The spleen contributes importantly to myocardial infarct exacerbation during post-ischemic reperfusion in mice via signaling between cardiac HMBG1 and splenic RAGE

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Online resource Table I

Genes	Primer sequences
IL-1β-F	AAATGATGGCTTATTACAGTGGC
IL-1β-R	GCTGTAGTGGTGGTCGGAGATT
INF-γ-F	GATGTAGCGGATAATGGAAC
INF-γ-R	GCAGGCAGGACAACCAT
TNF-α-F	CCGAGTGACAAGCCTGTAGCCC
TNF-α-R	CCCTTGAAGAGGACCTGGGAGTAGAT
MCP-1-F	CTCATAGCAGCCACCTTCAT
MCP-1-R	CTTCTTTGGGACACTTGCTG

Online resources Figure I



The infarct size exacerbation effects of different concentration of IHH. (n=3-4 in each group)

Mice underwent 20'/60' IR injury. We tested 3 different concentrations of IHH combined with sham heart homogenate. Sham and 1ug/g IHH had no IF exacerbation effect. 5 ug/g IHH lightly increased myocardial infarct size. The 10 ug/g IHH was found giving the most significant exacerbation effect. Risk Region (RR) were identical among groups. *: p<0.05 compared with control, sham IHH and 1ug/g IHH groups.

#: p<0.05 compared with other groups.

Online resources Figure II



Risk Regions were identical among groups in all experiments for TTC staining. A: Risk regions for Figure 1A; B: Risk regions for Figure 1C; C: Risk regions for Figure 2 D: Risk Regions for Figure 3C; E Risk Regions for Figure 3D; F Risk Regions for Figure 6

Online resource Figure III

Relative levels of mRNAs



40-IHH significantly increased mRNAs of cytokines in splenic leukocytes but not in bone marrow cells *: p<0.05 compared with Vehicle groups.

Online resources Figure IV



cFLFLF significantly reduced the myocardial infarct size in 40'/60' IR mice. RR (Risk region) had no significant difference between two groups. * p<0.05 compared with the 40'/60' IR group.