PAMPs and DAMPs: signal 0s that spur autophagy and immunity

Summary: Pathogen-associated molecular pattern molecules (PAMPs) are derived from microorganisms and recognized by pattern recognition receptor (PRR)-bearing cells of the innate immune system as well as many epithelial cells. In contrast, damage-associated molecular pattern molecules (DAMPs) are cell-derived and initiate and perpetuate immunity in response to trauma, ischemia, and tissue damage, either in the absence or presence of pathogenic infection. Most PAMPs and DAMPs serve as so-called ‘Signal 0s’ that bind specific receptors [Toll-like receptors, NOD-like receptors, RIG-I-like receptors, AIM2-like receptors, and the receptor for advanced glycation end products (RAGE)] to promote autophagy. Autophagy, a conserved lysosomal degradation pathway, is a cell survival mechanism invoked in response to environmental and cellular stress. Autophagy is inferred to have been present in the last common eukaryotic ancestor and only to have been lost by some obligatory intracellular parasites. As such, autophagy represents a unifying biology, subserving survival and the earliest host defense strategies, predating apoptosis, within eukaryotes. Here, we review recent advances in our understanding of autophagic molecular mechanisms and functions in emergent immunity.

Keywords: PAMPs, DAMPs, autophagy, apoptosis, immunity, inflammation

Introduction

Danger is everywhere. The host recognizes so-called danger signals with induction of an innate and then adaptive immune response (Fig. 1). In the setting of microbial infection, pathogen-associated molecular patterns (PAMPs), present in diverse organisms but absent in the host, provide exogenous signals that alert the immune system to the presence of pathogens, thereby promoting immunity (1–3). The notion that a ‘signal 0’ for innate immunity was necessary, as postulated by Janeway, distinguished it from later signals to drive adaptive immunity (2). This led to the first identification of so-called pattern recognition receptors (PRRs). In contrast, cells release damage-associated molecular pattern molecules (DAMPs) as endogenous danger signals that alert the innate immune system to unscheduled cell death, to microbial invasion, and in response to stress (4–7).

Autophagy is a process by which cytoplasmic components, including soluble macromolecules (nucleic acids,
proteins, carbohydrates, and lipids) and organelles (e.g., mitochondria, peroxisomes, and endoplasmic reticulum) are degraded by the lysosome (8, 9). It likely evolved as a cell stress response to starvation and subsequently to limit damage and maintain cellular homeostasis as a means to exert protein/organelle quality control (10). Autophagic dysfunction is linked to several human diseases. For example, it can exert tumor suppressing (11, 12) as well as tumor promoting functions (13, 14), in a context and cell type-specific manner. Several studies reveal a crucial role for autophagy in adaptive and innate immunity (15–23), with the term ‘immunophagy’ (22) referring to all such processes collectively (Fig. 1). Autophagy degrades microbes (such as viruses, bacteria, and protozoa) that invade and gain access to the cytosol (16, 24, 25). This process is termed ‘xenophagy’ (26). The precise membrane dynamics and mediators of xenophagy, however, are not fully understood. In this review, we provide a brief overview of the process and function of autophagy and focus on the complex relationship between immunophagy and the initiating signal 0s with specific emphasis on PAMPs and DAMPs.

Fig. 1. Signal 0s play critical roles in autophagy and immunity. Pathogen-associated molecular patterns (PAMPs) and damage-associated molecular patterns (DAMPs) serve as signal 0s, inducing autophagy and immunophagy in the emergent immune response before the later Signal 1 (antigenic peptide and major histocompatibility molecules), Signal 2 (costimulatory molecules such as CD80 and CD86), both present on the surface of DCs recruited by the signal 0s. Signal 3 represents the DC provided IL-6 family cytokine expression such as IL-12 and IL-23 which promote polarization of emergent T-cell response. Signal 4 represents the integrin expression on DCs, defining the origin of the DCs and driving specialized molecules on T-cells promoting T-cell traffic to tissues. LPS, lipopolysaccharide; HMGB1, high mobility group box 1; ATP, adenosine-5’-triphosphate; PRRs, pattern recognition receptors; TLRs, Toll-like receptors; NLRs, NOD-like receptors; RLRs, RIG-I-like receptors; RAGE, receptor for advanced glycation end products.
The processes and functions of autophagy: a brief overview

Lysosomes contain acid hydrolases, enzymes that function to break up waste materials and cellular debris and regulate cell death. Nutrient sensor complexes such as mTORC1 and mTORC2 sit within the lysosomal membrane, able to initiate anabolism and mitosis when the cell is nutritionally replete. Within the immune system, lysosomes link exocytosis, endocytosis, and phagocytosis (27). The ubiquitin-proteasome system (UPS) and autophagy are two functionally linked major degradation pathways. Impairment of the UPS is compensated by upregulation of autophagy (28). In contrast, impairment of autophagy prevents ubiquitinated protein delivery to the proteasome for degradation (29). Some proteins, such as histone deacetylase-6 (HDAC-6) (30), p53 (31), and p62/SQSTM1 (29), play roles in the cross-talk between the UPS and autophagy.

In mammals, three primary forms of autophagy have been described: chaperone-mediated autophagy, microautophagy, and macroautophagy, which differ in their mechanism and function (Fig. 2). Among these, macroautophagy has been the most extensively studied (8). However, the precise mechanisms mediating microautophagy in mammalian cells are still unclear (32). Macroautophagy most commonly (hereafter referred to as ‘autophagy’) is initiated by the formation of the phagophore, followed by a series of steps, including the elongation and expansion of the phagophore and the closure and completion of a double-membrane autophagosome, which sequesters cytosolic material. Autophagosome maturation includes several vesicular fusion events that originates from early and late endosomes (amphisomes) and lysosomes (autolysosomes), followed by breakdown and degradation of the autophagosome and amphisomes through acid hydrolases inside the autolysosome. Recycling of the resulting macromolecules is mediated through permeases. Several recent studies suggest that autophagosomes may also fuse with cell membranes, promoting exocytosis, secretion, or extrusion of autophagic contents (33–35).

Autophagy involves a series of dynamic membrane-rearrangement reactions mediated by a core set of proteins – the autophagy-related (ATG) proteins (36) and other autophagy interaction network components (37). ATG proteins are composed of four functional groups (Fig. 2), including a protein serine/threonine kinase complex that responds to upstream signals (Atg1/ULK1, Atg13, and Atg17),...
a lipid kinase signaling complex that mediates vesicle nucleation (Atg6/Beclin1, Atg14, Vps34/PI3KC3, and Vps15), and two ubiquitin-like conjugation pathways that mediate vesicle expansion (the Atg8/LC3 and Atg12 systems). Atg9 is a transmembrane protein and may provide lipids to the isolation membrane by cycling between distinct subcellular compartments (38). Beclin 1/Atg6 has an important role in autophagy and tumorigenesis (11). It interacts with several cofactors to regulate the class III phosphatidylinositol 3-kinase (PI3KC3) and promote formation of Beclin 1-PI3KC3 core complexes, thereby inducing autophagy (39, 40).

Autophagy can degrade substrates in a selective manner such as mitochondria, in a process termed mitophagy. Mitophagy is now a well-established mechanism necessary for elimination of dysfunctional mitochondria and regulation of mitochondrial quality by specific mediators such as Nix/BNIP3L, Parkin, and Atg32 (41). Cells may also remove damaged mitochondria to prevent the accumulation of reactive oxygen species (ROS). The induced ROS from mitochondria or NADPH oxidases have recently been shown to be important signals linking immunity with autophagy (42–44).

Autophagy and apoptosis are both tightly regulated biological processes (Fig. 3) that play a central role in cell survival and cell death (45). These two pathways are regulated by common factors such as Bcl-2 family members and various transcription factors (Fig. 3). In addition, expressions of autophagy gene products are required for clearance of apoptotic cells and the prevention of tissue inflammation (46). Compared with apoptosis (‘programed cell death’), autophagy is primarily a cell survival process (‘programed cell survival’). In some cases, when apoptosis is compromised, such as in the setting of Bax/Bak deficiency, activation of autophagy leads to cell death (47), presumably via self-cannibalization or bioenergetic failure.

**PAMPs: the exogenous signal 0s**

In 1989, Charles Janeway (1) proposed that the immune system evolved to protect the host, not against innocuous foreign antigens but rather against infectious pathogens, and postulated that receptors on antigen-presenting cells of the innate immune system recognize so-called signal 0s, now termed PAMPs. Shortly thereafter, when we (MTL) asked him about the role of signal 0s in the setting of sterile inflammation, including cancer, he indicated that their role was indeed a concern but was at that time unclear. PAMPs are essential functional components of microorgan-
isms that direct the targeted host cell to distinguish ‘self’ from ‘non-self’ (‘stranger hypothesis’) and promote signals associated with innate immunity (48). Major PAMPs are microbial nucleic acids, including DNA (e.g. unmethylated CpG motifs), double-stranded RNA (dsRNA), single-stranded RNA (ssRNA), and 5‘-triphosphate RNA, as well as lipoproteins, surface glycoproteins, and membrane components [peptidoglycans, lipoteichoic acid, lipopolysaccharide (LPS), and glycosylphosphatidylinositol]. They are recognized by Toll-like receptors (TLRs) and other PRRs, such as retinoid acid-inducible gene I (RIG-I)-like receptors (RLRs), AIM2 like receptors (ALRs), and nucleotide-binding oligomerization domain (NOD)-like receptors (NLRs) (48–50). Most of the TLRs are believed to be homodimers, although heterodimers exist in the cases of TLR1:TLR2 and TLR2:TLR6. The Toll/interleukin-1 receptor homologous region (TIR) adapter proteins [myeloid differentiation factor 88 (MyD88), TIR adapter protein (TIRAP)/MyD88 adapter-like (MAL), translocating chain-associated membrane protein (TRAM), and TIR-domain-containing adapter-inducing interferon-β (TRIF)] also appear to associate with one another and are often illustrated as homodimers or heterodimers (Fig. 4A).

TLR4 recognizes LPS (51), a major cell wall component of Gram-negative bacteria that activates the innate immune system. Recognition of LPS requires CD14 in addition to TLR4. The responsiveness of the TLR4 and CD14 complex to LPS is enhanced by MD2. While the recognition of extracellular DNA primarily involves TLR9 (52), recognition of cytosolic DNA appears to involve several sensors (Fig. 4A) such as DNA-dependent activator of interferon (IFN)-regulatory factors (DAI) (53), hematopoietic IFN-inducible nuclear protein with the 200-amino-acid repeat (HIN-200) family members [e.g. absent in melanoma 2 (AIM2), p202, p204 (IFN-inducible IFI16 protein)] (54–56), DNA-dependent RNA polymerase III (Pol-III) (57), and leucine rich repeat (in FLII) interacting protein 1 (LRRFIP1) (58). dsRNA is recognized by TLR3 (59), RIG-I (60), melanoma differentiation-associated gene 5 (MDA5) (61), and laboratory of genetics and physiology 2 (LGP2) (62) (Fig. 4A).

Following PAMP recognition, activated TLRs and other PRRs localized to the cell surface, the cytoplasm, and/or intracellular vesicles provide signals to the host indicating the presence of a microbial infection and trigger proinflammatory and anti-microbial responses by activating a multitude of intracellular signaling pathways, including adapter molecules, kinases, and transcription factors such as nuclear fac-
tor-κB (NF-κB), activator protein-1 (AP-1), and IFN regulatory factors (IRFs) (Fig. 4A). PAMP-induced signal transduction pathways ultimately result in the activation of gene expression and the synthesis of a broad range of molecules, including cytokines, chemokines, cell adhesion molecules, and immunoreceptors that direct the adaptive immune response to invading pathogens by sensing microorganisms.

**DAMPs: the endogenous signal 0s**

In 1994, Polly Matzinger (4) proposed that the immune system is more concerned with ‘danger’ or ‘damage’ than with the distinction between self and non-self. The model starts with the idea that the immune system defines danger as anything that causes tissue stress or destruction (63, 64). In this model, antigen-presenting cells are activated by PAMPs and DAMPs from stressed or damaged tissues or microbes (65). Matzinger’s ‘danger model’ suggests why potent immune responses are initially elicited in the setting of sterile inflammation.

DAMPs are cell-derived molecules that can initiate and perpetuate immunity in response to trauma, ischemia, cancer, and other settings of tissue damage in the absence of overt pathogenic infection (Fig. 4A). DAMPs are localized within the nucleus and cytoplasm (HMGB1), cytoplasm alone (S100 proteins), exosomes [heat shock proteins (HSPs)], the extracellular matrix (hyaluronic acid), and in
Fig. 3. Overview of the major signal transduction pathways that regulate autophagy and apoptosis. Common molecular regulators include gene products that affect both autophagy and apoptosis and the pathways that they influence. The process of apoptotic cell death is mediated by two central pathways: an extrinsic pathway involving cell surface receptors (the death receptor pathway), and an intrinsic pathway using mitochondria and the endoplasmic reticulum (the mitochondrial pathway). A third pathway is mediated by cytotoxic T and NK cells delivering perforin/granzymes to promote apoptosis. CD95 (also called Fas or APO-1) induces apoptosis by forming a death-inducing signaling complex (DISC) at the receptor that contains FADD, caspase-8, and the caspase-8 regulator. Autophagy can degrade active caspase-8. The FLICE inhibitory protein (FLIP), a caspase-like molecule without proteolytic activity, protects cells from CD95-induced apoptosis. FLIP can suppress autophagy. The intrinsic mitochondrial pathway is activated by diverse apoptotic signals such as DNA damage, growth factor deprivation, and oxidative stress. Cytoplasmic translocation of mitochondrial proteins [such as cytochrome c (cyt c) and Smac/DIABLO] lead to activation of the caspase cascade and initiate apoptosis. A pivotal event in the mitochondrial pathway is mitochondrial outer membrane permeabilization (MOMP), which is mainly regulated by Bcl-2 family members. Some of the Bcl-2 family members (e.g. Bcl-2, and Bcl-XL) are anti-apoptotic, whereas others (e.g. Bax, bid, Bad, Bim, PUMA, and Noxa) are pro-apoptotic. Mitophagy is a well established mechanism necessary for elimination of dysfunctional mitochondria and regulation of mitochondrial quality in yeast or mammalian cells associated with cytosolic mediators such as NIX, Atg3, optic atrophy 1 (OPA1), dynamin-related protein 1 (DRP1), unc-51-like kinase 1 (ULK1), Parkin, Pink1, voltage-dependent anion channel 1 (VDAC1), p62/Sequestosome 1 (SQSTM1), mitofusin 1, and mitofusin 2. Beclin 1, the mammalian ortholog of yeast Atg6, plays a central role in autophagy. It interacts with several cofactors (e.g. Arg14L, UVRAG, Bif-1, Rubicon, Ambr1, HMGB1, pIPIST, VMP1, SLAM, IP3R, PINK, and Survivin) to regulate the formation of Beclin 1–PI3KC3 complexes, thereby inducing autophagy. In contrast, the BH3 domain of Beclin 1 is bound to and inhibited by Bcl-2 or Bcl-XL. This interaction can be disrupted by phosphorylation of Bcl-2 and Beclin 1, or ubiquitination of Beclin 1. Interestingly, caspase-mediated cleavage of Beclin 1 promotes cross-talk between apoptosis and autophagy. Although apoptosis-associated cleavage of Beclin 1 and Atg5 inactivates autophagy, the cleavage of Atg4D by caspase-3 generates a fragment with increased autophagic activity. In the presence of growth factors, growth factor receptor signaling activates PI3KC1 at the plasma membrane. PI3KC1 activates the downstream target AKT, leading to activation of mammalian target of rapamycin (mTOR) by inhibiting the tuberous sclerosis complex 1/2 (TSC1–TSC2), which results in inhibition of autophagy. Overexpression of the phosphatase and tensin homolog (PTEN) gene, by an inducible promoter, antagonizes PI3KC1 to induce autophagy. Ras has a dual effect on autophagy. When it activates PI3KC1, autophagy is inhibited, but when it selectively activates the Raf1–mitogen-activated protein kinase (MEK)–extracellular signal-regulated kinase (ERK) cascade, autophagy is stimulated. AMPK monitors the energy status of the cell by sensing the AMP:ATP ratio. Several upstream kinases, including liver kinase B1 (LKB1, which is activated by energy depletion), calcium/calmodulin kinase kinase-β (CaMKKβ, which is activated by cytosolic Ca2+ levels), and TGFβ activated kinase-1 (TAK-1, which is also involved in IKK activation), can activate AMPK by phosphorylation. Activated AMPK promotes inhibition of mTOR kinase, which induces autophagy. NF-κB and p53 play a double role in regulating autophagy in a transcription dependent and/or independent fashion. In contrast, E2F1 and FOXO3 positively regulate autophagy in a transcription dependent fashion. Nuclear HMGB1 inhibits the p53-dependent transactivation from the Bax promoter. Accumulation of cytosolic HMGB1 sustains autophagy by liberating the Beclin 1 and PI3KC3 complexes. RAGE-induced autophagy is associated with decreased phosphorylation of mTOR and increased Beclin 1–PI3KC3 interactions.

plasma components such as complement (C3a, C4a, and C5a). Examples of non-protein DAMPs include ATP, uric acid, heparin sulfate, RNA, and DNA. DAMPs can be mimicked by release of intracellular mitochondria, consisting of formyl peptides and mitochondrial DNA (with CpG DNA repeats), to activate human polymorphonuclear neutrophils through activation of TLR9 (66), which reveals an important link between trauma and inflammation. Following interaction between DAMPs and DAMP receptors [e.g. TLRs and the receptor for advanced glycation end products (RAGE)], activation of mitogen-activated protein kinases (MAPKs), NF-κB, and PI3K/AKT signaling pathways ensues thus mediating a potent response to cell survival and cell death (Fig. 4A). Increased serum levels of these DAMPs are associated with inflammatory diseases, including sepsis, arthritis, atherosclerosis, systemic lupus erythematosus, Crohn’s disease, and cancer.

The chromatin-associated protein HMGB1 is considered to be one of the prototypical DAMPs. Release of HMGB1 extracellularly is a common denominator in the response to both cell or tissue injury including organ harvest and associated ischemia/reperfusion insults, and microbial invasion (67–69). The redox/thiol-reducing protein HMGB1 mediates the response to infection, inflammation, migration, proliferation, and differentiation (70–72). It is specifically recognized by several cell surface receptors including RAGE, TLR4, TLR2, triggering receptor expressed on myeloid cells-1 (TREM-1), and CD24 (Fig. 5). CD24 serves as a negative signaling molecule to limit DAMP- but not PAMP-mediated inflammation (73). The HMGB1 protein induces migration and activation of human dendritic cells (DCs), eosinophil, natural killer (NK)-DC cross-talk, and T-cell activation (70, 71). HMGB1 causes TLR4-dependent activation of NADPH oxidase as well as increased ROS production (74). Interestingly, a cysteine at position 106 (Cys106) within HMGB1 is required for binding to TLR4 and activation of cytokine release in macrophages (75). The oxidation of HMGB1 Cys106 alone is sufficient to block the immunogenic activity of HMGB1 for DCs (76). These findings suggest that redox regulates HMGB1 function in the setting of emergent immunity and inflammation (Fig. 5). In addition, HMGB1 is an essential component of DNA-containing immune complexes that stimulate cytokine production through a TLR9-MyD88 pathway involving RAGE (77). In contrast, HMGB1-containing
Fig. 4. The role of TLRs, RLRs, and NLRs in PAMP and DAMP recognition. (A) Signaling pathways triggered by pathogen-associated molecular pattern (PAMPs) and damage-associated molecular pattern molecules (DAMPs). Lipopolysaccharide (LPS) activates both the myeloid differentiation factor 88 (MyD88)-dependent and Toll-like receptor 4 (TLR4) pathways. The MyD88-dependent pathway is responsible for NF-κB and mitogen-activated protein kinase (MAPK) activation, which controls induction of proinflammatory cytokines. The TIR-domain-containing adapter-inducing interferon-β (TRIF)-dependent pathway activates IRF3 by TANK-binding kinase 1 (TBK1)/IKKe, which is required for the induction of IFN-inducible genes. TLR1-TLR2 and TLR2-TLR6 recognize bacterial triacylated lipopeptide or diacyl lipopeptide, respectively, and recruit TIR adapter protein (TIRAP) and MyD88 at the plasma membrane to activate the MyD88-dependent pathway. TLR5 recognizes flagellin and activates the MyD88-dependent pathway. TLR3, TLR7, TLR8, and TLR9 reside in the endosome and recognize dsRNA, ssRNA, CpG DNA, or mitochondrial DNA (Mit DNA), respectively. They recruit TRIF or MyD88 to activate the IRF3 or IRF7 pathway. All immunogenic nucleic acids bind indicated cytosolic DNA sensors or RNA sensors, including retinoid acid-inducible gene I (RIG-I)-like receptors (RLRs), which are required for subsequent recognition by specific pattern recognition receptors to activate innate immune responses. DAMPs such as HMGB1, S100 proteins (S100s), and heat shock proteins (HSPs) activate the receptor for advanced glycation end products (RAGE), TLR4, or triggering receptor expressed on myeloid cells-1 (TREM-1) and activate the MyD88-dependent pathway. ATP binding of the P2X7 receptor and uric acid, as well as asbestos and alum, increase activation of caspase-1 by the NALP3 inflammasome and other nucleotide-binding oligomerization domain (NOD)-like receptors (NLRs) to promote secretion of IL-1β and IL-18. PAMP and DAMP-mediated signaling and induction of an innate immune response usually results in resolution of infection, but may also cause chronic inflammation or autoimmunity by altering various cell death and survival mechanisms. (B) Mitochondria in mammalian cells are removed by autophagy via the NIX adapter during developmental elimination of mitochondria, or via PTEN-induced putative kinase 1 (Pink1) and Parkin-mediated ubiquitination (Ub) of voltage-dependent anion channel 1 (VDAC1), recognized by the adapter p62 for removal of stressed (e.g. depolarized or damaged) mitochondria. (C) Intracellular bacteria exposed to the cytosol are modified by ubiquitin (Ub) and recognized by the autophagic adapters p62 or nuclear dot protein 52 (NDP52)/TBK1 for sequestration into autophagosomes and subsequent elimination.
nucleosomes from apoptotic cells induce anti-double-stranded DNA (dsDNA) and anti-histone IgG responses in a TLR2-dependent manner (78). Indeed, HMGB family members function as universal sentinels for nucleic acids in innate immune signaling (79).

**PAMPs and autophagy**

A diverse array of pathogens interact with components of the autophagic pathway including *Brucella abortus* (80, 81), *Coxiella burnetii* (82), *Porphyromonas gingivalis* (83), *Salmonella enterica* (84), *Chlamydia trachomatis* (85), *Listeria monocytogenes* (86), Group A *Streptococcus* (87, 88), *Mycobacterium tuberculosis* (89), *Leishmania Mexicana* (90), *Shigella flexneri* (91), *poliovirus* (92), *herpes simplex virus* (93, 94), *sindbis virus* (95), *dengue virus* (96), and *coronavirus* (97). As such, autophagy is likely the most ancient of the immune mechanisms, predating integration of mitochondria, generation of reactive oxygen species, and all innate and adaptive immune responses (15–20, 24, 98, 99). Autophagy is not only a surveillance system to detect cytosolic microbes but also a mechanism for immune cells such as macrophages to capture, document, and digest microbes. Several of these components engage lipid rafts (96, 100). Recent studies have uncovered key ubiquitin-binding adapters, such as p62 (101–103), nuclear dot protein 52 kDa (NDP52) (104), neighbor of BRCA1 gene 1 (NBR1) (104, 105), and NIX (34, 106), in targeting bacteria (e.g. *Salmonella*, *Shigella*, *Streptococci*, *Listeria*, and *Sindbis virus*) or mitochondria to autophagosomes by binding to LC3 (Fig. 4B,C). Other molecular species such as diacylglycerol serve as lipid signals that can also target bacteria to the autophagosome by activation of protein kinase C (107). In addition, the mammalian Atg8/LC3 family has many confirmed or likely interactions with other proteins (37), suggesting that these novel partners may be involved in xenophagy or other forms of selective autophagy.

LPS, a prototypical PAMP, directly induces autophagy in macrophages by activating the p38 MAPK and PI3KC3 pathways (108). The major signaling target of PAMPs during infection is the transcription factor NF-κB. The relationship between NF-κB and its regulation of autophagy is an area of great interest, emerging as a negative regulator of autophagy induced by tumor necrosis factor (109). NF-κB p65 also directly regulates Beclin 1 expression (110). Other studies suggest that inhibitor of NF-κB kinase (IKK), a kinase upstream of NF-κB, is directly involved in the induction of autophagy (Fig. 3) and shows no strict NF-κB correlation with control of autophagy (111, 112). Moreover, TGFβ-activated kinase 1 (TAK1)-binding proteins 2 and 3 (TAB 2
ROS accumulates in mitochondria in Atg5 (113). Autophagy is required for the activation of NF-κB in mouse embryonic fibroblasts (114). Interestingly, LPS results in the activation of nuclear factor erythroid 2-related factor 2 (Nrf2), which controls autophagic degradation by p62 (115). p62 accumulation, in turn, results in hyperactivation of Nrf2 (116). This raises the intriguing question of whether the p62-Nrf2 pathway is an important means to activate autophagy during infection.

Selective viral autophagy plays a crucial role in antiviral host defense (117). Autophagy is essential for delivering cytoplasmic viral RNA to the endosomal pathway, extinguishing infection (118). The precise mechanisms underlying type I IFN production in autophagy are unknown but have been postulated to involve Atg5-Atg12 conjugation (119). ROS accumulates in mitochondria in Atg5−/− cells without autophagy, amplifying RLR signaling pathways (44). Atg5 also participates in immunity and intracellular killing of pathogens via autophagosome-independent processes, promoting immunity-related GTPase (IRG) trafficking (120). Autophagy is an anti-microbial effector of IRG. Murine IRG, Irgm1, promotes autophagy and sustains CD4+ T-cell viability (121). The human IRG (IRGM) can eliminate mycobacteria through induction of autophagy (122), requiring IRGM expression in mitochondria (123). Moreover, IRGM is a common target of RNA viruses-mediated autophagy, which regulates viral particle production (124). Recent study suggests that Rubicon, as part of a Beclin-1-Vps34-containing autophagy complex, positively regulates NADPH oxidase (NOX2) assembly for superoxide generation in TLR2 signaling, and negatively regulates CARD9/Becl10-MALT-1 complex and cytokine production in Dectin-1 and RIG-I signaling (125, 126), suggesting a direct impact of autophagy protein on pathogen-specific host defense.

Although autophagy serves as a potent cellular strategy to clear pathogens, several viruses have evolved to exploit autophagic signaling to promote their replication, including dengue virus (96), hepatitis C virus (127), and poliovirus (128), among others. The interaction between human immunodeficiency virus (HIV) and autophagy are indeed quite complicated. HIV-1 infects CD4+ T cells as well as macrophages. Autophagy promotes HIV-1 proliferation within macrophages (129, 130). HIV-1 is also targeted for elimination by autophagy, countered by the virus assembly proteins Nef (129) and Env (130). Autophagy is inhibited in HIV-1-infected CD4+ cells (130). HIV-1-infected cells inhibit autophagy in non-infected macrophages/monocytic cells through induction of Akt and STAT3 signaling (131). Further studies are required to better understand the contribution of autophagy to HIV pathogenesis.

Autophagy also plays a role in viral antigen processing and presentation, mediating major histocompatibility complex (MHC) class I or II presentation (132–134). This requires both TLR and Atg5 signaling in DCs (134–137). In contrast, Atg5, Atg7, and Atg16L1 are all required for NOD2-induced autophagy and antigen presentation (138). Stimulation of the B-cell receptor (BCR) by DNA-containing antigens results in the translocation of both the BCR and TLR9 to autophagosomes (139). It is not clear whether autophagy mediates TLR9 signaling following translocation of CpG DNA from the endoplasmic reticulum (ER)/Golgi apparatus into the lysosome or endolysosome (140, 141). However, Atg9a, but not Atg7, controls dsDNA-driven dynamic translocation of stimulator of IFN genes (STING) from the ER to the Golgi (142).

Recent evidence suggests that autophagy is likely to play a prominent role in the pathogenesis of Crohn’s inflammatory bowel disease (CD) (143). Genome-wide association studies have identified CD-associated susceptibility genes, such as Atg16L1, NOD2, and IRGM (144), which function to regulate autophagy. Mice engineered with deficiencies in the Atg16L1 gene displayed gut inflammatory phenotypes not previously associated with autophagy (145, 146). A causal relationship clearly exists linking environmental factors, ATG16L1 genetic susceptibility, and the development of CD (147), suggesting that the interaction between host defects in autophagy and environmental stressors such as infection may be crucial for the pathogenesis of certain inflammatory diseases. NOD2 variants have also been linked with CD (138). In mammalian cells, NOD1 and NOD2 signal to induce autophagy and functionally interact with Atg16L1 (138, 148). ATG16L1 genetic variation also modulates NOD2-induced pro-inflammatory cytokine responses (149). Autophagy is involved in other inflammatory disorders including cystic fibrosis (150), obesity (151), and sepsis (152).

**DAMPs and autophagy**

HMGB1 is one of the best characterized DAMPs, expressed largely in the nucleus as a chromatin-associated protein. HMGB1 release from the nucleus and from the cell is dependent on different types of stress (Fig. 5). In response to exogenous bacterial products (such as endotoxin or...
CpG-DNA) (67, 153) or endogenous inflammatory stimuli (e.g. TNF, IFN-γ, or hydrogen peroxide) (67, 154) or apoptotic cells (155), innate immune cells actively release HMGB1. In addition, HMGB1 can be passively released from necrotic cells (68) or cells infected by viruses (156), triggering an inflammatory response (Fig. 5). HMGB1 can also be released from chemotherapy drug-induced apoptosis at later stages of tumor development (157).

There is a complex reciprocal relationship between autophagy and HMGB1. Autophagy, not apoptosis, is a major regulator of HMGB1 localization and release by ROS in the early events following cell stress (158, 159). Notably, both endogenous and exogenous HMGB1 are important regulators of autophagy (158–161). Endogenous HMGB1 competes with Bcl-2 for interaction with Beclin 1 and orients Beclin 1 toward autophagosomes (159). Interaction between HMGB1 and Beclin1 relies upon the autophagic complex ULK1-mAtg13-FIP200 (162). In addition, HMGB1 may be involved in the regulation of Bcl-2 phosphorylation by the extracellular signal-regulated kinase (ERK)/MAPK pathway (159). The intramolecular disulfide bridge (C23/45) of HMGB1 is required for binding to Beclin 1 and sustaining autophagy (Fig. 6). Reducible HMGB1 binds to RAGE, induces Beclin 1-dependent autophagy (Fig. 5), and promotes resistance to chemotherapeutic agents or ionizing radiation (158). In contrast, oxidized HMGB1 increases the cytotoxicity of these agents and induces apoptosis via the mitochondrial pathway (158). There is a direct molecular interaction between HMGB1 and p53 in colorectal cancer to regulate apoptosis and autophagy (163). Loss of p53 increases cytosolic HMGB1 leading to increased binding to Beclin 1, thereby promoting autophagy, and decreasing apoptosis. In contrast, loss of HMGB1 increases cytosolic p53 and apoptosis and decreases autophagy. HDACs regulate HMGB1 nuclear versus cytosolic localization within monocytic cells (164). HMGB1 within the nucleus enhances DNA repair and chromatin modification following DNA damage (165). It is unknown whether HMGB1 mediates the HDAC-autophagy pathway in DNA-damage repair (166), but this represents a distinct possibility. In addition, HMGB1 and the downstream mediator heat shock protein β-1 (HSPB1/HSP27) modulates mitochondrial respiration and morphology by sustaining autophagy/mitophagy (167), as we have shown, suggesting that HMGB1 is essential for mitochondrial quality control. HMGB1 forms highly inflammatory complexes with ssDNA, LPS, IL-1β, and nucleosomes (168), which interact with TLR9, TLR4, IL-1R, and TLR2 receptors, respectively. Thus HMGB1 may mediate PAMP-induced autophagy, as a ‘universal’ factor important in host defense and immune homeostasis.

In addition to HMGB1, other DAMPs such as ATP, S100, and host-DNA induce autophagy in several cellular systems (139, 169–171). ATP-induced autophagy is associated with the rapid killing of intracellular mycobacteria within human monocytes/macrophages (169), thus supporting the notion that autophagy plays a key role in the control of mycobacterial infections. Moreover, ATP-induced autophagy is dependent on the mobilization of extracellular calcium and the P2X7 receptor (171) (Fig. 4A). Recent study suggests chemotherapy-induced autophagy causes the release of ATP from tumor cells, thereby stimulating anti-tumor immune responses including recruitment of dendritic cells and CD4+ and CD8+ T cells (172). However, autophagy can also limit T cell-mediated cytotoxicity (173). Thus, the process of autophagy plays dual roles in regulation of effective chemotherapy and the host-derived anti-cancer immune responses (174).

S100 proteins or calgranulins are a group of more than 20 related calcium-binding proteins. S100A8, S100A9, and S100A12 are all expressed by phagocytes and secreted at sites of inflammation. S100A8/A9 induced autophagy plays a key role in the removal of damaged mitochondria in the setting of apoptosis (170). Although there is no direct evidence that uric acid, the NALP3 inflammasome inducer, is involved in autophagy, a close link exists between autophagy and inflammasome activation (42, 146, 175).

**TLRs, NLRs, RLRs, and autophagy**

Crossstalk between autophagy and TLRs results in the activation of innate immune responses (176). TLRs promoting autophagy include the TLR2/TLR1 heterodimer (177), TLR3 (178), TLR4 (108, 178–180), TLR5 (180), TLR6 (180), TLR7/8 (178), and TLR9 (139, 180) in various cell types including macrophages, DCs, and neutrophils. Direct stimulation of TLR7, however, does not lead to induction of autophagy in plasmacytoid DCs (118). Conventional DCs demonstrate high levels of basal autophagy, and afford very little or no induction of autophagy on stimulation with other types of immunological agonists or TLR signals (146, 181). LPS stimulation increases the number of autophagosomes in primary human monocytes, although it fails to induce autophagy in primary mouse macrophages (108, 146). In addition, agonists of mouse TLR7 induce autophagy in RAW264.7 myeloid cell lines and weakly in murine primary bone marrow macrophages (178). These
studies suggest that induction of autophagy following TLR stimulation is a cell-type-specific response. TLR-induced autophagy appears to depend on both MyD88 and TRIF (178, 180) (Fig. 6). Moreover, TLR signaling enhances the interaction of MyD88 and TRIF with Beclin 1 by NF-κB. NF-κB inhibits TNF-α-induced autophagy. Multiple means to promote ROS production converge on the mitochondria or alternatively, NADPH oxidases such as NOX2 and NOX4. These in turn result in activation of autophagy through Atg4 activation, ultimately reducing the binding of Beclin 1 to Bcl-2. TRIF-dependent and/or MyD88-dependent TLR4 pathway is required for LPS-induced autophagy in macrophages. Moreover, TRAF6-mediated ubiquitination of Beclin1 amplifies TLR4-induced autophagy. HMGB1 links sterile injury and infection-induced immunity. Stimuli that enhance reactive oxygen species promote cytosolic translocation of HMGB1 and thereby enhance autophagic flux. HMGB1 directly interacts with Beclin1, displacing Bcl-2 requiring the cysteines at positions C23 and C45 within HMGB1. The HMGB1/RAGE interaction activates parallel signaling pathways, including ERK1/2 and NF-κB activation. Mitogen-activated protein kinases (MAPKs), such as JNK1 and ERK1/2, also phosphorylate Bcl-2 driving subsequent dissociation of the Beclin 1-Bcl-2 complex. Notably, DAPK phosphorylates Beclin 1, promoting the dissociation of Beclin 1 from Bcl-2 like proteins, which in turn induces autophagy. Activation of phagocytic NADPH oxidase (NOX2) by HMGB1 requires TLR4 expression; the activation of other NADPH oxidases and interaction with TLRs is less clear. Thus PAMPs (LPS) and DAMPs (HMGB1) drive autophagy/immunophagy, regulating immune, stromal, and tumor cell functions.

Fig. 6. Signaling pathways triggered by LPS and HMGB1 in autophagy/immunophagy. The inflammatory response, including the recruitment and migration of immune cells to the site of infection and release of cytokines, is mediated by lipopolysaccharide (LPS). LPS bound to LPS-binding protein (LBP) is presented to CD14. CD14 maneuvers the LPS-LBP complex to TLR4, and LPS, in combination with accessory molecule MD2, activates TLR4 signaling. LPS induces activation of the tumor necrosis factor receptor (TNFR)-associated factor 6 (TRAF6)-p38 MAPK pathway and induces the expression of TNF-α and Beclin 1 by NF-κB. NF-κB inhibits TNF-α-induced autophagy. Multiple means to promote ROS production converge on the mitochondria or alternatively, NADPH oxidases such as NOX2 and NOX4. These in turn result in activation of autophagy through Atg4 activation, ultimately reducing the binding of Beclin 1 to Bcl-2. TRIF-dependent and/or MyD88-dependent TLR4 pathway is required for LPS-induced autophagy in macrophages. Moreover, TRAF6-mediated ubiquitination of Beclin1 amplifies TLR4-induced autophagy. HMGB1 links sterile injury and infection-induced immunity. Stimuli that enhance reactive oxygen species promote cytosolic translocation of HMGB1 and thereby enhance autophagic flux. HMGB1 directly interacts with Beclin1, displacing Bcl-2 requiring the cysteines at positions C23 and C45 within HMGB1. The HMGB1/RAGE interaction activates parallel signaling pathways, including ERK1/2 and NF-κB activation. Mitogen-activated protein kinases (MAPKs), such as JNK1 and ERK1/2, also phosphorylate Bcl-2 driving subsequent dissociation of the Beclin 1-Bcl-2 complex. Notably, DAPK phosphorylates Beclin 1, promoting the dissociation of Beclin 1 from Bcl-2 like proteins, which in turn induces autophagy. Activation of phagocytic NADPH oxidase (NOX2) by HMGB1 requires TLR4 expression; the activation of other NADPH oxidases and interaction with TLRs is less clear. Thus PAMPs (LPS) and DAMPs (HMGB1) drive autophagy/immunophagy, regulating immune, stromal, and tumor cell functions.
TLR4-induced autophagy (182) (Fig. 6). In contrast, the de-ubiquitinating enzyme A20 reduces ubiquitination of Beclin 1 and limits the induction of autophagy (182). A similar process may be linked to p62 recognition of microbial targets (91), as TRAF6 binds to p62. In addition, phosphorylation by death-associated protein kinase (DAPK) of Beclin 1 on Thr 119 in the BH3 domain promotes autophagy (183). Other protein modifications have been identified in Beclin 1-PI3KC3 complex formation (40). Beclin 1 may play a critical role in TLR-mediated autophagy by post-translational modification.

Autophagy may also participate in the regulation of TLR-mediated inflammation. The most direct way in which autophagy influences inflammation is the breakdown of invading microorganisms such as the Bacille Calmette-Guerin (89, 184) or the centrally important adapter protein myeloid differentiation factor 88 (MyD88) (185). Phagocytosis is one of the basic tools of innate immunity. TLR signaling in macrophages links the autophagic pathway to phagocytosis (179). LC3-associated phagocytosis (LAP) is required for the clearance of dead cells (186). Phagocytosis of cells dying through autophagy evokes a pro-inflammatory response in macrophages (186, 187). Vitamin K3 reduces pancreatic inflammation in acute pancreatitis through inhibition of the autophagic pathway (188). Possible links between these two forms of cellular ‘eating’ represent a new dimension in host defense and inflammation, potentially accessible with novel therapeutics.

In mammalian cells, NLR family members such as NOD1 and NOD2 induce autophagy to control bacterial infection and promote antigen presentation (138, 148). In Drosophila, the NLR-type PRR PGRP-LE is crucial for the induction of autophagy of L. monocytogenes (189). Four NLR family members have been described as components of inflammasomes: NALP1, NALP3, NLRC4, and NAIP5 (175). Blockade of autophagy by genetic ablation of the autophagy regulators Atg16L1 or Atg7 enables LPS-dependent inflammasome activation including the processing of pro-IL-1β into IL-1β (146). Various inflammasomes stimuli trigger autophagy in macrophages by activating nucleotide exchange (the replacement of GDP by GTP) on RalB (190). Moreover, mitophagy/autophagy blockade leads to the accumulation of damaged, ROS-generating mitochondria, and this in turn activates the NLRP3 inflammasome (42). Conversely, autophagic proteins regulate NALP3-dependent inflammation by preserving mitochondrial integrity (191). These findings suggest that autophagy contributes to homeostatic regulation of the inflammasome through the clearance of dysfunctional mitochondria and ROS production. In addition, NLR members may negatively regulate maturation of the inflammasome through interact with Beclin1 (192). Interestingly, autophagy not only inhibits IL-1β release by targeting pro-IL-1β for p62-mediated lysosomal degradation (193, 194) but also promotes IL-1β release by unconventional secretory pathway (195), suggesting a dual roles of autophagy in regulation of IL-1β signaling including inflammasome activation.

The RLR family members recognize RNA viruses within the cytosol and induce the expression of potent antiviral factors, such as type I IFN and proinflammatory cytokines. RLRs are negatively regulated by Atg5-Atg12 (119) and can activate autophagy (196). Further analysis reveals that ROS play a key role in autophagy-mediated RLR signaling (44). In addition, the pathogen receptor CD46 (197) and the T-cell receptor CD40 on myeloid and other cells (198) can activate autophagy with microorganism recognition.

RAGE, AIM2/AIM2-like receptors, and DAI as DNA receptors and autophagy

RAGE induces cellular inflammation signaling events upon binding of a variety of ligands, such as glycated proteins, amyloid-β, HMGB1, and S100 proteins (199). Moreover, RAGE can directly bind to dsDNA and dsRNA in vitro (200), suggesting that RAGE may act as both a DNA and RNA receptor (Fig. 4A). In addition, interaction between RAGE and TLR9 contributes to autoimmune pathogenesis (77), whereas interaction between RAGE and TLR2 limits inflammation (201). RAGE is linked functionally to outcome in several infectious diseases including cancer, diabetes, and Alzheimer’s disease (199, 202). Blockade of RAGE suppresses inflammation, tumor growth, and metastasis in various tumor models (203, 204).

We have shown that RAGE positively regulates autophagy (Fig. 6). RAGE sustains autophagy and limits apoptosis, promoting pancreatic tumor cell survival (205–207). RAGE-sustained autophagy is associated with decreased phosphorylation of the mammalian target of rapamycin (mTOR) and increased Beclin 1-PI3KC3 interaction and ATG12-ATG5 conjugation (205–207). In addition, knockdown of RAGE but not TLR4 diminishes HMGB1-induced autophagy in cancer cells (158). In contrast, TLR4 is required for PAMPs such as LPS to induce autophagy in macrophages (108), suggesting that the induction of autophagy by DAMPs or PAMPs may have alternative receptor-dependent pathways. Another RAGE ligand, the heterodimer S100A8/A9, also induces autophagy (170), although it is
not clear whether RAGE mediates this process directly. RAGE is an important inflammatory mediator that modulates crosstalk between pro-survival pathways: IL-6/pSTAT3 and autophagy in pancreatic ductal adenocarcinoma tumor cells and contributes to early pancreatic intraepithelial neoplasia formation (205). RAGE also functions as a phosphatidylserine receptor and assists in the clearance of apoptotic cells by phagocytosis in macrophages (208). Further studies are required to explore the structural basis and protein modification(s) necessary for RAGE-mediated autophagy and phagocytosis in immunity.

Cytosolic DNA derived from vaccinia virus can be sensed by AIM2 in a complex with Asc and caspase-1, leading to the processing of pro-IL-1β to IL-1β (54). The AIM2-like receptors (ALR) including the recently identified IFI16 form a newly defined family activating a unique inflammasome. Their roles in regulation of autophagy are currently undefined. In certain cultured cell lines, AT-rich dsDNA can also be sensed by the protein DAI, which drives IFN-β production through activation of the protein kinase TBK-1 (53). The DNA sensor LRRFIP1 mediates the production of type I IFN via a β-catenin-dependent pathway (58). Cytosolic DNA-dependent RNA polymerase III (Pol-III) is the DNA sensor linking DNA release by pathogenic bacteria and viruses in the host cell cytosol to IFN-β production and innate immunity (57) (Fig. 4A). It is unknown whether these cytosolic DNA sensors, including p202 and p204 (IFI16), evoke a robust innate immune response in an autophagy-dependent pathway.

**HMGB1/RAGE-mediated autophagy and energy metabolism**

Autophagy functions in protein and organelle quality control under basal conditions and can be activated in response to stress. The breakdown products derived from autophagy have a dual role, providing substrates for both biosynthesis and energy generation (209). ATP is produced by cellular respiration (either glycolysis or oxidative phosphorylation/OXPHOS) in mitochondria. Mitophagy is important in maintaining mitochondrial homeostasis (41). Suppression of HMGB1 expression in fibroblasts and cancer cells significantly inhibits both OXPHOS and glycolysis, and ATP production is decreased in HMGB1-deficient cells (167). We demonstrated that HMGB1 modulates the expression of HSPB1. As a cytoskeleton regulator, HSPB1 is critical for dynamic intracellular trafficking during autophagy and mitophagy. Phosphorylation of HSPB1 (both Ser15 and Ser86) is required for HMGB1-dependent mitochondrial homeostasis. Loss of either HMGB1 or HSPB1 results in a phenotypically similar deficiency in mitophagy typified by mitochondrial fragmentation with decreased aerobic respiration and ATP production. In addition, the Pink1-Parkin pathway is required for HMGB1/HSPB1 mediated mitophagy. Recent findings indicate that mitochondrial STAT3 sustains the altered glycolytic and OXPHOS activities characteristic of cancer cells (210). We found that RAGE-mediated autophagy is required for IL-6-induced mitochondrial localization and function of STAT3 (205). Knockdown of RAGE by shRNA in murine and human pancreatic tumor cell lines significantly decreases IL-6/STAT3-mediated mitochondrial respiratory chain complex I activity and ATP production. These findings reveal a novel pathway coupling autophagy and cellular energy metabolism. Further studies involved with assessment of adaptive immune responses induced by chemotherapy (174) or immunotherapy with IL-2 (211) or cytolytic cells (212) suggest that there will be a complex interplay between innate factors such as DAMPs and PAMPs and autophagy.

**Conclusion**

Multiple positive feedback loops between DAMPs and PAMPs and their overlapping receptors temporally and spatially drive immune regulatory functions (Fig. 6). Interestingly, these exogenous and endogenous signal 0s all induce and increase autophagic flux in an ROS-dependent fashion. Interactions between immune and dying tumor cells likely determine the balance between immunity and tolerance to tumor cells. As a defense mechanism, autophagy limits damage, sustains cell viability, removes intracellular pathogens, and participates in antigen presentation. The organization of the cellular networks linking autophagy to other biologic processes is quite complicated (37). For example, mTOR (213), Beclin 1- (214), and Atg5/Atg7-independent (215) alternative autophagy-activating pathways have been discovered. Although the role of autophagy in host defense responses has been extensively investigated in vitro, it is now important to more broadly assess its role in vivo. Moreover, the relationship between autophagy and other membrane trafficking systems including phagocytosis, endocytosis, and exocytosis and their relation to host defense remains largely unknown and needs more intensive study. Whether autophagy is required for generation of immunological memory following inflammation initiated by signal 0s and the subsequent recruitment and maturation of inflammatory/immune
cells is unknown. Whether autophagy serves as a necessary feedback loop to allow emergence of innate and adaptive immune function and the recall response are similarly unclear. This knowledge is important in an era developing and applying autophagy-inhibiting drugs. In addition, DAMPs such as HMGB1 may have either a pro-tumor or anti-tumor immune effect, depending on the tumor type, established immune suppressor and effector cells, the state of oxidation extracellularly, and the overall nature of the tumor microenvironment. More studies are needed to confirm whether HMGB1, as a DNA binding protein, couples with cytosolic DNA sensors as well as RAGE to regulate autophagy and innate immunity in response to pathogen-derived DNA, mitochondrial DNA, or nuclear DNA damage. Clearly this evolutionarily ancient system of autophagy is connected to many emergent innate and adaptive immune responses, largely through the response to stress, DAMPs, and ROS.

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