Myeloid-derived suppressor cells promote tumor growth and sorafenib resistance by inducing FGF1 upregulation and fibrosis

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Abstract

Background

Considerable evidence implicates myeloid-derived suppressor cells (MDSCs) promote tumor progression and drug resistance. Sorafenib is the standard first-line therapy for advanced hepatocellular carcinoma (HCC). Clinical evidence indicates that sorafenib resistance is associated with increased MDSCs, by which MDSCs exerts these effects is obscure. This study aimed to investigate the mechanism of sorafenib resistance mediated by MDSCs.

Methods

A syngeneic mouse-liver cancer cell line BNL was subcutaneously injected to build a tumor-bearing mouse model, and syngeneic MDSCs were adoptive transferred into the tumor-bearing mouse. Tumor tissue was obtained, and transcriptomic analysis of the tumor was carried out on RNAseq data. A coculture system was used to verify the crosstalk between MDSCs and BNL cells.

Results

Adoptive MDSCs transfer into tumor-bearing mice induced an increase of tumor-infiltrating MDSCs, which led to tumor growth and impaired antitumor activity of sorafenib in BNL HCC models. MDSCs transfer contributed to tumor fibrosis and tumor-associated fibroblast (CAF) activation, associated with fibroblast growth factor (FGF1) upregulation. In contrast, MDSC depletion by anti-Ly6G antibody reduced fibrosis and increased sorafenib antitumor efficacy. Intriguingly, tumor-infiltrating MDSCs barely expressed FGF1. IL-6 derived from MDSCs increased FGF1 expression in BNL liver cancer cells, and anti-IL-6 attenuated this effect in vitro. MAPK pathway, one of the sorafenib targets, is the downstream signaling of FGF1 and is reactivated by MDSCs-mediated FGF1 upregulation.

Conclusions

Our finding demonstrated that MDSCs led to tumor growth and sorafenib resistance via FGF1 upregulation and subsequent indirect CAF activation. We offered a novel mechanism of MDSCs-driven HCC progression and sorafenib resistance.

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Keywords: Myeloid cells, Hepatocellular carcinoma, Sorafenib, Fibroblast cells, Combination therapy

Abbreviations: CAF, tumor-associated fibroblast; Col, collagen formation; CXCR4, C-X-C receptor type 4; ECM, extracellular matrix; FGF1, fibroblast growth factor 1; HCC, hepatocellular carcinoma; H&E, hematoxylin-eosin staining; IF, immunofluorescence; IHC, immunohistochemistry; LAMA1, laminin alpha 1; MAPK, Mitogen-activated protein kinase; MDSC, Myeloid-derived suppressor cells; PDGF, platelet-derived growth factor receptor; pERK, phosphorylated extracellular regulated protein kinases; SDF, stromal-derived factor 1; TME, tumor microenvironment; THBS 1, thrombospondin-1; VEGFR, vascular endothelial growth factor receptor; VTN, vitronectin; VWF, von willebrand factor.

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Introduction

Hepatocellular carcinoma (HCC) represents the sixth most commonly diagnosed cancer and the third leading cause of cancer-related mortality [1]. Furthermore, due to lack of effective treatment and frequent resistance, the long-term survival rate has not been significantly improved. Further emphasizing the urgent need to deepen our understanding of HCC progression and drug resistance. The tumor microenvironment (TME) comprises immune cells, stromal cells, fibroblasts and endothelial cells, which plays a complex and multidirectional interplay in remodeling characteristics of tumor cells. Therefore, understanding the interaction mechanism between nontumor cells and tumor cells is essential.

Myeloid-derived suppressor cells (MDSCs), a heterogeneous population of immature myeloid cells, induce local and possibly systemic immunosuppression and tumor progression [2]. There are two different types of MDSCs, as identified in studies in both mice and humans: polymorphonuclear MDSC (PMN-MDSC) and monocyted MDSC (M-MDSC) are similar to monocytes. The phenotypic characteristics of murine MDSCs have been described CD11b+Ly6G+ (PMN-MDSC) and CD11b+Ly6G− (M-MDSC). With potent immunosuppressive activity, both PNMM-DSC and M-MDSC have the ability to support tumor progression, although differences in the phenotypic characteristics. MDSCs correlate with early recurrence and poor prognosis in patients with HCC who underwent curative resection [3], radiotherapy [4], and hepatic arterial infusion chemotheroy [5]. Accompanied by tumor progression, MDSCs were also acceleratedly expanded, activated and released from the molecular from TME [2]. MDSCs drive HCC progression by suppressing CD4+ T cells, CD8+ T cells, natural killer cells, and the paracrine release of cytokines and chemokines with angiogenesis and recruitment [6]. Little attention, however, has been devoted to analyzing the direct effect of MDSCs on the tumor cells.

Although it offers very moderate survival improvements, Sorafenib is one of the most effective single-drug therapies for advanced HCC [7]. Sorafenib inhibits Raf, as well as the kinase activity of vascular endothelial growth factor receptor (VEGFR) and platelet-derived growth factor receptor (PDGFR) led to antiangiogenic effects. Immunotherapy is promising but depends on combination regimens which has a lower cost-effectiveness [8,9]. Therefore, it is crucial to understand the molecular mechanisms mediating sorafenib resistance to improve clinical outcomes for HCC patients. Here, we investigated the mechanism of MDSC-driven HCC progression and sorafenib resistance.

Methods and materials

Cell line and cell culture

BNL 1ME A.7R.1 (referred as BNL in this study), a babl/c mouse-derived HCC cell lines, were procured through the Shanghai Xin Yu Biotechnology(Shanghai, China) and were routinely maintained. They were tested for mycoplasma contamination by a Mycoplasma Detection Kit. MDSCs were isolated from the peripheral blood of BNL-bearing mice (21 days after inoculation). Then the purity of MDSCs were identified by flow cytometry analysis (FCAS)>90% purity, defined by CD11b+Gr1+ MDSCs). The expression of representative effector molecules of MDSC in Arginase1(Ang1), TGF-β, IL-6 (Fig. S1) and reactive oxygen species(ROS) (Fig. S2) were detected. To effectively simulate the tumor microenvironment, BNL were noncontact cocultured with MDSCs isolated from PBMC of tumor-bearing congenic mice through Transwell kit (Corning #3450). In particular, BNL cells(2 × 10^6 cells/well) were suspended in 1.5 ml serum-free medium in the top of insert well, and 5 × 10^4 well MDSCs were seeded in the lower chamber in 2.6 ml complete medium supplemented with 20 ng/ml GM-CSF. After 48 h cultivation, BNL cells were treated with DMSO or sorafenib 5 μM for 24 h.

Animals

Male babl/c mice (6 weeks) were purchased from the Animal Center of Southern Medical University and were maintained under specific pathogen-free conditions. Additional information on methods is available in the supplement.

Flow cytometric analysis

Single cells were stained by fluorecence-conjugated antibodies (Biolegend) as previously described [10]. Stained cells were analyzed using the FACS Analyzer (ECS Calibur, BD). The results were analyzed with FlowJo10.0 software. Gating strategy for the MDSC subsets are presented in Fig. S3. Additional information on methods is available in the supplement.

Hematoxylin-eosin staining(He&E), Mason’s trichrome staining, immunohistochemistry(IHC), immunofluorescence(IF)

HE staining was conducted according to standard protocols. Mason’s trichrome staining were performed by kit(Proteinbio, China)according to the instruction. For IHC, the slices were disposed by Elivision™ plus Polyer HP (Mouse) IHC Kit(MXB, China) and incubated with CD31 (1:2000, abcam ab182981). For immunofluorescence, the tissue slices were incubated with α-SMA (1:500, abcam ab124964) and VEGFA primary antibody (1:500, abcam ab52917) and with Alexa Fluor 488/Dylight-594 conjugated secondary antibody (1:400 Abbkine, Wuhan, China).

RNA-seq

Total RNA was extracted from the frozen tumor tissue using TRIzol® Reagent following the manufacturer’s instructions (Invitrogen). Genomic DNA was removed by DNase I (TaKara Jan). RNA quality was examined by 2100 Bioanalyzer (Agilent) and quantified using the ND-2000 (NanoDrop Technologies). The methods of library preparation, Illumina Hiseq xten/Nova seq 6000 Sequencing, read and Differential expression and functional enrichment analysis were described in the supplement.

Western blotting and qRT-PCR

The protein of cells and tumor tissue were extracted, quantified, separated, transferred and blocked as described previously [11]. The membranes were incubated with primary antibodies (β-actin, Affinity; GAPDH, Proteintech; FGF1, Ras, ERK, pERK, VEGFA, PDGFB and α-SMA, Abcam) overnight at 4 °C. Protein bands were quantified using ImageJ software with β-actin or GAPDH as the internal control. For qRT-PCR, total RNA was extracted from cells and tumor specimens using trizol reagent (RighGene, Guangzhou, China) following the manufacturer’s instructions. The expression of mRNA was measured via qRT-PCR using a SYBR PrimeScript RT-PCR Kit (Takara Bio, Shiga, Japan) in accordance with the manufacturer’s instructions. GAPDH was used as an internal control. Primer sequences for various genes are listed in Table 1.

MILLIPLEX® MAP array

Cytokines in the supernatant of cell culture were measured by Mouse High Sensitivity. Magnetic Bead Panel MILLIPLEX® MAP kit (Merck, Germany) following the manufacturer’s instructions.
Table 1

| Gene  | Forward primer             | Reverse primer             |
|-------|----------------------------|----------------------------|
| IL-6  | TAGTCTCTTCTCACCACATTCC     | TGGTCCTTAGCCACTCTCC       |
| TGFβ  | CTCCCGTGCGTTCTAGTGCGC     | GCTTTAGTTTGGCAGGATCTCG   |
| Arg1  | CTCCAGGCAAAGCTCTTAGAGAG  | AGGAAGCTGTCAATAGGACATCG  |
| FGF1  | CCGTCGAGGAGGTCTTCAAC      | GTCCCTGTGCCCCATCCAG      |
| PDGF  | CATCGGCTCTTGTGATGATCTTT   | GTGTCGGGCGTTGCTGCTAAGT   |
| GAPDH | AGGTCCGTTGTGAACGGATTG     | TGTAAGACCATGTAGGGTCA     |

Statistical analysis

The variances between the groups that are being statistically compared were similar. Quantitative analyses were done using Graphpad prism 5 (GraphPad Software, Inc. USA). Bubble chart and heatmap was carried out in R software. For experiment in vitro, all data were collected from at least three independent experiments. All data was expressed as mean ± SEM. ANOVA and t-test were used when data accorded with normal distribution and homogeneity of variance. Tukey test was used multiple comparisons. A P-value < 0.05 indicated that the difference was statistically significant.

Results

MDSC promoted tumor growth and attenuated the efficacy of sorafenib in vivo

We tested the effect of MDSC in a murine subcutaneous HCC model using BNL HCC cells. Syngeneic MDSCs were generated from peripheral blood of tumor-bearing mice. They were initiated coinjection with BNL cells (1:10) and re.injected into the tumor at the same amount weekly for three weeks (Fig. 1A). To confirm the mice were successfully transferred MDSC, we examined the number of MDSC in the tumor by Flow cytometric analysis. PMN-MDSC(CD11b+Ly6G+) was significantly increased in tumors of mice with transferred MDSCs, while the number of M-MDSC(CD11b+Ly6G+) was unchanged (Fig. 1B,C, P<0.0001), suggesting that transferred MDSCs promoted PMN-MDSC infiltrating in tumor. In contrast, a reduction of MDSCs was observed in the tumor of mice with sorafenib treatment. Then we analyzed the tumor growth based on tumor volume and weight. As shown in Fig. 1D and E, sorafenib significantly suppressed tumor growth (P = 0.0071) but caused only marginal inhibition in mice transferred MDSCs (P = 0.5097).

MDSCs boosted synthesis of extracellular matrix (ECM)

For mechanistic insights, the histopathology of tumor tissue was examined. With H&E staining, we noted a dramatic increase of fibrosis in the tumor from mice with transferred MDSCs (Fig. 2A). In the tumor of mice with transfer MDSCs, we observed a number of collagen fibers diffused around the tumor cells by Masson staining. Similar phenomena were also observed in the mice transferred MDSCs and treated with sorafenib (Fig. 2B), indicating that MDSCs might induce tumor fibrosis. To further understand this finding, we employed RNA-seq to analyze the gene expression profile of tumors from two major groups’ mice (treated with sorafenib and transferred with MDSCs vs. non-treated by sorafenib). As a result, the Kyoto encyclopedia of genes and genomes enrichment analysis showed significant enrichment for genes of the ECM-receptor interaction (Fig. 2C). And genes upregulated were significantly enriched for signatures related to collagen formation (Col), ECM organization (Col, Thbs, Lam, Vtn, Vwf), and cell motility (ITGA) (Fig. 2D). Genes encoding ECM-related proteins, including thrombospondin-1 (THBS 1), laminin alpha 1 (LAMA1), vitronectin and collagen IV were significantly enriched in tumor transferred MDSCs (Fig. 2E P = 0.0003). CAF is the synthetic machine of ECM [12], the genes of whose defining markers α-SMA (encode by Acta2 gene) and FAP were also enriched in the MDSC+ Sorafenib group [12,13]. Overall, these data suggested transferred MDSCs could cause the synthesis of ECM.

MDSCs induced CAF activation by upregulating FGF1 levels

Next, we examined the expression of α-SMA in tumor tissue, and the mice transferred MDSCs expressed increased α-SMA versus controls (Fig. 3A P<0.001). TGFβ, platelet-derived growth factor (PDGF), and FGF are reported the key mediators of CAF activation [13]. We examined the expression of FGE, TGFβ, and PDGF in tumor. The transcript expression level of FGF(FGF1, FGF18) significantly upregulated, whereas PDGF and TGFβ of that did not show an observable difference in mice transferred MDSCs (Fig. 3B,C). This result was further verified in vitro by the coculture of MDSC and BNL (Fig. 3D,E,F). Taken together, we demonstrated that MDSCs might induce CAF activation, which was associated with FGF1 upregulation.

IL-6 was a potential mediator of cross-talk between BNL HCC cells and MDSCs

We next questioned whether MDSCs secreted FGF1 and activated CAF based on the findings described above. It was consistent with previous studies, we found that FGF was hardly expressed in MDSC but in BNL (Fig. 4A) [14]. Emerging research suggests that tumor cells are inescapably responsible for CAF activation and proliferation, but the exact mechanism is unclear [15]. We presumed that MDSC-induced FGF1 upregulation of BNL tumor cells drove CAFs activation. To verify this inference, BNL were cultured in the presence (coculture) or absence of MDSC for 48h. As shown in Fig. 4B, the BNL cells cocultured with MDSC showed altered morphology compared to the control. Then, FGF1 expression of BNL cells was analyzed. Sorafenib decreased FGF1 expression, which could be partly reversed by the presence of MDSC (Fig. 4C P = 0.0002). To understand the mediator contributing FGF1 upregulation, we studied cytokines upon coculture system and found IL-6, one cytokine that can be secreted by MDSCs, significantly increased in the coculture system and MDSC supernatant (Fig. 4D). As shown in Fig. 4E, the IL-6 neutralizing antibody (10 ng/ml) blocked cells from MDSC-induced FGF1 upregulation (P<0.0001).

Ly6G+ myeloid cells depletion reduced fibrosis and enhanced sorafenib efficacy

To confirm whether MDSCs was responsible for tumor fibrosis, neutralizing antibody(Anti-Ly6G, a in vivo antibody) was injected into
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Fig. 1. MDSC promoted tumor growth and attenuated the efficacy of sorafenib in vivo. (A) Schematic diagram of the establishment of MDSCs transfer model. In short, homologous HCC cell line BNL(3 × 10^6/100 μL) was subcutaneously injected in balb/c mice to build tumor-bearing mice model. 3 weeks later, MDSCs were isolated from the PBMC of tumor bearing mice. Then, MDSCs were mixed with BNL cells and co-injected subcutaneously. MDSCs were injected into the tumor at the indicated dosage weekly for 3 consecutive weeks. (B) Representative flow cytometric plot of PMN-MDSC and M-MDSC. (C) Quantification of the number of fluorescently labelled MDSC from 100mg tumor tissue (n = 3). (D) Tumor volume was measured using a Vernier caliper, V = a × b^2/2, where a is the long diameter and b is the short diameter (n = 5, two-way ANOVA). (E) Tumor weight at experimental end stage (n = 5). (C), (D), (E) MDSCs condition was normalized to the control condition. Data are reported as mean ± SE. Specific n values of biological independent animals. ANOVA for four groups comparisons and Tukey test for multiple comparisons. Asterisks indicate comparison with the control group. *p < 0.05, **p < 0.01, ***p < 0.001.

mice to decrease (PMN-MDSC)s. The addition of anti-Ly6G mAb significantly improved the ability of sorafenib to suppress tumor growth (P<0.0001), while anti-Ly6G mAb alone did not present significant suppression for tumor (Fig. 5A P>0.05). Anti-Ly6G did not significantly decrease IL-6 level in the tumor (Fig. 5C), although with an MDSC depletion (Fig. 5B P<0.0001). The reason is that a variety of cells in the tumor microenvironment can secrete IL-6, yielding conflicting results. Moreover, Ly6G⁺ myeloid cells depletion reduced collagenous fiber (Fig. 5D P<0.01) and showed a correlation with the number of MDSCs.
Fig. 2. MDSCs boosted synthesis of extracellular matrix (ECM). (A) Representative images of H&E staining of tumor (Scale bar = 200 μm). (B) Masson staining of tumor tissue, collagen fibers were stained in blue (Scale bar = 100 μm). Collagen volume fraction was analyzed by ImageJ (n = 3). (C) Bubble chart showed kyoto encyclopedia of genes and genomes enrichment analysis for the differential expression of genes between Sorafenib group and MDSCs+Sorafenib group. (D) Heatmap displaying quantification of ECM-receptor interaction gene. (E) The gene expression of THBS1, LAMA1, vitronection and collagen IV, which encode ECM. (F) The gene expression of Acta2 (encode α-SMA protein) and Fap, which are the makers of CAFs. Data are expressed as mean ± SEM of n = 5 or 3 independent animals. t – test for two groups comparisons. *p < 0.05, **p < 0.001.
Myeloid-derived suppressor cells promote tumor growth

MAPK signaling re-activated led to angiogenesis and resistance to sorafenib

Mitogen-activated protein kinase (MAPK) might exert a central role in these findings, for the reason that which is the downstream of FGF1R activation as well as one of the targets of sorafenib [16,17]. Thus, we detected the expression of major molecules (FGF1, Ras, ERK and VEGFA) of the MAPK pathway. We showed sorafenib drastically suppressed the expression of phosphorylated extracellular regulated protein kinases (pERK) in vivo and vitro (Fig. 6A,B), and MDSCs impaired this effect in vitro (Fig. 6B). A similar result was shown by gene set enrichment analysis (GSEA) (Fig. 6C FDR = 0.0374). CD31, an endothelial cell marker, revealed that microvessel density decreased in tumor tissues of the sorafenib group and significantly increased in that derived from mice with the adoptive transfer MDSCs, whether or not treated with sorafenib (Fig. 6D P<0.0001). VEGFA staining showed a similar result (Fig. 6E P<0.0001). These results suggested MDSC re-activated MAPK pathway which mediated angiogenesis and sorafenib resistance.
Discussion

Growing evidence implicating MDSCs mediated the progression, poor prognosis, and therapeutic resistance of HCC, which are mainly linked to its immunosuppressive function [18]. Erewhile, several studies suggest that MDSCs directly impact tumor cells’ characteristics [19, 20] while underlying mechanisms are still unknown. Our study indicated that MDSCs promoted the FGF1 expression in BNL liver cancer cells and IL-6 is a mediator.

In a mouse xenograft model, we observed increased tumor growth and angiogenesis accompanied by tumor fibrosis and improved synthesis of ECM in tumors derived from the mice transferred MDSCs. With the depletion of MDSCs by anti-Ly6G, sorafenib significantly inhibited tumor growth, accompanied by reduced tumor fibrosis and angiogenesis. Several studies provided evidence that MDSC reduction positively correlated with sorafenib response and prognosis [21–23]. Targeting Ly6G improves sorafenib efficacy [24, 25]. However, models of HCC have not yet conclusively clarified the involvement of MDSC for tumor fibrosis. It is reported that MDSCs regulate fibrosis in the lungs, liver, and kidneys and help repair the central nervous system in non-neoplastic inflamed organs [26]. In HCC mice models, MDSCs accumulated in fibrotic livers, which significantly correlated with reduced tumor-infiltrating lymphocytes and increased tumorigenicity [25]. Chen’s research found that sorafenib caused tumor hypoxia and an increase in MDSCs, which promoted tumor fibrosis via the stromal-derived factor 1 (SDF) alpha/C-X-C receptor type 4 (CXCR4) axis [27]. We found sorafenib reduced the number of MDSCs, the different results of which might be attributable to the subcutaneous HCC model [24]. HIF, SDF, and CXCR4 were consistently elevated in the tumors of mice transferred MDSCs in this study.

FGF1, PDGF, and TGFβ pathways are the primary mechanisms contributing to CAF activation [13]. In this study, FGF1 significantly differed between groups. FGF1 activation plays a positive role in the expansion of CAFs through transcriptional repression of TP53 and escapes from p53-dependent stroma cell senescence [28] and section with induction of several key CAF effector genes(MMP1, HGF) [29]. However, no study answers whether MDSCs directly act on fibroblasts. Our finding showed that MDSC barely expresses FGF1, as described in another study. As previously speculated, the stimulation of MDSC-induced fibrosis of tumor tissue may be an indirect effect [14]. In TME, tumor cells and macrophages can mediate MDSC-induce fibrosis. Tumor cells are the central member of tumor tissue and can be effect by MDSC. A study in head and neck squamous cell carcinoma shows that tumor cell-secreted FGF1 increases mitochondrial oxidative phosphorylation and promotes lactate consumption of CAFs [30]. We carried out a series of experiments to support the conjecture. In several disease models, MDSC differentiated into macrophage-like phenotypes. Nevertheless, references to the role of macrophages on tumor fibrosis are contradictory [31]. More research is needed to understand this complex question. All in all, our findings and previous research showed that MDSCs triggered tumor fibrosis, facilitating enhanced growth, angiogenesis, and resistance to sorafenib.

Cytokines are essential mediators linking inflammation and cancer and exert vast immunoregulatory actions to tumor cells [32]. To deeper understand the interaction between MDSCs and BNL, we analyzed the most common mouse 18 cytokines. Four cytokines (GM-CSF, IL-4, TNF-alpha, and IL-6) secreted more in coculture than in monoculture. Among them, IL-6 was shown to have the most dramatic and specific increase(more than 3-fold) in secretion. Anti-IL6 neutralized the effect of MDSC. Our data showed that
MDSC-secreted IL-6 was essential for FGF1 upregulation of BNL cells. IL-6 is a prototypical cytokine that exerts a pro-tumorigenic role in inflammation-associated cancers [33]. Serum IL-6 concentrations were significantly higher in patients with HCC than healthy donors [34]. IL-6/IL-6 receptor promotes epithelial-mesenchymal transition progression, cancer stenosis of triple-negative breast cancer [35]. Anti-IL-6/IL-6 receptor monoclonal antibodies developed for tumor-targeted therapy have demonstrated promising results in preclinical studies and clinical trials [36]. We provided the first evidence that MDSCs secreted IL-6, a potential mediator of tumor progression and resistance to sorafenib in HCC. Notably, we only analyzed the most common mouse 18 cytokines and did not verify other cytokines secreted by MDSCs. In addition, we found IL-1, CXCL2, CCL2, CXCL5 were decreased in the coculture system compared with tumor cells in mono-culture, which might result from the negative regulation of GM-CSF [37].

Our findings are presented with a limitation that only a single cell line was carried out to investigate the crosstalk among MDSC, HCC, and CAFs. Further study will be performed in multiple cell lines. Nevertheless, a recent study supports the validity of our findings. This study showed that FGF1 upregulated in CAF of HCC patients at all stages and high levels of tumor-associated neutrophils infiltrated in the tumor, which indicated a poor prognosis among patients in middle and late stage. Considering nomenclature of immunosuppressive neutrophils, multiple populations of neutrophils with immunosuppressive features might be PMN-MDSCs. In addition, another clinical and preclinical study demonstrated that tumors with sorafenib-acquired resistance were enriched with FGF signaling cascades and FGF blockade delayed tumor growth and improved survival in sorafenib-resistant tumors, although variations of MDSCs was not mentioned.

In conclusion, we demonstrated that MDSCs facilitated CAF activation led to tumor growth, angiogenesis, and sorafenib resistance by inducing FGF1 upregulation. Our findings offer some insights into the crosstalk among MDSC, HCC and CAFs.
Fig. 6. MAPK signaling re-activated led to angiogenesis and resistance to sorafenib. (A) WB for MAPK pathway in tumor tissue (n = 5). (B) WB for MAPK pathway in BNL cells (n = 3). (C) GSEA of tumor tissue involved in MAPK pathways (n = 5). The normalized enrichment scores (NES) and tests of statistical significance (FDR) are shown. (D) Representative images (Scale bar = 200 μm) of tumor sections stained IHC with CD31 for sprouting neomicrovessels. Positive area was analyzed by ImageJ (n = 3). Specific n values of biological independent experiments. ANOVA for four groups comparisons and Tukey test for multiple comparisons. Asterisks indicate comparison with the control group. **p < 0.01, ***p < 0.001.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

CRediT authorship contribution statement

Xue Deng: Writing – original draft, Methodology, Formal analysis, Data curation, Writing – review & editing. Xueyan Li: Writing – review & editing, Methodology, Writing – review & editing. Xuan Guo: Methodology, Writing – review & editing. Yantong Lu: Methodology, Writing – review & editing. Yingjie Xie: Formal analysis, Data curation, Writing – review & editing. Xuhui Huang: Funding acquisition, Writing – review & editing. Juze Lin: Funding acquisition, Writing – review & editing. Wei Tan: Funding acquisition, Writing – review & editing. Changjun Wang: Conceptualization, Writing – review & editing, Funding acquisition.

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Ethics statement

The animal study was reviewed and approved by the Animal Care and Use Committee of Southern Medical University (Guangzhou, China).

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Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.neo.2022.100788.

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