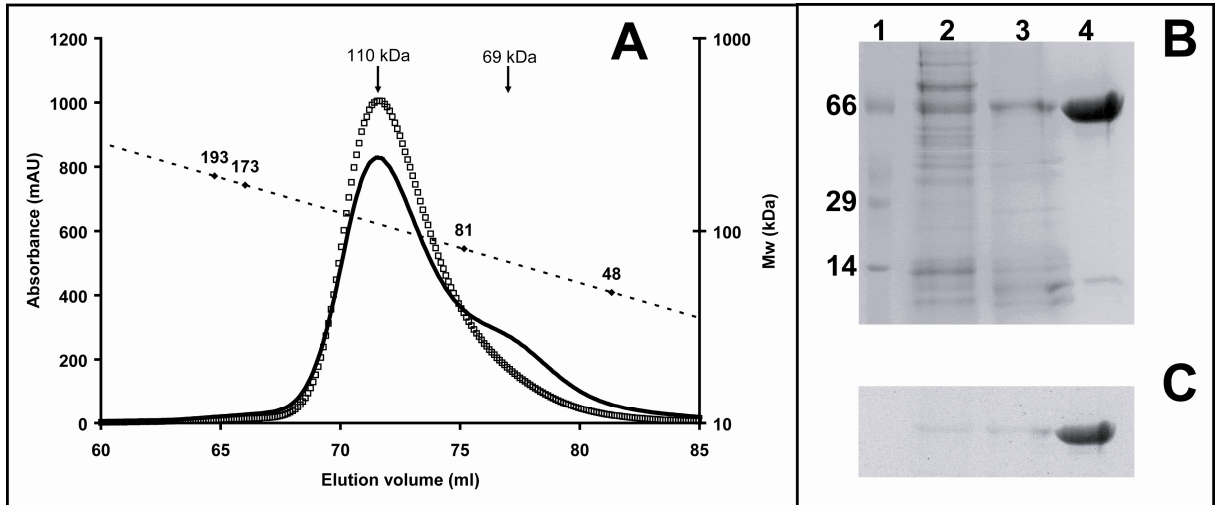


Supplementary Material



Supplementary Figure 1

(A) Size-exclusion chromatography (HiLoad 16/60 prep grade Superdex 200 column; GE Healthcare) of Cph1 Δ 2 in the Pr state recorded at 280 (-) and 660 nm (\square). Dotted line: column calibration with marker proteins (kDa). (B) SDS-PAGE gel after Coomassie staining, (C) SDS-PAGE gel demonstrating zinc acetate fluorescence of the supernatant after cell lysis (2), after Ni²⁺ affinity chromatography (3) and after SEC (4). Molecular weight markers are indicated in (1).

Supplementary Table 1

Data collection for Cph1Δ2 crystals

Data processing	blue light & glycerol	IR & cryo salts
Beamline	X-13, EMBL, Hamburg	ID14-3, ESRF, Grenoble
Wavelength (Å)	0.8015	0.9330
Detector	MAR CCD 165 mm	ADSC Q4 CCD
Space group	$P4_321$	$P4_321$
<i>a</i> , <i>b</i> , <i>c</i> (Å)	75.69, 75.69, 246.58	77.18, 77.18, 249.00
Maximal resolution (Å)	3.15 x 3.15 x 2.8	2.70 x 2.70 x 2.20
Total reflections	76347	183551
Unique reflections	14707	26369
Completeness ^a	0.789 (0.166); 0.976	0.672 (0.067); 0.985
$\langle I \rangle / \sigma \langle I \rangle$ ^a	16.5 (3.8)	23.0 (3.5)
R_{merge} ^b	0.072 (0.324)	0.049 (0.361)
Mosaicity (°)	0.42	0.35
Wilson <i>B</i> -factor (Å ²)	56.6	17.3

^a Values in parentheses correspond to the highest resolution shell; 2nd completeness corresponds to range 25-3.15 Å and 25-2.70 Å, respectively.

^b $R_{\text{merge}} = \frac{\sum_{\text{hkl}} \sum_i (I_i(\text{hkl}) - \langle I(\text{hkl}) \rangle)}{\sum_{\text{hkl}} \sum_i I_i(\text{hkl})}$.