

Secretory versus Degradative Autophagy: Unconventional Secretion of Inflammatory Mediators

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Key Words

Autophagy · Alarmins · Inflammasome · Calpain

Abstract

Autophagy (macroautophagy) is often defined as a degradative process and a tributary of the lysosomal pathway. In this context, autophagy carries out cytoplasmic quality control and nutritional functions by removing defunct or disused organelles, particulate targets and invading microbes, and by bulk digestion of the cytoplasm. However, recent studies indicate that autophagy surprisingly affects multiple secretory pathways. Autophagy participates in extracellular delivery of a number of cytosolic proteins that do not enter the conventional secretory pathway via the Golgi apparatus but are instead unconventionally secreted directly from the cytosol. In mammalian cells, a prototypical example of this manifestation of autophagy is the unconventional secretion of a major proinflammatory cytokine, IL-1 β . This review examines the concept of secretory autophagy and compares and contrasts the role of autophagy in the secretion of IL-1 α and IL-1 β . Although IL-1 α and IL-1 β have closely related extracellular inflammatory functions, they differ in intracellular activation, secretory mechanisms and how they are affected by autophagy. This example indicates that the role of autophagy in secretion is more complex, at least

in mammalian cells, than the simplistic view that autophagosomes provide carriers for unconventional secretion of cytosolic proteins.

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Introduction: Secretory Autophagy

Our current view of protein secretion from eukaryotic cells is dominated by the well-established classic paradigm of conventional secretion (fig. 1, right). This paradigm rests on the extensively characterized biosynthetic pathway utilized by proteins endowed with sorting signals (leader peptides) authorizing them to enter the lumen of the endoplasmic reticulum (ER). From there, they undergo modifications and transport through the Golgi apparatus (G), and are finally secreted upon fusion of post-Golgi carriers with the plasma membrane (PM). A new protein secretion paradigm has been recently defined in yeast [1–3] as autophagy-based unconventional secretion in eukaryotic cells. In principle, this phenomenon (secretory autophagy; fig. 1, left) may enable any cytosolic protein to be exported from the cell where it may exert either its primary or a cryptic biological function. A direct function of autophagy in unconventional secretion by providing carriers for secretion (fig. 1, left) has been

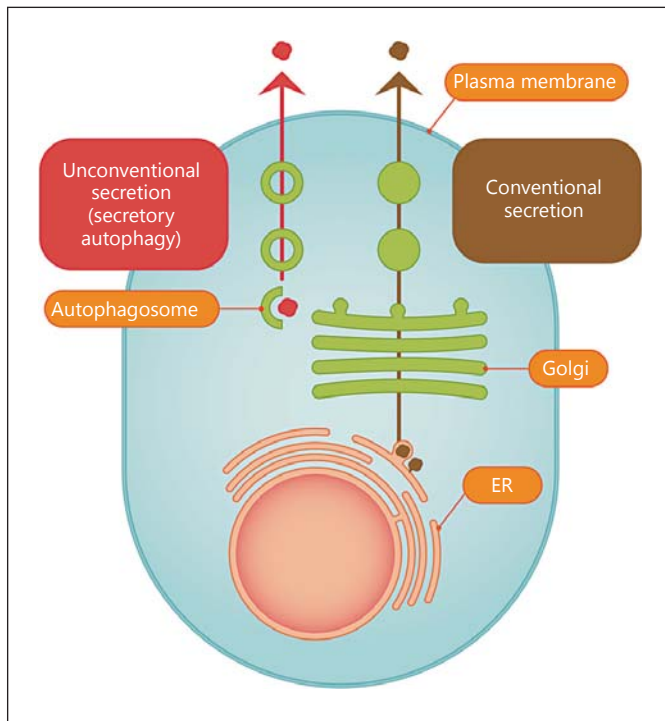


Fig. 1. Protein secretion: conventional biosynthetic pathway versus unconventional secretion through secretory autophagy. The well-developed paradigm of conventional protein secretion through the ER, Golgi and post-Golgi trafficking (right arrow) versus the process (left arrow) of autophagy-dependent unconventional secretion of cytosolic proteins (secretory autophagy). The proteins destined for conventional secretion enter ER via signal peptides, whereas cytosolic proteins destined for secretory autophagy are sequestered into autophagosomes to be exported from the cell.

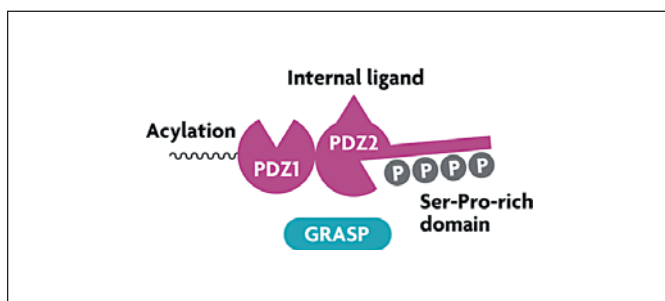


Fig. 2. GRASP domains and salient features. GRASP contains two PDZ domains, an internal protein sequence ('internal ligand') that binds to one of the PDZ domains allowing homo-oligomerization, N-terminal acylation that facilitates GRASP association with membranes, and a Ser-Pro-rich domain that is a substrate for phosphorylation by Ser/Thr kinases. Phosphorylation modulates GRASP interactions and the oligomerization state.

described in yeast, using a combination of genetic and cell biological tools [1–3]. Secretory autophagy has also been implicated in mammalian cells [4–6]. At present, there are two principal features defining secretory autophagy: (1) participation of Atg factors and contribution of autophagy as a pathway, and (2) dependence on Golgi re-assembly and stacking protein (GRASP; fig. 2) known as GRASP55 and GRASP65 in mammalian cells [7], dGRASP in *Drosophila*, GrpA in *Dictyostelium* and Grh1 in yeast [6, 8–11].

Degradative Autophagy

Autophagy is currently best known for its role in metabolism as well as a form of organelle and protein turnover and quality control [12, 13]. Canonical degradative autophagy (macroautophagy) allows cells to digest their cytoplasmic components as an endogenous source of nutrients and energy at times of starvation or as a mechanism for the clearance of defective organelles and toxic intracellular aggregates [14]. The canonical autophagy pathway depends on a suite of autophagy-specific factors (termed Atg and followed by a number) responsive to upstream signaling by TOR, AMPK and other inputs [12, 15, 16]. The Atg factors are responsible for the execution of autophagy and the formation of specialized double membrane organelles, termed autophagosomes. The Atg factors organize in several subcomplexes, including the Atg5-Atg12/Atg16 complex that acts as a conceptual equivalent of E3 ligases in the ubiquitin system. It regulates C-terminal lipidation of Atg8 (or its mammalian equivalents LC3s and GABARAPs) with phosphatidylethanolamine essential for autophagosomal membrane growth [12]. Atg8-phosphatidylethanolamine can effect membrane tethering and fusion [17, 18], albeit autophagic organelles are also subject to conventional membrane fusion via SNAREs, the canonical regulators of membrane fusion in most compartments of the eukaryotic cell [19]. The membrane precursors for the formation of autophagosomes originate from transient domains on the ER, termed omegasomes. There are also potential contributions from other compartments such as the plasma membrane, mitochondria and Golgi. To carry out degradative functions, autophagosomes fuse with lysosomes to form autolysosomes where the captured cargo is eventually degraded.

As stated above, autophagy is assumed to represent primarily a catabolic, lysosomal degradative pathway. The notion of autophagy as a purely degradative pathway

was recently challenged by the emergence of three independent reports on the role of autophagy in unconventional secretion of Acb1 in yeast [1–3] and IL-1 β and other cargo (e.g. HMGB1 and IL-18) in mammalian cells [4, 5]. These new insights assign a nondegradative function to autophagy manifested as unconventional protein secretion (fig. 1). Furthermore, it has become apparent that autophagy broadly intersects with protein trafficking and secretion to include effects on the constitutive biosynthetic pathway [20], regulated exocytosis [21] and alternative sorting of integral membrane proteins to the plasma membrane [22].

Secretory Autophagy and Unconventional Secretion

The information regarding how autophagy promotes unconventional secretion comes, at present, from a handful of studies [1–4, 22]. It is most completely defined in yeast, using the yeast prototypical cargo for unconventional secretion, Acb1 [1–3]. The role of autophagy in unconventional protein secretion and trafficking is a recently recognized function of the autophagic machinery. Not all unconventionally secreted proteins are secreted by the same mechanism. Unconventional secretion is a catch-all term for a spectrum of diverse processes [10]. This collection of often unrelated secretory processes delivers, via unique mechanisms, specific protein cargo localized in the cytosol to the extracellular environment or assists in the trafficking of integral membrane proteins to the plasma membrane without having to pass through the ER-G-PM pathway [10]. Only a subset of unconventional secretion is dependent on autophagy. When autophagy provides specific membrane carriers to execute unconventional secretion, this subset is referred to as autosecretion [5], type III secretion [6] or secretory autophagy, a term preferentially used in this review.

Secretory versus Degradative Autophagy Precursors

Are the precursors for secretory and degradative autophagy related? A membrane structure in yeast, termed compartment for unconventional protein secretion (CUPS) [2] has been identified as the source of organelles or trafficking intermediates for autophagy-based unconventional secretion. CUPS, the term used in yeast, may be related (although this has still to be established) to the mammalian cell omegasome, which acts as a cradle for

generating nascent autophagosomes normally associated with degradative autophagy [23]. CUPS forms in the general vicinity of but does not completely coincide with the ER exit sites marked by Sec13 [2]. The organizers of the ER exit sites (Sec12 and Sec16) and COPII components (Sec23 and Sec24, but not Sec13 and Sec31) are required for the canonical autophagosome formation in yeast [24]. Participation of the regulators of the early secretory pathway in autophagy has also been noted in mammalian cells [25, 26]. Starvation induces both CUPS [2] and omegasomes [23]. These structures are morphologically similar, PI3P-positive, and are decorated by autophagy (Atg) factors. As CUPS is reportedly not induced by rapamycin, it has been proposed to be distinct from omegasomes [2]. However, the original description of the omegasome and its role in autophagy along with its PI3P positivity did not report the use of rapamycin and was based solely on the starvation-dependent induction of autophagy [23]. The relationship of CUPS to the well-known preautophagosomal structure (PAS) in yeast also needs to be clarified since Sec23 of COPII is required for autophagy via PAS [24], but its mutation does not interrupt unconventional secretion of Acb1. Similarly, Sec12 matters for autophagy [24] but appears to be dispensable for CUPS formation in yeast [2]. The subtle differences outlined above may be technical but could indicate some subcompartmentalization of domains within precursor for autophagosomal organelles (CUPS, PAS and omegasomes) destined for degradative or secretory autophagy.

Secretory versus Degradative Autophagy: Cargo Selection

How is the cargo destined for secretory autophagy selected and separated from degradative autophagy cargo? Thus far, this has not been explored. Of potential relevance is that CUPS contains Vps23 [2]. Vps23, in mammalian systems known as TSG101, is a member of the endosomal sorting complex required for transport (ESCRT)-I complex associated with multivesicular endosomal sorting but also participating in a number of other trafficking processes. ESCRT components other than Vps23 are however not found in CUPS [2] and Vps23 itself is not necessary for CUPS formation, indicating that Vps23 may be a component of sorting elements associated with CUPS. Of further interest is that VPS23/TSG101, in its sorting function, can bind to ubiquitinated cargo, a typical recognition tag for autophagic adaptors

[27]. A role in secretory autophagy for conventional autophagy adaptors, e.g. p62, NBR1, NDP52 and optineurin, remains to be addressed.

GRASP in Unconventional Secretion and Secretory and Canonical Autophagy

The protein named GRASP (fig. 2), with orthologs in yeast [2, 3, 8], metazoans [28] and mammalian cells [29–31], is the only specific marker thus far known to be essential for unconventional protein secretion [10]. However, how it functions in autophagy-based unconventional secretion is not understood. A recently published study [4, 5] indicates that GRASP55 (one of the two mammalian GRASPs) is also necessary for canonical, degradative autophagy [4]. Thus, the early secretory and degradative autophagosomes may originate from the common ancestral membrane domains, with subsequent divergence and specialization for the secretion or digestion of captured substrates.

The yeast GRASP (Grh1) redistributes to CUPS. It is not known what signaling events regulate redistribution of the yeast GRASP orthologs to CUPS upon starvation. In mammalian cells, GRASP55 and GRASP65 localize to the Golgi ribbon, where they interact with Golgin-45 and GM130 (fig. 3). GRASP translocation in mammalian cells has been detected [4] and it could mirror the events in yeast, although this has still to be established. There are two types of factors that could be expected to control GRASP55 localization during degradative and secretory autophagy: (1) kinase(s) that may affect the GRASP55 homo-oligomerization state and thus the tethering of adjacent membranes (as in the Golgi ribbon), and (2) compartment-specific interacting partners binding to the PDZ domains of GRASPs, thus tethering them to the appropriate membranes that eventually coalesce. GRASPs have the ability to link membranes through their homotypic interactions between PDZ domains and internal PDZ-binding motifs ('internal ligand'; see fig. 2). This is morphologically best recognized in the formation, maintenance and disruption of Golgi ribbons, where GRASPs are known to play a role in the linking of Golgi cisternae [30, 31] (fig. 3). The ability of GRASPs to undergo homo-oligomerization depends on the phosphorylation state of their Ser-Pro-rich C-terminal domains and can be disrupted, e.g. during mitosis (causing Golgi dispersal) following complex phosphorylation patterns by several kinases including ERK, CDK1 and PLK1 [30, 31]. In addition to homo-oligomerization via the internal ligand,

GRASP55 and GRASP65 interact in the Golgi with Golgin-45 and GM130, respectively. Although phosphorylation of GRASPs can disrupt their homo-oligomerization (resulting in events such as Golgi disruption in mitosis), it is not known whether similar events could lead to GRASP translocation and modulation of interactions with other putative partners involved in secretory autophagy and unconventional secretion in general.

GRASP carries out its function in conventional autophagy as well, at least in mammalian cells [4]. The yeast GRASP equivalent is important for Atg9 relocalization upon starvation [2], mirroring Vps23 localization to CUPS. The effect of the yeast GRASP ortholog Grh1 on the sole integral membrane autophagy factor (Atg9) indirectly corroborates with the reported role of the mammalian GRASPs (GRASP55) in control of autophagy initiation [4].

Regulation of Secretory versus Degradative Autophagy

In mammalian cells, there is an involvement in autophagy of two closely related but distinct isoforms of the small GTPase Rab8: Rab8a and Rab8b [4, 32]. There are indications of specialization between Rab8a and Rab8b in secretory versus degradative autophagy (fig. 3). It has been shown that Rab8a, and potentially exocyst components, play a role in secretory autophagy [4]. In contrast, Rab8b affects the maturation stages of degradative autophagy [32].

Rab8b interacts with the Ser/Thr protein kinase TBK-1. TBK-1, in turn, phosphorylates two autophagy adaptors: optineurin [33] and the classic autophagy adaptor p62 (sequestosome 1) [32]. The latter is a key autophagy factor [34] and the founding member of a new subfamily of pattern recognition receptors termed sequestosome-like receptors [27]. This function of TBK-1 is likely superimposed on the tonic level of basal p62 phosphorylation carried out by casein kinase 2 at the same Ser-403 residue [35]. Of note, TBK-1 is a member of the IKK family of kinases, the well-known regulators of innate immunity [36]. IKKs can be either canonical (IKK- α and IKK- β) or atypical/IKK-related kinases (IKK ϵ and TBK-1). IKK α and IKK β play a role in the induction of autophagy [16]. A role for TBK-1 in autophagy maturation indicates a sequential action in autophagy of the canonical IKKs at the initiating stages and TBK-1 at the completion end of the process. Whether TBK-1 regulates autophagy at the point of divergence between secretory and degradative autophagy (by, for example, promoting the latter) remains to be determined.

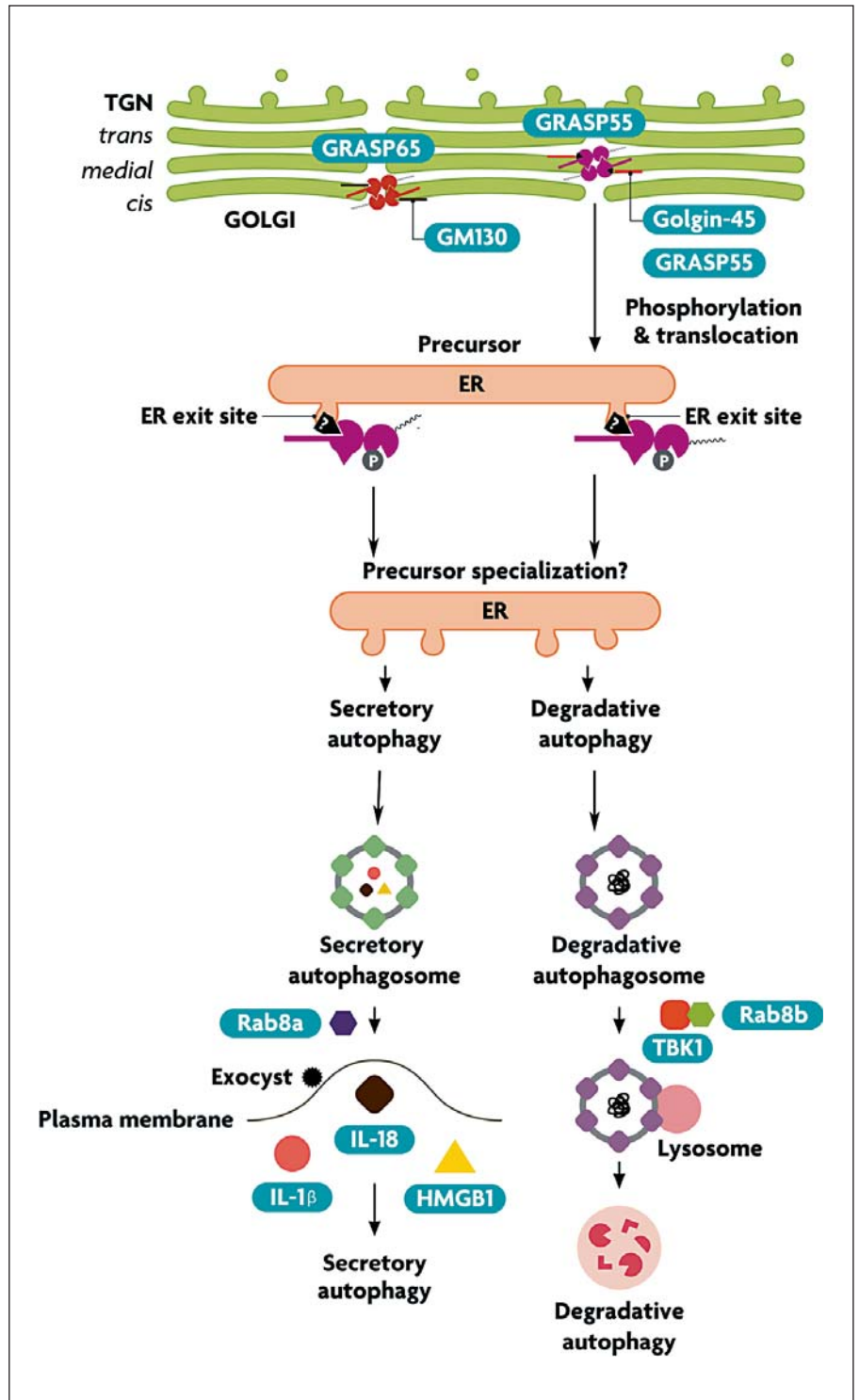


Fig. 3. Hypothetical model and the proposed putative points of divergence between degradative and secretory autophagy. The steps are discussed in the text.

Autophagy Role in IL-1 β Secretion and Inflammasome Activation

In mammalian cells, several cytosolic proteins are secreted through different pathways of unconventional secretion [6, 10]. The most prominent candidate to be unconventionally secreted via autophagy is the proinflammatory cytokine IL-1 β , as proposed [37] immediately following the initial reports on the molecular machinery of secretory autophagy in yeast [1, 3]. However, a number of converging reports focussing on IL-1 β from the immunological perspective have shown, through the observations in mice lacking Atg16L1 [38] followed up by mechanistic *ex vivo* and *in vivo* analyses [4, 39–42], that autophagy dampens IL-1 β activation by the inflammasome. This negative effect of autophagy on IL-1 β activation is a dominant effect of autophagy under steady-state conditions. Nevertheless, a contribution of secretory autophagy to the release of IL-1 β from the cells has been detected as a transient autophagy-dependent secretion of IL-1 β during the early, acute phase of inflammasome activation [4]. The latter observation supports a role (albeit only transiently detectable) of secretory autophagy in the transport of cytosolic IL-1 β from the cells via autophagy-dependent processes.

IL-1 β is synthesized as a leaderless cytosolic protein secreted from cells via membranous organelles [43, 44]. Secretory autophagy may assist in these unconventional secretion processes [4]. Once secreted, IL-1 β binds to the IL-1 receptor (IL-1RI) to initiate a number of inflammatory processes that contribute to innate immunity but can also cause tissue damage [45]. Pro-IL-1 β is processed from its inactive cytosolic precursor by inflammasome, a multiprotein complex responding to microbial products referred to as PAMPs (pathogen-associated molecular patterns) and endogenous molecules signaling damage or danger termed alarmins or DAMPs (damage-associated molecular patterns) [46]. An inflammasome typically consists of three parts: a PAMP- or DAMP-detecting module in the form of a Nod-like receptor (e.g. NLRP3) or an endogenous DNA (released from mitochondria)-detecting module such as AIM2, the hub adaptor protein ASC and caspase-1 that, in turn, enzymatically processes substrates, e.g. pro-IL-1 β into mature, biologically active products such as IL-1 β . Autophagy inhibits IL-1 β activation indirectly, by lowering the endogenous sources, such as reactive oxygen species (ROS) or DAMPs such as mitochondrial DNA released by depolarized or otherwise unkempt mitochondria accumulating in autophagy-deficient cells [39, 40] and recognized by the inflammasome

components. Autophagy may also act via the proposed autophagic degradation of inflammasome constituents [41, 42, 47]. In principle, the primary role of autophagy in the inflammasome context is to keep the cellular interior clean of endogenous DAMPs, lest these inflammasome agonists cause sterile inflammation and pathological tissue damage. Autophagy also plays a membrane-trafficking role contributing to acute unconventional secretion of IL-1 β , but this contribution is masked under steady-state conditions [4]. Secretory autophagy also contributes to the release of other cytosolic inflammasome substrates, e.g. IL-18 and other alarmins such as high-mobility group protein B1 (HMGB1) [4]. Thus, autophagy plays a dual role in inflammasome activation and secretion of IL-1 β and the release of other alarmins from the cell. It is important to recognize this duality as follows: (1) the positive contribution of autophagy to acute delivery of IL-1 α and alarmins from a cell is a manifestation of secretory autophagy as a membrane-trafficking process [4], and (2) the tonic suppression of inflammasome activation is a manifestation of degradative autophagy in its function as a quality control process in the cytoplasm.

Is the Role of Autophagy in IL-1 α Activation and Secretion Different from the Effects on IL-1 β ?

Another proinflammatory pathway, linked but distinct from inflammasome activation, is affected by autophagy. This pathway involves IL-1 α , a proinflammatory cytokine that shares two features with IL-1 β : they are both are cytoplasmic proteins and they both engage the same receptors and downstream signaling following extracellular release and binding to IL-1RI. However, IL-1 α is activated and secreted in a manner that differs from pro-IL-1 β processing and activation. IL-1 α is produced as a cytosolic proform that can be processed by calpain or other proteases [48] and actively exported out of the cell [49, 50] or be passively released upon cell death [51]. IL-1 α secretion has been reported to involve both caspase 1-independent and caspase 1-dependent pathways [52]. Caspase 1^{-/-} macrophages stimulated with LPS show a 20–30% reduction in IL-1 α secretion compared to controls [53], thus indicating that a major route of IL-1 α secretion is independent of caspase 1. Interestingly, despite this divergence, autophagy-defective cells (e.g. Atg5^{fl/fl} LysM-Cre⁺ macrophages) also generate excess IL-1 α . This is consistent with the existence of another major, inflammasome-independent activation pathway [50] that is also augmented in the absence of the cytoplasmic housekeeping func-

tions of autophagy [4, 39–42]. This pathway shares with the inflammasome-IL-1 β the very upstream triggers – e.g. ROS emanating from unempt mitochondria in autophagy-deficient cells. However, it differs in the lower, execution part of the pathway. ROS can activate calpain [54, 55], which in turn activates IL-1 α [50].

The findings with the ROS-calpain axis in IL-1 α activation [50] and reports regarding the ROS-RLR signaling [56] expand proinflammatory phenomena to noninflammatory pathways downstream of the accumulation of dysfunctional mitochondria and ROS in autophagy-deficient cells. Other changes with inflammatory consequences have been noted in mice with Atg5-deficient macrophages [57, 58]. It is not known whether IL-1 α is a substrate for the alternative secretory pathway, as may be the case for IL-1 β [4]. Thus far, only the negative role of autophagy in activation processes upstream of the IL-1 α secretion has been studied [50]. Kinetic studies of IL-1 α release may help discern a possible positive contribution of autophagy to the physical act of secretion of IL-1 α versus the dominant role of autophagy in keeping the upstream activation triggers down.

Concluding Remarks

Thus far, genetic studies support the existence of secretory autophagy. It remains to be defined whether and how secretory autophagy provides specific membrane carriers to capture cytosolic proteins and secrete them outside of the cell. The precise cell biological mechanisms executing secretory autophagy, the adaptors for cargo selection, the switches that separate degradative and secretory autophagy as well as other aspects of vectorial mem-

brane transports and protein cargo sorting have still to be delineated. The intersections of this pathway with the role of GRASP need to be understood. The spectrum of cytosolic proteins externalized by secretory autophagy and the scope of their extracellular functions also remain to be catalogued, in order to appreciate in full the biological and medical significance of this potential pathway.

We caution the reader that the models depicted in the figures of this review are hypothetical and should serve not as a firm summary of what is known but as a speculative starting point and a platform to test and – if need be – disprove these models, in the hope that this will stimulate further research. Regardless of the details, secretory autophagy is an area of investigation that is already expanding the sphere of influence of autophagy from a purely intracytoplasmic domain to the extracellular space. Furthermore, the existence of secretory autophagy argues that autophagy is not only a lysosomal degradative process but may play important biogenesis roles in the extracellular delivery of bioactive molecules. Even if this expanded concept is accepted, a common denominator for both processes will still be the removal of macromolecules and organelles from the cytoplasm, which can occur by degradation or secretion.

Acknowledgments

V.D. acknowledges recent and current support by AI042999, AI069345 and ARRA RC1AI086845 from the National Institutes of Health Colitis Foundation of America CCFA2053 and the Bill and Melinda Gates Grand Challenge Explorations grant. E.F.C. was supported by a supplement 3R01AI042999-13S1. This work was supported in part by the NCRN and the National Center for Advancing Translational Sciences NIH Grant UL1 TR000041.

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