

The fungal phytochrome FphA from *Aspergillus nidulans*

Sonja Brandt¹, David von Stetten², Mina Günther², Peter Hildebrandt², Nicole Frankenberg-Dinkel^{1,*}

¹ Physiologie der Mikroorganismen, Ruhr-Universität Bochum, Universitätsstr. 150, D-44780 Bochum, Germany.

² Technische Universität Berlin, Institut für Chemie, Sekr. PC14, Straße des 17. Juni 135, D-10623 Berlin, Germany.

SUPPLEMENTAL FIGURES

Figure S1. RR spectra of FphAN753 in the meta-Rc state in comparison to DrBphP. FphAN753 (top – abbreviated as FphA) and DrBphP were trapped in the meta-Rc state at -30°C during irradiation with red light. The spectra display the marker band region (1500-1700 cm⁻¹) from samples dissolved in H₂O and D₂O. The spectra of DrBphP were taken from ref. 19.

Figure S2. Absorbance spectra of FphAN753_H504A-BV and BV control sample. Absorbance spectra of 10 μM apo-FphAN753_H504A after incubation with 20 μM BV for 30 min in the dark (solid line) and 20 μM BV in EW buffer (dotted line).

Figure S 1. Brandt, S *et al.*

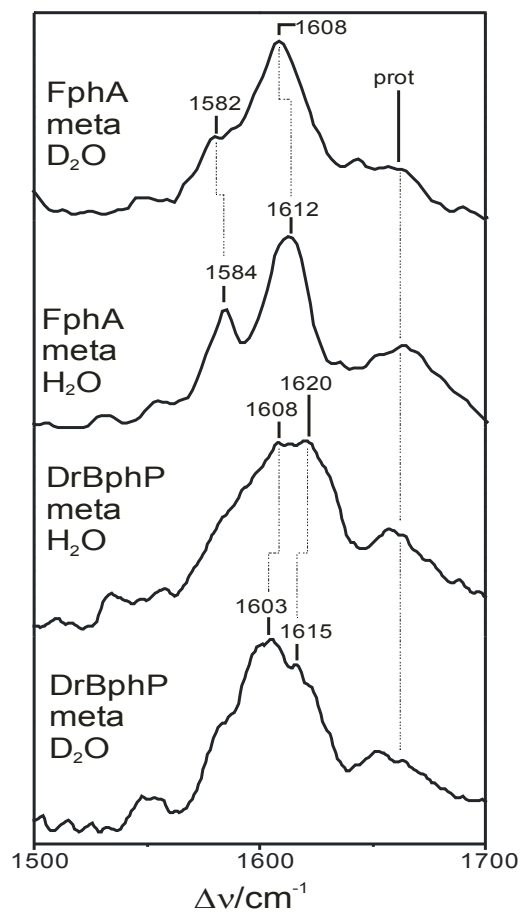


Figure S 2. Brandt, S *et al.*

