencing. We bred selected founder mice to identify germine transmission of the knockin allele, and further confirmed the F1 generation by PCR sequencing. Subsequent generations of mice carrying the \textit{Erbb3-K740M} allele were maintained on the B6 background, and genotyping was carried out by agarose gel electrophoresis of the PCR product after NolI digestion. The research ethics regulation and guidelines were approved by the Institutional Animal Care and Use Committee (IACUC) of Ewha Womans University.

2.2 | Cell staining

Both BrdU and TUNEL staining were undertaken as described previously.\textsuperscript{30} The number of BrdU- or TUNEL-positive cells per crypt was counted from 20 well-shaped crypts per mouse. The number of BrdU- or TUNEL-positive cells in polyps was counted in a given area at 400× magnification and the average number of stained cells was obtained from 2 to 8 fields of size-matched polyps from each mouse.

2.3 | Dextran sulfate sodium treatment and histological scoring

Mice aged 2–3 months were treated with 2.5% (w/v) DSS (Cat No. 160 110; MP Biomedicals) in drinking water. After 5 days, 2.5% DSS was replaced by drinking water for recovery. The body weight, stool consistency, and rectal bleeding were recorded daily. To evaluate the function of ERBB3 kinase, mice stayed in recovery for either 5 or 25 days. At the end of the experiment, the small intestine and colon were isolated and fixed in neutral buffered formalin, followed by embedding in paraffin and sectioning. We used H&E-stained sections for histological scoring using the previously described criteria (described in Data S1).\textsuperscript{31}

2.4 | Disease activity index measurement

We analyzed the body weight, stool consistency, and rectal bleeding daily in DSS-treated mice. The disease activity index (DAI) was the combined scores of weight loss, stool consistency, and rectal bleeding adapted as previously described.\textsuperscript{32} Each condition was graded on a scale of 0 to 4: weight loss percentage (0, none; 1, 0%-10%; 2, 10%-15%; 3, 15%-20%; 4, more than 20%); stool consistency (0, normal; 2, loose stool; 4, diarrhea); and rectal bleeding (0, normal; 2, hemorrhcruit; 4, cross bleeding).

2.5 | Quantitative RT-PCR

Total RNA preparation and quantitative RT-PCR (qRT-PCR) were carried out as described previously.\textsuperscript{30} The primer sequences used
for qRT-PCR are detailed in Data S1. Reactions were run on an ABI 7900HT Fast Real-Time PCR system (Thermo Fisher Scientific).

2.6 | In vivo phosphorylation analysis

We genotyped the neonatal pups (postnatal day 1). Next day, the male neonatal pups were injected s.c. with recombinant human NRG1-β1 (1 µg/g body weight) (396-HB-050; R&D Systems) or PBS (10 µg/g body weight) as the vehicle. After 10 minutes, skin and liver tissues were harvested and frozen in liquid nitrogen. We homogenized the tissues (50-100 mg) in 10 volumes of RIPA buffer supplements with proteinase and phosphatase inhibitors by means of a glass-Teflon homogenizer on ice. We measured protein concentration by using a Bradford Assay (Bio-Rad Lab).

2.7 | Western blot analysis

Equal amounts of protein extracts were separated by SDS-PAGE (7.5%), then were transferred to PVFD membranes. The detailed procedure for western blotting was described previously. The primary Abs are listed in Data S1. The protein-band densities were quantified by image analysis software ImageJ (https://imagej.nih.gov).

2.8 | Small intestinal crypt isolation and organoid culture

As described previously, we isolated small intestinal crypts from the ileum region. Briefly, the distal ileum was washed with ice-cold PBS. The lengthwise opened intestine was cut into pieces (less than 5 mm) and incubated in chelating buffer (2 mmol/L EDTA) with horizontal shaking for 30 minutes at 4°C. Ileum pieces were shaken by hand to dissociate the crypt from the basement membrane in dissociation buffer (D-sorbitol 54.9 mmol/L, sucrose 43.4 mmol/L). The crypt suspension was filtered by means of a cell strainer and pelleted by centrifuge. Five hundred whole crypts were suspended in 50 µL of a mixture containing 50% Matrigel (BD Biosciences) and 50% IntestiCul Organoid Growth Medium (Corning) and slowly ejected into 24-well plates. After polymerization, 750 µL IntestinCul media was added and replaced completely every 3 days. For organoid passageing, 7-9 days after seeding, the primary organoids were disrupted by being passed through a 26.5-gauge needle-equipped syringe in cold PBS. The dissociated organoids solution was centrifuged and embedded in Matrigel as above.

2.9 | Organoid measurement, treatment, and analysis

To evaluate the efficiency of organoid growth, we counted total organoid numbers/well under a light microscope at day 5 of culture in double-blinded fashion. We measured the size of organoids using the ImageJ program. The structure of the organoids was evaluated by counting the number of buds: sphere, no budding; organoid 1, 1 bud; organoid 2, 2 buds; organoid 3+, 3 or more buds. To examine the outgrowth of organoids by ligand treatment, we treated organoids with recombinant human epidermal growth factor (EGF; 10 ng/mL) (236-EG-200; R&D Systems) and recombinant human NRG1-β1 (10 ng/mL) for 3 days. For phosphorylation analysis, we homogenized organoid pellets in 50 µL RIPA buffer supplemented with proteinase and phosphatase inhibitors followed by western blotting. To examine the short-term effect of ligands on phosphorylation, we treated organoids with ligands for 10 minutes before preparing the protein extract.

2.10 | Macroadenoma counts and histopathological evaluation

We removed fecal material from the colon and small intestine of 2-month-old mice by flushing with PBS. Longitudinally opened small intestine and colon were splayed flat on 3MM paper. We counted polyp (>0.3 mm) numbers and diameters using a dissecting microscope and in-scope micrometer in double-blinded fashion. For histopathological evaluation, Swiss-rolled ileum was fixed, embedded in paraffin, and then stained in 5-µm sections with H&E. Histopathology of the polyps was classified as low grade, high grade, and adenocarcinoma following a previous report.

2.11 | Statistical analysis

We undertook statistical analysis by using GraphPad Prism 7.0 software. For significant analysis, we used Student’s t test or 2-way ANOVA with Sidak’s multiple comparisons test. P < .05 was considered statistically significant.

3 | RESULTS

3.1 | Erbb3<sup>K740M/K740M</sup> knockin mice show normal development

Based on previous reports for EGFR and ERBB2 kinase-inactive mutation, we selected a conserved lysine 740 amino acid of mouse ERBB3 as a site for mutagenesis. In vitro, mutant human ERBB3 with this lysine replaced by a methionine had strongly reduced autophosphorylation. We introduced knockin of the K740M (Lys-AAA to Met-ATG) sequence in mouse Erbb3 using the CRISPR-Cas9 system in 1-cell embryos. Of note, the codon sequence Ile-ATC (I739) was replaced by Ile-ATT to generate the Nall restriction enzyme site in the knockin allele (Figure 1A). Nall digestion of the PCR product of genomic DNA showed a 151-bp fragment for WT, 151-bp, 75-bp and 76-bp for Erbb3<sup>+/-K740M</sup> (Erbb3<sup>+/-M</sup>) heterozygous, and
75-bp and 76-bp for \textit{Erbb3} \textsuperscript{K740M} (\textit{Erbb3} \textsuperscript{M/M}) homozygous mice (Figure 1B). In addition, qRT-PCR analysis to compare the relative \textit{Erbb3} mRNA levels in the liver and colon tissues revealed that the \textit{Erbb3} mRNA level was the same in both WT and \textit{Erbb3} \textsuperscript{K740M/K740M} mice (Figure 1C). Western blot analysis showed that the ERBB3 proteins in \textit{Erbb3} \textsuperscript{K740M/K740M} mice were not significantly different from those in the WT mice, suggesting that the half-life of mutant ERBB3 protein is similar to that of WT ERBB3 (Figure 1D).

In contrast to the embryonic lethality of \textit{Erbb3} knockout mice,\textsuperscript{16,17} \textit{Erbb3} \textsuperscript{K740M/K740M} mice were born in the Mendelian ratio and had no obvious abnormal phenotype. Overall histology was similar in the ileum and colon between WT and \textit{Erbb3} \textsuperscript{K740M/K740M} mice (Figure 2A). The \textit{Erbb3} \textsuperscript{K740M/K740M} mice showed normal length of the small intestine and colon compared to the WT mice (Figure S1). After BrdU injection, the BrdU-positive cells per crypt were counted to evaluate the effect on intestinal cell proliferation. The number of BrdU-positive cells per crypt in the ileum and colon were not significantly different between the 2 groups (Figure 2B). The TUNEL assay revealed that apoptosis was also not significantly different in the ileum or colon of WT and \textit{Erbb3} \textsuperscript{K740M/K740M} mice (Figure 2C). Overall, kinase-inactive \textit{Erbb3} mutant mice showed normal intestinal homeostasis similar to that of intestinal epithelial cell (IEC)-specific \textit{Erbb3} knockout mice.\textsuperscript{24}

### 3.2 | In vivo tyrosine phosphorylation of the ERBB family is not affected in \textit{Erbb3} \textsuperscript{K740M/K740M} mice

In vitro, the isolated ERBB3 intracellular domain was able to autophosphorylate, and the endogenous full-length ERBB3 significantly upregulated ERBB3 tyrosine kinase activity when stimulated with NRG1.\textsuperscript{14,15} In order to validate the ERBB3 kinase activity in vivo, WT and \textit{Erbb3} \textsuperscript{K740M/K740M} littersmates were treated with either PBS or NRG1, and tissue extracts from skin were analyzed with western blotting. The NRG1-induced overall tyrosine phosphorylation levels (PY20) were similar between the WT and \textit{Erbb3} \textsuperscript{K740M/K740M} mice (Figure 3). Moreover, tyrosine phosphorylation of ERBB3 (pERBB3-Tyr1197), EGFR (pEGFR-Tyr1068), and ERBB2 (pERBB2-Tyr877) were not different between the 2 groups (Figure 3). Total levels of EGFR, ERBB2, and ERBB3 were similar between the 2 groups. There was little attenuation of ERBB3 phosphorylation in \textit{Erbb3} \textsuperscript{K740M/K740M} mice compared to WT in the liver (Figure S2).

### 3.3 | \textit{Erbb3} \textsuperscript{K740M/K740M} mice show normal recovery following DSS-induced colitis

The IEC-specific \textit{Erbb3} deletion mice show greater histological damage following DSS treatment and delayed recovery.\textsuperscript{24,25} To
clarify the contribution of ERBB3 kinase activity in those processes, Erbb3K740M/K740M mice were treated with 2.5% DSS for 5 days to induce colitis and then allowed to recover on normal drinking water for either a 5-day period (short-term recovery) or a 25-day period (long-term recovery). The body weight loss and DAI showed no significant difference between the WT and Erbb3K740M/K740M mice (Figure 4A,B). After a 5-day recovery, the small intestinal length or colonic length of WT and Erbb3K740M/K740M mice were similar (Figure 4C). Histological analysis of DSS-induced colitis by H&E staining of proximal, middle, and distal colonic sections showed no significant difference (Figure 4D), and all of the inflammation scores, epithelial regeneration scores, crypt damage scores, and total histological scores were similar between the WT and Erbb3K740M/K740M mice in short-term recovery (Figure 4E). There were also no significant differences in body weight loss (except on days 10 and 12), DAI, or histological scores in long-term recovery (Figure S3), indicating that ERBB3 kinase activity is not likely necessary for regeneration following intestinal damage.

3.4 | Neuregulin-1-β1-induced outgrowth of ileum organoids is inhibited in Erbb3K740M/K740M mice

Although ERBB3 is dispensable for the proliferation and survival of IEC in vivo, the abundant expression of ERBB3 in the IEC might have some function in intestinal homeostasis. To investigate the function of ERBB3 kinase activity in the intestine, we isolated and cultured the ileum crypts in a 3-D environment to form organoids, which are an ideal model to track morphology change under ligand treatment. In accordance with the in vivo proliferation result, the organoids from the ileum of Erbb3K740M/K740M mice showed organoid growth, size, and budding numbers similar to those of WT mice (Figure 5A,B). Exogenous treatment with NRG1-β1 for 3 days, but not EGF, promoted outgrowth of organoids and disrupted the architecture of organoids from WT mice, consistent with a previous study. However, Erbb3K740M/K740M organoids treated with NRG1-β1 failed to increase in size and complexity (Figure 5C,D; Figure S4). Also, the phospho-ERBB3(Tyr1197) protein level was significantly reduced in organoids from Erbb3K740M/K740M mice compared to WT
FIGURE 3 Neuregulin-1 (NRG1)-induced tyrosine phosphorylation of ERBB3 in vivo. A, Two-day-old WT (n = 3) and Erbb3<sup>K740M/K740M</sup> pups (n = 3) were treated with either PBS (−) or NRG1-β1 (+) for 10 minutes before harvesting skin tissues. Tissue extracts were analyzed by western blotting using indicated Abs. Photographs are representative images of 3 independent experiments. B, Relative phosphorylated (p) band intensities were normalized with total protein levels (n = 3). ns, statistically insignificant.

FIGURE 4 Effects of dextran sulfate sodium (DSS) on Erbb3<sup>K740M/K740M</sup> mice. Wild-type (n = 8) and Erbb3<sup>K740M/K740M</sup> (n = 8) adult mice were treated with 2.5% DSS for 5 d and switched to normal drinking water; mice were killed for analysis at day 10. A, Body weight loss during DSS treatment is expressed as a percentage of the weight on the day of first exposure to DSS. B, Disease activity index (DAI) score was graded daily in WT and Erbb3<sup>K740M/K740M</sup> mice. C, Small intestinal (top) and colonic (bottom) length of WT and Erbb3<sup>K740M/K740M</sup> mice. Data are represented as mean ± SEM of each genotype. D, Colon and ileum sections were stained with H&E. E, Inflammation score, epithelial regeneration score, crypt damage score, and total histological score of the colon are shown. Each dot represents the score of 1 individual mouse, and horizontal lines are means for each genotype. ns, statistically insignificant.
mice after NRG1-β1 treatment (Figure 5E). Neuregulin-1-β1 has poor effects on phospho-EGFR(Tyr1068) levels. In contrast, short-term (10-minute) treatment with NRG1-β1 resulted in similar activation of phospho-ERBB3(Tyr1197) protein levels in both WT and Erbb3 K740M/K740M mice (Figure S5), indicating that ERBB3 kinase activity affects phosphorylation of ERBB3 only in long-term NRG1-β1 treatment.

3.5 | Effect of ERBB3 kinase activity on Apc Min tumor development

Because IEC-specific knockout of Erbb3 results in a dramatic reduction of the number and size of polyps in Apc Min−/− tumor mice,24 the WT and Erbb3 K740M mice were crossed with Apc Min−/− mice to find out whether ERBB3 kinase activity contributes to intestinal tumorigenesis. The polyp numbers in the small intestine, but not in the colon of Erbb3 K740M/K740M mice were significantly reduced more than in the control mice (Figure 6A,B). The tumor sizes in the small intestine of control and Erbb3 K740M/K740M mice were not significantly different (Figure 6C). But the fraction of tumors less than 1 mm in diameter were 78.0 ± 2.1% in Erbb3 K740M/K740M, significantly higher than the 71.0 ± 2.4% in control mice (Figure 6D). Conversely, the fraction of tumors between 1 mm and 2 mm in diameter was smaller in Erbb3 K740M/K740M mice than in control mice (21.3 ± 2.1% and 27.7 ± 2.3%, respectively), and the fraction of tumors between 2 mm and 3 mm in diameter was also smaller in Erbb3 K740M/K740M mice than in control mice (0.46 ± 0.1% and 1.1 ± 0.19%, respectively), indicating that ERBB3 kinase activity contributes to polyp growth from the early stages. In control mice, high-grade adenomas showed more distorted glandular structures. In contrast, most high-grade adenomas in Erbb3 mutant mice showed less distorted glandular structures.
with multiple crypts and scattered goblet cells (Figure 6E). The number of low-grade adenomas was superior to high-grade adenomas in both control and Erbb3 mutant mice (Figure 6F). The number of low-grade adenomas was not significantly different between the control (10.2 ± 2.2) and Erbb3 mutant mice (9.5 ± 2.1), but the high-grade adenomas showed a significant decrease in the Erbb3 mutant (1.8 ± 0.7) compared to the control mice (4.7 ± 1.2). Adenocarcinoma was not found in either group. The number of BrdU-positive cells in intestinal polyps of Erbb3<sup>K740M/K740M</sup> mice showed no significant difference compared with those of control mice (Figure 6G). In contrast, the number of TUNEL-positive cells in intestinal polyps of Erbb3<sup>K740M/K740M</sup> mice was higher than that of control mice (12.8 ± 3.7 and 9.6 ± 1.9, respectively) (Figure 6H), showing that loss of ERBB3 kinase activity leads to increased apoptosis in intestinal polyps.
In this study, we investigated the role of ERBB3 kinase activity in embryonic development, intestinal homeostasis, and tumorigenesis using Erbb3 kinase-inactive mutant mice. Various mutant phenotypes were observed in the hypoplastic EGFR kinase mutant mice, $Egfr^{wa2/wa2}$, $Egfr^{wa5/wa5}$, and kinase-dead Erbb2 $K750M$ knockin mice.26,28,29 A dramatic impairment of mammary epithelial outgrowth in mice homozygous for the Erbb3 allele lacking the 7 known PI3K-binding sites indicates that ERBB3-associated PI3K activity is critical for mammary development.41 However, we observed that kinase-inactive Erb3 knockin homozygous mice were born and survived normally without any obvious defects, indicating that the kinase activity of ERBB3 is not crucial for embryonic development or survival. It is not surprising that the total phosphorylation and tyrosine phosphorylation of the ERBB family did not attenuate in kinase-inactive Erbb3 homozygous mice. Contrary to the essential role of the ERBB family in development and disease regulation, ERBB3 kinase activity itself does not seem to participate in normal homeostasis.

ERBB3 is required not only for its protective role of the intestinal epithelium during DSS-induced colitis but also for epithelium regeneration in both short-term and long-term recovery periods.24,25 Both ERBB2 and ERBB3 were shown to be reparative elements to protect against colitis injury by means of promoting cell survival.25 In this study, unlike in IEC-specific Erbb3 knockout mice, kinase-inactive Erbb3 mice showed normal recovery from the DSS-induced colitis, perhaps because kinase-inactive ERBB3 still forms a heterodimer with ERBB2 or other EGFR family members that in turn trigger signals pathways for cell proliferation and survival regardless of ERBB3 kinase activity. Indeed, the dual inhibition of ERBB2/ERBB3 is known to be required to fully inhibit AKT activity.42

In the intestinal organoid model, it is known that NRG1-β1 is superior to EGF in promotion of the hyperproliferation and dysregulated shape of ileal organoids.40 Epidermal growth factor is one of the essential factors for organoid culture,33 and the EGF concentration in the culture medium seems to be saturated as organoid outgrowth is not affected by the additional EGF treatment. Interestingly, the severe attenuation of organoid outgrowth in kinase-inactive Erbb3 mutant mice suggests that ERBB3 kinase activities contribute to outgrowth of epithelial cells under NRG1-β1 exposure, which is usually overexpressed in colon cancers.43 The phospho-ERBB3 level of kinase-inactive ERBB3 mutant was not reduced under short-term stimulation by NRG1-β1 either in vivo or ex vivo, but it failed to prolong this activated state in long-term stimulation ex vivo. The kinase activity of ERBB3 seems to take a distinct role in maintaining its phosphorylation state instead of transphosphorylation by dimerization with other receptors in the presence of abundant NRG1-β1. Epidermal growth factor receptor will form a heterodimer with ERBB3 only under an unsaturated level of EGF when EGF-free EGFR molecules are available to interact with ERBB3.44 Our data showed that NRG1-β1 induces a low EGFR phosphorylation, which indicates that EGFR phosphorylation is not advantageous in EGFR/ERBB3 heterodimers, because ERBB3 kinase activity is not strong enough to transphosphorylate EGFR.45,46

When ERBB3 is heterodimerized with other ERBBs by NRG1 binding, ERBB3 itself is transphosphorylated and led to downstream activation of PI3K/AKT/mTOR and RAS/RAF/MEK/ERK signaling pathways.3,8,47 The oncogenic regulation of NRG1 and kinase activity of ERBBs in the intestinal epithelium are becoming strongly evident, and ERBB3 is a principal element in the regulation of the ERBB signaling pathway and a potential cancer-drug target.48,49 Oncogenic ERBB3 mutations in human cancer showed different patterns in ligand dependency, in which several mutants were oncogenic in the absence of NRG1, more stimulated by NRG1 treatment, or less sensitive to NRG1.50 Expression of NRG1 has been found to be higher in tumors than in normal colonic epithelial cells and contributes to malignant transformation.43,47 These findings shed light on how ERBB3 kinase activity might have some functions in the presence of NRG1 ligands. The IEC-specific Erbb3 knockout mice showed fewer and smaller polyps when crossed with ApcMin mice, emphasizing the critical roles in tumor formation and progression by ERBB3 signaling.24 Our finding fewer polyps and more small polyps in Erbb3 kinase-inactive mice suggests that intrinsic ERBB3 kinase activity also contributes to early growth of polyps in ApcMin mice. Furthermore, an increase of apoptosis in intestinal polyps of Erbb3 kinase-inactive mice similar to the IEC-specific Erbb3 knockout mice indicates that ERBB3 kinase activity seems to support early tumor growth and survival given NRG1 overexpression in adenomas of ApcMin mice.

Taken together, we uncovered that the ERBB3 kinase activity contributes to intestinal adenoma formation in ApcMin mice and helps to induce ERBB3 phosphorylation, which is relevant to the outgrowth of ileal organoids in the presence of overexpressed NRG1. Several therapeutic mAbs targeting ERBB3 are under clinical evaluation,51 and the possibility of ERBB3 autophosphorylation in EGFR/ERBB2-targeted therapeutic resistant tumors suggests that the development of kinase inhibitors for the pan-EGFR family could also be an important therapeutic strategy in colorectal cancer.

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DISCLOSURE
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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