A new twist to the emerging functions of ceramides in cancer: novel role for platelet acid sphingomyelinase in cancer metastasis

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It is now appreciated that sphingolipids constitute a rich class of bioactive molecules that include ceramide, sphingosine, and sphingosine 1-phosphate whose formation is controlled by a network of highly regulated enzymes (Hannun & Obeid, 2008). Notably, several stress stimuli induce the production of ceramide, which, as a single entity, has been traditionally associated with apoptotic and growth suppressive functions. However, recent data clearly suggest that this simplistic formulation is no longer tenable.

See also: A Carpinteiro et al (June 2015)

The actions of distinct enzymes introduce variations in the structure of ceramide (chain length, fatty acid saturation, and hydroxylation), resulting in several individual ceramides. Furthermore, the production of sphingolipids is compartmentalized within the cell and controlled by distinct enzymes (Fig 1A). Among the best studied are the sphingomyelinases (SMases), a family of at least four different gene products that show distinct subcellular localization and distinct mechanisms of regulation (i.e., they respond to different stimuli). Among them, acid SMase (ASM in humans, Asm in mice), the product of the SMPD1 gene (sphingomyelin (SM) phosphodiesterase 1), modulates SM homeostasis but also participates in the response to a diverse set of cytokines or apoptotic stimuli (Jenkins et al, 2010). Indeed, several tumor types are known to have low levels of ASM. However, recent findings (Petersen et al, 2013), including the study by Carpinteiro et al (2015) in this issue of EMBO Molecular Medicine, are challenging this notion and redefining the study of ASM in tumor biology.

Under normal physiologic conditions, the SMPD1 transcript is translated into immature pro-ASM that is post-translationally modified to produce secretory ASM (S-ASM) and lysosomal ASM (L-ASM). Deficiency of L-SMase results in Niemann-Pick disease types A and B (NPD A and NPD B, respectively). NPD A, with < 1% of normal ASM activity, is characterized by lipid accumulation in the lysosomes resulting in organ failure and severe neurologic deficits, whereas NPD B patients have ~4% normal ASM activity and survive well into adulthood. S-ASM is stimulated by cytokines and mediates specific responses such as induction of chemokines, but also acts on SM in lipoproteins. Thus, even with one enzyme of sphingolipid metabolism, there could be more than one subcellular site of action. The study by Carpinteiro et al (2015) adds another layer by demonstrating cross-tissue action of S-ASM in the setting of tumor metastasis in vivo.

Carpinteiro et al (2015) were interested in defining the role of Asm and ceramide in hematogenous spread of tumor cells. They found that the number of lung metastases formed after injecting melanoma cancer cells into syngeneic mice was significantly decreased in the Asm knockout (KO) mice. Similar results were obtained using amitriptyline, an indirect inhibitor of Asm. These effects were not due to growth suppression once in the lung but rather to effects on earlier stages of the metastatic process. Moreover, ablation of Asm in the tumor cells had no effect on their ability to seed the lung. Together, the results suggest a role for host Asm in metastasis. In probing this role further, Carpinteiro et al (2015) turned to platelets presumably because of the role for platelets in the early phases of metastasis where platelet-rich thrombi form around tumor cells and create a favorable microenvironment (Fig 1B). Furthermore, treatment of platelets with thrombin induces Asm secretion (Romiti et al, 2000), raising the possibility for a role of Asm in platelet/tumor interactions. Indeed, the authors found that transfusion of wild-type (WT) platelets into the Asm KO restored the metastatic potential of melanoma cells, thus demonstrating that the defect in metastasis is largely due to the absence of platelet Asm.

One exciting corollary from this study relates to observations in the heterozygous Asm mouse (Asm<sup>−/−</sup>). Several studies have relied heavily on mouse models of NPD with total KO of the Smpd1 gene (Asm<sup>−/−</sup>). These mice have no detectable Asm activity, and they display many of the phenotypes seen in NPD A patients, including rapid and progressive neuropathology, accumulation of foam cells in viscera, and dramatically shortened lifespan. This severe clinical phenotype...
complicates the interpretation of results, even when conducted at early ages (e.g., 7 weeks in this study); e.g. foamy macrophages are present in bronchial alveolar lavage of Asm−/− animals at 3–4 weeks of age (Dhami et al., 2001). Therefore, it becomes difficult to discern primary effects due to loss of ASM from consequences of organ failure and cell pathology, many of which, such as dysfunction of the endolysosomal system, are shared among different lysosomal storage disorders and thus do not signify specific roles for the target enzyme. This may underlie some of the conflicting results on Asm. For example, Asm-deficient animals develop more liver metastases following intra-splenic injection of colon carcinoma cells (Osawa et al., 2013), but it was not clear whether this was secondary to liver dysfunction or due to specific actions mediated by Asm. In this regard, there has been a paucity of studies employing the heterozygous Asm mouse as an Asm-attenuated model. Asm+/− mice have 40–70% activity, and in contrast to nullizygous mice, no developmental phenotype has been observed (Horinouchi et al., 1995). Notably, Moles et al. (2010) employed heterozygous animals to define a role for Asm-mediated cathepsin activation in liver fibrosis. In this context, Carpinteiro et al. (2015) employed Asm−/− mice and found significant attenuation of metastasis, strongly adding credence to a pro-metastatic role for host Asm.

Mechanistically the authors show that Zn2+-dependent Asm activity (a signature of S-ASM) and ceramide levels increased in the media following co-incubation of B16F10 tumor cells with platelets and that platelets are the source of the secreted Asm. Indeed, treatment of B16F10 cells with recombinant Asm or with ceramide restored tumor formation in KO mice.

Using an antibody that recognizes ceramide, the authors suggest a role for ceramide raft formation by demonstrating that ceramide and integrins co-localized on the surface of B16F10 cells after treatment with Asm or with WT platelets. The authors also show that Zn2+-dependent L-Asm has sufficient activity at neutral pH to alter the lipid content of the plasma membrane, confirming and extending previous work by Schissel et al. (1996). This would constitute a mechanism for the trans-cellular action of ASM.

Several important questions are generated from this study. A critical biophysical property of proposed ceramide rafts is the threshold level at which plasma membrane ceramide begins forming rafts. From Fig 3

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Yusuf A Hannun & Benjamin Newcomb Platelet acid sphingomyelinase in cancer metastasis
in Carpinteiro et al (2015), it appears that increasing the total ceramide content to double the physiologic concentration induces raft formation and integrin activation. However, other work has shown that plasma membrane ceramide can induce apoptosis of tumor cells. This leaves the possibility that non-apoptotic metabolites of ceramide, such as sphingosine 1-phosphate, may also play a significant role in integrin activation. This possibility is not investigated in the current study, and the conclusion that ceramide rafts induce integrin clustering is primarily based on co-localization of integrins with ceramide, the latter detected by immunostaining with anti-ceramide antibodies. However, anti-ceramide staining has not been fully validated, and ceramide antibodies are known to immunostain other lipids including sphingomyelin (Cowart et al, 2002). Validation of the ceramide antibody should be established (e.g., by artificial production of ceramide at the plasma membrane, with bacterial SMase, for example, or by preventing ceramide production with selective inhibitors or accelerating ceramide removal).

Since several tumor types are known to express high levels of sphingomyelin and ceramide-producing enzymes (Komori et al, 1999), it would be interesting to determine whether this is associated with a higher intrinsic metastatic rate and whether pharmacologic inhibition of Asm could help reduce tumor metastasis. Asm inhibition has traditionally relied on the use of tricyclic antidepressants and other cationic amphipathic drugs (CADs), lysosomotropic molecules that induce loss of ASM (but also other enzymes such as acid ceramidase) (Hurwitz et al, 1994). Given their lack of specificity, these reagents can be used judiciously in cell biology and in in vivo studies to rule out a role for ASM if the results are negative. If the results are positive, however, they can only be considered as supporting evidence. For therapeutic purposes, it is thus unlikely that this class of molecules can become a class of viable inhibitors of ASM due to both specificity and potency issues.

Moreover, the use of CADs has resulted in some conflicting results. For example, Petersen et al utilized c-srcY527F-transformed NIH 3T3 murine embryonic fibroblasts to demonstrate that lysosomotropic agents, which inhibit Asm, selectively kill transformed cells. Using mammary fat pad injections of MCF7 breast carcinoma cells in SCID mice, they showed reduced xenograft growth in mice treated with siramesine (Petersen et al, 2013). On the other hand, Osawa et al (2013) showed that Asm KO mice had increased hepatic colonization following splenic injection of SLA colon carcinoma cells. As such, we believe that there is a pressing need to develop more specific and potent inhibitors of Asm and, if possible, target L-ASM selectively compared to S-ASM.

In conclusion, Carpinteiro et al (2015) provide compelling evidence of cross talk between platelets and tumor cells accomplished by trans-cellular signaling mediated by Zn2+ dependent Asm, which they show is required for lung seeding of melanoma cells. Although much still needs to be done, their work presents a new paradigm in the investigation of sphingolipids in cancer biology.

References

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