Effective treatment by using herbal extractive medicine to helpless liver cancer, ovarian cancer and mucinous breast carcinoma as III/IV line treatment

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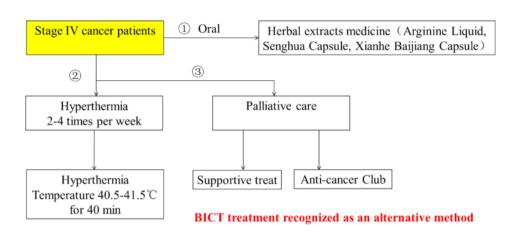
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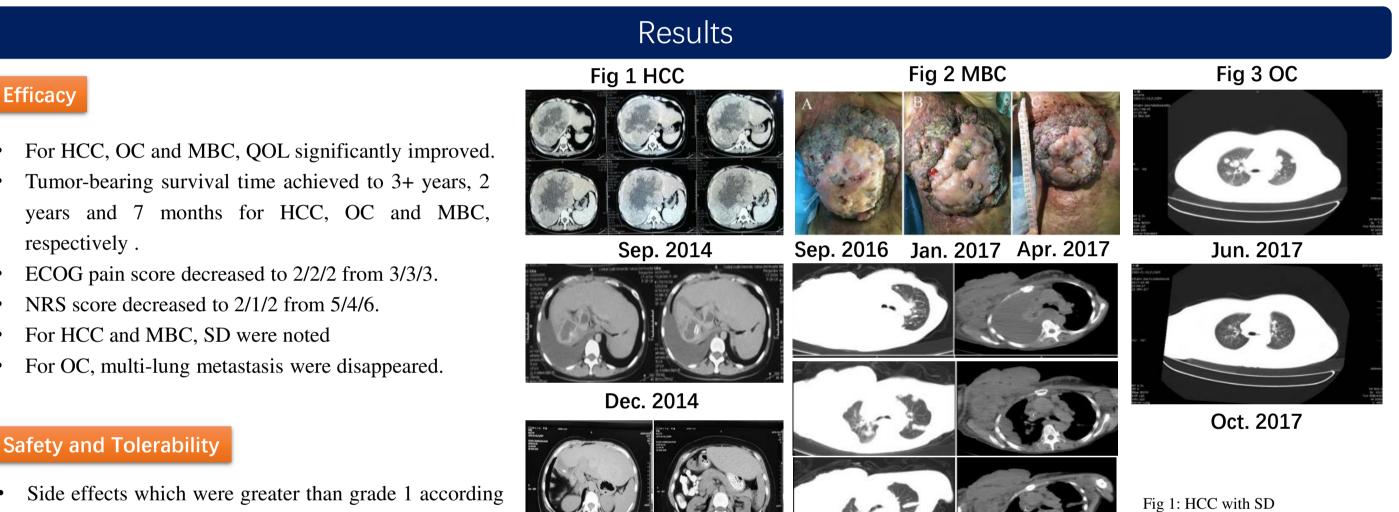
Background

No standard guideline for III/IV line treatments were determined, so we pay much attention on the such patients who need salvage treatments after I/II line treatment failure. Here, we reported terminal-stage giant hepatocellular cancer(HCC), ovarian cancer(OC) and mucinous breast carcinoma(MBC) cases who used herbal extractive preparations as palliative treatment that contain ginseng, Herba Agrimonia, White Flower Patrinia Herb etc.

Method

- Giant HCC (D=12cm) patient with intrahepatic metastasis, OC patient with multi-lung metastasis, and bilateral pure BMC patient with metastasis on bilateral lung, chest wall were enrolled.
- QOL, ECOG, NRS pain score and survival time were evaluated.
- Herbal extract preparations and arginine were given four times a day by oral, accompanied with nutrition medicine and hyperthermia.





to the CTCAE 3.0 were not observed.

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Fig 2: MBC with SD Fig 3: OC with PR

Drug analysis

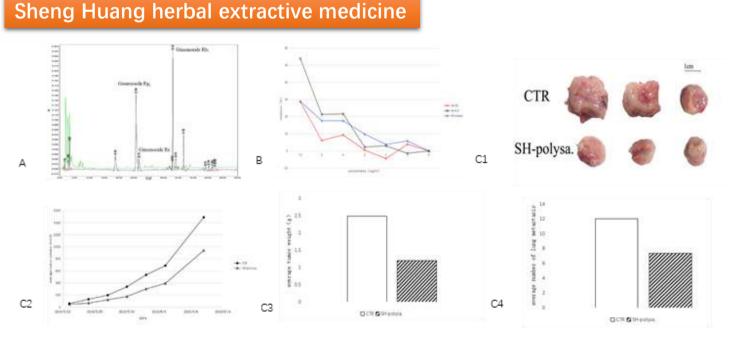


Fig. 5 A: Saponins (SH-ZG) and polysaccharides (SH-polysa.) were seperated from Shenhuang Drug and analyzed by HPLC. Ginsenoside Rg_1 , ginsenoside Re and Ginsenoside Rb_1 were detected in the sample. B: Proliferation of SH-ZG and SH-polysa were tested by MTT using 4T1 cells. The direct inhibition rate was not up to 30%. C: Xenograft 4T1 breast cancer were treated with SH-polysa. C1-3:The average tumor volume and average tumor weight of SH-polysa group significantly decreased compared to control group (P<0.05). C4: The lung metastasis numbers were decreased.

Conclusion

- Herbal extractive medicine(HEM) demonstrated an efffective for patients who failed at standard treatment regiment.
- HEM was acceptable with safety profile
- HEM provide a palliative care treatment opion for patients as III/IV line therapy.

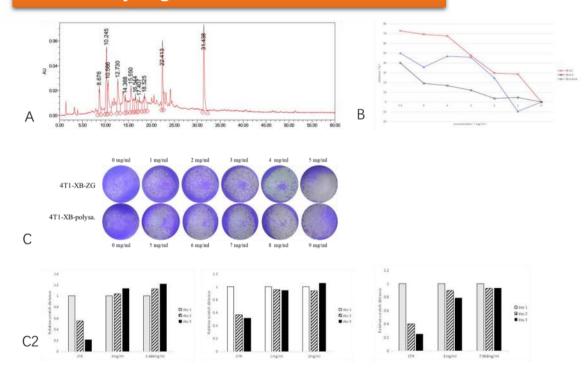


Fig. 6 A: (XB-ZG) and polysaccharides (XB-polysa.) were seperated from Xianhe Baijiang drug. After HPLC analysis, a component having a special ultraviolet absorption peak was founded and we used semi-preparative chromatographic system to separate this component. HPLC-MS analysis indicated that the component may flavonoids. **B:** 4T1 breast cancer cell viability could be inhibited by XB-ZG and XB-polysa., IC50 of XB-ZG to 4T1 is 3.367 mg/ml by MTT. **C:** C1: 4T1 cell line was treated with XB-ZG and XB-polysa. Cell clon was tested by crystal violet staining. XB-ZG and XB-polysa inhibited proliferation were dose dependence. C2-4: The relative scratches distance among groups.



Xianhe Baijiang herbal extractive medicine