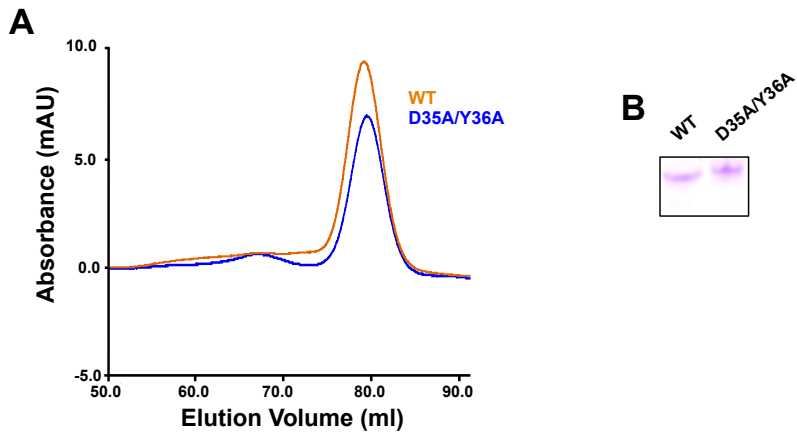


**Supplementary table 1.** Sequences of the primers used for the generation of CrmD mutants. The substituted bases are underlined.

Primer	Mutation	Sequence (5'-3')
CrmD37	K44A	GGTACAGACTATAACAGTAATAATCTATGTTGT <u>G</u> CACAATGCAATCCTGGAAT
CrmD38	K44A	ATTCCAGGATTGCATTGT <u>G</u> CACAACATAGATTATTACTGTTATAGTCTGTACC
CrmD39	P68A	TACAAAATGTGACAAGT <u>G</u> CGCAGATGACACCTTTACATC
CrmD40	P68A	GATGTAAAGGTGTCATCT <u>G</u> CGCACTGTGCACATTTTGTA
CrmD41	N77A	AGATGACACCTTTACATCCATT <u>C</u> TCTCATAGTCCCGCGTG
CrmD42	N77A	CACGCGGGACTATGAG <u>C</u> AGGAATGGATGTAAAGGTGTCATCT
CrmD43	N77F	AGATGACACCTTTACATCCATT <u>C</u> TTTCATAGTCCCGCGTG
CrmD44	N77F	CACGCGGGACTATGAA <u>A</u> AGGAATGGATGTAAAGGTGTCATCT
CrmD45	Q93A/E95A	AAGTTGTCGAGGCAAATGTAGCAGTAAT <u>G</u> CAGTAG <u>C</u> GACTAAATCGTGTAGTAAC
CrmD46	Q93A/E95A	GTTACTACACGATTTAGT <u>C</u> GCTACT <u>G</u> CATTACTGCTACATTTGCCTCGACAACCTT
CrmD47	Q103A/D104A/R105A	GTAATCAAGT <u>F</u> AGAGACTAAATCGTGTAGTAACACAG <u>C</u> GG <u>C</u> CGAGTATGTGCTGTGCATC
CrmD48	Q103A/D104A/R105A	GATGCACAGACACATACT <u>G</u> CG <u>G</u> CGCTGTGTTACTACACGATTTAGTCTCTACTTGATTAC
CrmD49	Y141A/S142A/S143A	GTGGTCTGGTTACGGTGTATATGGC <u>C</u> CG <u>C</u> AGCTAAAGGAGATGTAATATGTA AAAAGT
CrmD50	Y141A/S142A/S143A	ACTTTTACATATTACATCTCCTTTAG <u>C</u> TG <u>C</u> GGCGCCATATACACCGTAACCAGAACCAC
CrmD51	D35A/Y36A	CCCATTAATGGGAAATGTAACGGTACAG <u>C</u> CGTAACAGTAATAATCTATGTTGTAACAA
CrmD52	D35A/Y36A	TTGTTTACAACATAGATTATTA <u>C</u> TGTTAG <u>C</u> GGCTGTACCGTTACATTTCCCATTAATGGG
CrmD53	F72A/T73A/S74A	GACAAGTGCCAGATGACAC <u>C</u> G <u>T</u> G <u>C</u> AGCCATTCCTAATCATAGTCCCG
CrmD54	F72A/T73A/S74A	CGGGACTATGATTAGGAATGG <u>C</u> TG <u>C</u> AGGGTGTATCTGGGCACTTGTC
CrmD55	H78A/S79A	CCCAGATGACACCTTTACATCCATTCCTAAT <u>G</u> C <u>T</u> GCTCCCGCGTGTCTAAGT
CrmD56	H78A/S79A	ACTTAGACACGCGGGAG <u>C</u> AG <u>C</u> ATTAGGAATGGATGTAAAGGTGTCATCTGGG
CrmD57	R86A/K86A	TAGTCCCGCGTGTCTAAGTTG <u>T</u> G <u>C</u> AGGGCAGTGTAGCAGTAATCAAGTAGAG
CrmD58	R86A/K86A	CTCTACTTGATTACTGCTACAT <u>G</u> CGCCTG <u>C</u> ACAACCTTAGACACGCGGGACTA
CrmD59	S90A/S91A	GTGTCTAAGTTGTCGAGGCAAATGT <u>G</u> CG <u>C</u> TAATCAAGTAGAGACTAAATCGTGT
CrmD60	S90A/S91A	ACACGATTTAGTCTCTACTTGATTAG <u>C</u> GGCAGATTTGCCTCGACAACCTTAGACAC
CrmD61	E116A/F117A/E118A	TGCATCCGGATACTACTGCG <u>C</u> AG <u>C</u> TG <u>C</u> AGGATCAAACGGTTGCAGGC
CrmD62	E116A/F117A/E118A	GCCTGCAACCGTTTGATCCTG <u>C</u> AG <u>C</u> TG <u>C</u> GCAGTAGTATCCGGATGCA
CrmD63	D146A/I148A	GTATATGGCTACTCATCTAAAGGAG <u>C</u> TGTAGCATGTAAAAAGTGTCCGGGTAATATA
CrmD64	D146A/I148A	TATATTACCCGGACACTTTTTACAT <u>G</u> C <u>T</u> AC <u>G</u> CTCCTTTAGATGAGTAGCCATATAC



**Figure S1. CrmD D35A/Y36A mutant displays the same oligomeric state than wild type CrmD.** The oligomeric state of wild type CrmD (WT) and the CRD1 mutant D35A/Y36A was compared by size exclusion chromatography (A) and electrophoresis in native gel (B). In A, 100  $\mu$ g of WT (orange) and D35A/Y36A (blue) CrmD mutant in 1 ml of PBS were eluted in a Superdex 200 gel filtration column. In B, 500 ng of WT and D35A/Y36A were analyzed in a native gel using the PhastGel System. Proteins were loaded in a 12.5% homogeneous gel with native buffer strips. Subsequently, the gel was stained with coomassie blue.