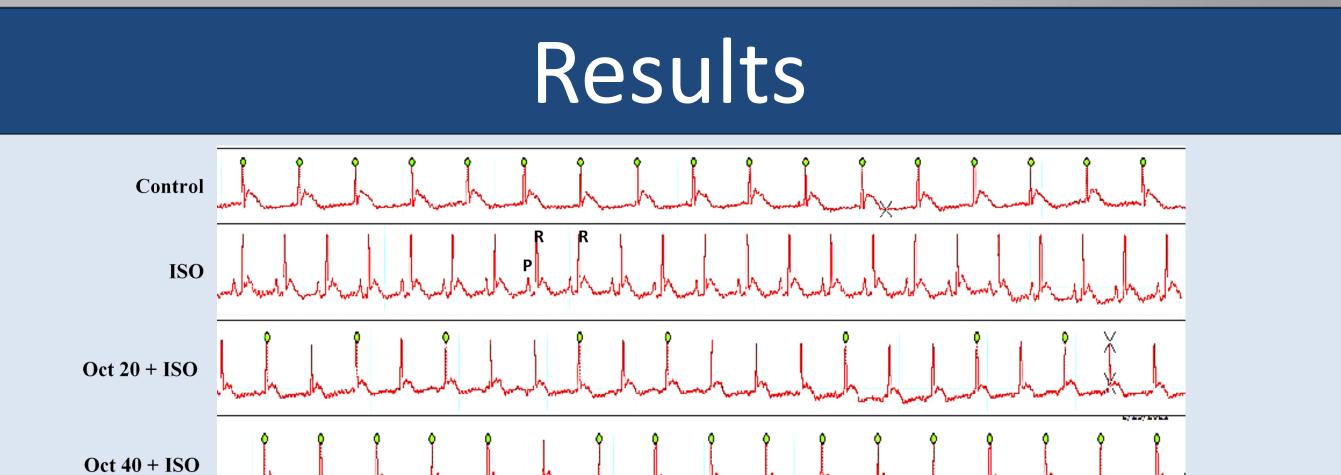
Prospective cardioprotective effect of octreotide in isoproterenol-induced myocardial infarction in rats

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Introduction

Myocardial infarction (MI) is a global health care problem, which instigates irreversible cardiac tissue damage and sudden death,

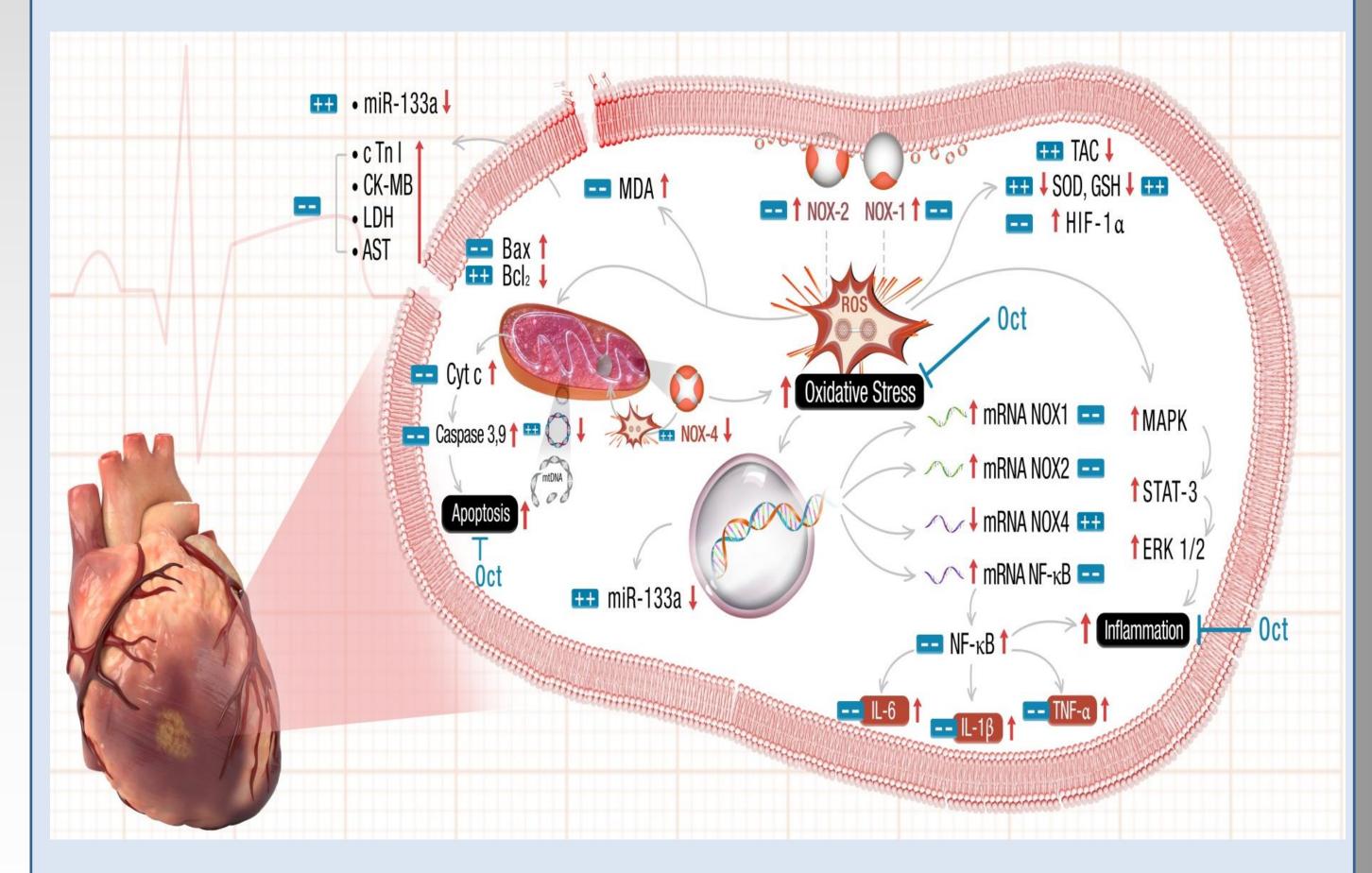


necessitating new prevention and management strategies. Hence, The current study is designed to investigate the cardioprotective effect of two different doses of octreotide on ISO-induced MI in rats with tackling the possible underlying trajectories involved including miR-133a, HIF-1 α , NOX-1/-2/-4, mitochondrial morphology, cardiac mtDNA copy number, and the inflammatory p38-MAPK/Erk1/2/STAT3/NF-κB pathway.

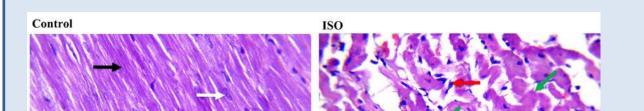
Materials and Methods

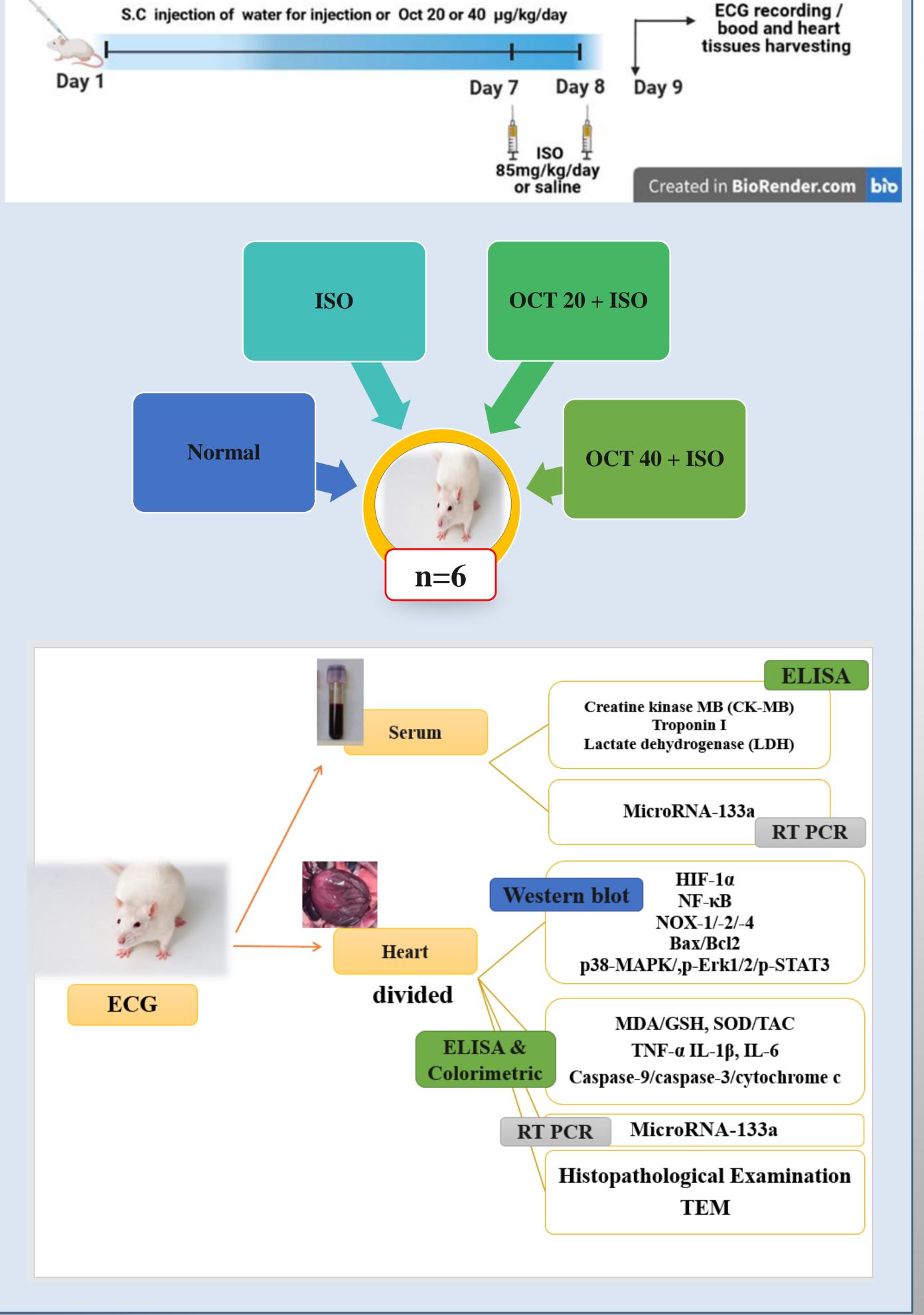
S.C injection of water for injection or Oct 20 or 40 µg/kg/day

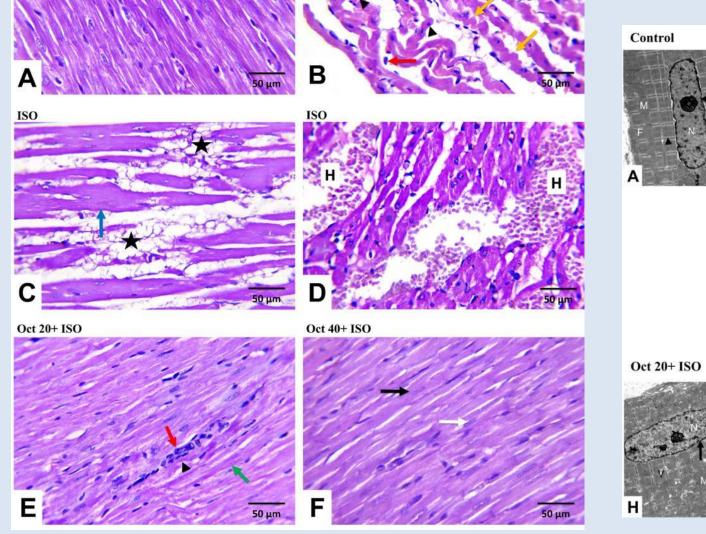
Effect of octreotide on electrocardiographic (ECG) pattern in myocardial infarcted rats

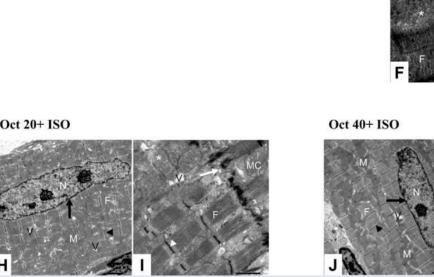


Graphical visualization showing the molecular changes in the cardiomyocyte during myocardial infarction and effect of octreotide on different measured parameters.









Representative photomicrograph demonstrating the histopathological changes of cardiac tissues in different groups (H&E, ×400).

photomicrographs Representative electron demonstrate the ultrastructural morphology of myocardium in different groups

Conclusions

Octreotide promising possesses cardioprotective capabilities against deleterious injuries induced by MI due to its antioxidant, anti-inflammatory, and antiapoptotic properties. Octreotide dose dependently modulated extremely vital molecular targets, which extensively participate in the MI associated injuries including the NOX-1/-2/-4, mitochondrial morphology, cardiac mtDNA copy number, and MAPK/Erk1/2/STAT3/NFkß pathway.

References

Braunersreuther, V., Montecucco, F., Asrih, M., Pelli, G., Galan, K., Frias, M., Burger, F., Quinderé, A.L., Montessuit, C., Krause, K.H., Mach, F., Jaquet, V., 2013. Role of NADPH oxidase isoforms NOX1, NOX2 and NOX4 in myocardial ischemia/reperfusion injury. J Mol Cell Cardiol 64, 99-107.

D'Oria, R., Schipani, R., Leonardini, A., Natalicchio, A., Perrini, S., Cignarelli, A., Laviola, L., Giorgino, F., **2020.** The Role of Oxidative Stress in Cardiac Disease: From Physiological Response to Injury Factor. Oxid Med Cell Longev 2020, 5732956.