

Original Article

Analysis of the Interactions of Botanical Extract Combinations Against the Viability of Prostate Cancer Cell Lines

Lynn S. Adams, Navindra P. Seeram, Mary L. Hardy, Catherine Carpenter and David Heber

Center for Human Nutrition, David Geffen School of Medicine, University of California at Los Angeles, Los Angeles, CA 90095, USA

Herbal medicines are often combinations of botanical extracts that are assumed to have additive or synergistic effects. The purpose of this investigation was to compare the effect of individual botanical extracts with combinations of extracts on prostate cell viability. We then modeled the interactions between botanical extracts in combination isobolographically. *Scutellaria baicalensis*, *Rabdosia rubescens*, *Panax-pseudo ginseng*, *Dendranthema morifolium*, *Glycyrrhiza uralensis* and *Serenoa repens* were collected, taxonomically identified and extracts prepared. Effects of the extracts on cell viability were quantitated in prostate cell lines using a luminescent ATP cell viability assay. Combinations of two botanical extracts of the four most active extracts were tested in the 22Rv1 cell line and their interactions assessed using isobolographic analysis. Each extract significantly inhibited the proliferation of prostate cell lines in a time- and dose-dependent manner except *S. repens*. The most active extracts, *S. baicalensis*, *D. morifolium*, *G. uralensis* and *R. rubescens* were tested as two-extract combinations. *S. baicalensis* and *D. morifolium* when combined were additive with a trend toward synergy, whereas *D. morifolium* and *R. rubescens* together were additive. The remaining two-extract combinations showed antagonism. The four extracts together were significantly more effective than the two-by-two combinations and the individual extracts alone. Combining the four herbal extracts significantly enhanced their activity in the cell lines tested compared with extracts alone. The less predictable nature of the two-way combinations suggests a need for careful characterization of the effects of each individual herb based on their intended use.

Keywords: botanical – isobologram – PCSPES – prostate cancer

Introduction

In Chinese medicine, formulas such as Huangquin-tang (1) which contain combinations of natural ingredients are prescribed for the treatment of prostate cancer. These combinations are intended to attain increased potency due to synergistic interactions between the individual components, such that the combination is superior to individual herbal treatment alone. Creating herbal mixtures as opposed to utilizing a single, active herb can present the benefit of lower

doses of each individual herb, thus reducing the incidence of dose-related side effects at the same bioeffective dose of a single herb. This treatment philosophy is consistent with the systemic theory of treating chronic diseases, which advocates the inclusion of many active organic substances in formulas to not only treat the disease but to create homeostasis within the body (2).

Interactions between organic substances have been tested in limited fashion in herbal combinations and have demonstrated the possibility of antagonism as well as synergism. In a recent report (3) of eight herbs from the PCSPES formula for prostate cancer, the most cytotoxic herb (*P. notoginseng*) was varied with respect to the rest of the mixture, and isobolographic analysis carried out on the resulting data. The analysis revealed

For reprints and all correspondence: Lynn S. Adams, University of California at Los Angeles, 900 Veteran Avenue, Los Angeles, CA 90095-1742, USA. Tel: +1-310-825-6938; Fax: +1-310-206-5264; E-mail: ladams@mednet.ucla.edu

antagonism of *P. notoginseng* against the mixture of the other herbs in the formula, even though the complete PCSPEs mixture had been shown to effectively inhibit cancer cell viability in several prior studies (4,5). Therefore, the present analysis was designed to further assess the actions and interactions of plant extracts from the PCSPEs formula (*Scutellaria baicalensis*, *Rabdosia rubescens*, *Panax-pseudo ginseng*, *Dendranthema morifolium*, *Glycyrrhiza uralensis* and *Serenoa repens*) separately and in combination, in a prostate cancer cell model.

The plants that were acquired were chosen due to either their inclusion in traditional Chinese formulas for prostate cancer or based on published reports of their effects in prostate cancer cell lines and/or patients. Effects that have been reported for these extracts include: the inhibition of proliferation in prostate, oral, colon and breast cancer cells (6), a decrease in PSA and androgen receptor expression in LNCaP prostate cancer cells, induction of apoptosis via caspase-3 activation (7) and immune stimulating properties (8). Other observed effects show the induction of G₁ and G₂/M cell cycle arrest in prostate cancer, leukemia and hepatic cancer cell lines (9–12) and decreased COX-2 and Bcl₂ expression in prostate cancer cell lines (13). *S. repens*, in particular, is widely known for its ability to alleviate frequency of urination in men with benign prostatic hyperplasia (BPH) (14–16). Studies have also shown that treatment with *S. repens* is associated with prostatic epithelial contraction and decreases in tissue dihydrotestosterone (DHT) levels in men with BPH (17,18).

Three prostate cancer cell lines were studied including an immortalized prostate epithelial cell line derived from a histologically normal adult human prostate and two cell lines of differing invasiveness to assess the effects of the botanicals in a model of the multistep process of carcinogenesis. The interactions among the four most potent extracts in the 22Rv1 prostate carcinoma cell line were then analyzed using isobolographic analysis. This method was first introduced by Loewe (19) in 1928 and has been widely utilized in interaction studies of both drugs and herbs (3,20–22). In contrast to traditional means of statistical analyzing dose-effect data which merely shows an effect but does not distinguish between additivity and synergy, isobolographic analysis depicts this difference graphically thus allowing for a visual assessment of the interaction. We then calculated a combination index (CI) for the data, to mathematically compare the observed response with the expected, calculated response. In addition to the two-way combinations, a mixture of the four most potent extracts was also tested in the viability assay and compared with controls and the two-way mixtures.

Methods

Plant Materials and Extracts

Plant materials were collected, authenticated, extracted and standardized as previously reported by our laboratory (23).

Briefly, the aerial parts of the Chinese herbs (*S. baicalensis*, *R. rubescens*, *P. ginseng*, *D. morifolium* and *G. uralensis*) were collected and authenticated in China by Phytomedical Research, Inc. (Beijing, China), and extracted by Botanica Biosciences (Ojai, CA, USA), in 70% ethanol : water. Commercial standards of baicalin, baicalein, chlorogenic acid and fatty acids were purchased from Sigma (St Louis, MO, USA). Ginsenosides and wogonin were purchased from Indofine (Hillsborough, NJ, USA). Commercial standard of oridonin was purchased from Herbstandard, Inc. (Chesterfield, MO, USA). Dried berries of Saw Palmetto (Florida, USA) were extracted sequentially in cold hexane, acetone and then ethanol at the Phytochemistry Laboratory of the Center for Human Nutrition, UCLA, USA. Solvent was removed *in vacuo* at low temperature to yield hexane, acetone and ethanol extracts, respectively. The Saw Palmetto extract was analyzed for fatty acid content by gas chromatography (GC). The extracts of the Chinese herbs were analyzed for their respective chemical marker compounds (referenced above) by high-performance liquid chromatography as previously reported (23). For cell culture assays, herb extracts were dissolved in DMSO with the exception of *G. uralensis*, which was dissolved in 70% ethanol, and sterile filtered with a 0.22 µm Millex[®]-GP filter unit (Millipore Corporation, Billerica, MA, USA).

Cell Culture

All cell lines were acquired from American Type Culture Collection (Rockville, MD, USA). 22Rv1 prostate cancer cells were grown in RPMI 1640 containing 10% fetal bovine serum (FBS) in the presence of 100 U ml⁻¹ of penicillin and 0.1 g l⁻¹ of streptomycin. RWPE-1 and RWPE-2 prostate cells were grown in Defined Keratinocyte Serum Free Medium (DKSFM) containing epidermal growth factor (EGF), insulin and fibroblast growth factor (FGF). Cells were incubated at 37°C with 95% air and 5% CO₂. All cells were maintained below passage 20 and used in experiments during the linear phase of growth.

ATP Assay

Cells were plated in 96-well plates at a density of 5000 cells per well for 22Rv1 cells and 10000 cells per well for RWPE-1 and RWPE-2 cells and allowed to attach for 24 h. Cells were then treated with either media, vehicle control (≤0.30% DMSO, ≤0.15% ethanol individually for individual extract tests or in combination where extract mixtures were used) or the appropriate treatments for 24, 48 and 72 h. Viability was measured utilizing the CellTiter-Glo[®] Luminescent Cell Viability Assay (Promega, Madison, WI, USA). When added to cells, the assay reagent produces luminescence in the presence of ATP from viable cells. Results were read on the Orion Microplate Luminometer (Bertholds Detection Systems, Pforzheim, Germany). Values obtained were subtracted from blank wells containing media and vehicle (no cells) to control for the effects of vehicles on the luminescence

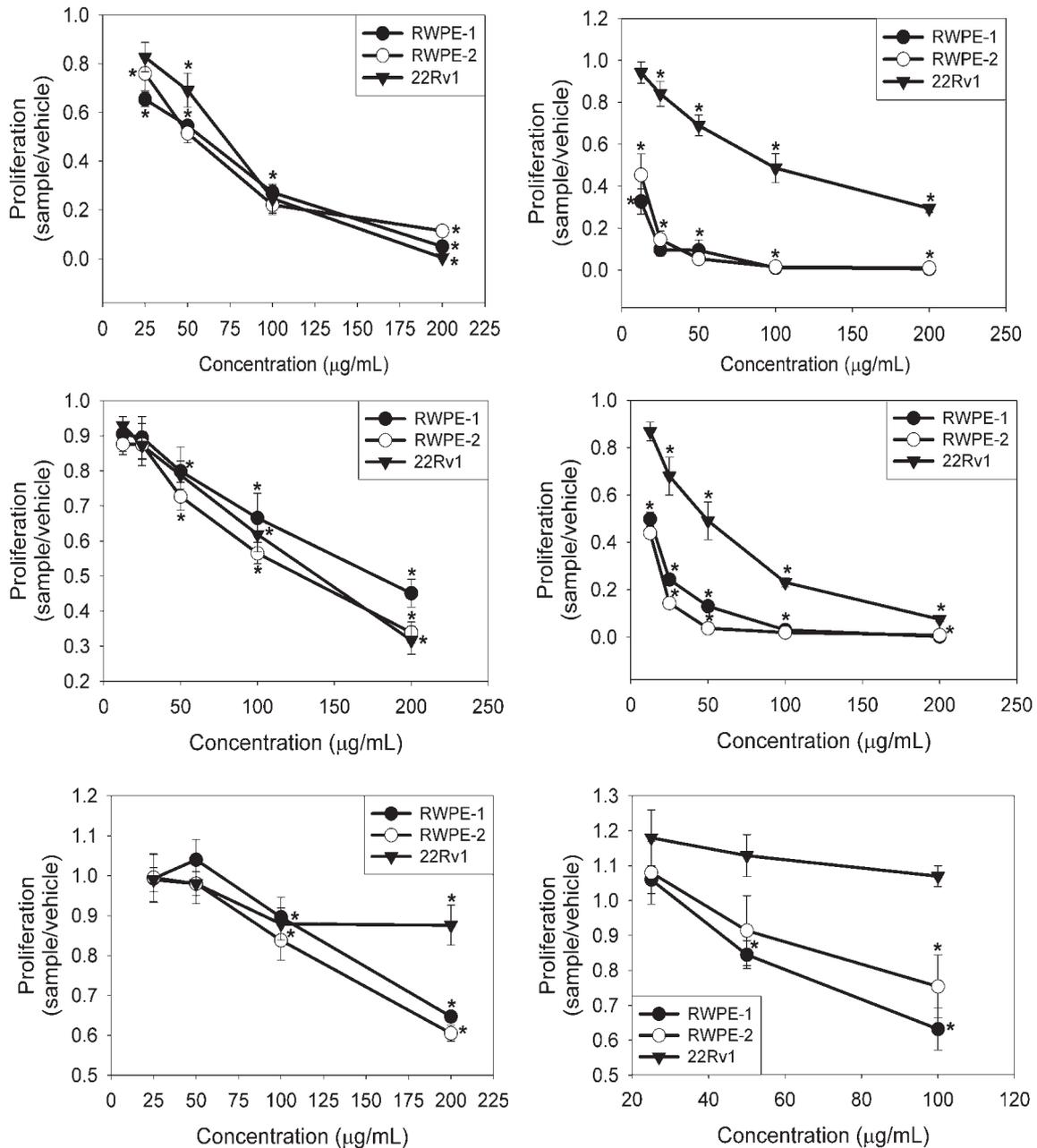


Figure 1. Herbal extracts inhibit proliferation in prostate cell lines. **I.** *S. baicalensis* **II.** *G. uralensis* **III.** *D. morifolium* **IV.** *R. rubescens* **V.** *P. ginseng* **VI.** *S. repens*. Cells were exposed to increasing concentrations of herbal extracts for 48 hours. Cell proliferation was determined via the CellTiter-Glo[®] Luminescent Cell Viability Assay. Data are expressed as a percentage of vehicle treated controls, mean \pm SE (n = 3). *P < 0.01 compared to vehicle controls.

reagents. Treatment values were then divided by vehicle control values to obtain percent change from control. Due to the large number of data collected from six extracts and three cell lines, only the concentration curve from the 48 h time point is shown.

Analysis of Antiproliferative Effect and Synergism

EC₅₀ values for each herbal extract were determined through linear regression with the GraphPad Prism 4 software program (GraphPad Software, Inc., San Diego, CA, USA). In combination experiments, extracts were added at equipotent quantities

and at $\frac{1}{2}$ and $\frac{1}{4}$ fractions of their IC₅₀ value. To assess synergy and antagonism, experimentally derived data was plotted on an isobologram and a CI was determined according to the method of Mertens-Talcott (18).

Statistical Analysis

Results are presented as mean values \pm SE. Data were analyzed by either one-way ANOVA followed by Dunnet's test ($\alpha = 0.05$) to compare experimental means with controls using GraphPad Prism 4 software. To compare statistical differences between theoretical and experimental means in the

Table 1. EC₅₀ values of individual herbal extracts in prostate cell lines

Extract	EC ₅₀ value (µg ml ⁻¹)		
	22Rv1	RWPE-1	RWPE-2
<i>S. baicalensis</i>	61.55	59.00	53.00
<i>G. uralensis</i>	94.64	9.00	11.00
<i>D. morifolium</i>	130.96	181.00	139.00
<i>R. rubescens</i>	45.00	12.50	11.00
<i>P. ginseng</i>	>200	>200	>200
<i>S. repens</i>	>200	121.00	134.00

isobolograms, Student's *t*-test was performed and $P \leq 0.05$ was accepted.

Results

Inhibition of Viability with Individual Extracts

Each of the individual extracts significantly inhibited viability of the three prostate cell lines tested in a dose-dependent manner, with the exception of *S. repens* in the 22Rv1 cell line ($P \leq 0.001$) (Fig. 1, I–VI). There was no significant difference between media and vehicle-treated controls as confirmed by two-tailed Student's *t*-test (ethanol: RWPE-1 $P \leq 0.2008$, RWPE-2 $P \leq 0.848$, 22Rv1 $P \leq 0.6098$; DMSO: RWPE-1 $P \leq 0.787$, RWPE-2 $P \leq 0.766$, 22Rv1 $P \leq 0.6098$). The EC₅₀ values obtained for each extract treatment are shown in Table 1. The four extracts that are most effective in inhibiting cell viability in these three cell lines include *S. baicalensis*, *G. uralensis*, *D. morifolium* and *R. rubescens*. Therefore, these extracts were chosen for further analysis as combinations of two herbs and all four together.

Two-way and Four-way Combination Treatments

Two-way combination studies of the four herbs exhibited varying inhibitory activities in the 22Rv1 cell line (Fig. 2). Inhibition of viability >50% of controls was taken to be indicative of additive or enhanced inhibition of growth since all botanical extracts were added at the concentration known to result in 50% inhibition when tested individually. No significant difference between media and vehicle-treated controls was observed as confirmed by two-tailed Student's *t*-test ($P = 0.530$). The mixture of *D. morifolium* and *S. baicalensis* was the most effective combination treatment, resulting in 72% inhibition of cell viability compared with vehicle control ($P \leq 0.001$). Similarly, the combination of *D. morifolium* and *R. rubescens* inhibited cell viability 57% compared with vehicle controls ($P \leq 0.001$). All other two-extract combinations resulted in <50% inhibition of cell viability. Nonetheless, all but one of the combinations tested resulted in significant inhibition of cell viability compared with controls. However, when the four herbs were combined into a single treatment (Quad), the result was an 88% inhibition of viability ($P \leq 0.001$) compared with vehicle controls (Fig. 2).

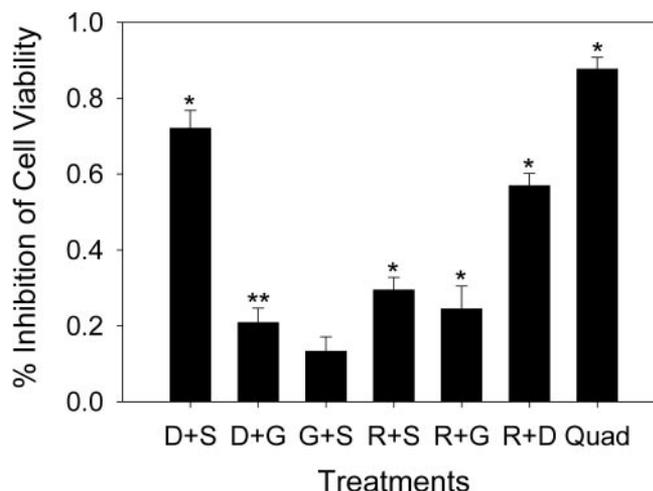


Figure 2. Herbal combinations inhibit proliferation in 22Rv1 prostate cancer cell line. Cells were exposed to two by two and a four way mixture (Quad) of the four herbs tested at EC₅₀ levels for the 22Rv1 cell line for 48 hours. Data are expressed as a percentage of vehicle treated controls, mean \pm SE ($n \geq 3$). Asterisks indicate significance from vehicle controls (single asterisk: $P \leq 0.001$, double asterisk: $P \leq 0.05$).

Isobolographic Analysis

Isobolographic analysis was performed utilizing the EC₅₀ values of each botanical extract at a 1 : 1 ratio and at appropriate dilutions of this dose ratio. The dose-effect data and statistics for the isobolograms are shown in Table 2. Plotting the EC₅₀ value of one herb of a combination on the *x*-axis and the other on the *y*-axis yielded the isobolograms shown in Fig. 3. The line connecting the two points on the graph demonstrates all combinations that would theoretically result in additivity. The combination of *D. morifolium* and *R. rubescens* falls on the line of additivity, which indicates an additive reaction with this combination. The *S. baicalensis* and *D. morifolium* combination fell below the line of additivity; however, the distance was not significant. The other four combinations all fell above the line of additivity; however, only one was not significant (*S. baicalensis* and *G. uralensis*). The remaining three combinations (*R. rubescens* and *G. uralensis*, *S. baicalensis* and *R. rubescens*) were significantly different from the line of additivity ($P < 0.05$) indicating antagonistic interactions.

Combination Index

Utilizing the EC₅₀ values of each individual herb and the EC₅₀ value of the combinations, the CI was derived to evaluate the level of interaction between the two herbs (18). Consistent with the results of the isobologram, the combination of *S. baicalensis* and *D. morifolium* was somewhat <1 (0.968), the combination of *D. morifolium* and *R. rubescens* was slightly >1 (1.299) and the CI for the *S. baicalensis* and *G. uralensis* combination was 2.034. These are the three combinations that were not significantly different from the theoretical additive dose. Alternatively, *S. baicalensis* and *R. rubescens* had a CI of 3.984, *G. uralensis* and *R. rubescens* was 7.98,

Table 2. Dose-effect data and statistics for isobolographic analysis

<i>S. baicalensis</i>		<i>D. morifolium</i>		<i>S. baicalensis</i>		<i>G. uralensis</i>	
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
25	0.1724	25	0.1262	25	0.1724	25	0.1585
50	0.3086	50	0.2116	50	0.3086	50	0.3098
100	0.7521	100	0.3800	100	0.7521	100	0.5133
200	0.9947	200	0.6826	200	0.9947	200	0.7051
Combination				Combination			
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
46.55	0.2619			39.02	0.2700		
69.83	0.3081			58.53	0.3100		
93.10	0.7206			78.04	0.4300		
Additive parameters		Experimental parameters		Additive parameters		Experimental parameters	
$f = 0.5$	$p(\text{sc}) = 0.3197$ $p(\text{den}) = 0.6803$	$Y = -2.198 + 1.428 \log(x)$		$f = 0.5$	$p(\text{sc}) = 0.3951$ $p(\text{gly}) = 0.6049$	$Y = -0.6436 + 0.557 \log(x)$	
Add = 96.22		$\log(\text{mix}) = 1.8890$ mix = 77.4462		Add = 77.86		$\log(\text{mix}) = 2.0530$ mix = 112.9796	
<i>S. baicalensis</i>		<i>R. rubescens</i>		<i>G. uralensis</i>		<i>D. morifolium</i>	
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
25	0.1724	25	0.3199	25	0.1585	25	0.1262
50	0.3086	50	0.3600	50	0.3098	50	0.2116
100	0.7521	100	0.7687	100	0.5133	100	0.3800
200	0.9947	200	0.9250	200	0.7051	200	0.6826
Combination				Combination			
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
27.84	0.1200			54.70	0.1200		
41.75	0.2000			82.05	0.2100		
55.67	0.2900			109.40	0.2200		
Additive parameters		Experimental parameters		Additive parameters		Experimental parameters	
$f = 0.5$	$p(\text{sc}) = 0.5777$ $p(\text{rap}) = 0.4223$	$Y = -0.7362 + 0.5933 \log(x)$		$f = 0.5$	$p(\text{gly}) = 0.4184$ $p(\text{den}) = 0.5816$	$Y = -0.4188 + 0.3126 \log(x)$	
Add = 53.25		$\log(\text{mix}) = 2.0840$ mix = 121.3389		Add = 112.58		$\log(\text{mix}) = 2.9390$ mix = 868.9604	
<i>D. morifolium</i>		<i>R. rubescens</i>		<i>G. uralensis</i>		<i>R. rubescens</i>	
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
25	0.1262	25	0.3199	25	0.1585	25	0.3199
50	0.2116	50	0.3600	50	0.3098	50	0.3600
100	0.3800	100	0.7687	100	0.5133	100	0.7687
200	0.6826	200	0.9250	200	0.7051	200	0.9250
Combination				Combination			
Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)	Dose ($\mu\text{g ml}^{-1}$)	Effect (% inhibition)
43.52	0.0340			35.99	0.1200		
65.27	0.1100			53.98	0.1800		
87.03	0.5700			71.97	0.2500		
Additive parameters		Experimental parameters		Additive parameters		Experimental parameters	
$f = 0.5$	$p(\text{den}) = 0.7443$ $p(\text{rab}) = 0.2557$	$Y = -2.777 + 1.674 \log(x)$		$f = 0.5$	$p(\text{gly}) = 0.6767$ $p(\text{rab}) = 0.3232$	$Y = -0.5187 + 0.4075 \log(x)$	
Add = 87.95		$\log(\text{mix}) = 1.9580$ mix = 90.7821		Add = 69.59		$\log(\text{mix}) = 2.4990$ mix = 315.5005	

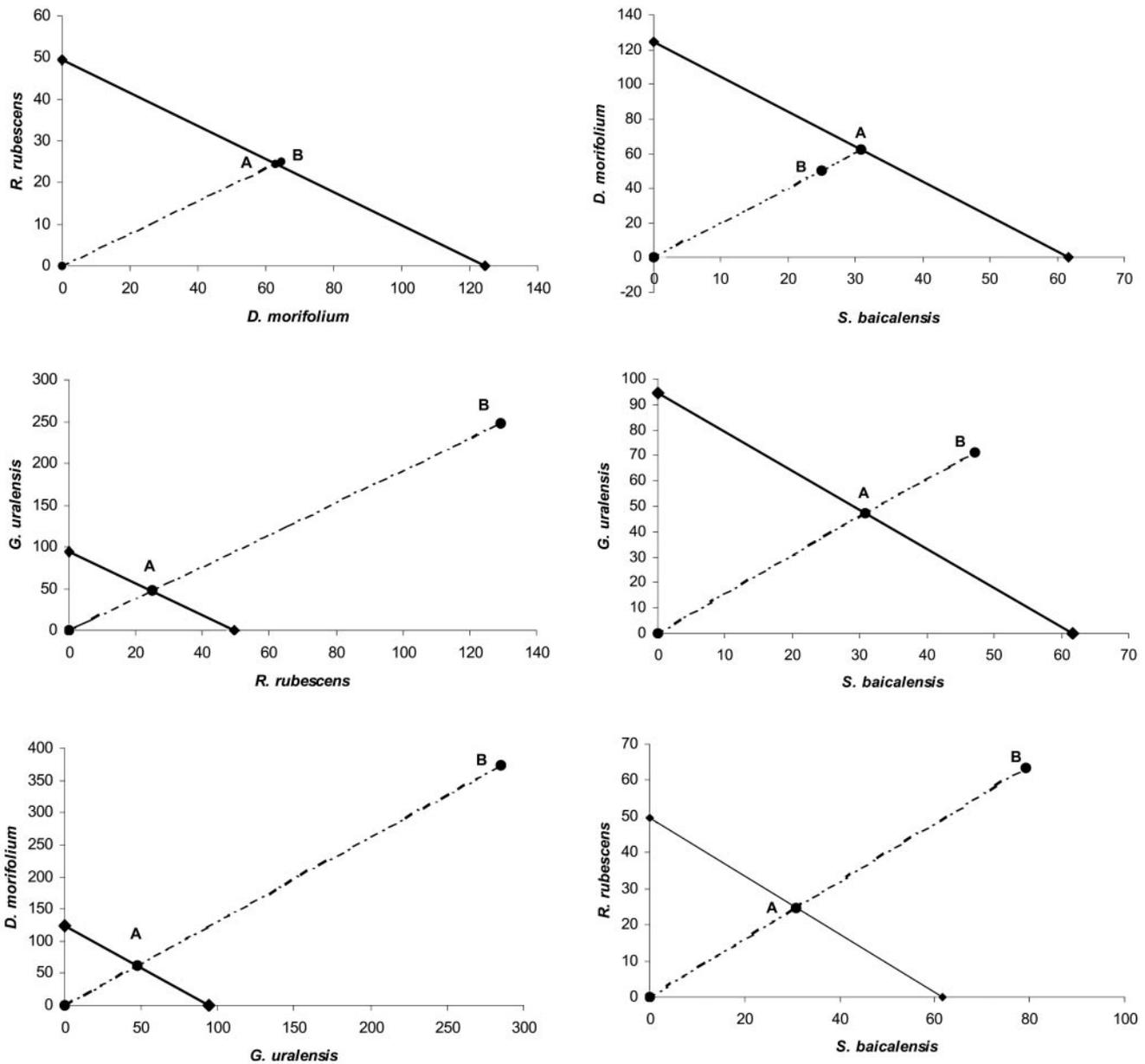


Figure 3. Isobologram demonstrating the interaction of two-by-two herbal combinations. The solid line represents the line of additivity (dose: $\mu\text{g/mL}$); the broken line represents proportions of the mixtures. Point A represents the calculated additive response where point B indicates the response achieved by testing.

Table 3. Interaction indices for herbal combinations

Combination	Combination index	P-value
<i>S. baicalensis</i> and <i>D. morifolium</i>	0.968	0.571
<i>D. morifolium</i> and <i>R. rubescens</i>	1.299	0.318
<i>S. baicalensis</i> and <i>G. uralensis</i>	2.034	0.242
<i>S. baicalensis</i> and <i>R. rubescens</i>	3.984	0.015
<i>G. uralensis</i> and <i>R. rubescens</i>	7.980	0.031
<i>G. uralensis</i> and <i>D. morifolium</i>	15.026	0.058

and the *G. uralensis* and *D. morifolium* combination had a CI of 15.026 (Table 3). These values reflect the isobologram showing that these combinations were significantly greater than the theoretical additive dose, indicating antagonism.

Discussion

The purpose of this investigation was 2-fold. First, we determined the activity of six widely used natural extracts from a well-known Chinese medicine combination (PCSPES) in different prostate cancer cell lines representing stages in the multistage process of prostate carcinogenesis. Second, we analyzed the interactions between the different herbs in combination, in a selected human prostate epithelial carcinoma cell line (22Rv1). These studies demonstrated varying effects of the four individual herbs on prostate cancer cells *in vitro*.

S. baicalensis, *R. rubescens*, *G. uralensis* and *D. morifolium* have been shown to inhibit viability in LNCaP prostate cancer cells *in vitro*. In this same report, *S. baicalensis* and

G. uralensis also decreased PSA and androgen receptor expression (24). Both a flavonoid in *S. baicalensis*, baicalin, and its aglycone form, baicalein, inhibited viability in DU145, LNCaP and PC-3 prostate cancer cell lines and induced apoptosis via caspase-3 activation (7), and induced G₁ and G₂/M cell cycle arrest in prostate cancer, leukemia and hepatic cancer cell lines (9–12). Oridonin, an extract of *R. rubescens*, decreased bcl₂ and increased bax expression in SPCA-1 lung cancer cells (25) and showed anti-angiogenic properties in an endothelial cell-based assay for angiogenesis (26).

The results of our initial viability studies confirm the effects observed with these same botanical extracts in previous studies of various prostate cancer cell lines (4,5). We found that botanical extracts from *S. baicalensis*, *G. uralensis*, *D. morifolium* and *R. rubescens* among the six that were tested showed substantial activity in these cell lines. *S. repens* and *P. ginseng* have been reported to reduce proliferation in LNCaP prostate cancer cells (13,27). These extracts were the least effective in RWPE-1, RWPE-2 and 22RV1 cells; therefore, the combination studies did not include these extracts. However, cell line specific effects are known to occur and natural compounds may act in either direct or indirect fashion. Indirect action of a compound may affect tissues or compounds outside of the cell and therefore not exhibit detectable activity in a cell culture model. An example of this would be the decrease of tissue DHT levels *in vivo* after treatment with *S. repens* (17,18).

Isobolographic analysis of the four most potent botanical extracts was studied in the 22Rv1 cell line which is the most aggressive prostate cancer cell line of the three tested. Our results indicate that some of the two-by-two combinations displayed additive effects suggesting that similar molecular targets or metabolic pathways are involved in their action. The combination of *S. baicalensis* and *D. morifolium* fell below the line of additivity indicating a trend toward synergy; however, this was not statistically significant.

Four of the combinations displayed clear antagonism. Of the four antagonistic combinations, three contain *G. uralensis* and two contain *S. baicalensis*. *S. baicalensis* and *G. uralensis* together inhibit each other to the point of rendering their action insignificant. It is unlikely that one extract would have an action in the cell that directly opposes inhibition of viability by another extract, as all of them individually result in the inhibition of cell viability. This observation affords us with opportunity for future studies into the mechanisms of these extracts.

More importantly, despite the interactions of the extracts in two-by-two combinations, the overall effect of the absolute mixture was to inhibit cell viability to 88%. This observation is consistent with the idea that combinations of botanical extracts are more effective than isolated components in the mixture. These interactions may depend upon the complimentary nature of different families of compounds in each of the extracts. This principle was demonstrated for food phytochemical extracts in a recent report which demonstrated that tomato powder containing lycopene, phytoene, phytofluene in a complex mixture

but not purified lycopene was able to inhibit prostate carcinogenesis in the NMU-testosterone mouse model (28).

The limitations in this work are related to the *in vitro* method used to evaluate cell viability, which may not reflect the conditions *in vivo* following absorption and metabolism of the extracts. As previously stated, while direct action may be observed in a cell culture model, indirect actions would unlikely be observed. Nonetheless, these findings provide valuable insight into the effect of the individual botanical extracts on prostate cancer cell viability and encourage investigation into the inhibitory targets and metabolic pathways affected by the botanical materials tested. The results of the isobologram suggest that unknown factors influencing the action of the extracts make their interactions unpredictable. The ratio of one extract to another, the mode of action and the necessity of common cellular resources are only some examples of variables that may determine the efficacy of a mixture. Therefore, comprehension of the activity of each extract when alone and in combination is essential to designing effective mixtures.

Acknowledgements

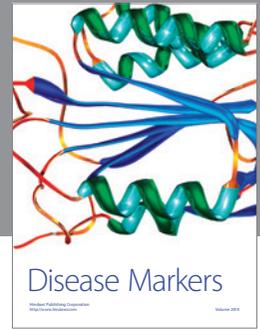
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