Electrostats and flexibility drive membrane recognition and early penetration by the antimicrobial peptide dendrimer bH1†

Harish K. Ravi, a Michaela Stach, a Thereza A. Soares, b Tamis Darbre, a Jean-Louis Reymond a and Michele Cascella* a

Molecular dynamics simulations of the polycationic antimicrobial peptide dendrimer bH1 (Leu)b(DapLeu)a(DapPhe)bDapLys-NH2 binding to membranes suggest that electrostatic interactions with the polyanionic lipopolysaccharide (LPS) and conformational flexibility of the 2,3-diaminopropanoic acid (Dap) branching units drive its selective insertion into microbial membranes.

Antibiotic resistant bacteria represent one of the major health care problems today, in particular Gram-negative pathogens such as Pseudomonas aeruginosa (PA), which are lethal to cystic fibrosis and immunocompromised patients. Recently, we reported the peptide dendrimer bH1 (Fig. 1) [1] as a new type of membrane disrupting antimicrobial peptide (AMP) [2] potentially suitable to address antibiotic resistance (Fig. 1). Dendrimer bH1 shows potent antimicrobial activity against Gram-negative E. coli (MIC = 1 µM) and P. aeruginosa (MIC = 5 µM) without showing significant hemolysis (MHC > 500 µM).

Most AMPs are polycationic amphiphilic sequences, which disrupt bacterial membranes by electrostatic binding to anionic groups of LPS or phospholipids followed by hydrophobicity driven membrane penetration, [3] which may lead to membrane lysis or cell penetration and subsequent interaction with an intracellular target. [4] To understand the selective interaction of bH1 with the LPS coated outer membrane of PA versus the eukaryotic cell membrane, we performed all atom molecular dynamics (MD) simulations of the dendrimer [5–8] interacting with membrane models constituted of a 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC) bilayer as a model of the euakaryotic membrane, or of an asymmetric LPS on top of a 1,2-dipalmytoyl-3-phosphatidyl-ethanolamine (DPPE) layer as a model of the PA outer membrane (structure in ESI†).

Standard MD protocols were followed using force field parameters and thermally equilibrated coordinates reported for POPC [9] and LPS. [10]

In both membrane systems bH1 approached the membrane surface in the first 10 ns of the simulation and established contacts via its multiple N-terminal ammonium groups. In bH1:POPC, the contacts involved a common terminal dendron Leu2Dap of bH1 binding to the phosphate groups of POPC present at the water–membrane interface. The dendrimer–membrane interface did not evolve significantly within the simulated time (500 ns) and the whole dendrimer structure remained fully solvated in the aqueous phase (ESI† Fig. S4).

In PA the outer phospho-saccharide chains of LPS assemble in a structured multilayer characterized by a high density of negative electrostatic charges, and a strong hydrophilic character. In our MD simulations, bH1 made contacts with the membrane surface within the first 15 ns and involved the positively charged Leu2Dap terminal dendron and the solvent-exposed LPS 0-β-D-glucose (0GB) and 0-α-D-glucose (0GA) moieties (Fig. 2E). Further penetration occurred around 18 ns by transient formation of hydrogen bonds between 6-α-D-glucose (6GA) in the LPS and the terminal leucines attached to the same Dap unit following a similar
The contacts of the Leu\textsubscript{2}Dap dendrons to 3H1 and PH2 were very stable, and no further penetration occurred within our simulation time (300 ns). Rather, the deep rooting of the first Leu\textsubscript{2}Dap dendron into the LPS, concomitant with deeper penetration of the leucines of the first Leu\textsubscript{2}Dap dendron to ~18 Å from the water–LPS interface, enabling its anchoring to the 1-glycero-\delta-manno-heptose-7-formamide (3H1) and phosphorylated 2\{2-hydroxyethyl\}-\delta-deoxy-\gamma-manno-heptose (PH2) of the LPS, whereby the three anionic phosphate groups of PH2 formed hydrogen bonds with the N-terminal ammonium groups of bH1 (Fig. 2H).

The contacts of the Leu\textsubscript{2}Dap dendrons to 3H1 and PH2 were very stable, and no further penetration occurred within our simulation time (300 ns). Rather, the deep rooting of the first Leu\textsubscript{2}Dap dendron into the LPS induced penetration of the other ones. In particular, after about 100 ns, one of the other three remaining Leu\textsubscript{2}Dap dendrons started to interact with the outer-core of the LPS, and then followed a similar hydrogen-bonding pattern observed for the initial anchoring Leu\textsubscript{2}Dap dendron. At the end of the MD simulation this second Leu\textsubscript{2}Dap dendron had penetrated well inside the outer core of the LPS and its terminal leucines were just 6 Å away from the inner core of the LPS.

Monitoring the insertion of the third and the fourth Leu\textsubscript{2}Dap dendrons, we observed a hydrogen-bonding pattern with the outer core identical to that of the first two dendrons. Translocation of the various terminal Leu\textsubscript{2}Dap dendrons of bH1 from the outer to the inner region of the LPS occurred on average in 100 ns by a diffusive mechanism in which the dendrimer built relatively weak and labile contacts with the various sugar moieties, in contrast to the initial insertion of the Leu\textsubscript{2}Dap dendrons into the LPS, which required some activation.

The Dap residue of the terminal Leu\textsubscript{2}Dap dendrons underwent significant conformational changes in the course of the MD simulation. Initially the dendrimer was fully hydrated and the dendrons assumed an extended conformation in which the Dap residue was present in the allowed \(\beta\)-sheet region of the Ramachandran plot (Fig. 2). Upon localization at the membrane–water interface, this extended conformation was destabilized and Dap assumed a coiled conformation typical for a right-handed \(\alpha\)-helix. The \(\beta\)-branch of the asymmetric Leu\textsubscript{2}Dap dendron is longer by one methylene group and conformationally more flexible than the \(\alpha\)-branch, and was always the first to penetrate the membrane, followed by Dap and the shorter \(\alpha\)-branch. After the Leu\textsubscript{2}Dap dendron had passed the region defined by the 0GB and 0GA saccharides and insertion into the LPS was complete, the secondary structure of Dap changed back to an extended conformation similar to that in the aqueous phase. Penetration of bH1 into the LPS thus seems to be favored by the conformational flexibility of the branching units, which allow local coiling and optimization of the interaction with the LPS at the water–membrane interface.

Electrostatic attraction between the positively charged terminal ends and complementary negatively charged phosphate groups provided the driving force inducing penetration of bH1 into the LPS during MD. The dendrimer insertion occurred through an entropy-driven process, with sequential binding of the different branches. The first layer of phosphate groups present in the LPS at about 18 Å from the membrane–water interface constituted the main basin of attraction for the terminal Leu\textsubscript{2}Dap dendrons. The simulated time (300 ns) was sufficient to observe localization of the dendrimer in this area of the LPS. The presence of a second
layer of phosphates roughly 26 Å deeper in the LPS may trigger further penetration in longer timescales.

The importance of electrostatic complementarity for the activity of bH1 was evidenced earlier by the observation that the uncharged, N-acetylated analog of bH1 shows no antibacterial or membrane disrupting effect. To further test this interaction, bH1 was exposed to 5(6)-carboxyfluorescein (CF) loaded large unilamellar vesicles (LUVs) composed of phosphatidylglycerol (PG) as head groups, and induced CF release at low concentration (1–30 μM) demonstrating an interaction and disruption of this membrane. In contrast, LUVs composed of phosphatidylcholine (PC) head groups showed only partial release of CF upon exposure to bH1 up to 200 μM mL⁻¹, showing that bH1 does not interact with zwitterionic groups (Fig. 3). bH1 also shows high potency against Gram positive Bacillus subtilis (MIC = 1 μM). Based on the present simulation we predict that bH1 can interact with negatively charged peptidoglycan elements, like teichoic acids, which are key to the structural stability of the outer cell wall.

In conclusion, the driving force facilitating penetration of the antimicrobial peptide dendrimer bH1 in the LPS of PA during MD was provided by electrostatic attractions exerted by the phosphate groups localized at the inner regions of the LPS layer on the external side of the membrane, which were established after initial membrane adsorption and sequential insertion of the four terminal Leu-Dap dendrons, an event requiring local coiling of the branching Dap amino acid to facilitate adaptation to the geometrical restraints at the membrane–water interface. The observation of these coiling events at the interface-crossing point suggests that flexible branching units are crucial to the antimicrobial activity of peptide dendrimers, a finding which may be useful in future designs. Note that in the simulation time presented here the LPS bilayer was still well organized and only marginally perturbed by the dendrimer. It is likely that the antimicrobial function of bH1 is associated with phenomena occurring at longer timescales, possibly comprising aggregation of multiple bH1 units into the LPS. A crucial role played by aggregation has been already described for membrane-binding antibacterial linear peptides. Future studies addressing longer MD simulations, also with a higher concentration of bH1 molecules, may be able to elucidate this other aspect of the biological activity of peptide dendrimers.

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Notes and references

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