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PARACRINE EFFECTS OF EARLY BONE MARROW CELLS TREATMENT IN EXPERIMENTAL MYOCARDIAL INFARCTION IN RATS: TISSUE EVALUATION OF INFLAMMATORY PROCESS, REDOX STATUS AND ECHOCARDIOGRAPHIC PARAMETERS

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Background: The redox unbalance and inflammation are associated with cardiac dysfunction postacute myocardial infarction (AMI). Transplant of bone marrow cells (BMC) can exert beneficial effects through paracrine actions on the host tissue. *Objective:* To assess cardiac function and its correlation with redox balance and inflammatory process in cardiac tissue 48 hours post-AMI treated with BMC. Methods: Male 8-week-old Wistar rats were randomized into four groups: Sham-operated (S); AMI; S + treatment (ST) and AMI + treatment (AMIT). Induction of AMI was accomplished through ligation of the left anterior descending coronary artery, with open-chest under mechanic ventilation. Determination of ejection fraction (EF) and infarcted area (%) were evaluated by echocardiography. Tumor necrosis factor (TNF-alpha) and Interleukin 6 (IL-6) were measured by western blot, and the oxidative stress (OE) was evaluated by reduced and oxidized glutathione ratio (GSH/GSSG) and measured by spectrophotometer. Results: Infarcted area was not different between groups AMI (52.8±5.7) vs. AMIT (54.2±4.3). EF (%) was lower in the infarcted groups: AMI (51±5%) vs. S (74±7%) (p=0.001) and AMIT (56±10%) vs. ST groups (73±3%) (p=0.001). The OE was increased in infarcted groups, AMI (8.21±3.8) vs. S (14.61±3.4) (p<0.05), AMIT (2.1±0.7) vs. ST (4.7±1.5) (p<0.05) and with treatment the OE was high, AMIT (2.1±0.7) vs. AMI (8.21±3.8) (p<0.005). However, it was observed that BMC treatment was able to minimize ventricular hypertrophy (mg/q) in AMIT (2.86±0.2) vs. AMI group (3.40±0.6) (p<0.001) and minimize TNF-alpha and IL-6 expression in infarcted treated group. We found a positive correlation between ventricular hypertrophy and cytokines' expression of TNFalpha (r=0.732; p=0.001), and IL-6 (r=0.720; p=0.001). Conclusions: Our data suggest that BMC treatment attenuated the ventricular hypertrophy and reduced the expression of pro-inflammatory cytokines through its paracrine effects, at least in this time point.