Enhancing effects on vacuole-targeting fungicidal activity of amphotericin B

Akira Ogita1,2, Ken-Ichi Fujita2 and Toshio Tanaka2*

1 Research Center for Urban Health and Sports, Osaka City University, Osaka, Japan
2 Department of Biology and Geosciences, Graduate School of Science, Osaka City University, Osaka, Japan

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INTRODUCTION

The polyene macrolide antibiotic amphotericin B (AmB; Figure 1A) was introduced in the mid-1950s as the first effective antifungal drug for treating systemic mycoses (Johnson and Einstein, 2007). AmB has been used for controlling life-threatening systemic fungal infections primarily caused by Candida albicans and Aspergillus fumigatus (Lemke et al., 2005). The susceptibility of fungal cells to AmB absolutely requires the presence of ergosterol in the fungal plasma membrane. Molecular modeling studies indicate that AmB creates aqueous pores consisting of an annulus of eight molecules linked hydrophilically to ergosterol embedded in phospholipid bilayers (Baginski et al., 2006). This type of ion channel structure results in altered plasma membrane permeability and leakage of vital cytoplasmic components, such as K+, ultimately inducing cell death (Kerridge, 1980, 1985). However, AmB-induced lethal effects cannot be simply explained by the disturbance in plasma membrane ion transport function, because the leakage of K+ has various secondary effects. Alonso et al. (1979) reported that AmB is a powerful inhibitor of both protein and ribonucleic acid synthesis in yeast cells at concentrations that do not affect the transport of amino acids or nucleosides. The induction of oxidative stress is also considered as an AmB-induced toxic event in C. albicans cells (Sokol-Anderson et al., 1986). The involvement of additional cytotoxic effects in AmB lethality is supported by the fact that the leakage of K+ is not necessarily accompanied by the loss of cell viability (Chen et al., 1978). Clearly, the exact mechanism of AmB lethality is very complex and has not been completely deciphered.

Recently, AmB-induced cytoplasmic events have been visualized as vacuole membrane fragmentation into numerous dotted particles in Saccharomyces cerevisiae and C. albicans, and such drastic structural damage has been considered as the major cause of AmB lethality (Ogita et al., 2006; Borjihan et al., 2009). Vacuole membrane fragmentation is observed concurrently with AmB-induced cell death when this antibiotic is used at a non-lethal concentration in combination with allicin (Figure 1B), an allyl-sulfur compound derived from garlic (Ogita et al., 2006). This Mini Review describes a possible mechanism underlying the vacuole-targeting fungicidal activity of AmB and its enhancement by allicin. This review also deals with chemosensitization of the vacuole-targeting fungicidal activity detected with a compound other than AmB.

ALLICIN AS AN ENHANCER OF THE FUNGICIDAL ACTIVITY OF AmB

Garlic (Allium sativum L.) has attracted much attention throughout human history as a medicinal panacea (Hobbs, 1992). A
the fungicidal activity of AmB (Ogita et al., 2006). AmB promotes K+ efflux during the process of cell death, but this ion efflux was not increased significantly in the presence of allicin (Ogita et al., 2010b). This means that AmB-induced K+ efflux could contribute to the lethality of AmB at least in part, but such an intracellular ionic imbalance is not a major cause in allicin-mediated enhancement of lethal cytotoxicity.

Fungal vacuole function is in osmoregulation, ion homeostasis, and cell volume regulation (Thumm, 2000; Wickner, 2000). Various hydrolytic enzymes, including proteases and nuclease, are thought to accumulate in vacuoles; hence, the damage to these organelles is considered a critical step in inducing cell death (Obara et al., 2001). In S. cerevisiae, vacuoles normally exhibit a swollen spherical architecture in untreated cells and are similarly observed in cells treated with either 120 μM allicin or a non-lethal concentration of AmB (0.5 μM; Ogita et al., 2006, 2007b, 2010b). Surprisingly, the organelles are visible as small discrete dots in the cytoplasm when cells are treated with a lethal concentration of AmB (5 μM), suggesting that vacuole membrane damage is the primary mechanism of AmB’s lethality (Ogita et al., 2006, 2007b, 2010b). Consistent with these findings is the detection of significant levels of AmB molecules in the cytoplasmic and plasma membrane fractions (Ogita et al., 2010b). Similar observations have been reported in studies on C. albicans (Borjihan et al., 2009).

**RELATIONSHIP BETWEEN AmB-INDUCED PLASMA AND VACUOLE MEMBRANE DAMAGE**

Although some controversy surrounds the lethal action of AmB, we believe that most researchers will agree that this antibiotic exerts its initial action at the fungal plasma membrane. According to the widely accepted hypothesis regarding the role of plasma membrane ergosterol, this neutral lipid is indispensable for the initial action of AmB (Ghamnoum and Rice, 1999; Lemke et al., 2005). This view is also supported by findings that AmB resistance results from lack of ergosterol biosynthesis (Ellis, 2002). Chemotherapeutic application of AmB is indeed based on its higher affinity and activity toward fungal plasma membranes containing ergosterol rather than cholesterol (Baginski et al., 2005). Brajtburg et al. (1980) discovered that the simultaneous addition of K+ and Mg2+ fully protects against the fungicidal activity of AmB. This fact supports the idea that AmB lethality is primarily dependent on the leakage of intracellular cations through the plasma membrane.

In contrast, Chen et al. (1978) reported that the aqueous channels or pores formed by AmB are not essential for its lethality for C. albicans, suggesting that the drug induces cell death by another mechanism. We have reported that the extracelluar addition of K+ and Mg2+ could restrict cytotoxicity as well as vacuole membrane fragmentation in S. cerevisiae treated with a lethal concentration of AmB alone (Ogita et al., 2010b). This may also suggest that AmB-induced leakage of these intracellular ions weakens the vacuole membrane or interferes with a mechanism of maintaining its normal rounded architecture. However, the simultaneous addition of allicin could still enhance AmB-induced cell death and the associated vacuole disruption. Our results reveal that, under certain conditions, AmB can cause vacuole disruption and cell death even when cells are protected against intracellular ion imbalance. Therefore, we conclude that allicin likely inhibits a mechanism required for maintaining vacuole membrane stability that cannot be protected by simply increasing the intracellular cation concentration.
INTRACELLULAR ERGOSTEROL-TRAFFICKING AS A POSSIBLE TARGET OF ALICIN

Yeast vacuoles normally undergo fission as well as fusion, and defects in vacuole fusion cause vacuole membrane fragmentation (Seeley et al., 2002). Ergosterol is required for proper fluidity as well as the relating function of plasma membrane (Zinser et al., 1991), and thus plays an important role in both endocytosis (Munn et al., 1999) and homotypic vacuole fusion (Kato and Wickner, 2001). This suggests the possibility that fungal cells can increase their vacuole membrane ergosterol content by provoking ergosterol-trafficking from the plasma membrane in response to the preliminary vacuole membrane damage such as that caused by the action of AmB at a non-lethal concentration (Ogita et al., 2009a). In support of such an urgent response, ergosterol-enriched S. cerevisiae cells are protected against the vacuole membrane damage induced by AmB alone at a lethal concentration as well as that caused by the combination of allicin and AmB at non-lethal concentrations (Ogita et al., 2009a). C. albicans responded identically to the lethal actions of AmB when the cells had been pretreated with exogenously added ergosterol (Borjihan et al., 2009).

Isolated vacuoles represent a much simpler system than intact cells for the study of vacuole membrane fission and fusion. They lack not only consecutive transport systems, but also the principal energy-producing systems, and most of the metabolic and biosynthetic systems of the cell (Boller et al., 1975). Although allicin alone had no effect on the morphology of the isolated vacuoles, AmB could disrupt their swollen spherical architectures regardless of whether allicin was added to the vacuole suspension (Borjihan et al., 2009; Ogita et al., 2009a). The vacuoles isolated from ergosterol-enriched cells are apparently resistant to the disruptive action of AmB when compared to those from untreated cells. The effect of allicin on the cellular localization of ergosterol could be visualized by filipin III. Ergosterol was, indeed, detected in the intracellular compartment corresponding to the vacuole membrane in addition to the plasma membrane when cells were treated with AmB alone at a non-lethal concentration (Ogita et al., 2009a). Unlikely, ergosterol remained in the plasma membrane without mobilization into the cytoplasm when allicin was added to the medium (Ogita et al., 2009a). These results suggest that the intracellular transport of ergosterol represents a protective reaction against the vacuole disruptive activity of AmB.

A model for the role of allicin in the vacuole-targeting fungicidal activity of AmB has been proposed based on these results (Figure 2). In response to the action of AmB, fungal cells increase the vacuole membrane ergosterol content by ergosterol-trafficking from the plasma membrane. As a result, at non-lethal concentrations of AmB, cells can maintain the large, spherical architecture of the organelle (Figure 2A). In the presence of allicin, however, AmB can cause vacuole membrane damage even at a non-lethal concentration, because the vacuole membrane cannot be suitably provided with ergosterol due to failure of its trafficking from the plasma membrane (Figure 2B). In contrast, in ergosterol-enriched cells, the vacuole membrane is protected against AmB-induced disruptive damage even if the intracellular ergosterol transport is inhibited by allicin (Figure 2C).

**FIGURE 2** | A proposed model for the role of allicin and ergosterol ( ◦) in the vacuole-targeting fungicidal activity of amphotericin B (AmB) ( ●). Untreated cells were incubated in the presence of AmB alone at a non-lethal concentration (A) and AmB at a non-lethal concentration and allicin (B). Ergosterol-enriched cells were incubated in the presence of AmB at a non-lethal concentration and allicin (C).
FUNGICIDAL ACTIVITIES OF OTHER COMPOUNDS THAT TARGET VACUOLES

Vacuole-targeting fungicidal activity has been markedly detected with the polyol macrolide antibiotic niphimycin (NM; Figure 3A) even in a nutrient medium (YPD) in which yeast cells are rather resistant to AmB, possibly because of their high ergosterol content derived from "yeast extract," a component of YPD medium (Subbiah and Abplanalp, 2003; Ogita et al., 2010b). The molecular structure of NM is characterized by a polyol lactone ring and alkylguanidium chain attached to the ring, suggesting the contribution of the alkyl side chain to the predominant lethality of NM even in ergosterol-rich medium (Nakayama et al., 2002).

N-Methyl-N′′-dodecylguanidine (MC12; Figure 3B) was synthesized and substituted for the alkyl side chain of NM, and was shown to strongly enhance the vacuole-targeting fungicidal activity of AmB against both S. cerevisiae and C. albicans (Usuki et al., 2000; Ogita et al., 2007a; Yutani et al., 2011b). The enhancing effect of MC12 on C. albicans pretreated with ergosterol was not observed, suggesting that it exerts an allicin-like inhibitory effect on intracellular ergosterol-trafficking.

The structure of the macrocyclic lactone ring of polyene macrolide compounds, in addition to the alkyl side chain, can also affect vacuole-targeting fungicidal activity. For example, nystatin possesses a polyene macrolide ring analogous to that of AmB, and its fungicidal activity results in disruption of vacuoles as is the case for AmB. The fungicidal activity of nystatin is also enhanced in the presence of allicin in parallel to an enhancement of vacuole disruption (Ogita et al., 2010a). In contrast, allicin is ineffective in enhancing the fungicidal activity of filipin III, a pentaene macrolide that exhibits toxic effects such as inducing oxidative stress but does not disrupt vacuoles (Bonneau et al., 2010; Ogita et al., 2010a).

Polymyxin B (PMB; Figure 3C) is a bactericidal cyclic peptide effective against Gram-negative bacteria. PMB passes through the outer membrane, interacting with plasma membrane phospholipids, and thus, its bactericidal action is proposed to depend on a marked increase in plasma membrane permeability. This antibiotic can also interact with the plasma membranes of S. cerevisiae and C. albicans, but its fungicidal activity is much lower than that detected against bacterial cells. In our chemical screening experiment, PMB exhibited fungicidal activity that is enhanced by allicin (Ogita et al., 2007b). As expected, the vacuole membrane damage was clearly observed in PMB-treated S. cerevisiae and could be enhanced along with cytotoxicity even in ergosterol-rich medium.

The vacuole-targeting fungicidal activity of PMB against C. albicans, A. niger, and Mucor mucedo is enhanced in the presence of ionophores like monensin and salinomycin (Ogita et al., 2009b). An organosulfur compound, zwiebelane A, isolated from onion bulbs, also enhances the vacuole-targeting fungicidal activity of PMB (Borjihan et al., 2010). Interestingly, the enhancing effects of ionophores and zwiebelane A were observed when S. cerevisiae cells were treated with PMB, but not with AmB, suggesting a difference in the mechanism of vacuole disruption between AmB and PMB.

In spite of its proven track record, the well-known side effects of AmB might prevent its clinical use. Clinical use of AmB has therefore improved with the aid of liposomal drug delivery systems that reduce its nephrotoxicity (Deray, 2002). A clinically effective dose of AmB may be also reduced when the antibiotic is used in combination with an agent like allicin. AmB resistance can be generated due to a genetic lack of plasma membrane ergosterol that plays an important role in the uptake of AmB into the fungal cytoplasm in addition to its participation in the formation of the plasma membrane's ion-permeable channel (Ogita et al., 2010b). Therefore, it seems possible to improve AmB-dependent chemotherapy by developing a method of enhancing its uptake into the cytoplasm of AmB-resistant fungal strains.
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