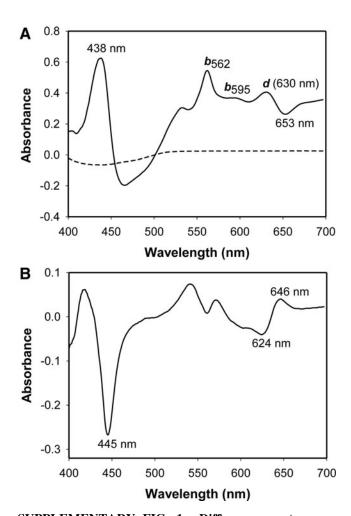
Supplementary Data



SUPPLEMENTARY FIG. 1. Difference spectra confirm the lack of cytochromes in the hemA mutant and their reconstitution. Cultures were grown as previously described (but without CORM-3 treatment) for 7–8 h (reconstituted cells) or 24 h (mutant cells), then harvested, washed thoroughly, and resuspended in 0.1 M KPi (pH 7). Cellular cytochrome content was assessed by dual wavelength spectrophotometry at room temperature. The resultant data were smoothed in SigmaPlot (Systat Software Inc.). Recorded wavelength positions are shown. (A) Reduced minus oxidized spectra of the hemA mutant (dashed line, 17.3 mg protein ml⁻¹) and after reconstitution with δ-ALA (solid line, 21.8 mg protein ml⁻¹). (B) CO-reduced minus reduced spectrum of cells after reconstitution with δ-ALA. Concentrations of cytochromes b (0.46 nmol mg protein⁻¹) and d (0.36 nmol mg protein⁻¹) were calculated from the reduced minus oxidized spectrum after reconstitution with δ-ALA.