RBT-1, a Pharmacologic Preconditioning Agent, Reduces the Incidence of Anemia, Blood Transfusion, and Use of Supplemental Iron in Patients **Undergoing Cardiac Surgery**

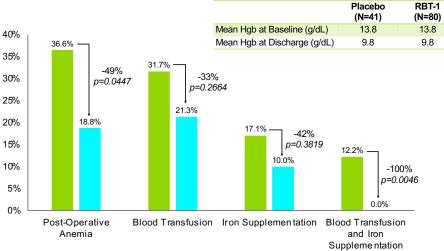


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Introduction	Methods	Discussion & Conclusions
 Patients undergoing cardiac surgery are at risk of anemia from blood loss, inflammation, and red blood cell lysis, necessitating blood product utilization and increasing the risk of post-operative complications RBT-1 (a combination of stannic protoporfin [SnPP] and iron sucrose [FeS]) is a novel preconditioning drug administered prior to surgery that upregulates anti-inflammatory, antioxidant, and iron-scavenging pathways 	 A randomized, double-blind, placebo-controlled trial Population: 121 patients scheduled to undergo non- emergent coronary artery bypass graft (CABG) and/or cardiac valve surgery on cardiopulmonary bypass (CPB) Treatment: Patients received one intravenous (IV) dose of RBT-1 or placebo (normal saline) 24 to 48 hours before surgery 	 Patients in placebo and RBT-1 groups had similar baseline characteristics A significant risk reduction in post-operative anemia was observed in response to RBT-1 When comparing all patients who required both blood transfusion and iron supplementation, a significant reduction was observed with RBT-1
 Activation of these pathways, along with the iron content of RBT-1, may improve iron levels, iron utilization, and reduce erythropoietin resistance A recent Phase 2 trial showed that the drug increased 	 Incidence of post-operative anemia, need for blood transfusion, and iron supplementation were recorded as reported by the investigators Hemoglobin (Hgb) levels were obtained at Baseline and 	 Hemoglobin levels at discharge in patients who received RBT-1 were comparable to those who received placebo, despite lesser rates of blood transfusion and iron supplementation
 cytoprotective protein levels and improved clinical outcomes Analyses of anemia incidence, transfusion, and iron usage 	followed through hospital discharge post-cardiac surgery	 Treatment with the novel preconditioning drug RBT-1 prior to cardiac surgery may reduce the risk of post-operative anemia and need for blood transfusion and iron

Results

Baseline Characteristics	Placebo (N=41)	RBT-1 (N=80)
Mean age (yrs)	65.4	65.6
Male sex (%)	73.2	75.0
White race (%)	92.7	86.3
BMI (kg/m ²)	29.7	31.5
Mean duration of surgery (hrs)	4.94	4.99
Mean time on CPB (hrs)	1.95	1.97
Comorbid Conditions	Placebo (N=41)	RBT-1 (N=80)
Age ≥65 years (%)	56.1	60.0
Congestive heart failure (%)	14.6	15.0
LVEF ≤35% (%)	4.9	11.3
Diabetes mellitus requiring insulin (%)	7.3	17.5
eGFR ≥20 to <60 mL/min/1.73m ² (%)	14.6	32.5
Pre-operative Hgb <10 g/dL (%)	2.4	1.3



Placebo (N =41) RBT-1 (N=80)



from the study are presented

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supplementation

