

Mechanisms Underlying Therapeutic Effects Of Traditional Chinese Medicine On Gastric Cancer

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Abstract: Gastric cancer (GC) is one of the most common malignant tumors in the world. It is the fourth most common cancer and has the second highest mortality rate globally. Metastasis is an important feature of gastric cancer and is the most common cause of death. Exploring the mechanism underlying the metastasis of gastric cancer and searching for new drug targets has become the focus of several studies. Traditional Chinese medicine may show promise for treatment of gastric cancer. In this review, we report the recent progress in research on the anti-metastasis activity of Chinese medicine, to facilitate clinical development of treatments for gastric cancer.

Keywords: Chinese medicine, metastasis, gastric cancer, cell cycle, angiogenesis

Introduction

Gastric cancer is the most common malignant tumor of the digestive system. China accounts for 42.6% of gastric cancer cases globally,¹ and one of the main causes of death is metastasis of malignant tumors.² With the continuous study of traditional Chinese medicine, numerous studies have illuminated that in cancer treatment it can be used to enhance the efficacy and diminish the side effects. Chen et al.³ Found that Huaier make the effective of HCC after curative liver resection which can extend RFS and reduce extrahepatic recurrence. In addition, Von Hagens et al.⁴ illustrated that in a phase I study, patients who have metastatic breast cancer with Long-term oral artesunate can consolidate the original curative effect and safety. Deeken et al.⁵ indicated that intravenous artesunate is 18 mg/kg which the treatment of solid tumor was well tolerated and diminishing the side effect. Compared to chemotherapy and radiotherapy in Western medicine, traditional Chinese medicine places considerable emphasis on overall regulation, and it has been proved to be an effective therapy for reducing side effects, improving the survival rate, and prolonging the survival period in cancer model systems. There are some anti-radiation traditional Chinese medicine in radiotherapy.⁶ Traditional Chinese medicine is proved to be efficacious in increasing the sensitivity to chemo- and radio-therapies.⁷ And evidence from the meta-analysis suggested that traditional Chinese medicine combined with chemotherapy has a positive effect on gastric cancer treatment.⁸ It can be seen that traditional Chinese medicine is of great significant for assitant-therapy to decline the side effect, safe and tolerable. The purpose of this review is to clarify the mechanism by which traditional Chinese medicine helps in the treatment of gastric cancer and prevention of metastasis as shown in Table 1.

Inhibition Of Proliferation Of Gastric Cancer Cells

The proliferation of tumor cells is the foundation of tumor invasion and metastasis. Excessive proliferation of tumor cells at the primary focal center increases pressure and promotes invasion and metastasis in the direction of low pressure, resulting in metastasis to neighboring tissue and distant organs. Traditional Chinese medicine and its components can suppress the proliferation of gastric cancer by arresting the cell cycle, inhibiting telomerase activity, inducing apoptosis via the mitochondrial pathway, and regulating inflammation-related factors.

Cell Cycle Arrest

Uncontrolled cell proliferation is one of the fundamental biological characteristics of tumors, based on dysregulation of the cell cycle. Various tumor components operate through cell cycle arrest. Cell cycle-dependent kinase (CDK2 and CDK4) plays a key role in cell cycle progression and is closely related to the entry of cells into S and G1 phase.⁹ Cell cycle arrest of tumor cells by traditional Chinese medicine is related to the loss of cell cycle-dependent protein kinase inhibitory factor activity, thereby inhibiting the CDK suppression signal and leading to cell cycle regulation disorder. Several studies have reported that^{10,11} matrine (including sophocarpine, oxysophocarpine, and sophoridine), astragalus polysaccharides (including ascorbic acid, glucose, and fructose), and cinobufotalin (an anti-cancer drug) can induce G1 cycle arrest. Sun et al.¹² found that Coleusin factor induced cell cycle arrest at the G0/G1 stage in SGC-7901 cells. Other studies^{13,14} reported that berberine and aianthone induced G2/M cell cycle arrest and inhibited the proliferation of gastric cancer cells.

Inhibition Of Telomerase Activity

Recent studies have discovered that telomerase is expressed in more than 85% of cancers, making telomerase an attractive and viable target for the development of anti-cancer therapeutics.^{15,16} Telomerase may play an important role in the process of tumor proliferation. Moreover, inhibiting tumor cell telomerase activity through traditional Chinese medicine can effectively inhibit tumor cell proliferation and interfere with metastasis of tumor cells.

Çalışkan et al.¹⁷ found that telomerase activity regulates the expression of human telomerase reverse transcriptase (hTERT) in gastric cancer cells, and suggested

Table 1 Traditional Chinese Medicine On Gastric Cancer

Ingredients of TCD	TCD	Experiment	Gastric cancer cell line and Experimental animal	Dosage	Event/Mechanism	Effect	Refs&PMID
Prescriptions	Xiaotan Sanjie recipe	In vivo	BALB/c nude mice (5×106 MKN-45)	1.496g	VEGF-C and VEGFR-3	Metastasis	37 21425575
Herbs	Luteolin	In vitro	MGC-803, HUVECs, His-746T	10, 20, 30 μM (Main 30 μM)	Suppressed Notch1-VEGF signaling	Angiogenesis and VM formation Metastasis	32 28655612
	Tanshinone IIA (TSN)	In vitro	SGC-7901	1, 5, 20 μg/L	Regulated NF-κB, AKT, and MMP-9 signaling	Proliferation and Migration	50 28184921
	Baicalein	In vitro	SGC-7901, MKN45	15, 30, 60 μmol/L	Downregulation of FOXM1	Growth arrest Apoptosis	54 26918059
		In vivo	BALB/c nude mice (1×106 SGC-7901)	15, 50 mg/kg			
	Baicalein	In vitro	SGC-7901 and MGC-803	<80 μM	p38	Invasion	44 25502212

(Continued)

Table 1 (Continued).

Ingredients of TCD	TCD	Experiment	Gastric cancer cell line and Experimental animal	Dosage	Event/Mechanism	Effect	Refs&PMID
Crude extracts	Coleusifactor (CF)	In vitro	BGC-823	25, 50, 100 μ M	G0/G1 Decreased cyclinD1, Bcl-2, and Bcl-xL, increased the p53, p21, and Rb.	Growth arrest Proliferation and Apoptosis	12 21093146
	Berberine hydrochloride (BER)	In vitro	AGS	15 μ M	G2/M Enhanced IL -8 expression Affected ERK1/2, SAPK/JNK, JAK2 and AP-1 pathway	Growth arrest Proliferation and Metastasis	13 24063987
	Allanthone	In vitro	SGC-7901	0.5, 1, 2, 4, 8 μ M	G2/M Bcl-2 and Bax	Growth arrest Apoptosis	14 None
	Celastrus Orbiculatus extract (COE)	In vitro In vivo	AGS and BGC-823 Nude mice(GFP GC cells)	160 mg/L 40 mg/kg	PI3K/Akt/mTOR/p70s6k pathway Induced the caspase-dependent mitochondrial pathway	Proliferation Apoptosis	20 28476678
	Xiaotan Sanjie decoction	In vitro In vivo	GCSCs and MKN-45 Sprague-dawley rats and nude mice (1 \times 106 CD44+ and CD44- cells)	1. 46, 2.92 and 5.84 g/mL 1. 46, 2.92 and 5.84 g/mL	Notch-1, Hes 1, VEGF and Ki-67	Metastasis	34 25278704
	Yiqi Huayu Jiedu decoction	In vitro In vivo	MGC-803 BALB/C nude mice (5 \times 106 MGC-803)	2 mg/mL 4 g/mL, 2 g/mL	TGF- β /Smad	Invasion and Metastasis	39 28539961
	Aqueous Huaier Extract	In vitro In vivo	SGC-7901 and MGC-803 Zebrafish (3 \times 102 SGC-7901 labeled with CM-Dil)	0.1, 0.2, 0.3, 0.5 mg/mL (main 0.5 mg/mL) 1 mg/m	Decreased N-cadherin and vimentin	Metastasis	48 29151976
	Celastrus Orbiculatus extract (COE)	In vitro	MGC-803	20, 40, 80 μ g/mL	Promoted the expression of maspin Phosphoinositide 3-kinase/Akt/mTOR and MAPK signaling pathways	Apoptosis Invasion and Migration	55 29387218

(Continued)

Table 1 (Continued).

Ingredients of TCD	TCD	Experiment	Gastric cancer cell line and Experimental animal	Dosage	Event/Mechanism	Effect	Refs&PMID
Compounds	Matrines	In vitro	SGC-7901	0.25, 0.5, 1.0, 2.0 mg/mL	G0/G1 Bax	Growth arrest Apoptosis	10 21519796
	Gambogic acid (GA)	In vitro	BGC-823	1.4 μM	Repressed hTERT transcriptional activity via c-Myc and Akt	Telomerase activity	18 18226852
	Crocodile choline (CCL)	In vitro	BGC-823, MGC-803, SGC-7901, MKN28 and GES1	10, 20, 40, 60 μg/mL	G2/M ΔΨ _m and ROS Notch pathway	Growth arrest Apoptosis Proliferation and Metastasis	22 28347224
		In vivo	BALB/C nude mice (1 × 106 BGC-823)	100 mg/kg			
	Ginsenoside	In vivo	BALB/C nude mice (5 × 106 NUCC-4-RFP)	1 mg/kg	VEGF-C, VEGF-D and VEGFR-3	Proliferation and Metastasis	36 28314255
	Matrine	In vitro	BCG-823	50 μg/mL	Affected vasodilator-stimulated phosphoprotein (VASP)	Migration	45 23685951
	Silibinin	In vitro	SGC-7901	50, 100, 200 μM	Decreased MMP-2 and MMP-9	Migration and Invasion	43 29344204
	Astragalus polysaccharide	In vitro	AGS	200 μg/mL	Inhibited phosphorylated AKT (p-AKT) and MMP-9	Metastasis	49 29428665
	Naringenin	In vitro	SGC-7901	20, 40, 80 μM	Regulated AKT pathway	Proliferation, Apoptosis, Migration, and Invasion	52 26960693

(Continued)

Table 1 (Continued).

Ingredients of TCD	TCD	Experiment	Gastric cancer cell line and experimental animal	Dosage	Event/Mechanism	Effect	Refs&PMID
Components	Piperlongumine (PL)	In vitro	AGS and HGC-27	5, 7.5, 10 μ M (Main 7.5 μ M)	G2/M Mediated by ROS and involves GADD45 α , XIAP, and CHOP. Suppressed STAT3 and TERT	Growth arrest Proliferation and Apoptosis	19 26874726
		In vivo	Nude mice (2 \times 106 HGC-27)	3.6 mg/kg			
	Pseudolaric acid B (PAB)	In vitro	BGC-823 and MKN-45	0.5, 2, 5, 10 μ M (10 μ M)	Suppressed PI3K/AKT, ERK1/2 and mitochondrial signaling	Apoptosis	21
		In vivo	BALB/C nude mice (1 \times 106 BGC-823)	30 mg/kg	Regulated MMP-9, HIF 1 α , VEGF, VEGFR2, E-Cadherin and Ezrin	Metastasis	28132880
	Curcumin	In vitro	BGC-823	15 μ mol/L	Regulated JNK-MAPK signal pathway	Growth arrest	61 25292057
	Curcumin	In vitro	MKN45, AGS, GES-1	3.125, 6.25, 12.5, 25.0, 50.0 μ mol/L	Regulated Cox-2 and NF- κ B pathway	Growth arrest	63 25884903
	(-) - patchouli alcohol (PA)	In vitro	GES-1	10 μ M	Decreased IL-4, IL-2, TNF- α Increased IL-13	Growth arrest	64 27017292
	Curcumin	In vitro	GC-MSCs	30 μ mol/L	Inhibited NF- κ B/VEGF signaling	Angiogenesis and Metastasis	33 29312505
	Green tea polyphenols	In vitro	MKN-28	10-4 M catechin equivalents	Inhibited MMP-2, MMP-9	Migration and Invasion	42 29450132
	Dehydroefusol	In vitro	SGC-7901 and AGS	12, 24, 48 μ M	Regulated VE-cadherin and MMP-2	Adhesion, Migration, and Invasion	53 25982451
		In vivo	BALB/C nude mice (4 \times 106 SGC-7901)	40, 120 μ g			

that inhibiting telomerase activity may be a potential anti-cancer therapeutic mechanism. Another study¹⁸ utilized the PCR-TRAP method to evaluate telomerase activity in BGC-823 human gastric carcinoma cells, and found that gambogic acid (GA) can inhibit telomerase activity by inhibiting the phosphorylation of Akt and reducing the expression of hTERT. Duan et al.¹⁹ demonstrated that piperlongumine (PL) acts as an anti-cancer agent by inhibiting telomerase activity and cell proliferation in vitro and in vivo.

Mitochondrial Apoptosis Pathway

Wang et al.²⁰ found that the ethyl acetate extract of *Celastrus orbiculatus* (*C. orbiculatus* extract, COE) could change the mitochondrial membrane potential of the cell, downregulate the apoptosis-related proteins Bax and caspase, and increase the expression of bcl-2 and PI3K/Akt. These data suggest that the COE may inhibit the proliferation and metastasis of gastric cancer cells by inducing apoptosis. Pseudolaric acid (PAB) has been shown to promote the apoptosis of various cells. Wang et al.²¹ evaluated the molecular mechanism of the antitumor activity of PAB in human leukemia U937 cells. U937 cells were observed to be activated by the Bcl-2-mediated mitochondrial pathway, and activation of the caspase-dependent pathway was regulated by PAB. In addition, the activity of caspase-3 increased after PAB treatment. This study thus showed that PAB can enhance U937 cell apoptosis, at least in part through the activation of the mitochondrial apoptotic pathway. Mao et al.²² found that crocodile bile accelerated cell apoptosis via the mitochondrial apoptosis pathway. Simultaneously, it reduced the mitochondrial membrane potential, and increased the production of reactive oxygen species, the Bax/Bcl-2 ratio, the levels of activated caspase-3, and the release of cytochrome C. These data indicate that crocodile choline is a potent inhibitor of gastric cancer cells via the mitochondrial apoptosis pathway.

Other Aspects

The literature shows that traditional Chinese medicine also exhibits anti-gastric cancer effects through autophagy.²³ Curcumin²⁴ may inhibit proliferation and induce the autophagy and apoptosis in GC cells through MTT assay and transmission electron microscopy (TEM). Lei²⁵ find that mulberry anthocyanins intervened SGC-7901 cells induced autophagy by TEM observation results.

Some studies have shown that traditional Chinese medicine can inhibit the effect of tumor cells through ROS. Kim DH et al.²⁶ reported for the first time that ISL induces apoptosis of renal cell carcinoma Caki cells by producing ROS, thus inducing p53 and inhibiting Stat3 signaling pathway. Silybin²⁷ can induce caspase-dependent cell death in vitro and inhibit the growth of glioma in vivo through Ca²⁺/ROS/MAPK-mediated pathway.

There are other natural products, bitter melon extract (BME), suppressing the cancer proliferation via different mechanisms. Bhattacharya et al.²⁸ shown that BME enhances NK cell-mediated HNSCC killing activity and reveals the potential immunomodulatory effects of BME. Muhammad et al.²⁹ found that BME attains anti-tumor activity by induction of autophagy and AMPK/mTOR pathway. Subsequently, they demonstrated that BME oral feeding effectively inhibited cancer cell growth in isogenic and xenograft mouse models. In addition, Bhattacharya et al.³⁰ shown that BME inhibited cell proliferation in BME orally-fed mouse tumors via lowering expression of proliferating cell nuclear antigen (PCNA) and c-Myc. Both of their studies indicates BME high potential clinical application.

Interference With Angiogenesis

By co-culturing endothelial cells with human or animal macrophages or supernatants, it was found that the angiogenic extracts caused chemotaxis of the peritoneal macrophages and human mononuclear cells in guinea pigs. These results imply that tumor extracts act indirectly to induce angiogenesis in vivo via their effect on host macrophages.³¹ Angiogenesis can be controlled by inhibiting the proliferation of vascular endothelial cells (VECs) and by regulating vascular growth factors.

Inhibition Of The Proliferation Of VECs

Angiogenesis is an essential pathological process in the metastasis of malignant tumors, and endothelial cell proliferation is the basis of angiogenesis. Therefore, the discovery and screening of drugs that inhibit the proliferation of VECs has become a focus of tumor clinical research. Zang et al.³² reported that Luteolin has an inhibitory effect on gastric cancer angiogenesis and Vasculogenic mimicry (VM) formation induced by inhibiting VEGF secretion, which is dependent on Notch1 expression. Huang et al.³³ observed that curcumin could inhibit GC-MSD driven angiogenesis, which inhibits the proliferation of vascular endothelial cells, producing an anti-tumor effect. Lynne M

Howells et al supported that in the treatment of patients with metastatic colorectal cancer. Curcumin is well safety and tolerated combining with FOLFOX chemotherapy. Bing et al.³⁴ reported that Xiaotansanjie decoction attenuates microvessel density by inhibiting the proliferation of vascular endothelial cells in gastric cancer.

Angiogenic Factors

Under conditions of local hypoxia, several molecular responses are induced during tumor growth which activate hypoxia-inducible factors. The transcription factor HIF accumulates in endothelial cells under hypoxic conditions and binds to the VEGF promoter to induce VEGF expression, resulting in angiogenesis.³⁵ Dai et al.³⁶ reported that Ginsenoside Rg3 inhibits tumor growth and reduces lymph node metastasis by inhibiting the expression of VEGF-C, VEGF-D, and VEGFR-3 in an orthotopic mouse model of human gastric cancer. Pang et al.³⁷ reported that xiaotan sanjie can inhibit proliferation of human gastric adenocarcinoma cells MKN-45 by attenuating tumor angiogenesis, which may be related to downregulation of the mRNA and protein expressions of VEGF-C and VEGFR-3.

Inhibition Of Gastric Cancer Cell Movement

Local invasion and distant metastasis are the most important biological features of malignant carcinoma. Tumors with invasive growth can not only continue to grow at the original site, but also spread directly to the surrounding tissue, and can also spread to other parts of the body through a variety of ways. Therefore, inhibiting the metastasis of gastric cancer cells is particularly important in the treatment of tumor, including inhibition of the migration and invasion of gastric cancer cells and the adhesion of gastric cancer cells.

Migration And Invasion

The migration and invasion capabilities of gastric cancer cells are associated with metastasis.³⁸ Once proliferating gastric cancer cells reach a certain density, they will invade the adjacent basement membrane and extracellular matrix, penetrate blood vessels, and enter the circulatory system. Via migration through blood vessels, a small metastasis focus will be formed on a target organ.³⁹ Other studies have also indicated that MMP-2 and MMP-9 facilitate cell migration and invasion. MMP-2 is involved in the metabolism of type IV collagen in the

basement membrane, which is often the initial step in cancer invasion and metastasis.^{40,41} In addition, MMP-9, which is the main enzyme for degrading extracellular matrix (ECM), is an important mediator of migration and invasion for gastric cancer cells.⁴² Lu et al.⁴³ reported that Silibinin acts by inhibiting the invasion and metastasis of human gastric cancer SGC-7901 cells, and by downregulating MMP-9 and MMP-2. Yan et al.⁴⁴ found that Baicalein could inhibit the proliferative, adhesive, invasive, and migratory abilities of SGC-7901 cells. The underlying mechanism was the inhibition of the expression of MMP-9 and MMP-2 by suppressing the activity of the p38 signaling pathway. Zhang et al.⁴⁵ demonstrated that alisol B can regulate phosphorylation of vasodilator-stimulated phosphoprotein (VASP) to inhibit gastric cancer cell adhesion and migration.

Adhesion

An increase in cell adhesion molecule expression and degradation of the extracellular matrix can control cancer metastasis and reduce migration of cancer cells.

Regulating The Expression Of Adhesion Molecules

Adhesion molecules regulate cell-cell adhesion, as well as adhesion of cells to the extracellular matrix. Reducing adhesion between malignant tumor cells and enhancing adhesion of tumor cells to host cells and the extracellular matrix are two of the basic mechanisms of malignant tumor metastasis. Adhesion is regulated by focal adhesion molecules including CD44 and E-Cad. One of the functions of CD44 is to regulate the adhesion and migration of tumor cells. In the active state, the cytoplasmic domain of CD44 interacts with the actin cytoskeleton, which promotes migration of tumor cells.⁴⁶ Sun et al.⁴⁷ found that HepG2 cells cultured with the serum of rats fed with Biejiajian Pills showed lower expressions of β -catenin protein both in the cytoplasm and the nuclei along with the inhibition of GSK-3 β phosphorylation and reduced expression of CD44v6 and VEGF. Biejiajian Pills suppress the proliferation and invasiveness of hepatocellular carcinoma. Xu et al.⁴⁸ found that Huaier could partly reverse the epithelial-mesenchymal transition (EMT) by decreasing the expression of the mesenchymal markers N-cadherin and vimentin and increasing the expression of the epithelial marker E-cadherin. Huaier extract significantly inhibited both the migratory and invasive abilities of gastric cancer cells. Ma et al Indicated that by Comprehensive clinical analysis, Huaier might enhance

clinical therapeutic effects and decline the side ones with Gastrointestinal cancers.

Inhibiting Extracellular Matrix Degradation

MMPs are the major zymogen family proteins that degrade tumor extracellular matrix (ECM). MMPs can promote tumor cell invasion of the ECM by degrading it, which permits tumor cells to migrate to other sites. Wu et al.⁴⁹ found that the combination of Apatinib and AsPs could inhibit the expression of phosphorylated AKT (p-AKT) and MMP-9 expression. Apatinib, in combination with AsPs, showed enhanced inhibitory effects on cell proliferation, migration, and invasion compared with Apatinib monotherapy. Jiao et al.⁵⁰ found that the migration and proliferation of the SGC-7901 cells were decreased after incubation with different concentrations of Tanshinone IIA (TSN) in a dose-dependent manner. Moreover, the expression levels of Ki-67, PCAN, MMP-2, MMP-9, and FOXM1 were decreased, and P21 was increased in the TSN-treated SGC-7901 cells.

Modulation Of Metastasis-Related Genes To Inhibit Metastasis

Inhibition of gastric cancer cell metastasis can be achieved by regulating metastasis-related gene expression in gastric cancer. In 2003, Ishikawa et al.⁵¹ reported that VEGF-C and -D genes may be associated with lymphatic metastasis of early gastric cancer, which suggested that VEGF-C and -D genes inhibit tumor metastasis. Dai et al.³⁶ reported that ginsenosides can target vegf-c, vegf-d, and vegfr-3 genes in mouse models of human gastric cancer cells and inhibit the progression and metastasis of malignant tumors. Bao et al.⁵² reported that naringenin may inhibit the migration and invasion of human gastric cancer SGC-7901 cells by reducing the expression of metastasis related genes MMP-2 and MMP-9. Liu et al.⁵³ reported that Dehydroeffusol can significantly inhibit the migration and invasion of gastric cancer cells in vivo and in vitro. Dehydroeffusol can inhibit gene promoter activity, influence the expression of the vasculogenic mimicry master gene VE-cadherin, and significantly reduce the expression of metastasis associated genes in gastric cancer cells (MMP2), thus effectively inhibiting gastric cancer metastasis. The proliferation of gastric cancer cells not only increases the pressure between gastric cancer cells, but also enhances metastasis. Most traditional Chinese medicines exhibit their anti-metastasis effect by regulating the relevant oncogenes and anti-oncogenes to further inhibit cellular proliferation and induce apoptosis.

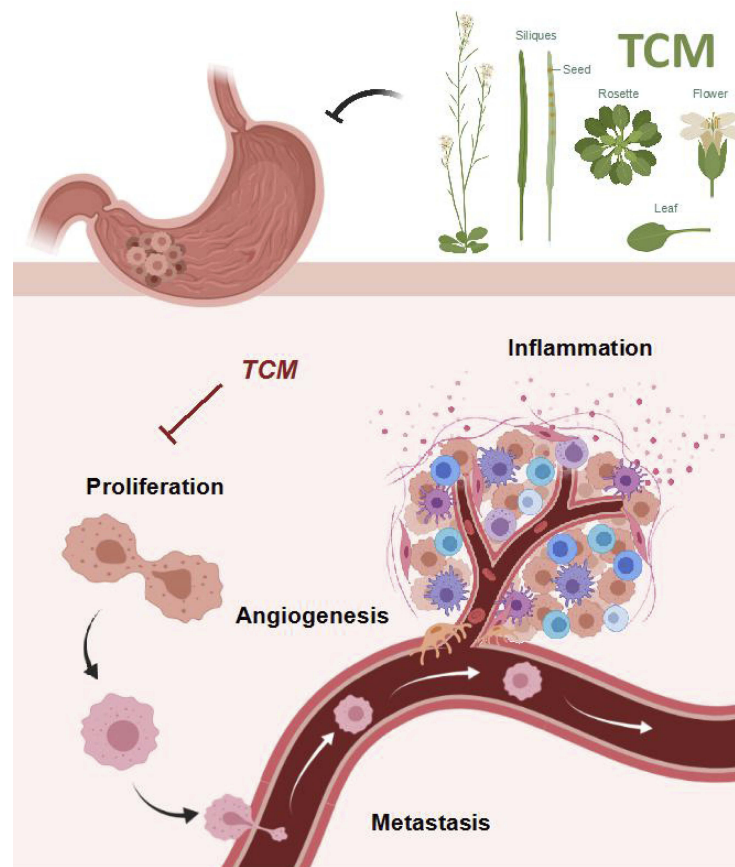
Recent studies have demonstrated that metastasis-associated genes include Bcl-2, Bax, maspin, and p53. Mu et al.⁵⁴ showed that baicalein has anti-apoptotic activity by down-regulating Bcl-2 and upregulating Bax in SGC-7901 cells. Qian et al.⁵⁵ indicated that COE could induce apoptosis of human gastric cancer MGC-803 cells overexpressing the maspin gene.

Regulation Of Inflammation-Related Factors

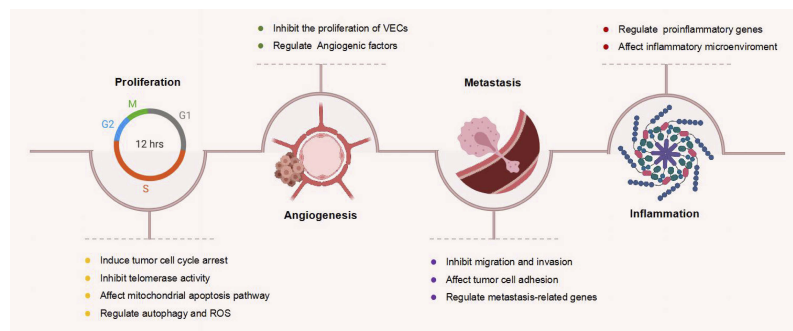
Cancer-related inflammation can promote tumorigenesis, tumor growth, and tumor metastasis in many types of cancers. Therefore, inhibiting cancer-related inflammation significantly improves the effects of cancer therapy. It is known that neoplastic cells often over-express proinflammatory mediators including proteases, eicosanoids, cytokines, and chemokines.⁵⁶ Moreover, 40 years ago, the phenomenon of TNF-mediated cell death in vitro and its physiological and pathophysiological effects were first described.^{57,58} Blocking TNF causes TNFR1 to no longer activate MAPK and NF- κ B signaling, resulting in down-regulation of proinflammatory genes.⁵⁹ Curcumin plays an anti-cancer role by inhibiting the occurrence, progression, and metastasis of various cancers, and inhibits cancer by two main processes that affect angiogenesis and tumor growth. Curcumin inhibits lymph node metastasis in gastric cancer, and its pleiotropic activity is mediated by modulating proinflammatory cytokines, and other signaling molecules, Human protein STAT3, NF- κ B, COX-2, 5-LOX, C-reactive protein, prostaglandin E (2), prostatic specific antigen, adhesion molecule, phosphorylase kinase, triglyceride, transforming growth factor beta, ET-1 HO-1, creatinine, ALT, and AST.⁶⁰⁻⁶³ Patchouli alcohol (PA), the main active ingredient of *Pogostemonis herba*, has anti-*Helicobacter pylori* and gastroprotective functions. In addition, PA significantly reduced the levels of pro-inflammatory cytokines such as interleukin -4 (IL-4), IL-2, and tumor necrosis factor (TNF-alpha), and increased the level of the anti-inflammatory cytokine IL-13 in GES-1 cells. To a certain extent, reducing the inflammatory reaction of the gastric mucosa can help block the precancerous lesions of gastric cancer.⁶⁴

Conclusion

This article summarizes how Chinese medicine can intervene at multiple levels and affect multiple targets to influence the metastasis of gastric cancer as shown in Figure 1.



A: General approach of mechanisms



B: Detailed approach of mechanisms

Figure 1 Mechanisms underlying therapeutic effects of traditional Chinese medicine on gastric cancer. **(A)** General approach of mechanisms. **(B)** Detailed approach of mechanisms.

However, it is difficult to confirm the active ingredients of traditional Chinese medicines due to their multimodal anti-tumor effects and complex composition, thus greatly limiting their potential. Additionally, clinical research for traditional Chinese medicine still lacks in-depth study and evaluation criteria, and as a result its authenticity is largely unrecognized. These findings suggest that the multi-effects of traditional Chinese medicine should be combined with those of Western medicine, allowing integrative medicine

to be used to control tumors. It is expected that further rigorous research will demonstrate that traditional Chinese medicine has clinical significance for treatment of the metastasis of gastric cancer and will lead to improvement of preventive measures.

With the continuous discovery and deeper exploration of the mechanism of gastric cancer and identification of the new molecular targets, the target of Chinese medicine in gastric cancer has been found, and the effects of

traditional Chinese medicine have been determined at the molecular level, which provides the basis for clinical applications of traditional Chinese medicine.

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Author contributions

All authors contributed to data analysis, drafting or revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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