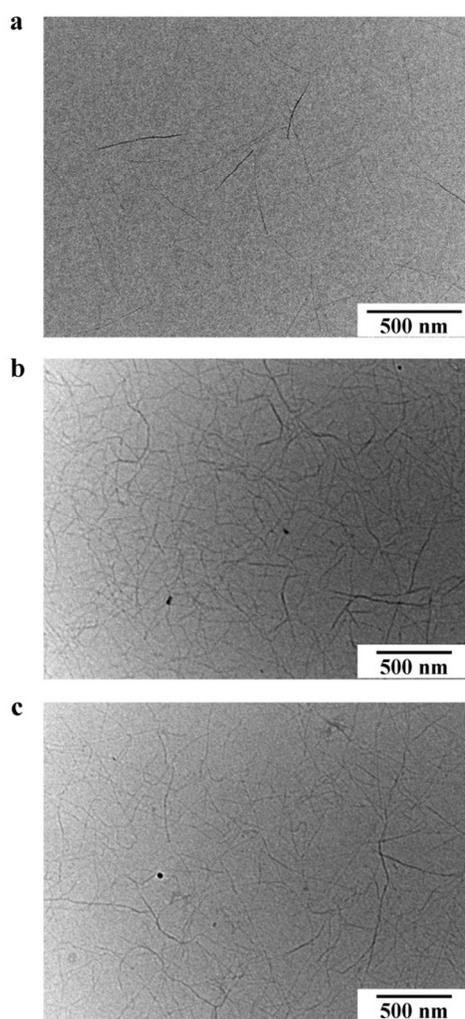


# A colloidal description of intermolecular interactions driving fibril-fibril aggregation of a model amphiphilic peptide

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## Transmission Electron Microscopy

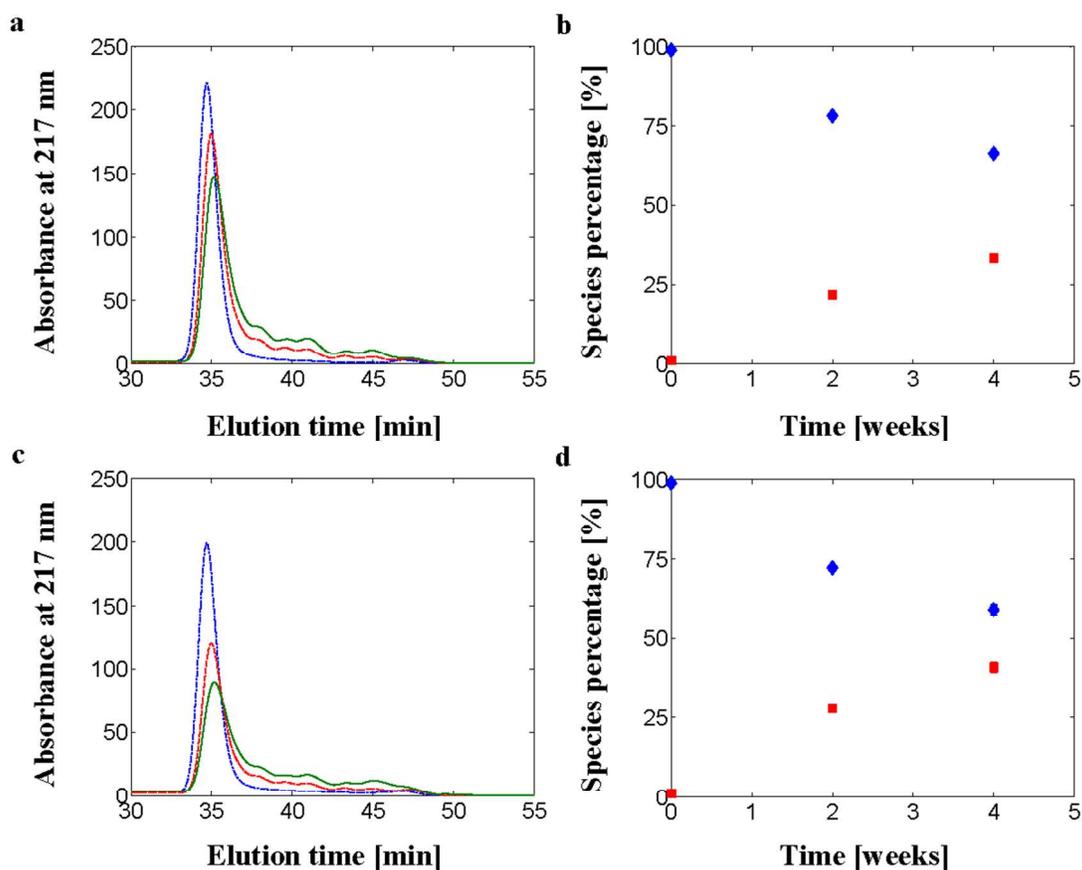
Peptide samples were recorded by using a FEI Morgagni 268 microscope. Peptide solutions were diluted to a final concentration of about 0.05 g/L, loaded on a carbon grid (Quantifoil, Jena, Germany), washed and stained with a 2% uranyl acetate solution.



**Figure S1. TEM pictures of RADA 16-I fibrils at 1 g/L in the presence of NaCl in 10 mM HCl at pH 2.0.** Representative TEM pictures of freshly prepared solution of RADA 16-I peptide (a) and aggregated solution of RADA 16-I peptide in the presence of 25 mM NaCl (b) and 100 mM NaCl (c).

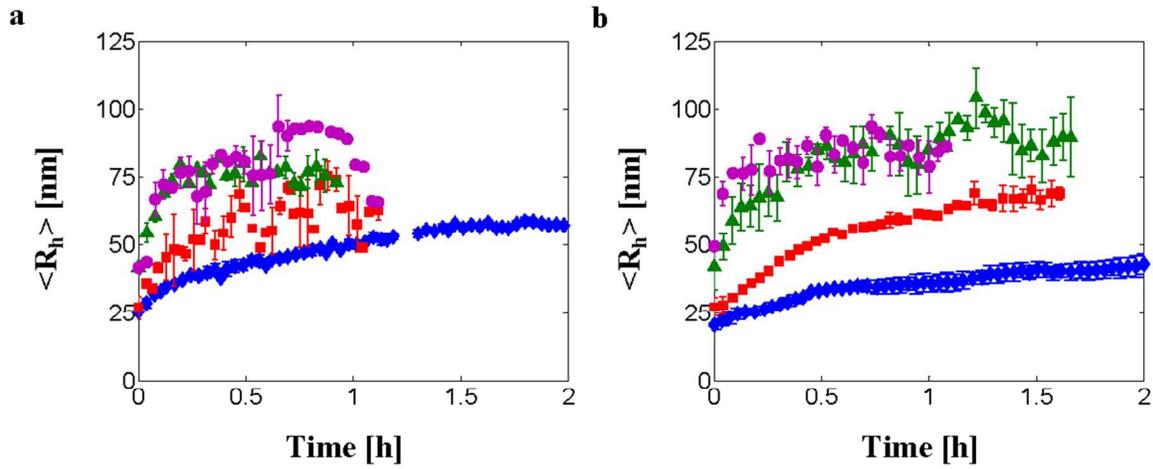
## Size Exclusion Chromatography

Size exclusion chromatography analysis was performed using a Superdex Peptide 10/300 GL, 10 mm×300 mm size-exclusion column (GE Healthcare, Uppsala, Sweden) mounted on a Agilent 1100 series HPLC unit (Santa Clara, CA, USA) consisting of an isocratic pump with degasser, an autosampler, a column oven, and a DAD detector. Each sample was eluted for 70 min at a constant flow rate of 0.4 mL/min using 10 mM hydrochloric acid at pH 2.0 as mobile phase. The fractionated samples were detected by UV absorbance at 217 nm.

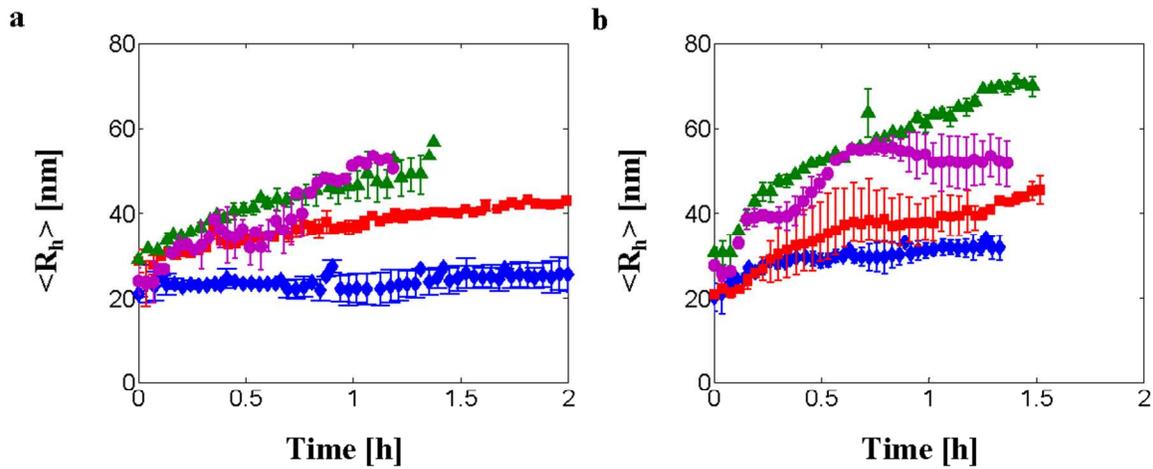


**Figure S2. The time evolution of monomer conversion in the presence of NaCl in 10 mM HCl at pH 2.0.** SEC chromatograms at zero time (blue, dash-dot line), after 2 weeks (red, dashed line) and 4 weeks (green, continuous line) of incubation in the presence of 25mM (a) and 75mM (c) NaCl. The time evolution of the percentage of the residual monomer (◆) and peptide fragments eluting at longer elution times (■) in the presence of 25 mM (b) and 75 mM (d) NaCl.

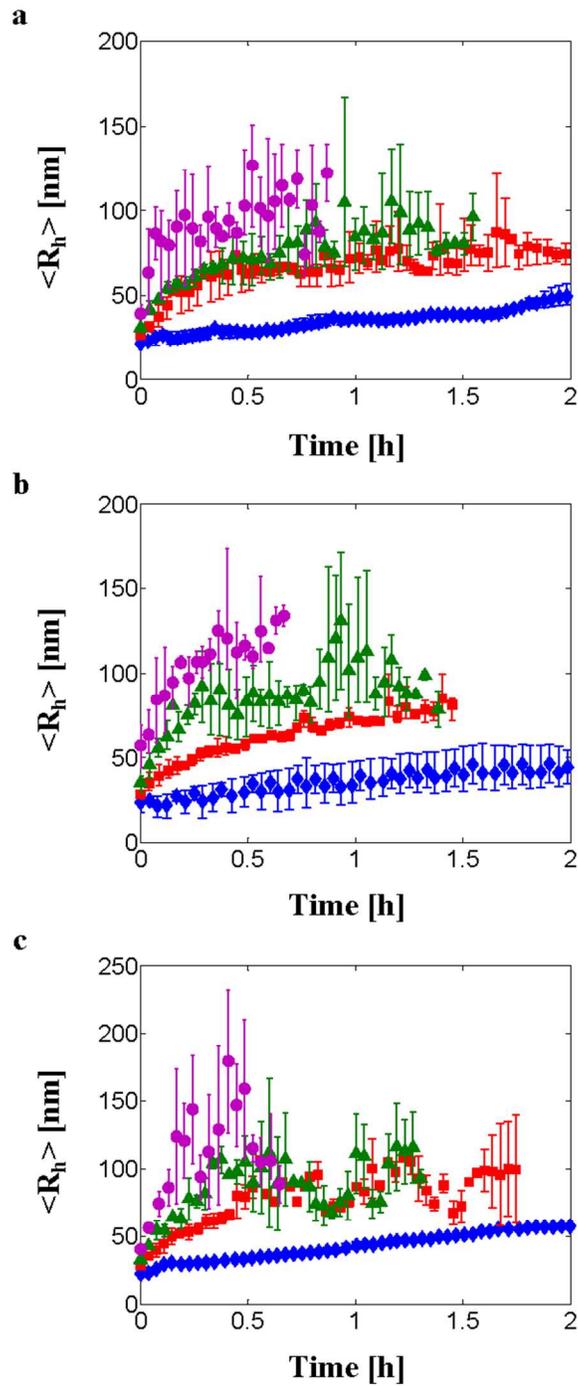
## Dynamic Light Scattering



**Figure S3. The anion effect on the aggregation of RADA 16-I peptide at 1 g/L in 10 mM HCl at pH 2.0.** The time evolution of hydrodynamic radius in the presence of NaNO<sub>3</sub> (a) and NaH<sub>2</sub>PO<sub>4</sub> (b) at ionic strength equal to 25 mM (◆), 50 mM (■), 75 mM (▲), 100 mM (●).



**Figure S4. The cation effect on the aggregation of RADA 16-I peptide at 1 g/L in 10 mM HCl at pH 2.0.** The time evolution of hydrodynamic radius in the presence of CaCl<sub>2</sub> at equal ionic strength (a) and CaCl<sub>2</sub> at equal concentration of Cl<sup>-</sup> ions (b) at ionic strength equal to 25 mM (◆), 50 mM (■), 75 mM (▲), 100 mM (●).



**Figure S5. The solvent effect on the aggregation of RADA 16-I peptide at 1 g/L in 10 mM HCl at pH 2.0.** The time evolution of hydrodynamic radius in the presence of 10% isopropanol (a), 10% ethanol (b) and 20% ethanol (c) in the presence of 25 mM ( $\diamond$ ), 50 mM ( $\blacksquare$ ), 75 mM ( $\blacktriangle$ ), 100 mM ( $\bullet$ ) NaCl.