CircHIPK3 promotes colorectal cancer growth and metastasis by sponging miR-7

Kaixuan Zeng1,2, Xiaoxiang Chen1,2, Mu Xu1, Xiangxiang Liu1, Xiujiu Hu1, Tao Xu1, Huiling Sun1, Yuqin Pan1, Bangshun He1 and Shukui Wang1,2

Abstract
Mounting evidences indicate that circular RNAs (circRNAs) have a vital role in human diseases, especially cancers. More recently, circHIPK3, a particularly abundant circRNA, was proposed to be involved in tumorigenesis. However, its role in colorectal cancer (CRC) has not been explored. In this study, we found circHIPK3 was significantly upregulated in CRC tissues and cell lines, at least in part, due to c-Myb overexpression and positively correlated with metastasis and advanced clinical stage. Moreover, Cox multivariate survival analysis showed that high-level expression of circHIPK3 was an independent prognostic factor of poor overall survival (OS) in CRC, hazard ratio (HR) = 2.75, 95% confidence interval [CI] 1.74–6.51, p = 0.009. Functionally, knockdown of circHIPK3 markedly inhibited CRC cells proliferation, migration, invasion, and induced apoptosis in vitro and suppressed CRC growth and metastasis in vivo. Mechanistically, by using biotinylated-circHIPK3 probe to perform RNA pull-down assay in CRC cells, we identified miR-7 was the only one microRNA that was abundantly pulled down by circHIPK3 in both HCT116 and HT29 cells and these interactions were also confirmed by biotinylated miR-7 pull-down and dual-luciferase reporter assays. Overexpression of miR-7 mimicked the effect of circHIPK3 knockdown on CRC cells proliferation, migration, invasion, and apoptosis. Furthermore, ectopic expression of circHIPK3 effectively reversed miR-7-induced attenuation of malignant phenotypes of CRC cells by increasing the expression levels of miR-7 targeting proto-oncogenes (FAK, IGF1R, EGFR, YY1). Remarkably, the combination of circHIPK3 silencing and miR-7 overexpression gave a better effect on tumor suppression both in vitro and in vivo than either circHIPK3 knockdown or miR-7 overexpression alone. Taken together, our data indicate that circHIPK3 may have considerable potential as a prognostic biomarker in CRC, and support the notion that therapeutic targeting of the c-Myb/circHIPK3/miR-7 axis may be a promising treatment approach for CRC patients.

Introduction
Colorectal cancer (CRC) is the third most common malignant disease and the fourth most frequent cause of cancer-related death worldwide1. Despite many advances in the diagnosis and therapeutic improvements of this disease, the prognosis of CRC patients remains poor, owing to late stage at initial diagnosis and high frequency of metastasis and recurrence2. Hence, the discovery of new potential biomarkers for prognosis predication and deeper elucidation of the exact molecular mechanisms underlying CRC malignancy may provide improved treatments of CRC patients.

Circular RNAs (circRNA) are highly conserved and stable covalently closed RNA transcripts generated by back-splicing of a single pre-mRNA with gene-regulatory potential3,4. Emerging evidences show that circRNAs possess closely related to human diseases, especially cancers, and may act as better biomarkers due to their abundance and stability5,6. Recently, circHIPK3, a particularly abundant circRNA7,8, have been verified to be involved in metabolic dysregulation9 and tumorigenesis10.
Zheng et al.\textsuperscript{10} showed circHIPK3 was significantly upregulated in liver cancer and promoted cell proliferation, implying that circHIPK3 might be an oncogene. On the contrary, a more recent study found circHIPK3 served as a tumor suppressor to inhibit bladder cancer growth and metastasis\textsuperscript{11}. Whether or not circHIPK3 as a suppressor or oncogene still remains under debate.

MicroRNAs (miRNAs), an evolutionarily conserved group of small regulatory noncoding RNAs, have been confirmed to be involved in various biological functions\textsuperscript{12}. Considerable studies reported that circRNAs as “miRNA sponges” to regulate gene expressions\textsuperscript{13–17}. The most well-known circRNA is CDR1as, which harbors more than 70 selectively conserved miR-7 target sites\textsuperscript{15,18}. However, the regulatory roles of circRNAs act as “miRNA sponges” in CRC are still largely unknown.

In this study, we identified that circHIPK3 was an oncogene, which was upregulated in CRC and increased circHIPK3 predicted poor prognosis. Furthermore, circHIPK3 could sponge endogenous miR-7 to sequester and inhibit miR-7 activity, thereby leading to increased FAK, IGFR1, EGF\textsubscript{R}, and YY1 expression. Our findings reveal a novel mechanism underlying circHIPK3 in CRC progression.

**Results**

**CircHIPK3 is significantly upregulated in CRC and increased circHIPK3 expression predicts poor prognosis**

CircHIPK3 (hsa_circ_0000284) is derived from the HIPK3 gene Exon2, whose spliced mature sequence length is 1099 bp. The result of Sanger sequencing confirmed the head-to-tail splicing in the RT-PCR product of circHIPK3 (Fig. 1a). We then evaluated the stability and localization of circHIPK3. Following Actinomycin D (an inhibitor of transcription) treatment, the half-life of circHIPK3 exceeded 24 h, while that of linear HIPK3 exhibited only about 3 h, indicating that circHIPK3 is highly stable (Fig. 1b). Resistance to digestion by RNase R further confirmed that circHIPK3 harbors a loop structure (Fig. 1c). Fluorescence in situ hybridization (FISH) assay showed that circHIPK3 predominately localized in the cytoplasm (Fig. 1d).

Next, we investigated the expression level of circHIPK3 in CRC cell lines and tissues. CircHIPK3 is significantly upregulated in CRC cell lines (HCT116, HT29, SW480, SW620, DLD1) compared with normal colon mucosal epithelial cell (FHC) (Fig. 1e). The similar results were also observed in CRC compared with matched normal tissues (Fig. 1f). Statistical analyses showed that increased expression of circHIPK3 was significantly associated with T status of tumor \((p = 0.028)\), lymph node metastasis \((p = 0.023)\), distant metastasis \((p = 0.006)\), and advanced clinical stage \((p = 0.004)\) (Table 1). Moreover, CRC patients with high expression of circHIPK3 had significantly shorter overall survival than those with the low expression of circHIPK3 by the analysis of Kaplan–Meier survival curve \((p < 0.001)\) (Fig. 1g). Further Cox multivariate survival analysis revealed that high circHIPK3 expression was an independent prognostic factor for poor survival of CRC patients (hazard ratio [HR] = 2.75, 95% confidence interval [CI] 1.74–6.51, \(p = 0.009)\) (Table 2). These results suggest that circHIPK3 upregulation as an early event in CRC development and a vital role in CRC progression.

The transcription factor c-Myb is an upstream regulator of circHIPK3 expression

Previous studies showed the enrichment for circHIPK3 transcribed by c-Myb in diabetes mellitus\textsuperscript{9,19}. Thus, we wonder whether c-Myb can also regulate the expression of circHIPK3 in CRC. We found c-Myb was significantly overexpressed in CRC cell lines (Fig. 2a) and tissues (TCGA database) (Fig. 2b), which is consistent with previous research. Then, HCT116 and HT29 cell lines were transfected with c-Myb siRNA, NC siRNA, Vector, and c-Myb, respectively. qRT-PCR results demonstrated that silencing of c-Myb decreased, but overexpression of c-Myb increased, the expression of circHIPK3 in both HCT116 and HT29 cell lines (Fig. 2c). Luciferase reporter assay showed that c-Myb overexpression noticeably enhanced the luciferase activity of the vector containing c-myb site within circHIPK3 promoter, whereas the luciferase activity of the vector with mutant c-Myb binding site was not affected (Fig. 2d). Moreover, ChIP assay also showed the amount of immunoprecipitated DNA from the circHIPK3 promoter was increased upon overexpression of c-Myb (Fig. 2e, f). Altogether, the above results indicate that c-Myb elevates the expression of circHIPK3 by directly binding to its promoter region.

Silencing of circHIPK3 inhibits CRC cells proliferation, migration, invasion, and induces apoptosis in vitro

In an attempt to investigate the biological functions of circHIPK3 in CRC, we designed three small interfering RNAs (siRNAs) targeting the junction sites of circHIPK3 to silence circHIPK3 expression in HCT116 and HT29 cell lines. These siRNAs obviously decreased circHIPK3 expression level, but had no effect on its liner isofrom (Fig. 3a). And we chose si-circHIPK3#1 for the subsequent experiment due to the highest inhibitory efficiency. The colony formation assay showed that circHIPK3 knockdown significantly suppressed colony-forming ability of HCT116 and HT29 cell lines (Fig. 3b). Cell proliferation was measured by the CCK8 (Fig. 3c) and EdU assay (Fig. 3d), and silencing of circHIPK3 significantly inhibited cell proliferation in these two cell lines. In addition, more apoptotic cells are
presented in si-circHIPK3 group as compared with si-NC group in HCT116 and HT29 cell lines, respectively (Fig. 3e). Moreover, transwell invasion assay without or with matrigel demonstrated that circHIPK3 silencing markedly impeded HCT116 and HT29 cells migration (Fig. 3f) and invasion (Fig. 3g) by 46% and 51%, respectively. These data collectively indicate that silencing of circHIPK3 can retard the progression of CRC cells.
CircHIPK3 can sponge miR-7 in CRC cell lines

To explore whether circHIPK3 can function as "miRNA sponge" in CRC cells, we selected the top ten miR-599, miR-93-3p, miR-365a-5p, miR-365b-5p, miR-421, miR-570-3p, miR-597-5p, miR-7, miR-1207-3p, miR-124-5p candidate miRNAs through CircNet database. A 3' terminal-biotinylated-circHIPK3 probe was designed to determine which miRNAs potentially interact with circHIPK3. The probe was verified to pull-down circHIPK3 in CRC cells and circHIPK3 overexpression increased the pull-down efficiency (Fig. 4a). qRT-PCR analyses revealed that miR-7 was the only one miRNA that was abundantly pulled down by circHIPK3 probe in both HCT116 and HT29 cells (Fig. 4b and c). To further consolidate the direct binding of miR-7 and circHIPK3, we utilized biotin-labeled miR-7 and its mutant mimics to pull-down circHIPK3 in HCT116 and HT29 cells with circHIPK3 overexpression, the results showed wild-type miR-7 captured more circHIPK3 compared with the mutant (Fig. 4d). Next, we carried out luciferase reporter assays and demonstrated that overexpression of miR-7 significantly decreased the luciferase activity of the vector containing the complete circHIPK3 sequence, but did not affect the luciferase activity of the vector with mutant miR-7-binding site in HCT116 and HT29 cells (Fig. 4e). Furthermore, we applied FISH to assess whether there is a co-location between circHIPK3 and miR-7, the result showed that circHIPK3 and miR-7 were co-localized in cytoplasm (Fig. 4f).

We then evaluated the expression of miR-7 in CRC tissues, qRT-PCR result showed miR-7 was significantly downregulated with the increasing of clinical stages (Fig. 4g). And overexpression of circHIPK3 decreased, but silencing of circHIPK3 increased, the expression of miR-7 in HCT116 and HT29 cell lines (Fig. 4h). Moreover, circHIPK3 expression was negatively correlated with the expression of miR-7 in CRC tissues (r = −0.453, p < 0.001) (Fig. 4i). Taken together, these data demonstrate that circHIPK3 acts as a miRNA sponge for miR-7 in CRC.

Overexpression of circHIPK3 effectively reverses miR-7-induced inhibition of CRC cells progression

miR-7, a well-known tumor suppressor, was involved in many human tumors development and progression including CRC. Our data also showed that overexpression of circHIPK3 decreased, but silencing of circHIPK3 increased, the expression of miR-7 in HCT116 and HT29 cell lines (Fig. 4h). Moreover, circHIPK3 expression was negatively correlated with the expression of miR-7 in CRC tissues (r = −0.453, p < 0.001) (Fig. 4i). Taken together, these data demonstrate that circHIPK3 acts as a miRNA sponge for miR-7 in CRC.

### Table 1 Correlations between circHIPK3 expression and clinical characteristics in CRC patients (n = 178)

| Clinicopathologic parameters | Total (n = 178) | CircHIPK3 expression | p value |
|------------------------------|----------------|-----------------------|--------|
|                              | Low (%) | High (%) |        |
| Age (years)                  |         |           |        |
| ≤65                          | 79      | 41 (51.9%) | 38 (48.1%) | 0.763 |
| >65                          | 99      | 48 (48.5%) | 51 (51.5%) |        |
| Gender                       |         |           |        |
| Male                         | 102     | 49 (48.0%) | 53 (52.0%) | 0.650 |
| Female                       | 76      | 40 (52.6%) | 36 (47.4%) |        |
| Tumor site                   |         |           |        |
| Colon                       | 115     | 62 (53.9%) | 53 (46.1%) | 0.210 |
| Rectum                       | 63      | 27 (42.9%) | 36 (57.2%) |        |
| Tumor size (cm)              |         |           |        |
| ≤5                           | 107     | 57 (53.3%) | 50 (46.7%) | 0.358 |
| >5                           | 71      | 32 (45.1%) | 39 (54.9%) |        |
| Pathological T category      |         |           |        |
| T1–T2                       | 48      | 31 (64.6%) | 17 (35.4%) | 0.028*|
| T3–T4                       | 130     | 58 (44.6%) | 72 (55.4%) |        |
| Lymph node metastasis        |         |           |        |
| N0                           | 102     | 59 (57.8%) | 43 (42.2%) | 0.0006**|
| N1–2                        | 76      | 30 (39.5%) | 46 (60.5%) |        |
| Distant metastasis           |         |           |        |
| M0                           | 155     | 84 (54.2%) | 71 (45.8%) |        |
| M1                           | 23      | 5 (21.7%)  | 18 (78.3%) |        |
| TNM stage                    |         |           |        |
| I–II                         | 121     | 71 (57.8%) | 51 (42.2%) | 0.004**|
| III–IV                      | 56      | 19 (33.3%) | 38 (66.7%) |        |
| Differentiation              |         |           |        |
| Well                         | 23      | 12 (52.2%) | 11 (47.8%) | 0.695 |
| Moderate                     | 66      | 31 (46.3%) | 35 (53.7%) |        |
| Poor                         | 26      | 11 (42.3%) | 15 (57.7%) |        |

*aUsing median circHIPK3 values as cutoff

**p < 0.01

*p < 0.05
Likewise, transwell invasion assay without or with matrigel revealed that miR-7 together with circHIPK3 overexpression obviously impeded HCT116 and HT29 cells migration (Fig. 5d) and invasion (Fig. 5e) compared with miR-7 overexpression alone. In addition, we also found that overexpression of miR-7 combined with knockdown of circHIPK3 displayed an additive suppressive effect on CRC cell malignant phenotype compared with miR-7 overexpression or circHIPK3 silencing alone (Supplementary Fig. 1a–c).

Accumulating evidence indicated that miR-7 could simultaneously target various oncogenes involved in diverse signaling pathways in different human tumors27. We then wonder whether circHIPK3 exert pro-tumor role by elevating the expression of miR-7 targeting oncogenes. A panel of growth and metastasis-related miR-7 targets was chosen for qRT-PCR analysis. The results showed that circHIPK3 status affected the expression of FAK, IGF1R, EGFR, and YY1, but not that of PAX6, XRCC2, and RAF1. Ectopic expression of circHIPK3 increased, but knockdown of circHIPK3 decreased, the expression levels of FAK, IGF1R, EGFR, and YY1 (Fig. 5f). Moreover, the expression of FAK, IGF1R, EGFR, and YY1 was markedly increased in CRC cells (HCT116 and HT29) co-transfected with miR-7 mimics and circHIPK3 vectors compared with the cells transfected with miR-7 mimics alone (Fig. 5g). These above results implicate that overexpression of circHIPK3 effectively reverses miR-7-induced attenuation of aggressive phenotypes of CRC cells by sponging miR-7 and subsequent promotion of FAK, IGF1R, EGFR, and YY1 expression.

Silencing of circHIPK3 combined with overexpression of miR-7 exhibits an additive inhibitory effect on CRC growth and metastasis in xenograft animal models

To further assess whether circHIPK3 exerts tumor-promoting effect in vivo, we established the xenograft mouse models by subcutaneously injecting equal amount of HCT116 cells (n = 8 for each group). After ~10 days, when the volume of the tumor reached about 100 mm³, si-circHIPK3 alone, agomir-7 alone or both were injected intratumorally every two days for two weeks. As shown in Fig. 6a, intratumorally injection of agomir-7 or si-circHIPK3 markedly reduced tumor volume and weight, respectively. More importantly, the combined si-circHIPK3 and agomir-7 group displayed less tumor volume and weight than that of si-circHIPK3 or agomir-7 alone (Fig. 6a). Likewise, miR-7 overexpression or circHIPK3 knockdown significantly decreased the expression of miR-7 targeting proto-oncogenes (FAK, IGF1R, EGFR, YY1) both at the mRNA and protein levels, and the combined group exhibited less expression of these oncogenes (Supplementary Fig. 2a, b). IHC analysis also showed that decreased Ki67, MMP9-positive cells, and microvascular density, and increased cleaved caspase-3-positive cells in si-circHIPK3 + agomir-7 group compared with si-circHIPK3 or agomir-7 group alone (Fig. 6b). We also established the metastasis models through tail-vein injecting HCT116 cells into nude mice (n = 8 for each group). After ~10 days, when the volume of the tumor reached about 100 mm³, si-circHIPK3 alone, agomir-7 alone or both were injected intratumorally every two days for two weeks. As shown in Fig. 6a, intratumorally injection of agomir-7 or si-circHIPK3 markedly reduced tumor volume and weight, respectively. More importantly, the combined si-circHIPK3 and agomir-7 group displayed less tumor volume and weight than that of si-circHIPK3 or agomir-7 alone (Fig. 6a). Likewise, miR-7 overexpression or circHIPK3 knockdown significantly decreased the expression of miR-7 targeting proto-oncogenes (FAK, IGF1R, EGFR, YY1) both at the mRNA and protein levels, and the combined group exhibited less expression of these oncogenes (Supplementary Fig. 2a, b). IHC analysis also showed that decreased Ki67, MMP9-positive cells, and microvascular density, and increased cleaved caspase-3-positive cells in si-circHIPK3 + agomir-7 group compared with si-circHIPK3 or agomir-7 group alone (Fig. 6b). We also established the metastasis models through tail-vein injecting HCT116 cells into nude mice (n = 8 for each group). After ~10 days, when the volume of the tumor reached about 100 mm³, si-circHIPK3 alone, agomir-7 alone or both were injected intratumorally every two days for two weeks. As shown in Fig. 6a, intratumorally injection of agomir-7 or si-circHIPK3 markedly reduced tumor volume and weight, respectively. More importantly, the combined si-circHIPK3 and agomir-7 group displayed less tumor volume and weight than that of si-circHIPK3 or agomir-7 alone (Fig. 6a). Likewise, miR-7 overexpression or circHIPK3 knockdown significantly decreased the expression of miR-7 targeting proto-oncogenes (FAK, IGF1R, EGFR, YY1) both at the mRNA and protein levels, and the combined group exhibited less expression of these oncogenes (Supplementary Fig. 2a, b). IHC analysis also showed that decreased Ki67, MMP9-positive cells, and microvascular density, and increased cleaved caspase-3-positive cells in si-circHIPK3 + agomir-7 group compared with si-circHIPK3 or agomir-7 group alone (Fig. 6b).
As illustrated in Fig. 6c, an average of 38 lung metastatic nodules per mice was observed in control group and 17, 19 were respectively observed in agomir-7 or si-circHIPK3 group, whereas only 6 were observed in the combined si-circHIPK3 and agomir-7 group. Overall, these data indicate that silencing of circHIPK3 suppresses
Fig. 3  circHIPK3 silencing suppresses CRC cells proliferation, migration, invasion, and induces apoptosis. a The interfering efficacies of three circHIPK3-targeting siRNAs on circHIPK3 and HIPK3 mRNA were measured by qRT-PCR. b Colony formation assay of HCT116 and HT29 cells transfected with control or circHIPK3#1 siRNAs. c, d CCK8 and EdU assays of HCT116 and HT29 cells transfected with control or circHIPK3 siRNAs were performed to evaluate cell proliferative ability. e Flow cytometry apoptosis analysis of HCT116 and HT29 cells transfected with control or circHIPK3#1 siRNAs. f, g Cell migration and invasion abilities of HCT116 and HT29 cells transfected with si-NC or si-circHIPK3#1 were evaluated by transwell migration and matrigel invasion assay. Data were represented as means ± S.D. of at least three independent experiments. *p < 0.05, **p < 0.01, ***p < 0.001. Scare bar = 50 μm.
CRC growth and metastasis in vivo and combined with miR-7 overexpression exhibits an additive effect on tumor repression.

Discussion

It is becoming increasingly clear that circRNAs have a crucial role in cancer development and progression and affect the hallmarks of cancer. However, their roles in CRC remain largely unknown. In this study, we found circHIPK3 was significantly upregulated in CRC, at least in part, due to c-Myb overexpression and increased circHIPK3 predicted poor prognosis. Mechanistically, ectopic expression of circHIPK3 could rescue the expression of miR-7 targeting oncogenes by sponging miR-7, thereby promoting CRC progression (Fig. 7). Moreover, the combination of circHIPK3 knockdown and miR-7 overexpression gave a better tumor-suppressive effect both in vitro and in vivo than did circHIPK3 silencing or miR-7.
Fig. 5 (See legend on next page.)
reintroduction alone. Thus, our study demonstrate that co-expressing miR-7 along with a circHIPK3 inhibitor may be a promising treatment approach for patients with CRC.

Previous studies indicated the enrichment for circHIPK3 transcribed by c-Myb in diabetes mellitus5,19. c-Myb overexpression occurring in many malignancies, including CRC, often marks poor prognosis30. And c-Myb have a pivotal role in promoting cell proliferation, survival, and metastasis by activating various signaling pathways31. We evaluated whether proto-oncogene c-Myb also regulated circHIPK3 expression in CRC. Luciferase reporter assay displayed that overexpression of c-Myb promoted the transcription of circHIPK3, and ChIP-qPCR analysis further confirmed c-Myb could directly bind to circHIPK3 promoter region in CRC cells. These indicated that transcription factor c-Myb is an upstream regulator of circHIPK3 expression, which were consistent with the previous reports.

Emerging studies show that circRNAs are abundant in the human transcriptome2,15,18. In addition, circRNAs are more stable than other types of RNA because of their covalently closed structures. Therefore, circRNAs are more suitable as potential cancer biomarkers than other RNAs, such as miRNAs, and lincRNAs, due to their abundance and stability. As expected, there are already many circRNAs recognized as cancer biomarkers. CiRS-7, circPVT1, circBRAF, circPRDM2, and circTCF25 were respectively identified to be a promising prognostic biomarker in colorectal cancer6, gastric cancer8, glioma32, hepatocellular carcinoma33, and bladder cancer34. Herein, we found that circHIPK3 was frequently upregulated in CRC and patients with circHIPK3 high expression had significantly shorter overall survival than those with the low expression of circHIPK3, implying that circHIPK3 might be a promising prognostic biomarker in CRC.

Up to now, numerous studies have shown that circRNAs exerted various biological functions by acting as competing endogenous RNAs (ceRNAs)3,5,36. Analogously, circHIPK3 could promote cell growth by sponging miR-12410. However, Shan et al.9 reported that circHIPK3 controlled diabetic proliferative retinopathy via sponging miR-30a, but not miR-124. Meanwhile, circHIPK3 was proposed to inhibit bladder cancer progression through sponging miR-55811. One likely explanation for these discrepancies is that circRNA have its biological roles in tissue/developmental-stage specific context3. Importantly, we identified that circHIPK3 could interact with miR-7 in CRC by biotinylated RNA pull-down and dual-luciferase reporter assays, which has not been reported by previous studies.

MiR-7, a well-known tumor suppressor, was proposed to participate in the development and progression of different types of human cancers27. In agreement with previous studies, our data indicated that miR-7 overexpression significantly inhibited CRC cells proliferation, migration, invasion and induced apoptosis, and these inhibitory effects were exceedingly similar to circHIPK3 silencing. Moreover, we found circHIPK3 could sponge endogenous miR-7 to sequester and reduce miR-7 activity, thus resulting in increasing the expression of miR-7 targeting oncogenes (FAK, IGF1R, EGFR, and YY1). FAK—a kinase regulated by cell—ECM interactions and known to be important for cellular migration, which promoted tumor invasion and metastasis by elevating the expression of VEGF, MMP2, and MMP938. Wealth of studies confirmed that IGF1R and EGFR could respectively activate PI3K/AKT and MEK/ERK signaling pathways to promote cancer progression and drug resistance39,40. And we also found that inhibition of circHIPK3 could reverse CRC cell resistance to EGFR inhibitor cetuximab (data not shown). Besides, YY1 was reported to promote cancer growth through inhibiting p53 and activating Wnt signaling pathways40. These oncogenes were frequently overexpressed in CRC and our data showed that circHIPK3 upregulated these proto-oncogenes expression to promote CRC progression, suggesting that circHIPK3 was an oncogene in CRC. Further studies will be required to investigate the role of circHIPK3 in other malignancies.

In summary, our findings provide robust evidence that c-Myb transcriptionally elevates circHIPK3 and circHIPK3 serves as a novel oncogenic circRNA by sponging miR-7, as well as a promising prognostic biomarker in CRC. Our data also suggest that targeting
Fig. 6 CircHIPK3 silencing retards tumor progression and combined with miR-7 overexpression exhibits a magnifying effect on tumor repression in vivo. 

**a** The images of tumor-bearing nude mice from four treatment groups (n = 8 for each group) on the 35th day. Tumor volumes were monitored with digital calipers during the time course of five weeks, and tumor weights were also measured at the end of this study. Scale bar = 1 cm.

**b** H&E staining for tissue morphology and IHC staining analysis of proliferation (Ki67), apoptosis (CC3, cleaved caspase-3), invasion (MMP9) and vasculature (CD34). Arrowheads denote blood vessels. Scale bar = 50 μm.

**c** Representative images of lung metastatic nodules in four treatment groups (n = 8 for each group). Arrowheads denote the lung metastatic nodules stained by H&E. Scale bar = 50 μm. The corresponding statistical plots were presented in right panel. **p < 0.01 vs control group, ##p < 0.01 vs agomir-7 or si-circHIPK3 group.**

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**RETRACTED ARTICLE**
the c-Myb/circHIPK3/miR-7 axis as a potential treatment strategy for fighting CRC.

Materials and methods

Cell cultures and patient tissues

Human normal colon epithelial cell (FHC) and colorectal cancer cell lines (HCT116, HT29, SW620, DLD1) were purchased from American Type Culture Collection (Manassas, VA, USA) and were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (Gibco, Vienna, Austria). Cells in this medium were placed in a humidified atmosphere of 5% CO2 at 37 °C. All cell lines were free of mycoplasma contamination.

The 178 samples of CRC and 40 adjacent normal formalin-fixed, paraffin-embedded (FFPE) tissues were obtained from patients during operation at Affiliated Nanjing First Hospital of Nanjing Medical University (Nanjing, China). The patient characteristics were showed in Table 1. None of these patients received preoperative chemotherapy or radiotherapy. This study was approved by the ethics committee of Nanjing First Hospital and written informed consent was obtained from each patient.

Quantitative reverse transcription polymerase reaction (qRT-PCR)

Total RNA was extracted from tissues and cells by using TRIzol reagent (Invitrogen, CA, USA). For miRNA, the expression was determined by stem-loop primer SYBR Green quantitative real-time PCR (RiboBio, Guangzhou, China). For circRNA and mRNA, Total RNA was reverse transcribed to cDNA and then qPCR was conducted by using a SYBR Green PCR Kit (Takara, Otsu, Japan). All primer sequences were designed and synthesized by RiboBio (Guangzhou, China). GAPDH was chosen as the reference gene for circRNA and mRNA. U6 was chosen as an internal control for miRNA. Gene expression was quantified using the $2^{-\Delta\Delta C_t}$ method.

Actinomycin D and RNase R treatment

The culture medium was added to Actinomycin D (2 mg/ml) or DMSO (Sigma-Aldrich, St. Louis, MO, USA) to assess the stability of circHIPK3 and its linear isoform. Total RNA (10 μg) was incubated with 40U RNase R (Epicentre Technologies, Madison, WI, USA) at 37 °C for 60 min. After treatment with Actinomycin D and RNase R, the expression levels of HIPK3 and circHIPK3 were determined by qRT-PCR.

Oligonucleotide transfection

si-circHIPK3, si-c-Myb, miR-7 mimics, and their respective control oligonucleotides were synthesized by Gene-Pharma (Shanghai, China). Transfections were performed with final concentrations of 50 nM of miRNA mimics and siRNAs using the Lipofectamine 2000 reagent (Invitrogen) according to the manufacturer’s protocol.

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The schematic cartoon of the c-Myb/circHIPK3/miR-7 axis in CRC. In CRC, circHIPK3 is frequently upregulated partly caused by the transcription factor c-Myb overexpression, then circHIPK3 can sponge more endogenous miR-7 to sequester and inhibit miR-7 activity, thereby leading to increased proto-oncogenes (FAK, IGF1R, EGFR, and YY1) expression, which promoting CRC development and progression.
Plasmids construction and stable transfection

The method of constructing stable circHIPK3 or c-Myb overexpression cell lines has been described in the previous study11. In brief, human circHIPK3 and c-Myb cDNA was synthesized and cloned into pcD-ciR and pcDNA3.1 vector (Geenseed Biotech, Guangzhou, China), respectively. Then, HCT116 and HT29 cells were transfected with these plasmids, followed by selected with G418.

Luciferase reporter assay

For the promoter of circHIPK3 luciferase reporter assay, the wild-type and mutant promoters of circHIPK3 were synthesized and inserted into pGL3-basic vectors (GeneCreat, Wuhan, China), then the pGL3-basic-circHIPK3-wt/mut vectors and pcDNA3.1-c-Myb vectors were co-transfected with pRL-TK into HCT116 and HT29 cells using Lipofectamine 2000 reagent. For circHIPK3 and miR-7 luciferase reporter assay, the circHIPK3 sequences containing wild-type or mutated miR-7 binding sites were respectively synthesized and inserted into pmirGLO luciferase vector (GeneCreat, Wuhan, China), and then co-transfected with miR-7 mimics into HCT116 and HT29 cell lines were accomplished using Lipofectamine 2000. Cells were harvested at 48 h after transfection and luciferase activity was detected by the dual-luciferase reporter assay system (Promega). Relative luciferase activity was normalized to the Renilla luciferase internal control.

Chromatin immunoprecipitation assay

Chromatin immunoprecipitation (ChIP) assay was conducted using the ChIP Assay Kit (Biotinyl, Shanghai, China) according to the manufacturer’s guidelines with slight modifications. In brief, HCT116 and HT29 cells were treated with 1% formaldehyde for 10 min and quenched with 0.125 M glycine. Then the lysate was fixed with 4% paraformaldehyde and stained with Apollo Dye Solution for proliferating cells, followed by mounted with Hoechst 33342. Then the EdU-positive cells were photographed and counted under an Olympus FSX100 microscope (Olympus, Tokyo, Japan) in five randomly selected fields.

Apoptosis analysis

Cell apoptosis was analyzed using the Annexin V-FITC/Propidium Iodide (PI) Apoptosis Detection Kit (BD Biosciences, #556547) according to the manufacturer’s protocol. After incubation for 24 h, the HCT116 and HT29 cells were collected and then analyzed by fluorescence-activated cell sorting using FACScan (BD Biosciences, San Jose, CA, USA). The cell apoptosis data were analyzed by Flowjo V10 software (Tree Star, San Francisco, CA, USA).

Transwell migration and invasion assays

The cell migration and invasion assays were conducted by using transwell chamber (Corning, NY, USA), which was coated with (invasion assay) or without (migration assay) the matrigel mix (BD Biosciences, CA, USA) according to the manufacturer’s protocol. After incubation for 24 h, the cells located on the upper surfaces of the transwell chambers were scraped with cotton swabs and the cells located on the lower surfaces were fixed with methanol for 10 min, followed by stained with crystal violet. Then the stained cells were photographed and counted in five randomly selected fields.

RNA fluorescence in situ hybridization

The RNA fluorescence in situ hybridization assay was performed by using Fluorescent In Situ Hybridization Kit (Ribobio, Guangzhou, China) according to the manufacturer’s guidelines. After incubation for 24 h, the cells located on the upper surfaces of the transwell chambers were scraped with cotton swabs and the cells located on the lower surfaces were fixed with methanol for 10 min, followed by stained with propidium iodide and Hoechst 33342. Then the EdU-positive cells were photographed and counted under a confocal microscope.

Biotinylated RNA pull-down assay

The pull-down assay with biotinylated RNA was performed as described11,41. In brief, for circHIPK3 pulled down miRNAs, the biotinylated-circHIPK3 probe was
incubated with C-1 magnetic beads (Life Technologies, Carlsbad, CA, USA) to generate probe-coated beads, then incubated with sonicated HCT116 and HT29 cells at 4 °C overnight, followed by eluted and qRT-PCR. For miR-7 pulled down circHIPK3, HCT116 and HT29 cells with circHIPK3 overexpression were transfected with biotinylated miR-7 mimics or mutant using Lipofectamine 2000. The cells were harvested, lysed, sonicated, and incubated with C-1 magnetic beads (Life Technologies, Carlsbad, CA, USA), followed by washed and qRT-PCR.

Western blotting

The HCT116 and HT29 cells were lysed in RIPA lysis buffer. Then, equal amounts of protein were resolved by SDS-PAGE analysis and electrotransferred onto a PVDF membrane (Millipore, Schwalbach, Germany), then blocked with 5% skim milk powder and incubated with primary antibody at 4 °C overnight. The primary antibodies used were anti-FAK (#3285, Cell Signaling Technology), anti-IGF1R (#ab39675, Abcam), anti-EGFR (#4267, Cell Signaling Technology), anti-YY1 (#66281-1-lg, Proteintech), anti-GAPDH (#ab181602, Abcam). Then the membranes were incubated with HRP-conjugated secondary antibody for 1 h at room temperature, the blots were visualized using an enhanced chemiluminescence kit (Pierce, Waltham, MA, USA).

Immunohistochemistry (IHC)

IHC was performed on formalin-fixed, paraffin-embedded tissue sections as described previously. Primary antibodies against Ki67 (#ab15580, Abcam), cleaved caspase-3 (#ab2302, Abcam), MMP9 (#ab3885, Abcam), and CD34 (#ab81289, Abcam) were used. The complex was visualized with DAB complex, and the nuclei were counterstained with hematoxylin. All sections were scored by the semi-quantitative H-score approach and validated by two experienced pathologists.

Animal experiments

For xenograft tumor model, 5-week-old male BALB/c nude mice were randomly divided into four groups (n = 8 for each group). HCT116 (5 × 10⁶/0.2 ml PBS) cells were subcutaneously inoculated into the right flank of each nude mouse. Tumor volumes were measured every 3 days with digital calipers and were calculated by the following formula: tumor volume = 1/2 × (length × width). After ~10 days, when the volume of the tumor reached about 100mm³, cholesterol-conjugated si-circHIPK3 alone, agomir-7 alone or both (10 nM) (Gene-Pharma, Shanghai, China) in 0.1 ml of saline buffer were locally injected into the tumor mass, respectively. The injections were performed seven times at an interval of two days between each injection (i.e., day 10, 12, 14…). Thirty-five days later, the mice were killed and the volume and weight of the tumors were measured. Then the tumor tissues were harvested for use in further hematoxylin and eosin (H&E) and IHC staining.

For metastasis assay, HCT116 (1 × 10⁶/0.2 ml PBS) cells with corresponding treatment were tail-vein injected into 32 five-week-old male BALB/c nude mice which were randomly divided into six groups (n = 8 for each group). Seven weeks later, the mice were killed, and all the lungs are surgically removed and the number of macroscopically visible pulmonary metastasis nodules per mouse was counted by two experienced pathologists. Then the lung tissues were fixed in 10% neutral phosphate-buffered formalin followed by HE staining. The animal experiments were approved by the Animal Care Committee of Nanjing Medical College (acceptance no.: SYXK20160006).

Statistical analysis

Statistical analyses were performed using SPSS 22.0 (IBM, SPSS, Chicago, IL, USA) and figures were produced using GraphPad Prism 6.0 or Originpro.9.0. Differences between the different groups were tested using the Student’s t-test or one-way ANOVA. Kaplan–Meier method was used to evaluate the survival rate and analyzed by log-rank test. The correlations were analyzed using Pearson’s correlation coefficients. The univariate and multivariate analyses were analyzed using Cox proportional hazards models. All experimental data were presented as the mean ± S.D. of at least three independent experiments. The differences were considered to be significant at p < 0.05.

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Author details

1General Clinical Research Center, Nanjing First Hospital, Nanjing Medical University, 210006 Nanjing, China. 2School of Medicine, Southeast University, 210009 Nanjing, China

Conflict of interest

The authors declare that they have no conflict of interest.

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