SEC-SAXS studies of double-stranded RNAs from trypanosome RNA editing

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SAXS studies of double-stranded RNAs are hindered by the presence of aggregates and the frequent presence of intermolecular interactions. Size exclusion chromatography SAXS can remove aggregated RNA prior to exposure of the RNA to the X-ray beam and thereby permit SAXS studies that would otherwise be difficult. We present several successful SEC-SAXS studies of double-stranded RNAs from trypanosome RNA editing that were done at APS BIOCAT ID-18 and SSRL BL 4-2. We also discuss the practical aspects of these studies and the remaining challenges.