Caspase-8 is activated by cathepsin D initiating neutrophil apoptosis during the resolution of inflammation

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In the resolution of inflammatory responses, neutrophils rapidly undergo apoptosis. We describe a new proapoptotic pathway in which cathepsin D directly activates caspase-8. Cathepsin D is released from azurophilic granules in neutrophils in a caspase-independent but reactive oxygen species–dependent manner. Under inflammatory conditions, the translocation of cathepsin D in the cytosol is blocked. Pharmacological or genetic inhibition of cathepsin D resulted in delayed caspase activation and reduced neutrophil apoptosis. Cathepsin D deficiency or lack of its translocation in the cytosol prolongs innate immune responses in experimental bacterial infection and in septic shock. Thus, we identified a new function of azurophilic granules that is in addition to their role in bacterial defense mechanisms: to regulate the life span of neutrophils and, therefore, the duration of innate immune responses through the release of cathepsin D.
the cysteine protease cathepsin B and the aspartic protease cathepsin D have been reported to be involved in apoptosis regulation (18–20). Genetic evidence for the role of cysteine cathepsins in apoptosis is provided by studies showing resistance against TNF-induced liver apoptosis in mice lacking cathepsin B (19), perhaps because of insufficient cleavage of Bid (21–23). Cathepsin D was shown to activate Bax in T cells (24) and to be involved in the release of cytochrome c from mitochondria in fibroblasts (20, 25). Moreover, pepstatin A (PepA), a pharmacological inhibitor of cathepsin D, blocked mitochondrial cytochrome c release and caspase activation in cardiomyocytes and fibroblasts (25, 26). Collectively, these data suggested a role for lysosomes and cathepsins in proapoptotic pathways proximal to mitochondrial activation in at least some forms of apoptotic cell death.

Because neutrophils rapidly undergo apoptosis after phagocytosis of bacteria (7, 27), we hypothesized that azurophilic granules, in which cathepsins are located and intracellular bacterial killing occurs, might be able to somehow trigger the normal apoptotic program in these cells. To resolve the issue of whether cathepsins are involved in neutrophil apoptosis pathways, we specifically inactivated cathepsin B and D, respectively, by both genetic and pharmacological means. Our studies revealed that cathepsin D is released from azurophilic granules during the initial phase of neutrophil apoptosis, leading to death receptor–independent activation of caspase-8. Importantly, this newly identified alternative proapoptotic pathway of caspase-8 activation observed in neutrophils is blocked under inflammatory conditions and is crucial for the resolution of innate immune responses.

RESULTS
Cathepsin D, but not cathepsin B, deficiency delays neutrophil apoptosis
Neutrophils are known to express cathepsin G in azurophilic granules (28). In initial experiments, we addressed the question of whether the apoptosis–relevant cathepsins B and D are expressed in normal blood neutrophils and obtained evidence that this is the case as assessed by immunoblotting (unpublished data). To determine the possible involvement of these cathepsins in the regulation of neutrophil apoptosis, we purified neutrophils from normal as well as cathepsin B– and cathepsin D–deficient mice. Neutrophils were isolated from bone marrow and blood, and then cultured. Mature neutrophil populations from cathepsin D–/– mice demonstrated more viable cells compared with those isolated from wild-type or cathepsin B–/– mice (Fig. 1 A). In this model, we additionally observed that blood neutrophils die more rapidly than bone marrow neutrophils, perhaps because the mean age of these cell populations is lower compared with blood neutrophils. The delayed death of cathepsin D–/– cells was more pronounced in blood compared with bone marrow neutrophils but was nevertheless also significant in the latter.

Similar to mouse neutrophils, human neutrophils also underwent spontaneous cell death in vitro. Cell death, however, occurred somewhat slower in the human compared with the mouse system (~40% vs. 88% mean cell death in 24-h cultures; Fig. 1, A and B). To investigate the effect of pharmacological inhibition of cathepsin B and D, we used the previously characterized specific cathepsin inhibitors CA-074-ME (CA), a cell-permeable inhibitor of cathepsin B and possibly other cysteine cathepsins (29), and PepA, which, as mentioned in the Introduction, blocks the aspartic protease cathepsin D (20).

In initial experiments, we confirmed the specificity of these two inhibitors (Fig. S1, available at http://www.jem.org/cgi/content/full/jem.20072152/DC1). In contrast to CA, PepA delayed spontaneous death of human blood neutrophils in a concentration–dependent manner (Fig. 1 B). The optimal concentration of PepA for death inhibition was 100 μM.

We next investigated whether the antideath effect mediated by PepA was caused by inhibition of apoptosis. PepA reduced phosphatidylserine redistribution, a characteristic feature of apoptotic neutrophils (12, 13), with the same efficacy as GM-CSF (Fig. 1 C, top). Agnostic anti–Fas antibody stimulation was used as an additional control and accelerated neutrophil apoptosis in this in vitro system, which was in agreement with previously published work (12, 13). We also analyzed DNA fragmentation, another hallmark of apoptotic cells. Again, PepA and GM-CSF significantly blocked apoptosis, whereas anti–Fas stimulation resulted in increased DNA fragmentation (Fig. 1 C, bottom). Although human and mouse neutrophils appeared to undergo cell death with different kinetics, these data collectively suggested that cathepsin D exerts proapoptotic activities in cultured mature neutrophils.

Cathepsin D translocation precedes cytochrome c release
To identify where cathepsin D is located within neutrophils, we performed double immunofluorescence analysis on mature neutrophils using confocal microscopy. Cathepsin D colocalized with myeloperoxidase (MPO) in freshly isolated neutrophils (Fig. S2 A, available at http://www.jem.org/cgi/content/full/jem.20072152/DC1). MPO is known to be stored in azurophilic granules of neutrophils (30). In contrast, cathepsin D did not colocalize with cytosolic caspase-3 (Fig. S2 B). These data suggested that cathepsin D is stored, like MPO, in azurophilic granules. Using an anti–cathepsin B antibody, we observed the same results (unpublished data). Upon culturing for 3 h, a subpopulation of neutrophils showed a diffuse staining pattern for cathepsin D and MPO (Fig. 2 A, left; see Fig. S2 A for higher magnification; the same data were observed regarding cathepsin B [not depicted]), suggesting the nonspecific permeabilization of azurophilic granules and subsequent cytosolic release of at least some of its components. Both GM-CSF and G-CSF preserved the punctate pattern in the majority of the cells, whereas the pancaspase inhibitor z-VAD and PepA had no effects. In agreement with these immunofluorescence data, GM-CSF and G-CSF, but not z-VAD, suppressed the enzymatic activity of cathepsin D in cultured neutrophils (unpublished data).

Mitochondria have been shown to be essential for optimal apoptosis induction in neutrophils (6, 8, 31). Therefore, we analyzed the kinetics of mitochondrial release of
Figure 1. Lack of expression and pharmacological inhibition of cathepsin D, but not cathepsin B, delays neutrophil apoptosis. (A) Viability assay. Mouse neutrophils from cathepsin D−/− (n = 7), cathepsin B−/− (n = 6), and wild-type (n = 6) mice were isolated from bone marrow (left) or blood (right) and cultured in the absence of cytokine support for 24 h before viability was measured. Values are means ± SD. *, P < 0.05; **, P < 0.01. (B) Viability assay. Concentration- and time-dependent effects of CA (left) and PepA (right) on human blood neutrophil death (n = 3). Values are means ± SD. *, P < 0.05; **, P < 0.01. (C) PS redistribution (top) and DNA fragmentation (bottom) assays. Human blood neutrophils were analyzed after 8-h (PS analysis) and 22-h (DNA analysis) cultures. (left) Values are means ± SD (n = 3). *, P < 0.05. (right) The quantitative analysis (percentages) of early and late apoptotic cells is shown. Representative results from three independent experiments.
Figure 2. Cathepsin D is released into the cytosol and proceeds mitochondrial cytochrome c liberation during neutrophil apoptosis. (A) Confocal microscopy, cathepsin D colocalization, and staining pattern analysis. Freshly isolated and cultured neutrophils were stained with anti-cathepsin D and anti-MPO antibodies. (left) Representative data. Correlation coefficient and statistical analysis are shown in Fig. S2 A. No detectable staining was observed using control antibodies (not depicted). Bar, 10 μm. (right) Quantitative and statistical analysis of time-dependent cathepsin D release experiments. Values are means ± SD (n = 3). The same data were observed if MPO and cathepsin B releases were investigated (not depicted). (B) Confocal

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two proapoptotic factors in cultured neutrophils in the presence and absence of PepA by immunofluorescence analysis (Fig. 2 B, left). As expected, cytochrome c and Smac demonstrated a punctate staining pattern and appeared to colocalize in freshly isolated neutrophils (Fig. S2 C). The mitochondrial location of these two proapoptotic factors was confirmed by the demonstration of colocalization with the mitochondrial marker protein cytochrome oxidase subunit I (unpublished data). After culturing for 12 h, we obtained a diffuse staining pattern in a significant proportion of neutrophils. In the presence of PepA and z-VAD, however, the numbers of cells turning into a diffuse pattern were significantly decreased. The same observation was made using GM-CSF and G-CSF.

We systematically counted the cells exhibiting a diffuse staining pattern and examined cathepsin D and cytochrome c release in a time-dependent manner (Fig. 2, A and B, right). Our data clearly indicated that the release of cathepsin D release occurs earlier than cytochrome c release. Moreover, because PepA and z-VAD blocked cytochrome c release, it can be concluded that both cathepsin D and caspase activities are required for the proapoptotic activation of mitochondria in neutrophils. Collectively, these data suggested that caspase-independent permeabilization of azurophilic granules occurs in early phases of neutrophil apoptosis. This process preceded the caspase-dependent liberation of mitochondrial proapoptotic factors and is blocked by survival cytokines.

To confirm that azurophilic granules undergo permeabilization and that the release of cathepsin D precedes cytochrome c release from mitochondria, we analyzed the kinetics of translocation of these two proteins using cytosolic extracts from cultured neutrophils and immunoblotting (Fig. S3, available at http://www.jem.org/cgi/content/full/jem.20072152/DC1). Although cathepsin D was already detected in cytosols from 3-h cultured neutrophils, we obtained evidence for cytochrome c at later time points only (6 and 9 h). These data further supported the concept that membranes of azurophilic granules are permeabilized before those of mitochondria during neutrophil apoptosis. Moreover, transmission electron microscopy of cultured neutrophils demonstrated damaged azurophilic granule membranes, suggesting the permeabilization of these organelles in this process (Fig. S4).

To demonstrate cathepsin D and B expression and their potential release in neutrophils under in vivo conditions, we analyzed neutrophils in tissue sections of patients suffering from inflammatory diseases. Leukocytoclastic vasculitis is a skin disorder known to be associated with apoptotic neutrophils present in skin lesions (32). We confirmed multiple apoptotic neutrophils in the dermis of these patients, as determined by regular histology and activated caspase-3 staining. Immunofluorescence analysis on these tissues demonstrated evidence for cathepsin D and MPO releases in these cells (diffuse staining pattern; Fig. 2 C). In contrast, we detected almost no apoptotic neutrophils in tissue sections from patients suffering from ulcerative colitis and acute appendicitis, in which neutrophils exhibited a punctate staining pattern, suggesting that azurophilic granules were intact. We also performed cathepsin B stainings and obtained the same results (unpublished data). In conclusion, permeabilization of azurophilic granules and the subsequent release of cathepsins and MPO in apoptotic neutrophils is not only an in vitro phenomenon but also occurs in vivo.

Caspase activation triggered by cathepsin D

Because the proapoptotic activation of mitochondria was both caspase D and caspase dependent, but the release of cathepsin D (and its enzymatic activation) was caspase independent, we addressed the question of whether active cathepsin D is able to induce caspase activation. In agreement with previous work (8, 12, 16), we observed caspase-8, caspase-3, and Bax cleavage in a time-dependent manner in association with spontaneous neutrophil apoptosis, as assessed by immunoblotting (Fig. 3 A). Caspase-8 cleavage proceeded caspase-3 cleavage both in spontaneous and Fas receptor-mediated apoptosis. Bax is cleaved by calpain-1 during neutrophil apoptosis (8), an event that is detectable before caspase-3 cleavage. PepA but not CA blocked cleavages of caspase-8, caspase-3, and Bax, confirming that pharmacological inhibition of cathepsin D but not cathepsin B delays neutrophil apoptosis.

Although we obtained evidence for the generation of active caspases in these experiments (a 15-kD fragment of caspase-8 and a 17-kD fragment of caspase-3), previous work suggested that cleavage does not necessarily indicate caspase activation (33). Therefore, we also analyzed the effect of pharmacological inhibition of cathepsin D on enzymatic caspase activities in cultured neutrophils. Spontaneous neutrophil apoptosis was associated with increases of both caspase-8 and -3 activities, which were completely blocked by PepA (Fig. 3 B). PepA did not directly block the activity of pure caspase-8 or -3 enzymes (Fig. S1 B), suggesting that cathepsin D is indeed able to trigger caspase activation in neutrophils.

Cathespin D directly activates caspase-8

Bid has been described as a target of cathepsin B in vitro (21–23). Therefore, we hypothesized that cathepsin D is also
able to cleave cellular substrates involved in apoptosis regulation. In an in vitro cell-free assay using cytosolic extracts of freshly isolated blood neutrophils, we observed that the addition of cathepsin D, but not cathepsin B, was followed by the cleavage of caspase-8 (15-kD fragment) and caspase-3 (17-kD fragment) in a concentration- and time-dependent manner (Fig. 4 A and B). As seen during spontaneous neutrophil apoptosis, caspase-8 cleavage was again detectable before caspase-3 cleavage in this cell-free system. In the presence of PepA, cathepsin D–induced caspase cleavages were completely prevented (Fig. 4 A). In addition, the caspase-8 fragment generated by incubation of cytosolic extracts with cathepsin D could be affinity labeled with the biotinylated caspase substrate VAD-fmk (bVAD-fmk; Fig. 4 B, right), suggesting that the 15-kD fragment of caspase-8 is enzymatically active (34). Additional enzymatic measurements confirmed that cathepsin D, but not cathepsin B, increased the enzymatic activities of both caspase-8 and -3 in the cytosolic extracts of neutrophils (Fig. 4 C).

Cathepsin D processed and activated both caspase-8 and -3 in vitro, but the cleavage of caspase-8 seemed to be earlier than that of caspase-3. Therefore, we hypothesized that caspase-8 is required for caspase-3 activation in neutrophils. Indeed, cathepsin D was unable to mediate caspase-3 cleavage (17-kD fragment) after immunodepletion of caspase-8 in cytosolic extracts of freshly isolated blood neutrophils (Fig. 4 D). We also used cytosolic extracts of the small cell lung carcinoma (SCLC) lines U1285 (caspase-8 containing) and SW2 (caspase-8 deficient) and confirmed that cathepsin D–mediated caspase-3 cleavage (17-kD fragment) is caspase-8 dependent (Fig. 4 E).

Although cathepsin D activated caspase-8, it remained unclear whether this is a direct or an indirect process. To address this question, we performed identical in vitro experiments using pure recombinant human caspase-8 and -3 proteins instead of cytosolic extracts. Cathepsin D cleaved caspase-8 (15-kD fragment) but not caspase-3 (Fig. 5 A). Moreover, the generated caspase-8 fragment was enzymatically active because it was trapped by bVAD-fmk (Fig. 4 B, right), suggesting that the 15-kD fragment of caspase-8 is enzymatically active (34). Additional enzymatic measurements confirmed that cathepsin D, but not cathepsin B, increased the enzymatic activities of both caspase-8 and -3 in the cytosolic extracts of neutrophils (Fig. 4 C).

Cathepsin D deficiency prolongs innate immune responses
To determine whether the newly identified prosapoptotic pathway resulting in caspase-8 activation in the absence of death receptor signaling is relevant in a more physiological context, we analyzed the role of cathepsin D in the regulation of innate immune responses in vivo. To this end, we used cathepsin D–deficient mice and compared their response to a bacterial challenge with that of wild-type controls. We found that cathepsin D deficiency significantly prolonged the innate immune response, indicating that cathepsin D is an important regulator of innate immune function.
Figure 4. Cathepsin D activates caspase-8, which is required for subsequent caspase-3 activation. (A) Immunoblotting. Concentration-dependent cleavage of caspase-8 and -3 proteins by cathepsin D, but not cathepsin B, in cytosolic extracts from freshly isolated human blood neutrophils. Results are representative of three independent experiments. (B) Time-dependent cleavage of caspase-8 and -3 proteins by cathepsin D, but not cathepsin B, in cytosolic extracts from freshly isolated human blood neutrophils. Cytosolic extracts were also used for affinity labeling with bVAD-fmk and analyzed by immunoblotting. Results are representative of three independent experiments. (C) Caspase activity assays in cytosolic extracts of freshly isolated human blood neutrophils after treatment with cathepsins. Values are means ± SD (n = 3). *, P < 0.05. (D) Immunoblotting. Cleavage of caspase-3 (17-kD fragment) mediated by cathepsin D in normal and caspase-8-immunodepleted cytosolic extracts from neutrophils. Results are representative of three independent experiments. (E) Immunoblotting. Cleavage of caspase-3 (17-kD fragment) mediated by cathepsin D in caspase-8-expressing and -deficient cytosolic extracts from SCLC lines. Arrowheads in A, B, D, and E indicate key cleavage products.
setting, we injected wild-type and cathepsin D−/− mice intraperitoneally with low-dose LPS. In this experimental model of bacterial peritonitis, we observed a moderate increase in blood neutrophil numbers peaking at 24 h in wild-type mice (P < 0.05). 32 h after LPS injection, neutrophil numbers started to decline. Compared with wild-type mice, cathepsin D−/− mice demonstrated much higher neutrophil numbers at 24 h that further increased 32 h after LPS administration (Fig. 6 A). We also analyzed the numbers of peritoneal lavage neutrophils 32 h after LPS injection. Although almost no neutrophils were detected in wild-type mice, cathepsin D−/− mice demonstrated evidence for peritoneal neutrophil infiltration (Fig. 6 B). In addition, although practically all peritoneal lavage neutrophils underwent rapid apoptosis, the neutrophils from cathepsin D−/− mice were apoptosis resistant (Fig. 6 C).

Macrophages have been shown to be key cellular players in the uptake of apoptotic neutrophils and in the resolution of inflammation (35, 36). Therefore, the possibility existed that the increased inflammation seen in cathepsin D−/− mice...
percentage of phagocytosed neutrophils was seen between wild-type and cathepsin D−/− mice (Fig. 6 D). We also observed no difference between wild-type and cathepsin D−/− mice when we intraperitoneally injected low-dose LPS in the mice before performing the same macrophage phagocytosis assays (unpublished data). Collectively, these data excluded a defect in the uptake of apoptotic neutrophils and suggested that the stronger and prolonged inflammatory response in cathepsin D−/− compared with wild-type mice is largely caused by a block of neutrophil apoptosis in this model of experimental bacterial infection.

Because experimental LPS administration in mice reproduces a human bacterial infection only in part, we investigated the newly identified cathepsin D–caspase-8 pathway in neutrophils of patients suffering from septic shock. Analysis of cathepsin D release in the cytosol in neutrophil cultures showed that at least 85% of the sepsis neutrophils maintained intact azurophilic granules with no evidence for permeabilization within the first 9 h. In contrast, at this time point, ~90% of normal neutrophils released cathepsin D in the cytosol (Fig. 7 A). Moreover, sepsis neutrophil cultures did not show evidence for caspase-8 cleavage within 9 h (Fig. 7 B), further supporting the concept that caspase-8 activation is mediated by cathepsin D in neutrophils. Consequently, spontaneous neutrophil death (Fig. S5, available at http://www.jem.org/cgi/content/full/jem.20072152/DC1) and apoptosis (Fig. 7 C) was markedly delayed in sepsis under ex vivo conditions. No difference was observed regarding cathepsin D expression between normal and sepsis neutrophils as assessed by immunoblotting (unpublished data).

**Reactive oxygen species (ROS) deficiency delays cathepsin D release and apoptosis**

The absence or inhibition of cathepsin D delays but does not prevent neutrophil apoptosis. This suggests that alternative mechanisms must exist for apoptosis induction. Genetic deficiency of nicotinamide adenine dinucleotide phosphate (NADPH) oxidase or its pharmacological inhibition results in delayed neutrophil apoptosis (12, 37). We therefore investigated the effect of combined inhibition of both ROS production and cathepsin D activity on neutrophil apoptosis. Inhibition of ROS production was performed by diphenyleneiodonium (DPI). Although pharmacological inhibition of both NADPH oxidase and cathepsin D demonstrated additive antidilative effects, the complete prevention of neutrophil death was not achieved even under these conditions (Fig. 8 A).

We next investigated whether ROS were required for the translocation of cathepsin D in the cytosol. To have a clean system, we used neutrophils from a patient with chronic granulomatous disease (CGD) that are unable to generate ROS based on a genetic defect in the NADPH oxidase. In the absence of ROS, we observed delayed cathepsin D release (a culture period of 7 h was required to obtain cathepsin D release in 50% of the neutrophils, whereas only 1 h was required in normal neutrophil populations; Figs. 2 A and 8 B). Interestingly, under ROS-deficient conditions, both GM-CSF
is inhibited by the neutrophil survival cytokines GM-CSF and G-CSF in vitro and in vivo. Because phagocytosis of bacteria triggers apoptosis in neutrophils (7, 27), it is also tempting to speculate that this previously reported death mechanism is initiated by the release of lysosomal enzymes from the azurophilic granules in the cytosol. The exact molecular mechanisms regulating the permeabilization of azurophilic granules under the different conditions remain to be investigated.

DISCUSSION

Lysosomal enzymes such as cathepsins and MPO are stored in the azurophilic granules of neutrophils and synthesized early during maturation of these cells in the bone marrow (38). These enzymes are used during phagocytosis and intracellular killing (30), as well as in the extracellular defense against bacteria (39). In this paper, we provide evidence for an important new function of azurophilic granules, namely, the induction of apoptosis in mature neutrophils. Azurophilic granules release cathepsin D, which directly activates the initiator caspase-8, leading to subsequent calpain (as indicated by Bax cleavage) and caspase-3 activation. This proapoptotic pathway and G-CSF had only modest additional inhibitory effects on cathepsin D release. The delayed cathepsin D release in ROS-deficient neutrophils was associated with no detectable cleavage of caspases (Fig. 8 C) and reduced apoptosis (Fig. 8 D). These data suggest that ROS are involved in the membrane permeabilization of azurophilic granules.

Figure 8. Delayed cathepsin D release and subsequent apoptosis defect in ROS-deficient neutrophils. (A) Viability assay. Neutrophils were isolated from human control individuals and cultured in the presence and absence of 20 μM DPI and 50 μM PepA for 24 h before analysis. Values are means ± SD from three independent experiments. *, P < 0.05. (B) Confocal microscopy. Blood neutrophils from a patient suffering from CGD were cultured and analyzed as shown in Fig. 2 A and Fig. S2 A. (C) Immunoblotting. In contrast to normal neutrophils (Figs. 3 A and 7 B), no active fragments were detectable in CGD neutrophils in up to 9-h culture periods. (D) PS redistribution assay. Blood neutrophils from a control individual and a CGD patient were cultured for 8 h, and analyzed by flow cytometry. The quantitative analysis (percentages) is shown in the subpanels.
caspase is activated in the absence of death receptor ligation in neutrophils. Because cathepsin D is ubiquitously present in lysosomes, it is possible that this pathway can also be used to activate apoptosis in other cells, such as SCLC cells.

To obtain original insights in the permeabilization process of azurophilic granules, we performed experiments in neutrophils derived from a CGD patient. We obtained evidence that ROS are involved, because CGD neutrophils demonstrated delayed cathepsin D release compared with normal neutrophils. These data are in agreement with recent work published by Blomgran et al., who showed that during microbe-induced apoptosis of human neutrophils, ROS-dependent lysosomal destabilization represent an early event (41). This destabilization provoked the release of cathepsin B, which then induced the cleavage of the proapoptotic Bcl-2 protein Bid, mitochondrial damage, and subsequent caspase activation and apoptosis. Our data, however, suggest that cathepsin B is functionally unimportant for apoptosis induction, because its genetic and pharmacological inactivation had no influence on neutrophil death. Clearly, additional work is required to better understand the molecular mechanisms leading to the membrane permeabilization of azurophilic granules during neutrophil apoptosis.

Cathepsin D is inactivated at the pH found in the cytoplasm (42). How then can caspase-8 be activated by cathepsin D in a cell? One possibility is that caspase-8 is cleaved because of the known endopeptidase activity of cathepsin D (i.e., at least partially retained at a neutral pH) (43, 44). On the other hand, because of the nonspecific membrane permeabilization of azurophilic granules, it is likely that acidification of the cytosol may occur, at least in close proximity to the granules, allowing cathepsin D to be completely active. Acidification of the cytosol has previously been noticed in apoptotic cells (43, 46). Although, this study cannot completely explain how cathepsin D maintains its activity in the cytosol, the delayed neutrophil apoptosis in cathepsin D−/− mice represents the most direct demonstration that it is indeed the case.

How is caspase-8 activated by cathepsin D? At present, we do not understand the molecular mechanism of any initiator caspase (47). Two models have been proposed. Initially, it was thought that initiator caspases autoprocess themselves when they are brought into close proximity with each other. Later studies suggested that caspase-8 aggregation serves primarily to facilitate dimerization of the enzyme and that this does not require interchain proteolysis. In contrast, interchain proteolysis, but not enforced dimerization of caspase-8 by Fas-associated protein with death domain, was a prerequisite for its activation by granzyme B or caspase-6 (34, 48). Therefore, the proteolytic processing of caspase-8 by cathepsin D might initiate its activation, but the exact mechanism remains to be investigated.

Most studies of neutrophil accumulation concentrate on adhesion and migration. In contrast, little information is available regarding the functional consequences of delayed neutrophil apoptosis in innate immune responses. Therefore, the demonstration that delayed apoptosis caused by cathepsin D deficiency amplifies and prolongs neutrophil inflammation in vivo is notable. Future studies may provide new insights into how GM-CSF and G-CSF stabilize the membrane of azurophilic granules, which results in “functional cathepsin D deficiency,” at least in terms of caspase-8 activation and subsequent apoptosis induction. As indicated in this study, the stabilization of azurophilic granules might play an important pathogenic role in sepsis, CGD, ulcerative colitis, and appendicitis. Besides ROS, another potential candidate regulating the permeability of azurophilic granules is Bax, which has been shown to regulate lysosomal membrane permeabilization (49). In addition, heat shock protein 70 has been reported to promote cell survival by inhibiting lysosomal membrane permeabilization (50). Independent from these pathogenic mechanisms, permeabilization of azurophilic granules may provide a new therapeutic strategy to induce neutrophil apoptosis and, consequently, to resolve exaggerated innate immune responses.

MATERIALS AND METHODS

Reagents. Both cathepsin B and D recombinant enzymes, the specific colorimetric cathepsin B substrate (N-benzoyloxycarbonyl (z)-Arg-Arg-para-nitroaniline), and DPI were obtained from EMD. The specific fluorogenic cathepsin D substrate [(7-Methoxyecoumarin-4-ylacetyle-L-Leu-Leu-Val-Asp (z-LEVD)-fmk (z-LEVD), z-Ile-Glu-Thr-Asp (IETD)-fmk (z-IETD), and z-Asp-Glu-Val-Asp (DEVD)-fmk (z-DEVD) were obtained from BD Biosciences. PepA, CA, sucrose, and LPS from Escherichia coli 0111:B4 were purchased from Sigma-Aldrich. Anti-Fas agonistic monoclonal antibody (CH11) was obtained from MBL International. bVAD-fmk was obtained from MP Biomedicals.

Mice. Cathepsin B−/−, cathepsin D−/−, and wild-type mice were generated in our laboratory as previously described (51, 52). For all experiments, mice with a C57BL/6j background that were 3–4 wk old were used. Mice were maintained under pathogen-free conditions. All animal experiments were reviewed and approved by the Animal Experimentation Review Board of the State of Bern.

Cells. Peripheral blood neutrophils were purified from healthy normal individuals and patients suffering from septic shock by Ficoll-Hypaque centrifugation (12, 13). Septic shock patients fulfilled the following inclusion criteria: (a) documented or suspected infection; (b) signs of systemic inflammation in response to infection; and (c) systolic arterial blood pressure <70 mm Hg, despite adequate fluid resuscitation, in the absence of other causes of hypotension (53). We also purified neutrophils from the blood of a patient with CGD. We obtained Institutional Review Board approval for the study from the Kantonale Ethikkommission Bern. The purity of the isolated human neutrophil populations was always >95%, as assessed by staining with Diff-Quik (Baxter) and light microscopy analysis. Neutrophils were also isolated from wild-type, cathepsin B−/−, and cathepsin D−/− mice. Mature mouse bone marrow (obtained from femur and tibia of the hind legs) and blood neutrophils (obtained by cardiac puncture) were positively selected using anti–Gr-1 monoclonal antibody (Miltenyi Biotec), as previously described (54).

The purity of the resulting mouse neutrophil populations was >90% (bone marrow) and >95% (blood), respectively. For macrophage phagocytosis assays, we injected 2–3 ml of ice-cold 30% sucrose solution in the peritoneal cavity of wild-type and cathepsin D−/− mice, gently palpated for ∼30–60 s, and aspirated the solution, which contained sufficient numbers of peritoneal macrophages. The SCLC cell lines U1285 (expressing caspase-8) and SW2 (lacking caspase-8) (55) were provided by Dr. U. Zangemeister-Wittke (University of Bern, Bern, Switzerland).
**Cell cultures.** Human blood and mouse bone marrow neutrophils were cultured at 10⁶ cells per milliliter. Mouse blood neutrophils were used at 25 × 10⁶ cells per milliliter. Neutrophils were cultured in complete culture medium (RPMI 1640 containing 10% fetal calf serum) in the presence and absence of 50 ng/ml GM-CSF, 25 ng/ml G-CSF, 1 µg/ml anti-Flt-3, 0.5–20 µM CA, 1–300 µM PepA, 20 µM DPI, and 50 µM z-VAD for the times indicated in the figures.

**Determination of cell death and apoptosis.** Neutrophil death was asayed by uptake of 1 µM ethidium bromide and flow cytometric analysis (FACS Calibur; Becton Dickinson) (12, 13, 56). To determine whether cell death was apoptosis, redistribution of phosphatidylserine (PS) in the presence of propidium iodide (PI) was measured by flow cytometry (12, 13, 56). Neutrophil apoptosis was also assessed by oligonucleosomal DNA fragmentation (12, 13, 56).

**Gel electrophoresis and immunoblotting.** 10⁶ cells per milliliter were washed with PBS supplemented with protease inhibitor cocktail (Sigma-Aldrich) and lysed with RIPA buffer (50 mM Tris-HCl [pH 7.4], 150 mM NaCl, 0.25% sodium deoxycholate, 1% Nonidet P-40, 1 mM EDTA supplemented with protease inhibitor cocktail). After a 10-min centrifugation to remove insoluble particles, equal amounts of the cell lysates, cell-free extracts, or subcellular fractionation extracts were loaded on gels (NuPage; Invitrogen). Separated proteins were electrotransferred onto polyvinylidene difluoride (PVDF) membranes (Immobilon-P; Millipore). The filters were incubated overnight at 4°C in TBS/0.1% Tween 20/5% nonfat dry milk with mouse anti-caspase-8 (1:1,000; Cell Signaling Technology), rabbit anti-Bax (1:1,000; BD Biosciences), rabbit anti-caspase-3 (1:1,000; Cell Signaling Technology); mouse anti-cathepsin D (1:1,000; Sigma-Aldrich), or mouse anti-cytochrome C (1:500; BD Biosciences) antibodies. For loading controls, stripped filters were incubated with anti-GAPDH (1:3,000; Chemicon) monoclonal antibody. Filters were washed in TBS/0.1% Tween 20/5% nonfat dry milk for 30 min at room temperature and incubated with the appropriate horseradish peroxidase–conjugated secondary antibody (GE Healthcare) in TBS/0.1% Tween 20/5% nonfat dry milk for 1 h. Filters were developed by an enhanced chemiluminescence technique (ECL kit; GE Healthcare) according to the manufacturer's instructions.

**Cell-free assays.** 20 × 10⁶ freshly isolated human blood neutrophils were washed with PBS containing 4% BSA and lysed for 35 min on ice in 60 µl of CEB buffer (20 mM Hepes [pH 7.2], 250 mM sucrose, 10 mM KCl, 1.5 mM MgCl₂, 2 mM EDTA, 1 mM dithiothreitol (DTT), 100 µM PMSE, 10 µg/ml pepstatin A). The lysate was incubated with a mechanical homogenizer by 25 strokes. After a 1-h centrifugation step at 21,000 g at 4°C, the supernatant (cytosol) was used for the cell-free assay. 10 µg of cytosolic fractions was incubated for the times indicated in the figures at 37°C in buffer C (340 mM NaOAc, 60 mM acetic acid, 4 mM EDTA, 0.1% CHAPS, 8 mM DTT [pH 5.5]) or buffer D (500 mM glycine–HCl [pH 3]) in the presence or absence of the indicated amounts of cathepsin B and D, as well as 100 µM PepA. In some experiments, caspase-8 was removed from neutrophil cytosolic extracts by immunodepletion using 5 µg of anti-human caspase-8 antibody (Cell Signaling Technology). In other experiments, we prepared cytosolic extracts from SCLC cell lines U1285 and SW2, which were incubated for 30 min at 37°C in the presence and absence of 0.6 U cathepsin B, 0.3 U cathepsin D, and 100 µM PepA for 30 min at 37°C. Activated caspase-8 was also detected by precipitation with bVAD-fmk (34).

**In vitro protease cleavage assays.** To investigate direct cleavage of caspases by cathepsin D, 2 µl of pure caspase-8 (generated from histidine-tagged human caspase-8 cloned in PET15b [57]; provided by K. Schulze-Osthoff, University of Düsseldorf, Düsseldorf, Germany) and caspase-3 (provided by C. Borner, University of Freiburg, Freiburg, Germany) recombinant proteins were incubated with 0.1 and 0.3 U cathepsin D in the presence or absence of 100 µM PepA for the times indicated in the figures at 37°C in buffer D. Recombinant activated human caspase-8 was analyzed by mass spectrometry immediately after purification. Samples for Edman sequencing were prepared by blotting the separated cleavage products onto PVDF membranes and cutting out the bands corresponding to the fragments of interest. Edman sequencing was performed at the Functional Genomics Center operated by the Swiss Federal Institute of Technology Zurich (ETH Zürich) and the University of Zurich, and mass spectrometry analyses were performed by the SVS MS-Plateform of the University of Geneva on a fee-for-service basis.

**Histological examination.** Tissue sections from leukocytoclastic vasculitis, ulcerative colitis, and acute appendicitis patients were fixed in 4% paraformaldehyde and embedded in paraffin. 5-µm sections were stained with hematoxylin and eosin and examined by light microscopy (Axiovert 35; Carl Zeiss, Inc.).

**Confocal laser scanning microscopy.** Cytopsins with 2 × 10⁶ cells per milliliter were made from freshly purified neutrophils or neutrophils cultured in the presence or absence of 100 µM PepA, 50 µM z-VAD, 25 ng/ml G-CSF, or 50 ng/ml GM-CSF on noncoated slides. Cells were fixed in 4% paraformaldehyde for 10 min at room temperature and washed three times in PBS (pH 7.4). Permeabilization of cells was performed with 0.05% saponin in buffer A (3% BSA in PBS) for 5 min at room temperature and with acetone for 15 min at −20°C. To prevent nonspecific binding, slides were incubated in blocking buffer (33% human immunoglobulin, 33% normal goat serum, 33% BSA) for 1 h at room temperature. Indirect immunostaining of caspase D, MPO, caspase-3, cytochrome C, CoxI, and Smac were performed by using the following primary antibodies diluted in blocking buffer: monoclonal anti–caspase D (1:100; Sigma-Aldrich), polyclonal anti–MPO (1:6,000; Dako), polyclonal anti–caspase-3 (1:100; Cell Signaling Technology), monoclonal anti–CoxI (1:400; Invitrogen), monoclonal anti-cytochrome C (1:100; BD Biosciences), and polyclonal anti-Smac (1:100; Inogenex). Mouse and rabbit control antibodies, respectively, were used at the same concentrations in each experiment.

Immunofluorescent stainings were also performed on 5-µm-thick paraformaldehyde–fixed paraffin-embedded tissue sections from leukocytoclastic vasculitis, ulcerative colitis, and acute appendicitis patients. Slides were dried for 2 h at 52°C and deparaffinized. Slides were blocked and stained as described in the previous paragraph. After incubation with primary antibodies, cells and tissues were incubated with the appropriate TRITC- and FITC–conjugated secondary antibodies (1:500) for 1 h in the dark at room temperature. For active caspase-3 staining, polyclonal anti–caspase-3 antibody (1:200; Cell Signaling Technology) was used. The–fading agent Mowiol (EMD) was added. Slides were covered with coverslips and analyzed by a confocal laser scanning microscope (LSM 510: Carl Zeiss, Inc.) equipped with Ar and HeNe lasers. To quantify the amount of cells with diffuse staining pattern (caspase D, MPO, cytochrome C, and Smac experiments), 100 cells were counted in randomly chosen regions, and the mean number of cells demonstrating diffuse staining was calculated.

For colocalization studies, unprocessed, unfiltered and undeconvoluted datasets were analyzed using Imaris software package (Bitplane AG), considering every singular layer of a stack separately. Quantitative data of colocalization events were determined by the statistics modules in the colocalization and Voxelshop software of the Imaris package. Intensities were given as the sum of all colocalizing voxels in a dataset, and a computer image was generated.
Experimental peritonitis. Cathepsin D\textsuperscript{−/−} and wild-type mice were infected intraperitoneally with 30 μg LPS. Blood was collected at the time points indicated in the figures, and neutrophil numbers were analyzed by differential counts using Turk solution (distributed by Dr. Gregg Chennae AG). After 32 h, mice were killed by CO\textsubscript{2} inhalation and infected intraperitoneally with 2 ml PBS. The peritoneal lavage fluid was collected and we performed differential counts according to standard morphological criteria on cytospin preparations stained with Giemsa-May-Grünwald solution (Sigma-Aldrich).

In addition, neutrophils were isolated, cultured for 24 h, and subsequently analyzed regarding PS redistribution.

Macrophage phagocytosis assay. Uptake of apoptotic mouse neutrophils by mouse macrophages was investigated as described previously (16, 35), with slight modifications. In brief, \(\sim 2 \times 10^9\) peritoneal macrophages of wild-type and cathepsin D\textsuperscript{−/−} mice were cultured in complete culture medium on a glass coverslip in 24-well tissue culture plates (VWR International AG) in the presence of 50 ng/ml of mouse GM-CSF. 10\textsuperscript{6} mature mouse bone marrow neutrophils of wild-type mice were cultured for the indicated times and added to macrophages at 37 °C for 30 min. After coincubation, cells were fixed with 1% acetone-formalin and stained for MPO activity with di-methoxybenzidine in the presence of hydrogen peroxide. Cells were lightly counterstained with Harris' hematoxylin. The proportion of neutrophils phagocytosed by macrophages was determined by two independent investigators by counting in at least five fields (a minimum of 100 neutrophils was evaluated).

Statistical analysis. Statistical analysis was performed using the Mann-Whitney U test. The figures show mean levels ± SD. For multiple comparisons, differences of the mean levels were analyzed using analysis of variance, followed by Tukey's HSD test. P < 0.05 was considered statistically significant. For colocalization studies, Pearson's correlation coefficient was determined in at least 10 different cells, and means ± SD were calculated.

Online supplemental material. Fig. S1 demonstrates the specificity of the pharmacological inhibitors PepA and CA. Fig. S2 demonstrates the localization of cathepsin D in azurophilic granules and its translocation in the cytosol. Results of quantitative colocalization analysis are provided. Fig. S3 demonstrates the time-dependent appearance of cathepsin D in the cytosol compared with cytochrome c, as assessed by immunoblotting. Fig. S4 shows a transmission electron micrograph suggesting membrane permeabilization of azurophilic granules during neutrophil apoptosis. Fig. S5 illustrates the delayed spontaneous death of blood neutrophils from septic shock patients compared with control individuals. Supplemental materials and methods describe subcellular fractionation and transmission electron microscopy. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20072152/DC1.

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