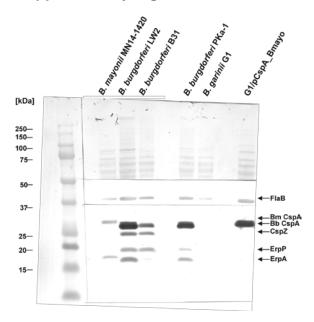
Supplementary Figure 5



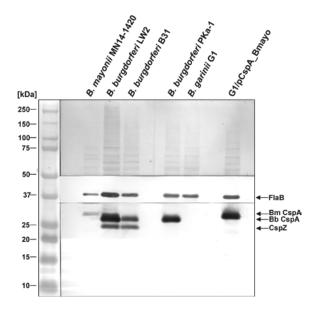
Uncropped gel and Western blots of figure 3A. Nitrocellulose membranes were scanned by a GS-710 Imaging Densitometer (Bio-Rad) and for image processing the Quantity One 4.2.1 software (Bio-Rad) was used. The general settings were as follows:

Application: Blot; HRP-Substrate (DAB)

Filter: Green

• Resolution: 42,3 * 42,3 microns = High for small gels with tiny features

Transform operations: high: 4095; low: 379; gamma: 0.71



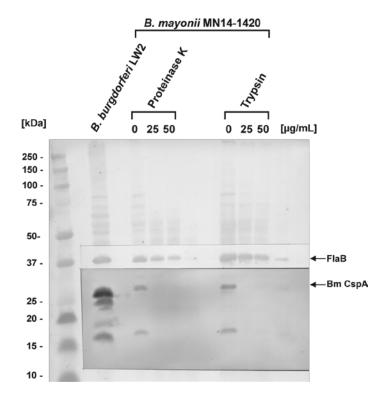
Uncropped gel and Western blots of figure 3B. Nitrocellulose membranes were scanned by a GS-710 Imaging Densitometer (Bio-Rad) and for image processing the Quantity One 4.2.1 software (Bio-Rad) was used. The general settings were as follows:

• Application: Blot; HRP-Substrate (DAB)

Filter: Green

Resolution: 42,3 * 42,3 microns = High for small gels with tiny features

Transform operations: high: 4095; low: 0; gamma: 0.85



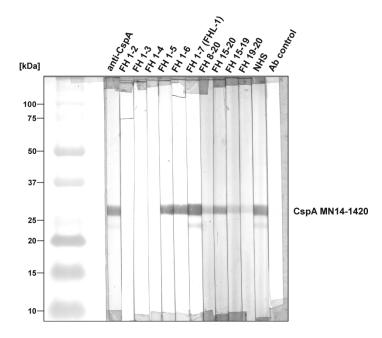
Uncropped gel and Western blots of figure 3C. Nitrocellulose membranes were scanned by a GS-710 Imaging Densitometer (Bio-Rad) and for image processing the Quantity One 4.2.1 software (Bio-Rad) was used. The general settings were as follows:

Application: Blot; HRP-Substrate (DAB)

• Filter: Green

Resolution: 42,3 * 42,3 microns = High for small gels with tiny features

Transform operations: high: 4095; low: 0; gamma: 1.0



Uncropped gel and Western blot of figure 4E. Nitrocellulose membranes were scanned by a GS-710 Imaging Densitometer (Bio-Rad) and for image processing the Quantity One 4.2.1 software (Bio-Rad) was used. The general settings were as follows:

Application: Blot; HRP-Substrate (DAB)

• Filter: Green

• Resolution: 42,3 * 42,3 microns = High for small gels with tiny features

Transform operations: high: 4095; low; gamma: 1.00