

Supplementary Table. Natural compounds and extracts as biofilm disrupting agents and QS inhibitors.

Natural agent	Bacterial species	Further information	Bibliographic reference
Chitosan	<i>Pseudomonas aeruginosa</i> PA01, <i>Streptococcus mutans</i> ATCC 35668, <i>Staphylococcus aureus</i> MW-2, <i>Acinetobacter baumannii</i> ATCC 19606, <i>Klebsiella pneumoniae</i> ATCC13883, <i>Enterococcus faecalis</i> ATCC51299	Chitosan derivatized with arginine and lactobionic acid could disaggregate a two-day old biofilm	[21]
Vitamin E	<i>Staphylococcus aureus</i> , <i>Staphylococcus epidermidis</i> , <i>Escherichia coli</i> , <i>Klebsiella pneumoniae</i> , <i>Proteus mirabilis</i> , <i>Acinetobacter baumannii</i> , <i>Pseudomonas aeruginosa</i> , <i>Pseudomonas putida</i>	An oily topical formulation to limit bacterial and fungal biofilm production was tested in a clinical trial involving 20 wounded patients	[21]
Abscissic acid	<i>Propionibacterium acnes</i> , <i>Staphylococcus aureus</i> , <i>Pseudomonas aeruginosa</i>	Abscissic acid, alone or in association with plant extracts ( <i>Morinda citrifolia</i> L., <i>Olea europaea</i> var. <i>sylvestris</i> Brot., <i>Curcuma longa</i> L.) was assayed for: The inhibitory activity against <i>Propionibacterium acnes</i> , <i>Staphylococcus aureus</i> , <i>Pseudomonas aeruginosa</i> and <i>Malassezia furfur</i> ; the inhibition of the synthesis of QS molecules; and the reduction of inflammation markers in keratinocytes infected with bacteria.	[21]

		In addition, a synergistic effect between abscissic acid (30 µg/mL-60 µg/mL) and plant extracts (2%-15% by weight) against <i>Staphylococcus aureus</i> and <i>Propionibacterium acnes</i> biofilm was assessed	
Aqueous and ethanolic extracts of <i>Rhamnus prinoides</i> L'Hér. stem and leaves and their main constituents	<i>Streptococcus mutans</i> , <i>Staphylococcus aureus</i> , <i>Pseudomonas aeruginosa</i> , <i>Bacillus subtilis</i>	Extracts (3 mg/mL) and single bioactives (4-hydroxy-4-methyl-2-pentanone, ethyl 4-ethoxybenzoate and benzoic compounds) were tested against polymicrobial biofilms in-vitro of <i>Staphylococcus aureus</i> and <i>Pseudomonas aeruginosa</i>	[21]
Mannose, Methyl α - Dmannopyranoside, 2-Deoxy-Dglucose, Methyl α - D-glucopyranoside	<i>Desulfovibrio vulgaris</i> ATCC 29579, <i>Desulfovibrio desulfuricans</i> DSM 12129	High concentrations (1 mM-500 mM) of each compound gave a consistent eradication of a mature biofilm if applied for 2 h-14 h. Mannose was the best biofilm dispersing agent for both species	[21]

Lipopolysaccharide	<i>Vibrio vulnificus, Pseudomonas aeruginosa, Staphylococcus aureus, Listeria monocytogenes</i>	Detoxified lipopolysaccharide from <i>Bacteroides vulgatus</i> MGM001 was effective in association with lipoteichoic acid to reduce biofilm formation on various materials, except acrylic matrices	[21]
Amorfrutin B	<i>Pseudomonas aeruginosa</i>	showed promising activity with inhibition	[1]
Coumarin and hydroxylated derivatives	<i>Chromobacterium violaceum</i> CV026	All the different coumarins tested with the exception of 4-hydroxycoumarin and dihydrocoumarin, inhibited the violacein production.	[22]