



Title	Berberine and Coptidis Rhizoma as novel antineoplastic agents: A review of traditional use and biomedical investigations
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3 **Berberine and Coptidis Rhizoma as novel antineoplastic agents: a review of**
4 **traditional use and biomedical investigations**
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8 Jun Tang^{a,b}, Yibin Feng^{a,*}, Saiwah Tsao^c, Ning Wang^a, Robert Curtain^d, Youwei Wang^b
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12 ^a *School of Chinese Medicine, The University of Hong Kong, 10 Sassoon Road, Pokfulam, Hong Kong,*
13

14 *PR of China*
15

16 ^b *College of Pharmacy, Wuhan University, Wuhan 430072, PR of China*
17

18
19 ^c *Department of Anatomy, The University of Hong Kong, 21 Sassoon Road, Pokfulam, Hong Kong, PR of*
20

21 *China*
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23 ^d *Genome Research Center, The University of Hong Kong, 21 Sassoon Road, Pokfulam, Hong Kong, PR*
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25 *of China*
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59 * Corresponding author: Tel.: +852 25890482; fax: +852 28725476
60 *E-mail address: yfeng@hku.hk.*
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Abstract

Ethnopharmacological relevance: Coptidis Rhizoma (*Huanglian*) and its major component, berberine, have drawn extensive attention towards their antineoplastic effects in the recent years. The antineoplastic effects are related to the Chinese Medicine (CM) properties of Huangliang in treating diseases by removing *damp-heat* and purging *fire* and counteracting *toxicity*.

Aim of the review: To trace the long history of the traditional use of *Huanglian* from folk medicines, especially from Chinese medicine, to recent pharmacological studies of *Huanglian* and berberine, with an emphasis on their antineoplastic effects and the promise as novel antineoplastic agents.

Methods: A total of 7 databases were extensively searched for literature research. The terms and keywords for searching included *Huanglian*, berberine, Coptis, Coptidis Rhizoma, anticancer, antiinvasion, antimetastasis and mechanism. The papers including ours with studies on anticancer and mechanism, pharmacology and toxicology of *Huanglian* and/or berberine were focused.

Results: In view of traditional use, the anticancer effects of *Huanglian* can be ascribed to its CM trait by removing *damp-heat*, *fire* and *toxicity*. From modern biomedical studies, anticancer effects have been demonstrated in both *Huanglian* and berberine. The underlying molecular mechanisms involve cell cycle arrest, apoptosis induction and antiinflammation. Berberine is an essential anticancer compound in *Huanglian*. In some studies, the use of *Huanglian* was shown to be more effective and beneficial than the use of berberine alone. The presence of other protoberberine-type alkaloids in *Huanglian* might give synergistic effects for the anticancer effects. Berberine also demonstrates effects of antiangiogenesis, antiinvasion and antimetastasis in some cancer cell lines, however, more investigations are required to unravel the underlying mechanisms involved.

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Conclusions: The modern evidences of treating cancer with *Huanglian* and berberine have a strong linkage with traditional concept and rules of using *Huanglian* in CM practice. As anticancer candidates with low toxicity, berberine and its altered structure, as well as *Huanglian* and its formulae, will attract scientists to pursue the potential anticancer effects and the mechanisms by using technologies of genomics, proteomics and other advanced approaches. On the other hand, relatively few *in vivo* studies have been conducted on anticancer effects of *Huanglian* and berberine. The clinical application of berberine or *Huanglian* as novel cancer therapeutic agents requires *in vivo* validations and further investigations of their anticancer mechanisms.

Keywords: Traditional Chinese medicine; Coptidis Rhizoma; *Huanglian*; Berberine; Antineoplastic agent; mechanisms of anticancer action

1
2 *Abbreviations:* AP-1: activator protein 1; ATF3: activating transcription factor 3; Bcl-2: B
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4 cell lymphoma/leukemia-2; BCD: berberine chloride dihydrate; BCL: berberine hydrochloride;
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6 CDK: cyclin-dependent kinase; COX-2: cyclooxygenase-2; FasL: factor associated suicide
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8 ligand; GADD153: growth arrest and DNA damage inducible gene 153; GAP: good
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10 agricultural practice; HIF: hypoxia inducible factor; *Huanglian* extract: the water extract of
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12 *Huanglian*; IC₅₀: fifty percent inhibitory concentration; ID₅₀: fifty percent inhibitory dose;
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14 IFN: interferon; IκB: inhibitor of NF-κB; IKK: IκB kinase; IL: interleukin; JNK, c-Jun *N*-
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16 terminal kinase; LOAEL: lowest observed adverse effect level; MAPK: mitogen-activated
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18 protein kinase; MDR: multidrug resistance; MLD: minimal lethal dose; MMP: mitochondrial
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20 membrane potential; MMPs: matrix metalloproteinases; NAG-1: nonsteroidal anti-
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22 inflammatory drug activated gene; NCI: National Cancer Institute; NDEA: *N*-
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24 nitrosodiethylamine; NF-κB: nuclear factor-κB; NOAEL: no observed adverse effect level;
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26 PARP: poly ADP-ribose polymerase; ROS: reactive oxygen species; CM: traditional Chinese
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28 medicine; TNF: tumour necrosis factor; Urokinase-type plasminogen activator: u-PA.
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1. Introduction

Coptidis Rhizoma (*Huanglian* in Chinese) has been used for more than two thousand years by CM physicians for treatment of *damp-heat* syndromes (Chinese Herbals Editor Board, 1999). Berberine (Fig. 1) is an isoquinoline alkaloid commonly used in both China and other countries as a botanic drug which is also a major active compound in *Huanglian*. It is widely present in the plant kingdom: Berberidaceae Ranunculaceae and Papaveraceae (Felter and Lloyd, 1898; Galle et al., 1994; Lenka et al., 2007). In this review, we focused on *Huanglian* of Ranunculaceae, because traditional use and ethnopharmacological relevance of *Huanglian* is link to biomedical investigations of *Huanglian* and berberine. As specified in the Chinese Pharmacopoeia (The State Pharmacopoeia Commission of the P.R. China, 2005), *Huanglian* may contain three *Coptis* species: *Coptis chinensis* Franch. (*Weilian* in Chinese), *Coptis deltoidea* C. Y. Cheng et Hsiao. (*Yalian* in Chinese) and *Coptis teetoides* C. Y. Cheng (or *Coptis teeta* Wall., *Yunlian* in Chinese) (Fam. Ranunculaceae). There are a few other native *Coptis* species distributed in other areas of the world which are also used as *Huanglian*, e.g. *Coptis japonica* Makino. in Japan (Society of Japanese Pharmacopoeia, 2001) and *Coptis trifolia* Salisb. (or Goldthread) which were used as domestic medicines in North America (Lloyd and Lloyd, 1887). In China, *Weilian* has been extensively cultivated in eastern *Sichuan* and western *Hubei* provinces under good agriculture practice (GAP) for Chinese Crude Drugs (Xu et al., 2004) (Fig. 2). It is worthwhile to note that the wild *Yunlian* has been remarked as vulnerable with the conservation status ranking No. 2 in China. Nevertheless, its status as an herbal drug would not be greatly influenced due to a similar extensive cultivation to *Weilian* in the southwestern part of China. Berberine is an alkaloid component in *Huanglian*. The recent studies confirmed that *Huanglian* is a good producer of berberine. Berberine constitutes the most abundant alkaloid in the dried herb (5.20%-7.69%, w/w) (Iizuka et al., 2000a; Xu et

1 al., 2004). High production of berberine can also be achieved via modifications of metabolic
2 pathways in plants by genetic engineering (Sato et al., 2001).
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4 The main pharmacodynamic properties of berberine and *Huanglian* have long been
5 recognized on the treatment of intestinal infections including acute gastroenteritis, cholera and
6 bacillary dysentery which can be linked to their antibacterial, antiviral and anti-inflammatory
7 effects (Amin et al., 1969; Subhuti, 2005; Kyoko et al., 2007). The purification of berberine
8 from *Huanglian* has greatly facilitated investigation studies into the therapeutic applications
9 of *Huanglian*.
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18 The development of *Huanglian* and berberine in clinical applications can be broadly
19 divided into several stages:
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- 22 (a) Early clinical use of *Huanglian* in CM began at about 2000 years ago in the treatment of
23 diarrhea, eye inflammation and women's abdomen ailments caused by *damp-heat*, which
24 is still a common use of *Huanglian* today in CM (Xu et al., 2004; Chen and Gao, 1997a
25 and 1997b).
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33 (b) Early medical uses and development of berberine-containing botanic drugs in North
34 American folk medicine in the treatment of sub-acute and chronic inflammations
35 including gastric disorders, respiratory affections and cancer during the 1800s; but
36 gradually lost their application status in western medicine (Lloyd and Lloyd, 1887; Felter
37 and Lloyd, 1898).
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45 (c) The pre-clinical and clinical studies and development of berberine as a natural antibiotic to
46 treat various bacteria-associated diarrheas in the late 1900s (Dutta and Panse, 1962;
47 Subbaiah and Amin, 1967; Khin et al., 1985; Rabbani et al., 1987).
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53 It is worth highlighting that the antineoplastic activities of berberine have been reported
54 intermittently (Hano, 1959). The initial positive results from these studies together with the
55 information from CM led to reconsideration of the use of berberine and *Huanglian* in the
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1 treatment of cancer diseases. After 1990s, there were more studies on the antineoplastic
2 effects of berberine and *Huanglian* in diverse cancer cell types, and the underlying
3 mechanisms of the antineoplastic action were also investigated especially on those of
4 berberine. Meanwhile, many other pharmacological benefits have also been reported
5 including lowering of blood cholesterol (Kong et al., 2004), anti-inflammatory effects (Kuo et
6 al., 2004), therapies of experimental colitis (Zhou and Mineshita, 2000) and treatment of
7 diabetes (Zhou et al., 2007) for berberine, hepatoprotective (Ye et al., 2009) and anticachectic
8 effects (Iizuka et al., 2000b) for *Huanglian*.

19 In this review, the history of the early development of *Huanglian* and berberine as botanic
20 drug was traced at first and recent publications on their effects on various cancer cell types
21 and the underlying antineoplastic mechanisms, genomics, proteomics and computer-aided
22 molecular design as well as *in vivo* studies, general pharmacology and toxicology were then
23 focused in English and Chinese databases. No restrictions regarding the language of
24 publication were imposed, but most relevant studies were published in English and Chinese
25 bibliographies, the electronic literature searches were conducted in the following databases (6
26 English databases and 1 Chinese database): MEDLINE (1966 to date, via Pubmed), EMBASE
27 (1974 to date), CINAHL (Cumulative Index to Nursing and Allied Health Literature, 1982 to
28 date), AMED (Allied and Complementary Medicine Database, 1985 to date), The Cochrane
29 Central Register of Controlled Trials (CENTRAL),EBM Reviews Full Text - CDSR, ACP
30 Journal Club, and DARE (1991 to date) and China Journals Full-text Database (1915 to date,
31 via CNKI). The following search key words or terms were used: *Huanglian*, berberine, Coptis,
32 Coptidis Rhizoma, anticancer, antineoplastic mechanisms, genomics, proteomics and
33 computer-aided molecular design as well as *in vivo* studies, general pharmacology and
34 toxicology. Further manual searches were conducted in our own files as well as in the
35 reference lists of all located articles. After systematic review, we did not find any review

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paper for the anticancer efficacy and safety of *Huanglian* and berberine until this paper was completed.

All articles were read in full by all authors and data relating to anticancer, genomics, proteomics and computer-aided molecular design as well as *in vivo* studies, general pharmacology and toxicology of *Huanglian* and/or berberine and results were extracted by authors. The authors agreed to a consensus on the assessed data and cases of discrepancy would be settled by discussion. Articles which are included in the study had to meet all of the following criteria: (a) experimental studies on anticancer effects of *Huanglian* and/or berberine; (b) explore underlying mechanisms for anticancer, antiinvasive and antimetastatic effects of *Huanglian* and/or berberine; (c) new approach such as genomics, proteomics and computer-aided molecular design to studies on anticancer effects of *Huanglian* and/or berberine; (d) animal studies and clinical trails for anticancer effects of *Huanglian* and/or berberine; (e) general pharmacological and toxicological effects of *Huanglian* and/or berberine; (f) traditional use and ethnopharmacological relevance to anticancer effects of *Huanglian* and/or berberine; (g) articles in English were preferred, but some information which can be only found in Chinese (partially information in d, e and f) were also included. Articles out of the above criteria were excluded. As a result, over 100 peer-reviewed international papers in English as well as 21 peer-reviewed papers or books in Chinese and 5 web information have been cited (136 reference papers in total) This is a first review paper which provided a whole picture and related information on topic of antineoplastic effects of *Huanglian* and berberine. The potential values of berberine and *Huanglian* as novel antineoplastic agents for future clinical application are also discussed. The plant sources, biosynthesis, pharmacological actions and phytochemistry of berberine and cultivation of *Huanglian* can be referred to other review articles published (Li et al., 2008a; Lenka et al., 2007; Yu et al., 2006; Xu et al., 2004).

2. The early history of *Huanglian* and berberine development as botanic drugs

2.1. The early years of *Huanglian* development

The earliest record of *Huanglian* can be dated back to *Shen Nong's Herbal Classic* compiled in the Han Dynasty of China (about A.D. 200), in which *Huanglian* was regarded as one of the top-grade drugs (tonic and non-toxic) (Xu et al., 2004; Wu et al., 1982). Since then, the use of *Huanglian* was frequently cited in many other official or un-official medical compilations e.g. *Shang Han Lun* (Treatise on Cold-induced Febrile Diseases) and *Jin Kui Yao Lue* (Synopsis of Prescriptions of the Golden Chamber, about A.D. 210, Han Dynasty), *Wai Tai Mi Yao* (Medical Secrets from the Royal Library, A.D. 752, Tang Dynasty), and *Ben Cao Gang Mu* (Compendium of Materia Medica, A.D. 1596, Ming Dynasty) (Chen and Gao, 1997a and 1997b). *Huanglian* is an important component in many traditional Chinese compound formulae (*Fufang* in Chinese). At least 1,760 *Huanglian* formulae have been recorded in Chinese *Herbals* until Song Dynasty (A.D. 1279), and about 108 of which are still in use today (Chen and Gao, 1997a). The well-known and useful ones, such as *Xie Xin* Decoction from *Jin Kui Yao Lue* and *Huanglian Jiedu* Decoction from *Wai Tai Mi Yao*, have received great attention in evidence-based studies (Feng, 2000; Ohta et al., 2004).

Huanglian can remove *damp-heat*, *fire* and counteract *toxicity*, both of which are related to inflammatory or tumour pathogenesis in CM (Chinese Herbals Editor Board, 1999; Liao et al, 2003). Its traditional use in the treatment of diseases such as stuffiness and fullness of abdomen, diarrhea, vomiting, high fever, fidgetiness and insomnia, eye inflammation, diabetes, carbuncles and sores as specified in Chinese Pharmacopoeia may be results of multiple pharmacological properties including anti-inflammatory and anticancer activities of

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Huanglian. The CM physicians may not have recognized the pathological nature of some diseases as clearly as western medicine. For example, in CM, tumours were described as ‘lumps’ or ‘carbuncles’ in the body and were usually believed to be caused by *heat*-induced *toxicity* and stagnation of *qi* and *blood*. There were indeed some approaches to treat ‘tumours’ in CM practices by clearing *blood stagnation*, *lumps*, to remove *heat* and to counteract *toxicity* (Liao et al, 2003).

In other traditional medicines such as Ayurvedic medicine and early western medicine, in the year of 1800s, the use of berberine-containing botanic drugs was also dominant including the Indian Barberry (*Berberis aristata* DC., Berberidaceae), European Barberry (*Berberis vulgaris* L., Berberidaceae), American Hydrastis (*Hydrastis canadensis* L. or Goldenseal, Ranunculaceae) and Goldthread. The American Hydrastis was largely used both as a medicine and as a coloring material among certain aboriginal tribes of North America. In his first edition of ‘Collections for an Essay Toward a *Materia Medica* of the United States’ (1798), B. S. Barton had referred to the Cherokee use of Hydrastis as a cure for cancer, which may represent the oldest recorded use of berberine-containing botanic drugs for cancer treatment. Some homoeopathic practitioners even suggested long and continued use of Hydrastis to retard *scirrhus* of the breast (Felter and Lloyd, 1898). However, evidence of pre-clinical or clinical studies for these herbs was scanty at that stage. The official use of these botanic drugs gradually diminished in the late 1800s.

2.2. *The early years of berberine development*

Berberine was firstly isolated and identified as a plant isoquinoline alkaloid (Lenka et al., 2007) in the early nineteenth century (Felter and Lloyd, 1898). In the succeeding years from 1910s to 1960s, the chemistry and synthesis of berberine was thoroughly studied (Johannes

1 and Rose, 1930; Huffman and Miller, 1960) and total synthesis of berberine was achieved in
2 1969 (Kametani et al., 1969). In the early 1960s, berberine and its salts, such as berberine
3 sulphate, had been demonstrated to be valuable to treat cholera, severe diarrhea and
4 amoebiasis by Indian researchers (Dutta and Panse, 1962; Subbaiah and Amin, 1967). Clinical
5 trials of berberine to treat bacterial diarrhea were reported in 1985 and 1987 (Khin et al., 1985;
6 Rabbani et al., 1987). Berberine itself may not be an effective antibiotic; however, its anti-
7 microbial activity is potentiated by the MDR inhibitor, 5'-methoxyhydnocarpin, which does
8 not have antimicrobial activity on its own (Stermitz et al., 2000). This study showed that
9 *Huanglian* extract may have higher therapeutic effects than berberine alone and may be the
10 results of synergistic actions of other components present in *Huanglian*. In China, berberine
11 was isolated and used as a botanic drug in the late 1950s (Subhuti, 2005). The initial clinical
12 application was for treatment of intestinal infections.

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The early development of berberine into an anticancer agent was not as promising as its application as a natural antibiotic agent. However, later screening of the Native American anticancer Hoxsey formula, a mixture of botanic drugs containing Barberry (*Berberis vulgaris* L.), resulted in the identification of berberrubine as the antitumour principal (Patricia et al., 2002; Hoshi et al., 1976). Interestingly, berberrubine is not present as the natural form in the botanic drug (Lenka et al., 2007). It is most likely an *in vivo* metabolite of berberine that is active for cancer therapy (Zuo et al., 2006).

2.3. *The link between traditional use and modern research of berberine and Huanglian for cancer therapies*

Even though there are few indications of *Huanglian* in early history directly related to treatment of cancer, clues and implications can be derived from traditional uses of *Huanglian*

1 in the treatment of diseases with similar symptoms of tumours, as well as basic research
2 especially on its main active principal, berberine. In modern concepts, cancer is characterized
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4 by a rapid and uncontrolled formation of abnormal cells, which may mass together to form a
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6 growth or tumour, proliferate throughout the body and initiate abnormal growth at other sites.
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8 Therefore, anticancer drugs involve pharmacological effects of cytotoxicity, antiproliferation,
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10 antiinflammation, cell cycle arrest, apoptosis induction, antiinvasion and antimetastasis etc..
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14 There is no concept of cancer in ancient CM, but some symptoms may relate to cancer.
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16 Symptoms such as diarrhea, vomiting, fever, and bleeding are not specific symptoms, cancer
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18 may also show similar symptoms. *Huanglian*, firstly recorded in *Shen Nong's Herbal Classic*
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20 (the oldest herbal compilation in China) to treat diarrhea, inflammation of the eye, and
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22 women's abdomen ailments caused by *damp-heat*, also has long been used for the treatment
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24 of these diseases in CM practice (including its formulae). The syndromes with the trait of
25
26 *damp-heat* or *heat-toxicity* can be easily linked to the modern concept of infection or
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28 inflammation. Therefore, the early pharmacological study of *Huanglian* and its formulae were
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30 found to be beneficial in anticontagious and antiinflammatory effects. In current studies, many
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32 cytokines released by cancer cells can be directly translated to be a similar target of anti-
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34 inflammatory drugs (Lee et al., 2007a). While the precise mechanisms underlying antitumour
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36 effects remain to be investigated, there is increasing evidence showing that *Huanglian* and/or
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38 berberine may inhibit the inflammation-triggered tumorigenesis (Izuka et al., 2000; Kuo et al.,
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40 2004; Lee et al., 2007b; Pandey et al., 2008). Inflammation and cancer have been viewed as
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42 vicious partners in crime and their relationship has become a hot topic for cancer research
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44 (Balkwill and Mantovani, 2001; Coussens and Werb, 2002; Mantovani and Balkwill, 2006;
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46 Lu et al., 2006). During inflammation, cytokine, growth factors and free radicals are produced,
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48 to activate transcription factors including NF- κ B and downstream cancer-related genes
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50 (Hussain and Harris, 2007). The link between anticancer and antiinflammation, can be
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1 established after antiinflammation, anticachectic, antioxidant, antiproliferation and
2 antiinvasion properties of berberine, *Huanglian* and/or its formulae have been discovered *in*
3 *vitro* and *in vivo* (Fukuda et al., 1999a, 1999b; Izuka et al., 2000b; Anis et al., 2001; Wang et
4 al., 2002; Jantova et al., 2003; Kuo et al., 2004; Peng et al., 2006; Lee et al., 2007b; Ye et al.,
5 2009, Tsang et al, 2009).

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11 In many modern Chinese herbal books about tumour treatment, *Huanglian* is chosen as one
12 of the anticancer herbal drugs and is widely used in routine CM practice (Tse, 1977; Qian,
13 1980; Xu et al, 1997; Chang, 1998; Bo, 2002, Feng et al, 2008). In view of CM, the anticancer
14 effect of *Huanglian* can be ascribed to one of the CM traits by clearing *damp-heat* or
15 removing *heat* and *toxicity* (Chinese Herbs Editor Board, 1999; Liao et al, 2003), while from
16 biomedical studies, cytotoxic, antiproliferative, antiinflammatory, antiinvasive and
17 antimetastatic effects should all contribute to anticancer effects of *Huanglian* and berberine
18 (Izuka et al., 2000; Jantova et al., 2003; Kuo et al., 2004; Peng et al., 2006; Lee et al., 2007b,
19 Tsang et al., 2009). In view of historical development, modern studies and clinical
20 applications, anti-inflammatory effect was frequently mirrored in *Huanglian* application in
21 many classic herbal books, for example, four formulae containing *Huanglian* (Coptidis
22 Rhizoma) such as *Gegen HuanqQin Huanglian* decoction (Puerariae Radix, Scutellariae
23 Radix, Coptidis Rhizoma, Glycyrrhizae Radix), *Banxia Xiexin* decoction (Pinelliae Rhizoma,
24 Scutellariae Radix, Coptidis Rhizoma, Zingiberis Rhizoma, Giensen Radix, Ziziphi Jujubae
25 Fructus, Glycyrrhizae Radix), *Huanglian* decoction (Coptidis Rhizoma, Pinelliae Rhizoma,
26 Cinnamoni Ramulus, Zingiberis Rhizoma, Ginseng Radix, Ziziphi Jujubae Fructus,
27 Glycyrrhizae Radix) from *Shang Han Lun* and *Jin Kui Yao Lue*, *Huanglian Jiedu* decoction
28 (Coptidis Rhizoma, Scutellariae Radix, Phellodendri Cortex, Gardeniae Fructus) from *Wai Tai*
29 *Mi Yao* and *Zuo Jin Wan* (Coptidis Rhizoma and Evodiae Fructus) from *Dan Xi Xin Fa* (A.D.
30 1481, Ming Dynasty) are usually used to treat stomach ache, vomiting, fever, bleeding and
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1 diarrhea, which are strongly related to infections, inflammations and even cancers (Feng et al,
2 2008).

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4 As a classic formula, *Huanglian Jiedu* decoction has been prescribed to treat gastroenteritis,
5 ulcer, acute febrile illnesses and tumours nowadays, in which a pharmacological basis was
6 studied (Sun et al., 2006). This decoction is the aqueous extract of a mixture of four herbs
7 with *Huanglian* as a major component. All herbs have some similar nature to *Huanglian* for
8 *heat* removal, suggesting an integrated anti-inflammatory effect, which may be more easily
9 understood from a traditional view of CM. Our clinical observations have shown that
10 *Huanglian*-containing formulae were also effective for treating adenoma and carcinoma of the
11 large intestine, Hilar Cholangiocarcinoma (Low and Feng, 2004; Feng et al, 2008). It could be
12 hypothesized that Chinese medicines with a similar druggable nature to *Huanglian*, may also
13 have effective antitumour effects in CM practice. This idea forms one basis for our future
14 effort to establish links between CM and western counterparts.

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16 In reality regarding CM practice, practitioners rarely use a pure compound or single herbal
17 drug to treat diseases, but use a combination of single herbs to mix together (so called *Fufang*,
18 or composite formula). It is a common model for many Chinese medicines studies in recent
19 years, the goal of research is to “dig out” major active single herb(s) in a composite formula
20 for some diseases and further explore active compound(s) in the single herb(s). To the
21 contrary, CM practitioners always integrate traditional use with results derived from modern
22 research. For example, a CM practitioner may use *Huanglian* combined with other herbs or
23 *Huanglian Jiedu* decoction to treat cancer, but not berberine. Berberine is one of the active
24 compounds from *Huanglian* but they are not equal with each other. The effects and
25 mechanisms of *Huanglian* should include those of berberine; however, berberine would not
26 fully represent that of *Huanglian*. Thus, *Huanglian* and berberine in this review have been
27 described separately in order to establish differentiation and their innate correlation.

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Numerous experimental studies of *Huanglian* and berberine on antitumour, as well as in modern CM practice, have been carried out in recent years, the topic of antiinflammation and anticancer is also a most recent issue. Therefore, *Huanglian* and berberine are defined in the title as the novel antineoplastic agents. From views of traditional use and pharmacological studies, *Huanglian* and berberine inhibit inflammation-triggered tumorigenesis, whereby clues have been found from traditional use, but other anticancer effects such as cytotoxicity, antiproliferation, cell cycle arrest, apoptosis, antiinvasion and antimetastasis were discovered in the last ten years, which can be considered as new findings from scientific research.

3. The pharmacological studies of berberine and *Huanglian* for cancer therapies

3.1. Cancer cell lines used and antiproliferation properties of berberine

Just as mentioned above, intensive studies on antineoplastic effects of berberine have occurred since the 1990s. Numerous cancer cell lines, either human or murine, have been used to examine the cytotoxic effects of berberine. Berberine could inhibit the growth of many cancer cell lines *in vitro* at a relatively low IC₅₀. The cancer cell types, which were sensitive to berberine, are broad including both hematological and non-hematological cancer arising from leucocytes, liver, lung, stomach, colon, skin, oral, esophagus, brain, bone, breast and sexual organs (either hormone-dependent or independent). In some cases, the IC₅₀ of berberine was lower than 4 µg/mL, which is below the safety limit established by National Cancer Institute (NCI) (Letašiová et al., 2006), indicating the potential of berberine as an antineoplastic drug on human cancer e.g. human cervical cancer, leukemia or colon cancer. Recently, we have confirmed the cytotoxicities of berberine on various cancer cell lines including HONE 1 (nasopharyngeal carcinoma), HK1 (nasopharyngeal carcinoma), HeLa (cervical carcinoma)

1 and HepG2 (hepatocellular carcinoma) cancer cell lines in a dose- and time-dependent manner,
2 this confirms the broad spectrum of antineoplastic activity of berberine (unpublished data).
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7 *3.2. Underlying mechanisms involved in the cytotoxicities of berberine*

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11 Recent advances in the study of cell cycle checkpoint controls and DNA repair, have
12 identified key small-molecule inhibitors of the cell cycle relevant to understanding the
13 mechanisms of cancer chemotherapeutic agents (Leland and Michael, 1994; Crews and
14 Shotwell, 2003). Berberine can induce DNA topoisomerase I poisoning and apoptotic cell
15 death (Kettmann et al, 2004a). Many more recent findings, stapled into the antineoplastic
16 story of berberine, are based on the knowledge of cell cycle regulation and the downstream
17 effects or phenotypes related to cell-cycle inhibition, particularly the caspase(s)-dependent
18 apoptosis and associated cell signaling pathways. The intrinsic apoptotic pathways are vital
19 for many normal physiological processes and initiated by a series of apoptotic regulatory and
20 effector proteins such as caspases. Aberrant or dysregulated apoptotic response contributes to
21 the development of cancer (Reed, 2002). There are two main pathways for cell death by
22 triggering the activation of the caspase(s) cascade. One pathway induces apoptosis via
23 interaction of the Fas/FasL system (the extrinsic cell death pathway). Fas is a member of the
24 tumour necrosis factor (TNF) death receptor family, which plays a key role in linking drug-
25 induced damage to apoptotic cell processes. Another one is the activation of the mitochondrial
26 pathway (the intrinsic pathway). Studies on treatment of cancer by berberine indicated that
27 apoptosis is involved in the many cancer cell lines examined. The underlying mechanisms for
28 apoptosis induction by berberine, in more or less degree of diversity, are summarized below.
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56 More details can be referred to in Fig. 3.
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(a) Cell cycle arrests at G₀/G₁, G₁ and/or G₂/M, and coordinated suppression of cyclin-dependent kinases (e.g., CDK 2, 4, 6) and cyclins (e.g., cyclin B, D, E) (Eom et al., 2008; Mantena et al., 2006a and 2006b; Lin et al., 2006b; Lin et al., 2006c; Kang et al, 2005; Jantova et al., 2003). Our current studies on human nasopharyngeal carcinoma cells (HONE1 cells), suggest that berberine could induce apoptosis via G₂ arrest which demonstrated by increased levels of cleaved-PARP, cleaved caspase 3 and cleaved caspase 9 (Tsang et al., 2009). In some cases, the induction of cell cycle arrest was relied on p53 status, for example, G₁ arrest was induced in HCO and U2OS cells (p53 functional), but G₂/M arrest in Saos-2 cells (p53 deficient) (Liu et al., 2009a); in others, the cell cycle arrest in G₀/G₁ phase was much higher in p53 expressing SK-N-SH cells than those in p53-deficient SK-N-MC cells (Choi et al., 2008), and also the case in p53 positive LNCaP prostate cells than those in p53 negative PC-3 cells (Choi et al., 2009).

(b) The mitochondria/caspase-dependent and/or Fas/FasL signal pathways, evidenced by the altered ratio of anti-apoptotic (Bcl-2 proper, Bcl-x_L) and pro-apoptotic (Bax, Bid) members of the Bcl-2 family proteins (Eom et al., 2008; Hsu et al., 2007; Mantena et al., 2006a and 2006b; Lin et al., 2006a; Lin et al., 2006c), loss of mitochondria membrane potential and/or release of cytochrome c (Lin et al., 2007a; Jantova et al., 2007; Hwang et al., 2006; Lin et al., 2006a; Lin et al., 2006c; Mantena et al., 2006a and 2006b), induction of reactive oxygen species (ROS) and/or Ca²⁺ production (Meeran et al., 2008; Lin et al., 2007a and 2007b; Jantova et al., 2007; Hsu et al., 2007; Lin et al, 2006a; Lin et al., 2006c), activation of Fas or FasL (Hsu et al., 2007; Hwang et al., 2006) as well as activation of a series of caspases, especially the downstream executioner, caspase-3.

(c) Many other cell signaling pathways have also been found to be associated with the anticancer effects of berberine. Hsu et al. (2007) have reported a new mechanism for berberine-induced apoptosis in SW620 human colonic carcinoma cells by targeting the

1 redos/ROS pathway and JNK/p38 pathway. Another study on the same cancer type
2 showed that berberine could induce cell growth arrest and apoptosis of HCT-116 and
3
4 SW480 cells by targeting two pro-apoptotic proteins ATF3 and NAG-1, in which the p53-
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6 dependent pathway and the PKC, ERK, and GSK-3b pathways were involved,
7
8 respectively (Rojsanga et al., 2007). Recent research further unraveled a vital role of p53
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10 related pathway in the berberine-mediated tumour suppression. Berberine could induce
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12 apoptosis of both A549 (p53+/+) and H1299 (p53-/-) human lung cancer cells, which
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14 effect was more pronounced in the A549 in both *in vitro* and *in vivo* studies (Katiyar et al.,
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16 2009). Similarly, the cells with functional p53 were more susceptible to berberine than the
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18 p53-deficient cells, such as in SK-N-SH than SK-N-MC and in LNCaP than PC-3 cells
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20 (Choi et al., 2008 and 2009).
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- 26 (d) Berberine induces apoptosis via positive or negative regulation of various cytokines
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28 functioned in cellular network. These include up-regulation of GADD153 (Lin et al.,
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30 2007b), inhibition of cyclooxygenase-2 (COX-2) and Mcl-1 (Kuo et al., 2005) and down-
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32 regulation of nucleolar phosphoprotein nucleophosmin/B23 and telomerase (Wu et al.,
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34 1999). A recent study demonstrated that nucleophosmin plays an important role in the
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36 regulation of apoptosis as one of Bax chaperons (Kerr et al., 2007). Also, this chaperon
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38 was found to be crucial and critical in regulation of the tumour suppressor p53 (Mauguel et
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40 al., 2004). GADD153, as a leucine zipper transcription factor, was found to mediate
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42 berberine-induced apoptosis in human cervical cancer Ca Ski cells, shown by elevated
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44 ratio of p53 and Bax/Bcl-2 proteins and increased levels of ROS and Ca²⁺ release from
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46 endoplasmic reticulum (ER). The proposed mechanism is that enforced expression of
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48 GADD153 sensitizes the cells to ER stress, which involves down-regulation of Bcl-2 and
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50 enhanced oxidant injury (McCullough et al., 2001).
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In addition to direct apoptosis induction, a large number of studies demonstrated that berberine may exert indirect effects against cancers by involving the pro-inflammatory or NF- κ B pathway (Kuo et al., 2004; Lee et al., 2007b; Pandey et al., 2008), antioxidant defense system (Thirupurasundari, et al., 2009), anti-metastasis pathway (Lin et al., 2008; Kim et al., 2008; Ho et al., 2009a) as well as synergistic effects with other chemotherapeutic agents (Liu et al., 2009b; Peng et al., 2008). The aforementioned berberine-induced ROS generation, may play a key role in modulating inflammatory responses through the inhibition of transcription factors such as NF- κ B, which mediates expression of many pro-inflammatory mediators (Lin et al, 2006a; Jantova et al., 2007; Hsu et al., 2007). One recent study dissected the possible role of berberine in the NF- κ B signaling pathway (Pandey et al., 2008). Some key components and downstream gene products in this pathway were mediated for berberine-induced apoptotic and anti-inflammatory activities. The mechanism in part, was ascribed to the direct modification of cysteine residue 179 in IKK, resulting in the stabilization of I- κ B α and suppression of phosphorylation and nuclear translocation of p65, and finally inhibition of NF- κ B reporter activity. Moreover, NF- κ B-regulated gene products were also suppressed by berberine including those functional in antiapoptosis (Bcl-xL, Survivin, IAP1, IAP2, and cFLIP), proliferation (cyclin D1), inflammation (COX-2), and invasion (MMP-9). These results established a strong link of enhanced apoptosis induced by TNF- α and chemotherapeutic agents and inhibition of TNF-induced cellular invasion with targeted NF- κ B signaling pathway by berberine.

Berberine could inhibit motility and invasion ability of the highly metastatic lung cancer cell line, A549 cells, under non-cytotoxic concentrations via down-regulation of nuclear transcription factors e.g. c-fos, c-jun and NF- κ B (Peng et al., 2006). Other explorations for the anti-metastasis function indicated that berberine could inhibit TNF- α -induced MMP-9 expression and cell invasion via mediating AP-1 DNA binding activity in MDA-MB-231

1 human breast cancer (Kim et al., 2008), induce down-regulation of MMP-1, -2, and -9 and
2 related gene expressions in human SNU-5 gastric cancer cells (Lin et al., 2008), and down-
3 regulate u-PA, MMP-2 and -9 expressions through MAPK and NF- κ B signaling pathways in
4 human SCC-4 tongue squamous carcinoma cells (Ho et al., 2009). We found that anti-
5 invasion of berberine on HONE1 cell is a different mechanism which through inhibition of
6 RhoA signaling pathway (Tsang et al., 2009). As regards the control of hematopoietic tumor
7 cell, berberine could inhibit the migration of HL-60 cells, primary acute myeloid leukemia
8 (AML) cells and leukemic stem cells (LSCs) induced by chemokine stromal cell-derived
9 factor-1 (SDF-1), showing to be promising for prevention of leukemia (Li et al., 2008b).
10 Berberine could also inhibit AP-1 (a complex of c-fos and c-jun) activation in human
11 hepatoma cells (Fukuda et al., 1999b), oral cancer cells (Kuo et al., 2004) and breast cancer
12 cells (Kim et al., 2008), as well as COX-2 expression in colon cancer cells (Fukuda et al.,
13 1999a).

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31 Several studies showed that berberine could confer synergistic effects against cancer in
32 combination with ER antagonists, irradiation and other chemotherapeutic agents. A recent
33 study by Liu et al. (2009b) demonstrated that the combined treatment of ER antagonists with
34 berberine enhanced the growth inhibitory effect on MCF-7 cells (ER⁺) but not on MDA-MB-
35 231 cells (ER⁻). Peng et al. (2008) found that 5 and 10 μ M concentrations of berberine
36 combined with irradiation significantly strengthened the radio-sensitivity on A549 cells,
37 which molecular basis involved the induction of autophagic cell death as well as cell cycle
38 G2/M arrest (only at 5 μ M berberine). Arsenic trioxide (As₂O₃) is usually indicated for the
39 treatment of acute promyelocytic leukemia *in vivo*, and also induces apoptosis in various
40 tumor cells *in vitro*. Lin et al. (2008) investigated the co-effects of berberine on As₂O₃-
41 mediated inhibition of cancer cell metastasis using rat and human glioma cell lines. As a result,
42 berberine at 10 μ M enhanced the As₂O₃ (5 μ M) mediated reduction in motility and invasion of
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glioma cells, which synergistic effects were achieved by the inhibition of PKC- α activity and downstream transcriptional factors, myc and jun, and MT1-MMP and MMP-2 in the PKC-mediated signaling pathway involved in cancer cell migration. The synergistic anticancer effect using both As₂O₃ and berberine was also found in human neuroblastoma SH-SY5Y cells. The mechanism was based on the mitochondria/caspase-3 dependent signaling pathway (Kim et al., 2007). The synergy researches promote largely the development of multimodality or adjuvant chemotherapy in cancer treatment. However, the idea can be further derived from the principle of CM practice, in which different combination of the Chinese herbs could be used to treat the different syndromes under various diseases, and the research can be tuned fine by referring to the efficacious or proprietary Chinese formulae. A recent study (Wang et al., 2008) was based on such a formula, *Zuo Jin Wan*, which has a long-standing use for the treatment of gastrointestinal disorders especially inflammations. Berberine and evodiamine, another natural antitumour alkaloid, composing of the two major active ingredients of the formula, were combined to treat human hepatocellular carcinoma SMMC-7721 cells. The cell apoptosis, cell cycle distribution, and the level of TNF- α were remarkably elevated.

It is worthwhile to note that berberine induced apoptosis in U937 cells without affecting the cell cycle or induction of necrosis in B16 or HeLa cells (Letašiová et al., 2006). Berberine could also antagonize the hypoxia-induced angiogenesis in gastric adenocarcinoma via HIF-1 α repression (Lin et al., 2004b). Therefore, the antineoplastic effects of berberine against various cancer cells may involve different mechanisms and is dependent on cell types and concentrations of the agents used.

Based on the above studies and our ongoing research results on berberine, we can conclude that berberine has multiple antineoplastic effects mainly through cytotoxicity, anti-proliferation, antiinvasion and antiinflammation mechanisms (Fig. 3). The main mechanism underlying induction of cytotoxicity, growth inhibition of berberine on cancer cells, could be

1 mediated through DNA topoisomerase I inhibition and cell cycle arrest which eventually
2 induces apoptosis via the caspase-3 or Fas/FasL signal pathways. Berberine-induced
3 anticancer effects were strongly correlated with its anti-inflammatory properties through
4 cross-talk between different signaling pathways, such as NF- κ B pathway for anti-
5 inflammatory effects and caspase-dependent pathways for apoptosis. The mechanisms
6 underlying anti-invasion of berberine are relatively less understood at present. In addition to
7 down-regulate u-PA, MMP-2 and -9 expressions, the signaling pathway involved in cell
8 motility and invasion, such as the RhoA GTPase/ NF- κ B pathway, may be considered (Fig. 3).
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21 *3.3. Antiproliferation properties and underlying mechanisms of Huanglian*

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26 *Huanglian* has also been used in many anticancer studies in such cancer types as human
27 breast cancer (Li et al., 2000; Kang et al, 2005), human colon cancer (Li et al., 2000), human
28 esophageal cancer (Iizuka et al., 2000a), human gastric cancer (Li et al., 2000), human
29 leukemia (Lin et al., 2004a), human liver cancer (Lin et al., 2004a) and human pancreatic
30 cancer (Hara et al., 2005). We have also reported that *Huanglian* had the strongest
31 cytotoxicity among a selection of sixteen anticancer Chinese herbs in rat leukemia L-1210
32 cells (Luo et al., 2002). A much higher inhibitory activity for growth of tumour cells was
33 present in the water extract of *Huanglian* (or *Huanglian* extract, the most clinically applied
34 form in CM), compared to extracts using other solvents for extraction. Cells treated for 3
35 hours with 5 μ g/ml of the *Huanglian* extract showed 89 % inhibition, while those treated with
36 ethanol extract showed only 28 % inhibitory activity.
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53 In some studies, the antitumour effects of *Huanglian* extract were compared with berberine,
54 in which either IC₅₀ or ID₅₀ values were used for assessment of *in vitro* cytotoxicity (Iizuka et
55 al., 2000a and 2003; Lin et al., 2004a; Hara et al., 2005). Iizuka et al. (2000a) found that only
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Huanglian had potent antitumour effects on various esophageal cancer cells among all ingredients of the complex prescription ‘Oren-to’ (*Huanglian Jiedu* Decoction), which may be due to its major component, berberine. In another study by Lin et al. (2004a), purified berberine and coptisine (Fig. 1) were shown to be the active components, which were more potent than *Huanglian* extract against hepatoma and leukaemia cells. However, other studies (Kang et al, 2005; Hara et al., 2005; Iizuka et al., 2003; Li et al., 2000) on human pancreatic and gastric cancer cell lines demonstrated that *Huanglian* may not necessarily be less potent than berberine alone after normalization to levels of berberine present in the herb extract. The underlying mechanism may be related to genes or cellular pathways resistant to berberine’s action that were overcome or regulated by other components in *Huanglian* (Fig. 1).

On the other hand, *Huanglian* has also been found to mediate cancer cell apoptosis via similar pathways to berberine, including: a) the regulation of cytokines e.g. up-regulation of IFN- β and TNF- α (Kang et al, 2005); b) indirect effects against cancers involving the anti-inflammatory pathways. *Huanglian* could inhibit IL-6 secretion in human esophageal tumour cells (Iizuka et al., 2000b); c) the anti-angiogenesis activity (Wang et al., 2004) and/or d) inhibition of DNA topoisomerase I (Kobayashi et al., 1995). These results have mirrored the diversity of *Huanglian* against various cancers. Even though the anti-inflammatory effects of *Huanglian* have demonstrated the art of CM for cancer therapies by removing *heat* and *toxicity* (inflammatory elements in cancerous pathogenesis) and promoting virtue (body immunity) simultaneously, there are still many uncertainties ahead about the underlying mechanisms need to be resolved in the future.

3.4. Future approaches

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Current studies broadly indicate the involvement of cell cytotoxicity, cell cycle regulatory machinery and apoptosis signaling pathways as targets of anticancer by berberine and *Huanglian*. The apoptotic cell death induced by berberine and *Huanglian* in cancer cells may be the result of cell cycle arrest. The detailed molecular mechanisms remain to be elucidated, especially for *Huanglian*, whose specialty for cancer prevention or therapy has been poorly studied. Similar to the antimicrobial action (Stermitz et al., 2000), synergism may play a key role in treatment of various tumours with *Huanglian*. The nature and identity of these synergistic components as well as the types for combination however, remain to be unveiled. Furthermore, detailed involvement of pro-apoptotic genes (e.g. *p53*, *c-Myc* and *pRb*), anti-apoptotic genes (e.g. *Bcl-2* gene family members or other cancer-specific genes), the types of cell death (apoptotic, autophagic and necrotic) and the downstream gene transcription (transcriptome analysis), translation (protein expression profile) and microRNAs (miRNA) in cells treated by berberine and *Huanglian*, remain to be fully clarified. The differential responses of cancer cells and their corresponding normal counterparts to berberine and *Huanglian* also need to be investigated, since only a few studies have been conducted to compare the cancer cell and non-neoplastic cell and show no insult on normal ones (Meeran et al., 2008; Mantena et al., 2006a; Hwang et al., 2006). It is encouraging to see that some transcriptional or regulatory factors, receptors as well as non-apoptosis cell death pathways are being identified as new targets of berberine and/or *Huanglian* such as the aforementioned pro-apoptotic proteins, nucleophosmin, HIF-1 α , COX-2 and GADD153, RhoA/NF- κ B signaling pathway and autophagic cell death, leading to a better understanding of the underlying mechanisms.

4. Genomics and proteomics approaches to study antineoplastic effects of berberine and *Huanglian*

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2 Very few genomics (or more exactly transcriptome analysis) and proteomics approaches
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4 have been carried out to investigate the inhibitory effects of *Huanglian* and berberine on
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6 tumour cells. With a microarray-based mRNA approach, Efferth et al (2005) had examined
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8 the expression values of 31 ABC transporter genes and another 9,706 genes from 60 NCI
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10 cancer cell lines treated with berberine. Twenty genes were identified that significantly
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12 predicted sensitivity or resistance of the cell lines to berberine and expression of 8 ABC
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14 transporters correlated well with the IC₅₀ values for berberine. Among them, the ABCB1
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16 (MDR1) gene was validated to confer resistance to berberine in CEM/VCR1000 leukemia
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18 cells. Another study (Hara et al, 2005) used 8 human pancreatic cancer cell lines which
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20 examined the ID₅₀ of *Huanglian*. By hierarchical cluster analysis, the investigators identified
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22 27 *Huanglian*-related genes using DNA microarray data from 12,600 genes and suggested that
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24 berberine can account for the majority of the anti-proliferation activity of *Huanglian*.
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26 Previously, gene expression patterns explored by the same group using pancreatic cancer cells
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28 and oligonucleotide microarrays, also led to the identity of 12 genes from approximately 11,
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30 000 genes showing a strong correlation ($r^2 > 0.81$) with the ID₅₀ values of both agents,
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32 suggesting that these genes are associated with the anti-proliferative activity of berberine in
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34 *Huanglian* (Iizuka et al., 2003). These genomic approaches help to identify novel marker
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36 genes and cell signaling pathways affected by berberine and *Huanglian*.
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46 Similar to mRNA expression based on transcriptome analysis, proteomics also provides
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48 high-throughput analysis of expressed proteins in berberine treated cells in examining or
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50 predicting cell response to many external stimulus. However, this may not correlate with
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52 changes in mRNA levels due to complicated protein-protein or protein-DNA interactions (Fell,
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54 2001). A study conducted by Tan et al. (2006) investigated differential protein expression in a
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56 human liver cancer cell line, HepG2, after berberine treatment. The results showed a network
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1 of proteins involved in the MAPK phosphor-relay systems, metabolism, and regulation of cell
2 cycle and DNA damage responses. This work serves as a paradigm for further studies on the
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4 molecular mechanisms of berberine or *Huanglian* against many tumours.
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8 9 **5. Computer-aided molecular design and prediction of cell response to berberine and** 10 **analogs** 11 12 13 14 15 16

17 Several studies have reported the structure and activity of berberine as a small molecule
18 ligand. In one study, the X-ray crystal structures of the human topoisomerase I-DNA
19 cleavable complex, combined with a Sybyl software package, have been used to analyse the
20 properties of berberine as a potent topoisomerase I poison for a next generation anticancer
21 drug (Kettmann et al, 2004a). Another study using NCI database by COMPARE analysis,
22 investigated the responses of 60 cancer cell lines to berberine (the IC₅₀ values) and examined
23 the correlation to another 43,177 compounds included in the database, in order to establish the
24 molecular determinants of response of tumour cells to berberine. Preliminary research results
25 were able to correlate the sensitivity or resistance of the cell lines with berberine, based on
26 microarray-mRNA expression assays (Efferth et al, 2005). Recent progress in structural and
27 functional aspects of telomeric G-quadruplex and telomerase, demonstrated that the preferable
28 chemicals to telomerase activity (another promising near-universal anticancer target) could be
29 designed with high selection, specificity and fewer side effects. One such research, targeting
30 telomerase, found that the 9-substituted derivatives of berberine displayed a stronger binding
31 affinity with the telomeric G-quadruplex and corresponding high inhibitory effect against
32 telomerase activity (Zhang et al., 2007). These molecular modeling studies, have enabled
33 exploration of the actions of small anticancer drugs such as berberine in a more precise and
34 selective fashion.
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6. *In vivo* studies in animals and humans

6.1. *Huanglian*

As aforementioned, *Huanglian* can remove *damp*-induced *heat* and counteract *toxicity*, both of which are related to tumour pathogenesis in CM (Chinese Herbals Editor Board, 1999; Liao et al, 2003). *Huanglian* is now recorded in some anticancer Chinese herbal books (Bo et al, 2002; Xu et al, 1997; Tse, 1977) and can be used in daily CM clinical practice for cancer treatment. *Huanglian* and *Huanglian* formulae (with *Huanglian* as the main herb) such as *Huanglian Jiedu* decoction have been described as anticancer agents and used for treating pancreas cancer, breast cancer, liver cancer and colon cancer in clinical Chinese herbal books (Qian, 1980; Chang, 1998). Owing to the potential *in vitro* results in the treatment of cancers, a phase I dose escalation study of *Huanglian* in patients with advanced solid tumours is currently ongoing at Memorial Sloan-Kettering Cancer Center (ClinicalTrials.gov, 2009).

An *in vivo* study showed that *Huanglian Jiedu* decoction can significantly inhibit the H22 tumour in mice in a dose-dependent manner, all of the *Huanglian Jiedu* decoction in different dosage groups could highly inhibit the proliferation of 4 human cancer cell lines: Swille, SPC-A-1, SGC-7901 and MCF-7 (Sun et al., 2006).

6.2. *Berberine*

The effect of berberine on WEHI-3 leukemia cells *in vivo* was examined. The results showed that Mac-3 and CD11b markers, which indicate inhibition of differentiation of macrophages and granulocytes precursors, were down-regulated after treatment. There was no

1 affect on the expression of CD14 marker, except for the CD19 marker, which indicated
2 promotion of differentiation of the B-cell precursors. The spleen sizes from mice treated with
3 berberine, were found to be lower when compared to those from untreated animals (Yu et al.,
4 2007). The treatment with berberine was also found to be able to inhibit the leukemia growth
5 induced by Friend murine leukemia virus in BALB/c mice and further support its potential for
6 virally induced cancer chemoprevention (Harikumar et al., 2008). On the other hand, *in vivo*
7 studies involving solid tumours were usually conducted with the murine xenograft animal
8 models. The anticancer activity of berberine was examined in a transplantable B16 tumour
9 system at a range of doses from 1 mg/kg to 10 mg/kg. Significant reduction of tumour volume
10 was observed on day 16 at doses of 5 and 10 mg/kg. The dose of 1 mg/kg stimulated increase
11 of tumour mass, but other tested concentrations, 5 and 10 mg/kg, reduced the tumour mass
12 (Letašiová et al., 2005a). In other studies, the treatment with berberine (10mg/kg of body
13 weight) inhibited significantly the human tongue cancer growth in xenograft mice, showing
14 the promise for tongue cancer prevention in clinic (Ho et al., 2009b). The oral administration
15 of berberine inhibited the growth of A549 (p53+/+) and H1299 (p53-/-) lung tumours
16 inoculated subcutaneously in athymic nude mice, especially more effective in A549 tumour
17 xenograft mice (58% tumour volume reduction vs. 35% in H1299 at a dose of 100 mg/kg)
18 (Katiyar et al., 2009). Berberine also strongly suppressed the growth of human prostate cancer
19 xenograft in mice, in which LNCaP xenografts with p53 expression were more easily affected
20 by apoptotic cell death than those inoculated with p53-deficient PC-3 cells (Choi et al., 2009).

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48 Furthermore, the chemomodulatory activity of *Alstonia scholaris* extract (ASE) was
49 investigated in combination with berberine hydrochloride (BCL) on Ehrlich ascites
50 carcinoma-bearing mice (Jagetia and Baliga, 2004). The tumour-bearing animals were
51 injected with various doses of ASE, and 8 mg/kg of BCL (one-fifth of the LD₅₀) was
52 combined with different doses of ASE (60-240 mg/kg). The combination of 180 mg/kg of
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1 ASE with 8 mg/kg of BCL showed the greatest antitumour effect; the number of tumour-free
2 survivors in the treated group was greater, median and average survival time increased to 47
3 and 40.5 days, respectively, compared to the non-treated group. Similarly, 180 mg/kg of ASE
4 in combination with different doses of BCL (2-12 mg/kg), showed a dose-dependent increase
5 in anticancer activity up to 8 mg/kg of BCL. The best effect was observed when 180 mg/kg of
6 ASE was combined with 6 or 8 mg/kg of BCL, where an increase in antineoplastic activity
7 was observed. Clinically, ionizing radiation is one of most commonly used strategies for lung
8 cancer therapies, however, how to maintain the radio-sensitizing effects still remains a
9 challenge. In the study by Peng et al. (2008), berberine being given intraperitoneally at 2.0
10 mg/kg twice weekly in combination with irradiation almost abridged the growth of implanted
11 Lewis lung carcinoma in C57BL/6 mice. The mechanism was in a good agreement with the
12 autophagy-mediated tumor diminishment. The results verified the synergistic cytotoxic effect
13 of berberine and irradiation.

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31 Berberine was also found to significantly inhibit carcinogenesis induced by 20-
32 methylcholanthrene (200 µg/0.1 mL/mouse) or *N*-nitrosodiethylamine (NDEA; 0.02% NDEA
33 in distilled water, 2.5 mL/animal by gavage, five days a week for 20 weeks) in a dose-
34 dependent manner using mice and rats. Administration of berberine (0.5, 2.5 or 5.0 mg/kg)
35 could significantly reduce the incidence of tumour in animals after injection of 20-
36 methylcholanthrene, with increased life spans observed compared with controls. When
37 berberine (10, 25 or 50 mg/kg) was administered simultaneously with NDEA, the markers of
38 liver injury (liver weight, gamma-glutamyl transpeptidase activity and glutathione *S*-
39 transferase level) were reduced significantly compared with animals treated with NDEA alone.
40 Similar decreases in the serum levels of lipid peroxide, bilirubin and glutamate pyruvate
41 transaminase were also noted. Morphology of liver tissue and levels of marker enzymes
42 indicated that berberine offered protection against chemical carcinogenesis (Anis et al., 2001).
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6.3. Uncertainties

Although *Huanglian* has been used clinically for treating cancers in China, a gateway to comprehensive clinical trial is just on the very preliminary stage (ClinicalTrials.gov, 2009). On the other hand, a note of caution was observed in a study using berberine and *Huanglian* to treat hepatoma cancer cells in which high cytotoxic effect in peripheral blood mononuclear cells was observed (Lin et al., 2004a). In addition, antagonistic actions of berberine on other chemotherapeutic agents e.g. paclitaxel-induced cytotoxicity in oral (OC2), gastric (SC-M1) and colon (COLO 205) cancer cells was also observed through up-regulation of MDR1 (Lin et al., 1999), which may compromise the actions of other chemotherapeutic agents. These issues should be validated in *in vivo* studies comparing with berberine, *Huanglian* and other chemotherapeutic agents.

7. Adverse effects under *Huanglian* and berberine treatments

7.1. *Huanglian*

According to unpublished data, *Huanglian* has no major side effects after being *p.o.* administrated 200 mg/kg daily for three weeks. The acute toxicity study showed that LD₅₀ of *Huanglian* was 4.89 g /kg in mice by *gavage* (Qiu et al., 2004a); and at a daily dose of 3 g/kg for 8 days *Huanglian* could markedly elevate AST level in serum (Qiu et al., 2004b). There were some reports suggesting that *Huanglian* may cause haemolytic jaundice and nuclear jaundice of the newborn infants with glucose-6-phosphate dehydrogenase deficiency (Yang and Gao, 1996). However, neither experimental studies nor clinical investigations found any

1 related evidence on the offsprings after administration directly or through pregnant mothers
2 with *Huanglian* or *Huanglian* formulae. In contrary, *Huanglian* is commonly used as a single
3 drug or *Huanglian* formulae in CM to prevent infections in newborn infants. Prescriptions of
4 *Huanglian* are not prohibited in Chinese medicines for pregnant women (Gao et al, 2002).
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17 Berberine sulfate also has not caused any histopathological changes in rat tissues and
18 organs when administered for 6 weeks in 500 mg/kg daily oral doses (Kowalewski et al.,
19 1975). The LD₅₀ values for berberine have showed great variation among different species, or
20 may be dependent on administrative routes. The *i.v.* administration of berberine and its
21 analogs may induce higher toxic effects than *p.o.*. In an earlier study, MLD of BCL in *s.c.*
22 administration to mice was found to be 10 mg/kg, while BCL was not toxic by oral
23 administration in mice (Aoyama, 1951). In another study, the LD₅₀ of berberine sulfate was
24 24.3 mg/kg in mice under *i.p.* administration but 392 mg/kg on oral route (Sabir and Bhide,
25 1971). In rats, LD₅₀ of berberine sulfate has been reported more than 1g/kg on *p.o.*, 88.5
26 mg/kg on *i.p.*, and 14.5 mg/kg on *i.m.* administration (Kowalewski et al., 1975). On the other
27 hand, LD₅₀ for *i.p.* administration to mice and rats were found to be 30 mg/kg and 205 mg/kg
28 respectively, allowing mice to be approximately six-fold more sensitive to toxicity from
29 berberine (Registry of the Toxic Effects of Chemical Substances, 1998). Berberine chloride
30 dihydrate (BCD) was evaluated for developmental toxicity in rats and mice (Jahnke, et al.,
31 2006). In this study, there were no maternal deaths, however adverse effects were noted in
32 maternal rat, but not in fetus. The lowest observed adverse effect level (LOAEL) in maternal
33 rat remained at 531 mg/kg/day based on the feeding study and developmental toxicity. The
34 no observed adverse effect level (NOAEL) was raised to 1000 mg/kg/day BCD based on the
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gavage study. In mice, 33% of the treated females died. Surviving animals had increased water intake, and average fetal body weight per litter decreased 5-6%, with no change in live litter size. Maternal toxicity LOAEL remained at 841 mg/kg/day BCD, based on increased water consumption, while developmental toxicity LOAEL was raised to 1000 mg/kg/day BCD based on decreased fetal body weight. Developmental toxicity NOAELs in both rat and mice studies are about 500 times higher than the amount of berberine taken for dietary supplements (e.g., goldenseal) by humans and about 60-100 times higher than a pharmacologic dose.

The most common adverse responses of berberine include constipation, laxative, anaphylaxis and other skin allergies such as dermatitis and rashes, and overdose may cause respiratory and circulatory system problems. Furthermore, berberine could displace bilirubin from serum-binding proteins, causing jaundice, kernicterus, and brain damage in infants (Bateman et al., 1998; Chan, 1993, 1994). Therefore, exposure of pregnant women to berberine is not recommended because it may lead to jaundice and kernicterus in the fetus (Chan, 1993).

8. Concluding remarks

Huanglian is a Chinese medicinal herb and berberine constitutes its major active component. It has been used clinically in certain CM formulae for treating inflammatory diseases and cancers. There are three distinct medicinal agents related to *Huanglian* which are used in CM, that is, Chinese composite formulae containing *Huanglian* (e.g., *Huanglian Jiedu* decoction), the single herb *Huanglian*, and single compound berberine, however, their physiochemical nature, pharmacological profiles and clinical indications in CM are somewhat similar to each other. *Huanglian* is the main active herb of *Huanglian Jiedu* decoction, both of

1 which contain the main active principal, berberine. Hence, in the standpoint of chemistry, in
2 that berberine is the major active compound against cancer, all three agents are strongly
3 correlated. Actually, their clinical uses in modern CM practice have largely been benefited
4 from basic research achievements on berberine. In return, the clinical successes further
5 promote research activities and the understanding of molecular mechanisms as addressed
6 above. In this review, recent progresses in research of berberine and *Huanglian* against
7 cancers are highlighted.

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17 In CM, there were no direct claims of *Huanglian* to cancer therapies in early history.
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19 However, there are many indications of anti-tumour effects in Chinese herbs with similar core
20 properties to *Huanglian* such as clearing *damp-heat*, *fire* and *toxicity* (they can be conceived
21 as characteristics of inflammatory diseases nowadays) and their perspective uses in treatment
22 of diseases with tumour symptoms, which can be derived from both traditional knowledge and
23 modern basic research. The early development of berberine and *Huanglian* as botanic drugs to
24 treat infections and inflammations may also be helpful to understand more clearly their
25 potential value as anticancer agents. The positive results of carcinostatic research of
26 berberine-containing plants in early 1900s and the purification of berberine have encouraged
27 researchers to explore antineoplastic properties of *Huanglian* and berberine in conventional
28 approaches. Even though many other pharmacological effects have been reported in berberine
29 and *Huanglian* including lowering of blood cholesterol and blood pressure and decrease of
30 blood glucose, the potential use of berberine and *Huanglian* as antineoplastic agents has
31 triggered new excitement and enthusiasm in investigations. The relatively low toxicities at
32 therapeutic level for both *Huanglian* and berberine also show additional benefit for their
33 further development.

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Berberine is an essential anticancer compound in *Huanglian*. However, it is interesting to
note in some studies that *Huanglian* extract may be more effective than berberine alone (Hara

1 et al., 2005; Iizuka et al., 2003; Li et al., 2000). One explanation is that other components in
2 *Huanglian* may act synergistically with berberine (Fig. 1). The identities of these components
3 and their mode of action are still unknown. This would suggest that berberine alone might not
4 fully represent the therapeutic properties of *Huanglian* used in CM practice. This point is also
5 best illustrated by the use of *Huanglian* formulae (with the main herb as *Huanglian*) in CM, in
6 which various ingredients follow the rule of drug synergism and compatibility (Qiu, 2007).
7 The synergy research in combination with berberine and other typical tumour-killing
8 approaches have drawn a large attention so far, however, it may be greatly promoted with the
9 full understanding of the intrinsic principle implied in CM.
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21 The anticancer effects of berberine and *Huanglian* mainly involve cytotoxicity, anti-
22 proliferation and anti-invasion. It is clear that berberine is able to inhibit the growth of various
23 types of cancer cells by inhibiting DNA topoisomerase I, inducing cell cycle arrest and
24 apoptosis mainly through caspase-3 or Fas/FasL signaling pathways. Current studies also
25 demonstrate that berberine-induced anticancer effects are strongly correlated with anti-
26 inflammatory properties through cross-talk between different signaling pathways targeted,
27 such as NF- κ B pathway for anti-inflammatory effects and caspase-dependent pathways for
28 apoptosis. This evidence may further establish a link with the art of CM to treat cancers from
29 its original standpoint by removing *heat* and *toxicity* (inflammatory elements in cancerous
30 pathogenesis). In addition to down-regulation of u-PA and matrix metalloproteinases (MMPs)
31 expression, the mechanism underlying anti-angiogenic, anti-invasive and anti-metastatic
32 effects of berberine is less understood at present, which warrant further investigation. We
33 propose that inhibition of the RhoA GTPase/NF- κ B pathway may be involved anti-invasive
34 effect of berberine according to our current study. *In vivo* studies in animals will also be
35 needed to validate the *in vitro* studies.
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1 All in all, most studies and discoveries on the anticancer properties of berberine and
2 *Huanglian* have just been conducted and found during the last ten years, many more details
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4 for their clinical potential, such as cancer types, molecular targets, dosages, and indication on
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6 different individuals, remain uncovered. Therefore, further studies are still warranted in order
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8 to transfer the knowledge for antineoplastic actions of berberine and *Huanglian* to clinical
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10 applications by using advanced approaches.
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Figure captions

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5 Fig. 1 The structures of berberine and other key protoberberine-type alkaloids contained in
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7 *Huanglian*

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12 Fig. 2 The natural habitat and cultivation of *Huanglian* (*Coptis chinensis* Franch.) in two
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14 famous endemic regions in China. Upper panel (pictures kindly provided by government of
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16 *Shizu* county of Chongqing city): A. An exhibition board showed the GAP authorized field for
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18 *Huanglian* cultivation in *Shizu* county of Chongqing city, and this board indicated the area has
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20 been certificated by the SFDA (state food and drug administration) of China for the
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22 cultivation and production of the official *Huanglian*, whose quality and safety can be
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24 guaranteed. The area with a exhibition board also plays an important role in the promotion,
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26 guidance, and paradigm of the herb cultivation in other appropriate areas; B. one GAP base
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28 for *Huanglian* cultivation, which demonstrated that ecological conditions should be humid,
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30 cool, shadowed, at high altitude around 1500-2300 m sea level and with high organic and
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32 weak acidic soil; C. A photo of one single plant of *Huanglian* taken in April. Lower panel
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34 (photo taken by ourselves): a. An exhibition board showed the GAP authorized field for
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36 *Huanglian* cultivation in *Lichuan* county of *Hubei* Province; b. A GAP base for *Huanglian*
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38 cultivation, with the same conditions as B; c. A photo of one single plant of *Huanglian* taken
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40 in April.

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51 Fig. 3 A schematic diagram of molecular machinery for antineoplastic properties of berberine.
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53 Berberine has multiple anticancer effects through direct cell toxicity, anti-proliferation and
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55 anti-invasion pathways. Bold arrows with number 1, 2, 3, 4, 5 and 6 show the different cell
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57 signaling pathways targeted by berberine.→: stimulation or positive regulation; ⇝: not yet
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completely demonstrated action; \perp : inhibition. 1. Cdk_i-Cdk-cyclin cascade; 2. Mitochondrial/caspase-dependent pathway; 3. Fas/FasL signaling pathway; 4. DNA damage via topoisomerase I inhibition, and followed by p53 pathway in response; 5. NF- κ B pathway; 6. RhoA GTPase/NF- κ B pathway

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Tang et al., Fig. 1

Figure 1.

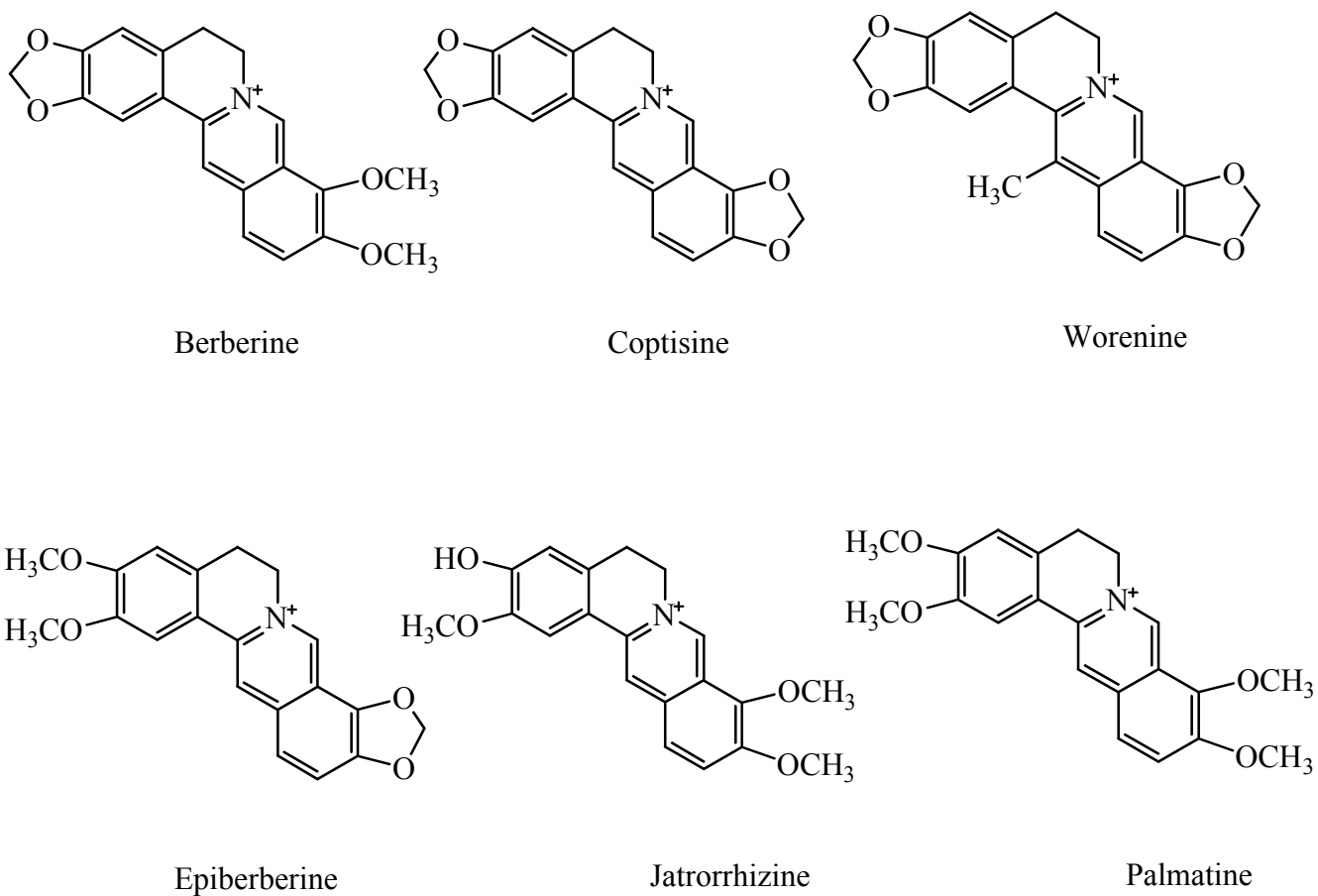


Figure 2.



Tang et al., Fig. 3

Figure 3.

