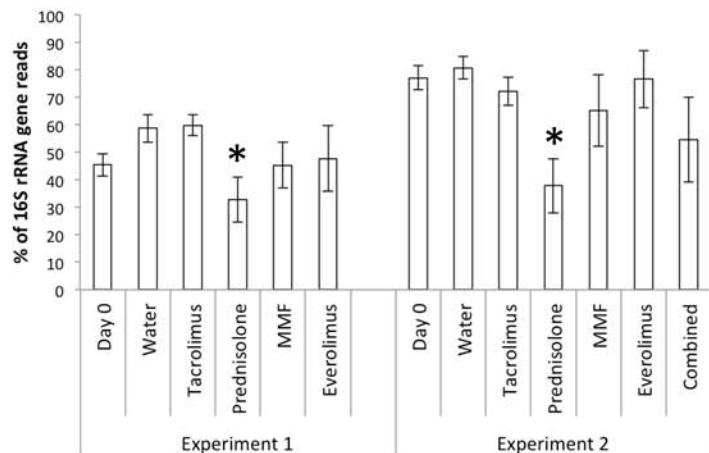


Table S1: Primers and cycling conditions used in this work

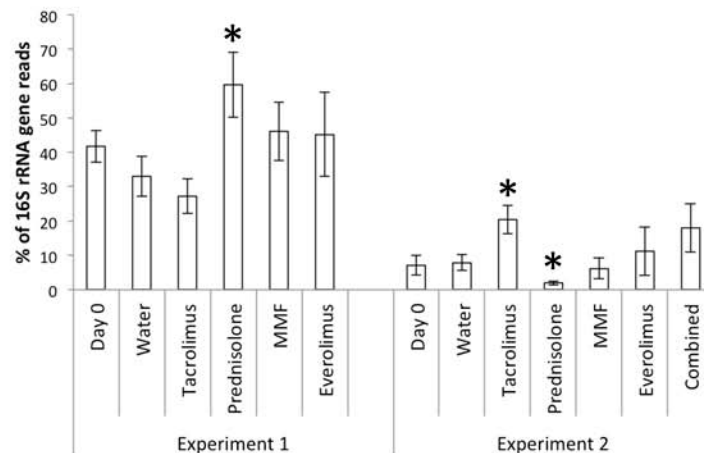
Target gene	Forward primer	Reverse primer
Reg3 β	GGCTTCATTCTTGTCTCCA	TCCACCTCCATTGGGTTCT
Reg3 γ	AAGCTTCCTTCCTGTCCTCC	TCCACCTCTGTTGGGTTTCA
IL-22	GCAATCAGCTCAGCTCCTGT	CGCCTTGATCTCTCCACTCT
GADPH	ATTGTCAGCAATGCATCCTG	ATGGACTGTGGTCATGAGCC

The reaction cycling conditions were: one enzyme activation step at 95 °C for 10 min followed by 40 cycles of 95 °C, 30 s, annealing for 30 s at 60 °C and extension at 72 °C for another 30 s. Fluorescence readings were taken at the end of each extension.

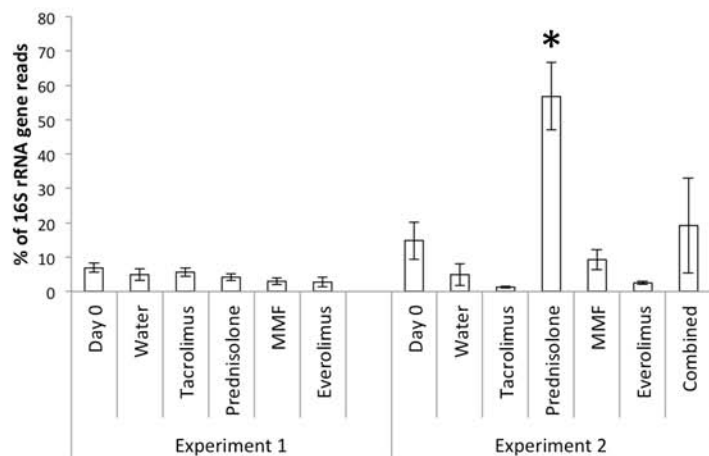
Bacteroidales



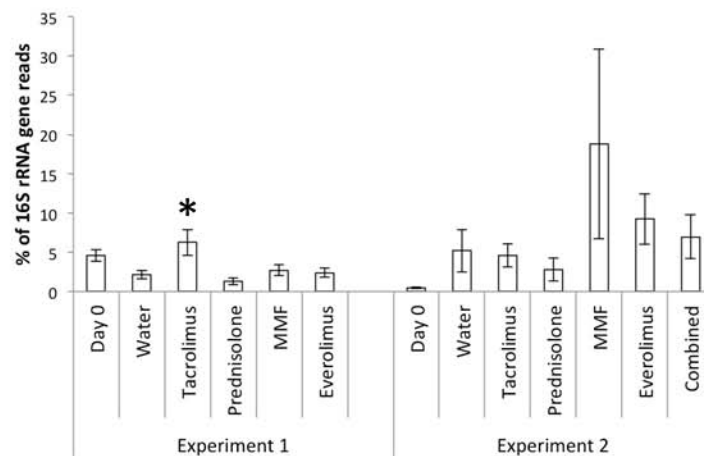
Clostridia



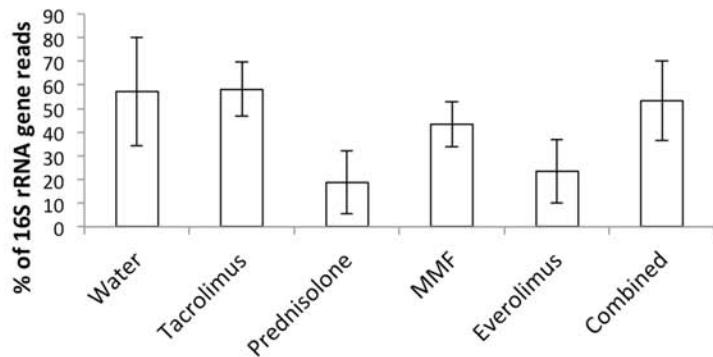
Bacilli



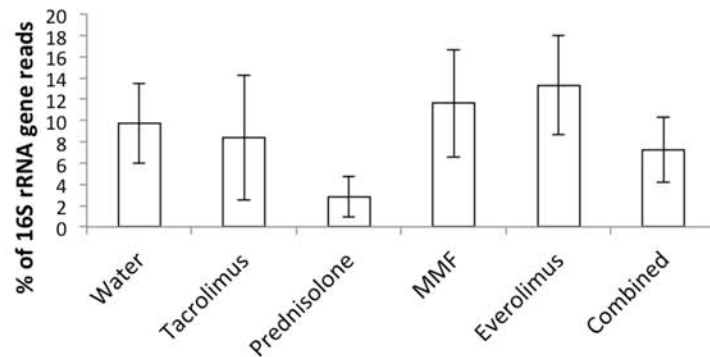
Erysipelotrichia



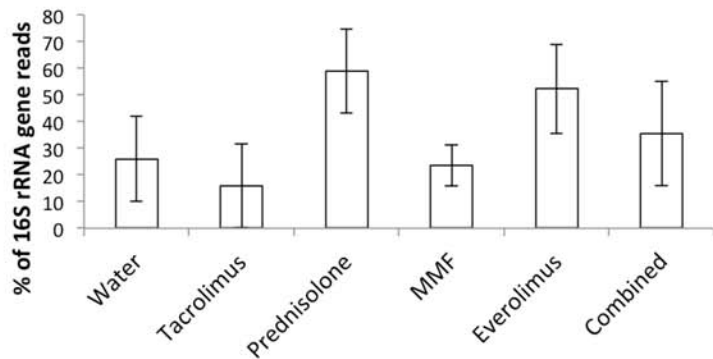
Erysipelotrichia



Clostridia



Bacilli



Bacteroidia

