Fluorescence-based Thermal Shift Assay (TSA)

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PALOMA VARELA[1]

[1]Institute for Integrative Biology of the Cell, CEA, CNRS, Université Paris-Saclay, Gif-sur-Yvette, France

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PALOMA VARELA

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PROTOCOL STATUS

Working

MATERIALS TEXT

Thermal stability of fully purified $ZitR_{MG}$ protein was calculated by induced thermal denaturation using a Q-PCR ABI Prism® 7900HT (CTPF platform, IMAGIF). We used Sypro Orange dye, which non-specifically binds to hydrophobic surfaces and can be measured at 488 nm. Purified $ZitR_{MG}$ protein (2.5 mg) was analyzed under different conditions of buffer (Tris, phosphate, HEPES and MES), pH (ranging from 9.2 to 4), salt (NaCl up to 350 mM), and glycerol concentrations (up to 10 %) (data not shown).