

Figure S1. Typical EPR spectra of the homogenate. The arrows show the magnetic field for $g = 4.3$ and the peak to peak intensity (I_s) used to measure the relative amount of labile iron.

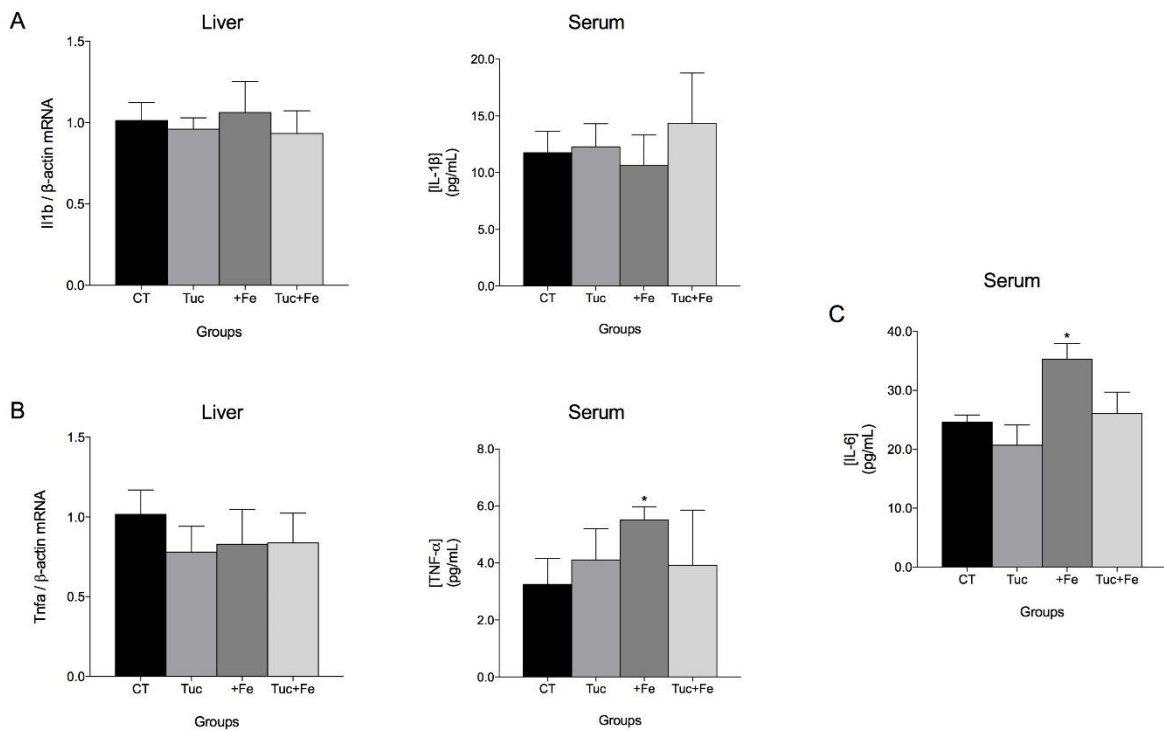


Figure S2. Hepatic mRNA levels quantification of interleukin 1 beta (Il1b) and tumor necrosis factor alpha (Tnfa) mRNA levels and serum protein levels of IL-1 β , IL-6 and TNF- α in rats fed a control diet, tucum-do-cerrado (Tuc), iron-enriched diet (+Fe) or tucum-do-cerrado iron-enriched (Tuc+Fe) diet, for 12 weeks. Data are the average \pm standard deviation ($n = 6$). Statistical differences compared to: * CT group, # Tuc group and $\text{\textcircled{S}}$ +Fe group ($p < 0.05$).

Table S1. Primer sequences used for Bmp6, Hamp, Hfe, Hfe2, Smad7, Trf1 and Actb real-time PCR assays.

Gene	Primers sequences (5' - 3')	GenBank accession number
Bmp6	GACAGCAGAGTCGCAATCG (forward)	NM_013107.1
	AGCTCACGTAAAGCTCATGC (reverse)	
Hamp	TGATGCTGAAGCGAAGGA (forward)	NM_053469
	TGTGTTGAGAGGTCAGGAC (reverse)	
Hfe	TGGGCAAGATCACCTTGAATT (forward)	NM_053301.4
	GGATCCTGTGCTCTTCCCACT (reverse)	
Hfe2	GTAGCATCGGGAGCCAAC (forward)	NM_001012080.1
Smad 7	TCAAAGG CTGCAGGAAGATT (reverse)	NM_030858.1
	AGAGGCTGTGTTGCTGTG (forward)	
	CATCGGGTATCTGGAGTAAGG (reverse)	
Trf1	GAGTTCCTGACATCATCAAGCA (forward)	NM_022712.1
	TCCAGCCTCACGAGGAGTAT (reverse)	
Actb	GTCGTACCACTGGCATTGTG (forward)	NM_031144
	CTCTCAGCTGTGGTGGTGAA (reverse)	

Table S2. Dilutions and companies of primary antibodies used for protein immunoblotting.

Protein	Dilution	Code/Company
pSMAD1/5/8	1:500	#9511 / Cell Signaling Technology
pSTAT3	1:500	#9131 / Cell Signaling Technology
β -actin	1:1000	#4967S / Cell Signaling Technology