

adrenergic receptor (AR) levels in the heart. Thus far, not much research has been done with regard to traditional Chinese medical treatment for CHF. We investigated the effect of Shexiangbaoxin pills (SXBP) on the function of the heart and the expression of α_1 -AR and β -AR subtypes in the messenger RNA (mRNA) levels and protein levels of non-infarction left ventricular tissue from rats with CHF induced by myocardial infarction.

Methods Models of CHF were established by left anterior descending coronary artery ligation. Fifty-four Wistar rats were randomly divided into five groups: normal control group (group A), sham operation group (group B), CHF model group (group C), positive medicine control group (group D), and small-dose SXBP group (group E) and large-dose SXBP group (group F), deployed intragastrically. Cardiac function was examined by echocardiography before and after therapy; mRNA expressed levels were measured by semiquantitative reverse transcription polymerase chain reaction (RT-PCR) for β_1 -AR, β_2 -AR, β_3 -AR, α_{1A} -AR, α_{1B} -AR, and α_{1D} -AR; protein levels were measured by western blotting analysis for β_1 -AR, β_2 -AR, α_{1A} -AR, α_{1B} -AR, and α_{1D} -AR in non-infarction left ventricular tissue.

Results There was no significant difference in the left ventricular ejection fraction (LVEF) between groups A and B. Compared to group B, LVEF of groups C, D, E, and F were significantly decreased ($p < 0.01$) before therapy. After therapy, compared to group C, LVEF of group F was significantly improved ($p < 0.05$). Compared to group B, β_1 -AR and α_{1B} -AR expressed levels were markedly decreased ($p < 0.05$), α_{1A} -AR and β_3 -AR were significantly increased ($p < 0.01$) in group C, and in both mRNA and protein expressed levels β_2 -AR had no significant difference between groups B and C ($p > 0.05$). α_{1D} -AR mRNA levels were unchanged in each group ($p > 0.05$), but α_{1D} -AR protein level was significantly decreased in group C ($p < 0.05$). After treatment, compared to group C, mRNA levels of β_1 -AR and α_{1B} -AR were significantly increased ($p < 0.05$ and $P < 0.01$), and α_{1A} -AR was markedly decreased in groups D, E, and F ($p < 0.05$). β_3 -AR level significantly declined in both groups D and F ($p < 0.01$), but β_2 -AR and α_{1D} -AR expressed levels remained unchanged in each group ($p > 0.05$). Protein levels, compared to group C, β_1 -AR was significantly increased ($p < 0.01$, $p < 0.05$, and $p < 0.01$) and α_{1A} -AR was markedly decreased in groups D, E, and F ($p < 0.05$, $p < 0.01$, and $p < 0.01$). β_2 -AR expressed level was significantly increased in group F ($p < 0.05$). α_{1B} -AR expressed level was significantly increased in both groups E and F ($p < 0.05$), and α_{1D} -AR was remarkably increased in both groups D and F ($p < 0.05$).

Conclusions After SXBP treatment, LVEF was increased and cardiac function was significantly ameliorated in rats with CHF. The therapeutic effect of SXBP may be related to better blood supply for myocardium and up-regulation of β_1 -AR and α_{1B} -AR, and down-regulation of α_{1A} -AR and β_3 -AR. The results show that SXBP can be used in treatment of CHF and the therapeutic effect of large-dose SXBP is superior to small-dose SXBP.

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EFFECTS OF SHEXIANGBAOXIN PILLS ON THE EXPRESSION OF CARDIAC α_1 - AND β -ADRENERGIC RECEPTOR SUBTYPES IN RAT HEARTS WITH HEART FAILURE INDUCED BY MYOCARDIAL INFARCTION

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Objectives Chronic heart failure (CHF) had been characterised as an activated sympathetic system leading to the alteration of