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Background {#Sec1} ===== Cells of the innate immune system (neutrophils, macrophages and monocytes) are the first responders to tissue injury and infection, in response to danger-associated molecular patterns (DAMPs) such as high-mobility group box 1 (HMGB1) and heat shock protein 70 (HSP70) \[@CR1--\@CR3]. Cell stimulation and activation occur via a series of membrane receptors, such as the Toll-like receptors (TLRs), the cytoplasmic NOD-like receptors (NLRs), or via engagement of receptors on the cell surface, such as the receptor for advanced glycation end-products (RAGE) \[@CR4, \@CR5]. DAMPs and activation of these receptors can activate the NF- κ B pathway, which initiates the production of cytokines and chemokines and increases the transcription of numerous genes encoding inflammatory mediators \[@CR6, \@CR7]. The role of HMGB1 in TLR signaling in macrophages is well established. HMGB1 from dying or activated cells can act as a potent cytokine/chemokine-inducing factor \[@CR8] or a 82157476af

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