

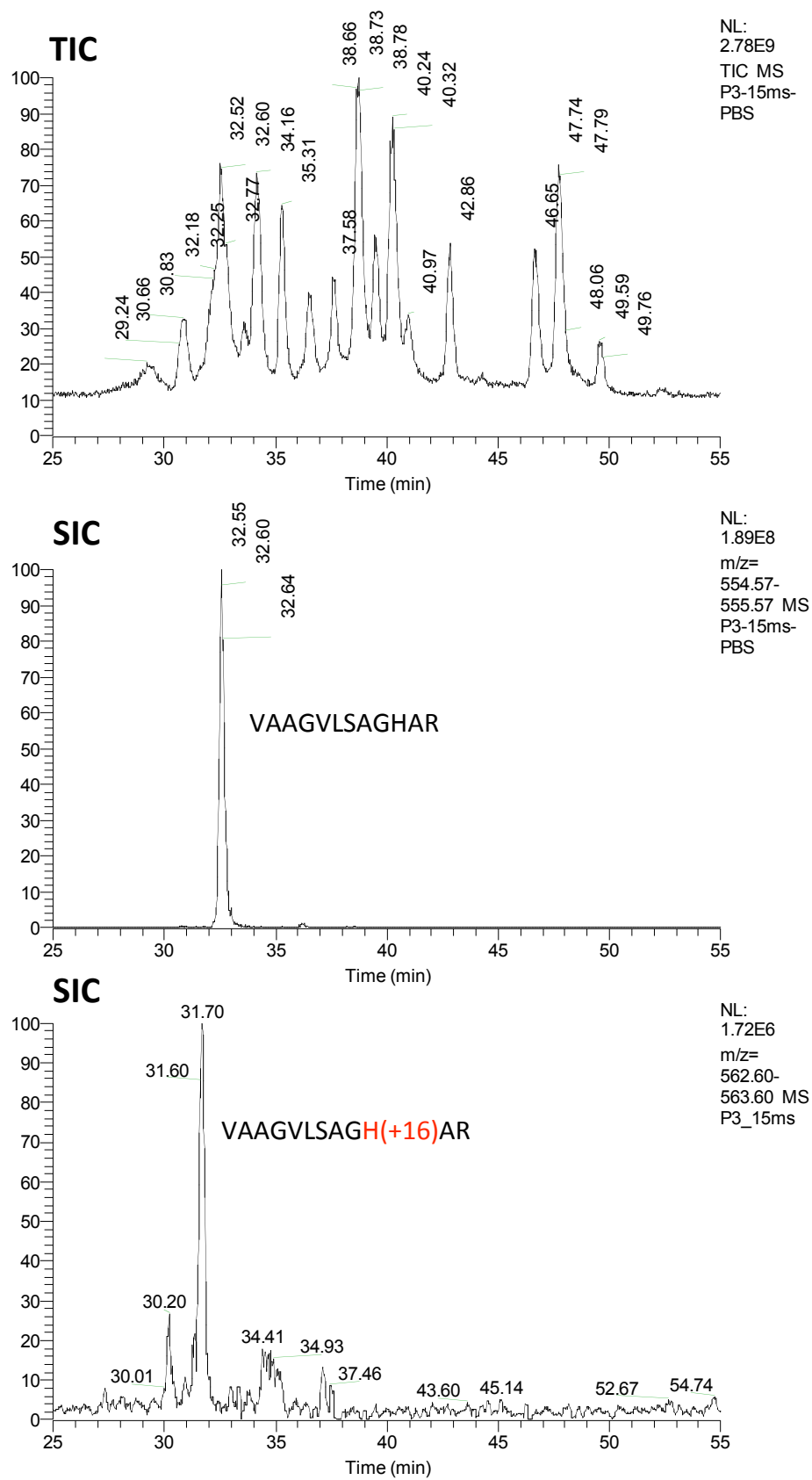
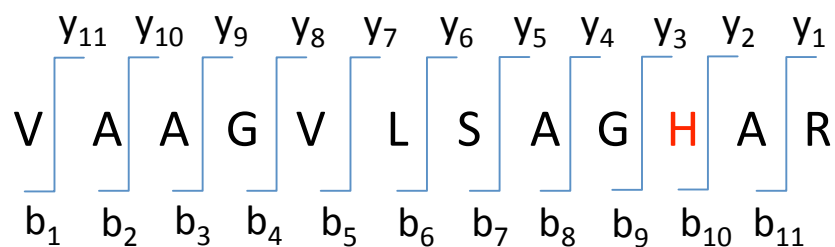
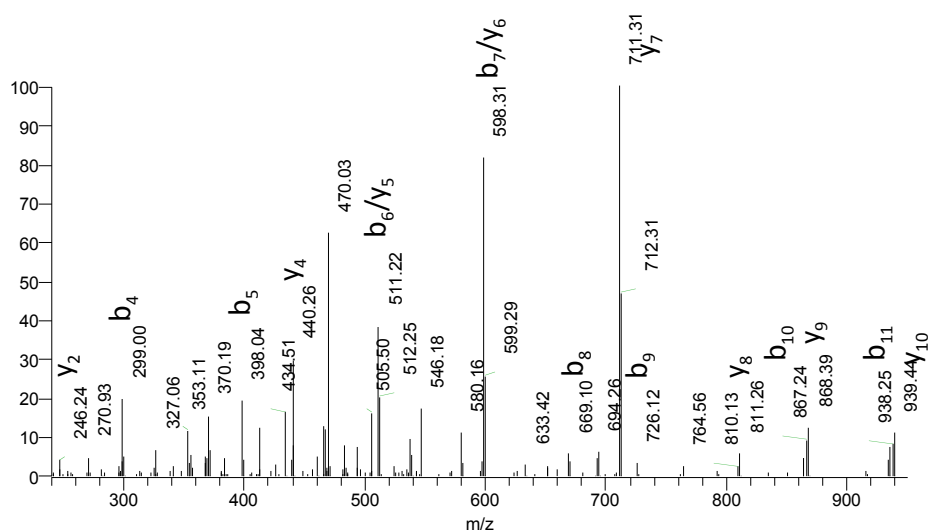


Figure S2.



VAAGVLSAGHAR



VAAGVLSAGH(+16)AR

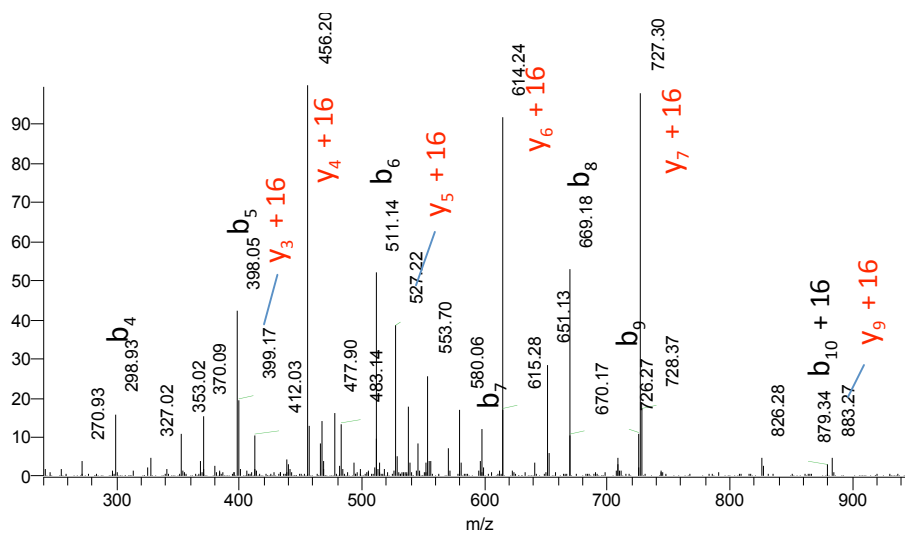


Figure S3.

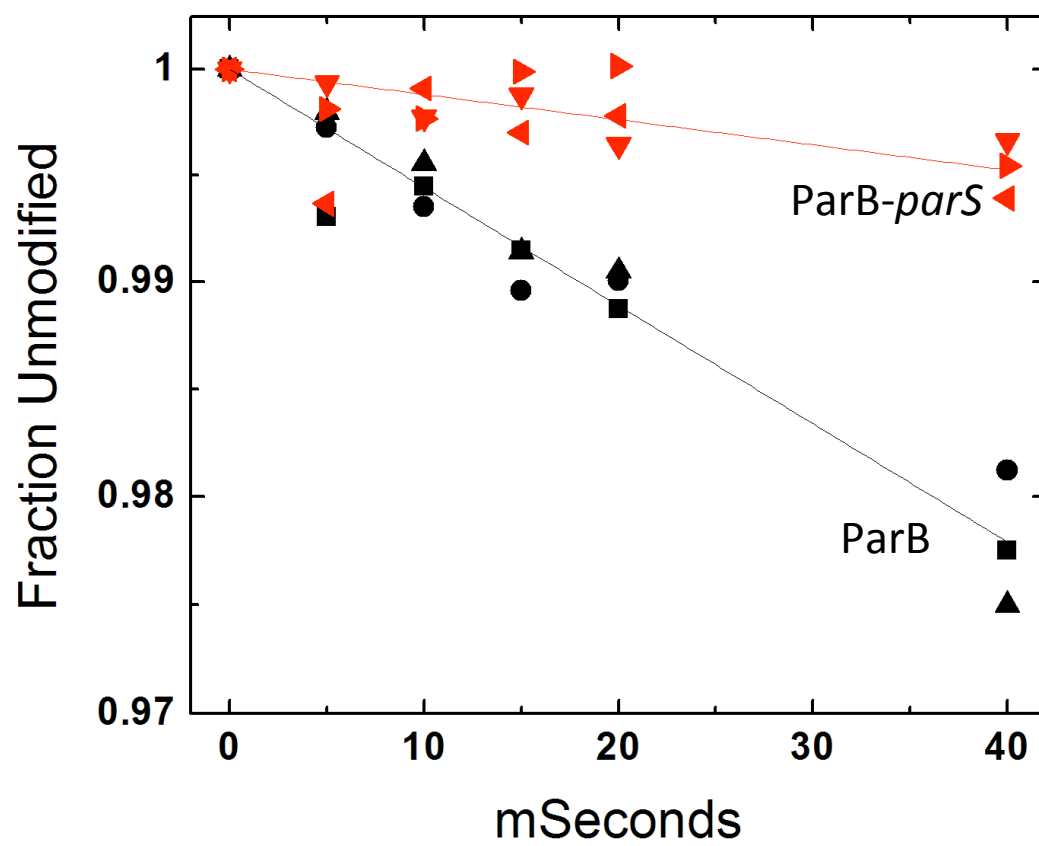


Figure S4.

Figure S2. The top panel shows a typical total ion current chromatogram (TIC) of a peptic digest of X-ray irradiated (15ms) ParB, represented by software XCalibur 2.01 (Thermo Electron). The middle panel shows the selected ion chromatogram (SIC) for the native (unmodified) and modified (+16 m/z shift, single peak) peptide 211-222 and their relative abundance after 15ms of X-ray irradiation.

Figure S3. MS/MS fragmentation of native and modified peptide 211-222, which shows +16 modification at H220.

Figure S4. A representative dose response plot for peptide, 211-222 VAAGVLSAGH(+16)AR. The fraction of unmodified peptide at any exposure is calculated by the following equation:

$$\text{Fraction Unmodified} = \frac{\text{Unmodified peak area}}{(\text{Unmodified peak area} + \text{Modified peak area})}$$

The peak areas are calculated from the selected ion chromatogram. A time evolution of SICs are used to calculate fraction unmodified vs. exposure time. Hydroxyl radical modification rate constants are determined from the single exponential curve fitting as described in Experimental Methods.