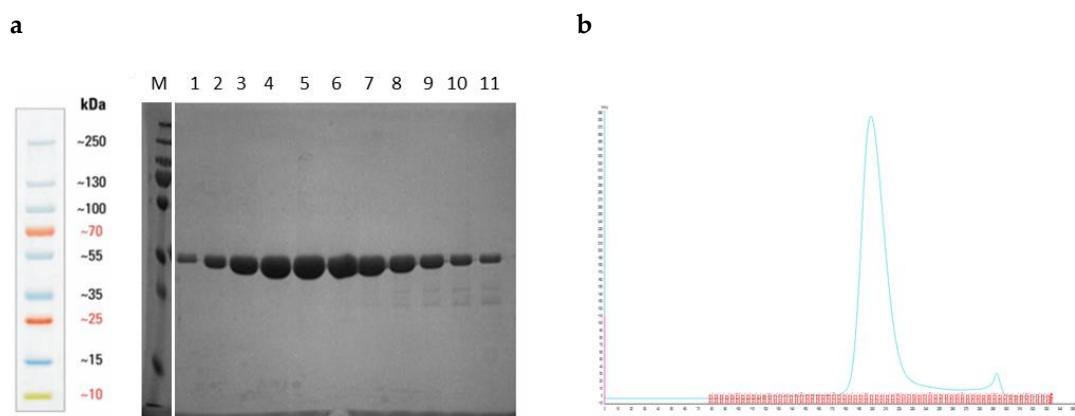


CLUSTAL O(1.2.4) multiple sequence alignment

EL11_EPI51560.1	SAGYDYTSVQNKVNELLGVKAYRKSVDLAREVIRGTTWGNNGSTRKQRLTQAGYDYDAVQK	300
EL3_WP_065189629.1	SAGYDYASVQNKVNELLGVKAYRKSVDLAREVIRGTTWGNNSMRKHRLTQAGYDYDAVQK	300
EL10_WP_064340486.1	SAGYDYASVQNKVNELLGVKAYRKSVDLAREVIRGTTWGNNGSTRKQRLTQAGYDYNQAVQK	300
EL12_WP_048730021.1	SAGYDYTSVQNKVNELLGVKAYRKSVDLAREVIRGTTWGNNGKTRKQRLTQAGYDYNQAVQK	300
EL8_EPI52092.1	QAGYDYTSVQNKVNKLGVKACRKSVDLAREVIRGTTWGNNGNERKNRLTQAGYDYDTVQK	300
EL5_EPI54247.1	QAGYDYTSVQNKVNKLGVKACRKSVDLAREVIRGTTWGNNGNERKNRLTQAGYDYDTVQK	300
EL7_WP_004104952.1	QAGYDYTSVQNKVNKLGVKACRKSVDLAREVIRGTTWGNNGNERKNRLTQAGYDYDTVQK	300
EL4_WP_004111234.1	SAGYDYASVQNKVNKLGVKAYRKSVDLAREVIRGTTWGNNGNERKNRLAQAGYDYDTVQK	300
EL6_EIK79883.1	QAGYDYASVAK-----	251
EL1_AEF31373.1	SAGYDYASVQNKVNELLGVKACRKSVDLAREVIRGAWGNGSTRKQRLAQAGYDYDTVQK	300
EL9_WP_004118369.1	SAGYDYASVAK-----	251
EL2_WP_065189413.1	SAGYDYASVQNKVNELLGVKACRKSVDLAREVIRGTTWGNNGNERKNRLTSAGYDYDTVQK	300
EL13_WP_076002731.1	SAGYDYASVQNKVNELLGVKACRKSVDLAREVIRGAWGNGSTRKQRLTSAGYDYDTVQK	300
EL14_WP_076002856.1	.*****:*** :	
EL11_EPI51560.1	RVNELL	306
EL3_WP_065189629.1	RVNELL	306
EL10_WP_064340486.1	RVNELL	306
EL12_WP_048730021.1	RVNELL	306
EL8_EPI52092.1	RVNELL	306
EL5_EPI54247.1	RVNELL	306
EL7_WP_004104952.1	RVNELL	306
EL4_WP_004111234.1	RVNELL	306
EL6_EIK79883.1	-----	251
EL1_AEF31373.1	RVNELL	306
EL9_WP_004118369.1	-----	251
EL2_WP_065189413.1	RVNELL	306
EL13_WP_076002731.1	RVNELL	306
EL14_WP_076002856.1	RVNELL	306

Supplementary Figure 1: Wild-type endolysins (EL1-EL14) encoded on prophage-like regions of *Gardnerella* genomes. A multiple sequence alignment was performed by Clustal Omega analysis (<https://www.ebi.ac.uk/Tools/msa/clustalo/>). The protein accession numbers are given after each endolysin name (e.g. EL1_AEF31373.1). The yellow background indicates the EAD (H-domain), the linker sequence is highlighted in grey, and the cell wall binding domain (B-domain) in blue. The domain regions were predicted by Interpro (<https://www.ebi.ac.uk/interpro/>).



Supplementary Figure 2. (a) PM-477 endolysin protein was recombinantly expressed and purified via Ni-NTA chromatography (FPLC). Fractions obtained from the final size exclusion chromatography (FPLC) were analyzed for purity by SDS-PAGE and Coomassie blue staining. MWM was used to confirm the expected protein size (left). Fractions 1-11 were pooled and used for further experiments. (b) FPLC chromatograph of PM-477 purification on a Superdex 75 column resulting in a defined peak from 182 mL to 222 mL. Red bar represents the fractions 1-11 that were pooled and concentrated to a final volume with a concentration of 0.616 mg/mL (determined by A280 measurement).