Imaging virus assemblies with in situ cryoEM

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With cryo-electron microscopy, structures of purified proteins and protein complexes can be routinely determined to near-atomic resolution using single particle analysis (cryoEM-SPA) method. Structures of macromolecular assemblies that are intrinsically flexible and dynamic, and often function in higher-order assemblies that are difficult to purify, have recently been analyzed to near near-atomic resolution using cryo-electron tomography and sub-tomogram averaging (cryoEM-STA). The study of these complexes and assemblies in situ using cryoEM-STA, coupled with cryoFIB and correlative and integrative imaging, opens a new frontier in structural cell biology. I will present technology development in cryoFIB and cryoEM-STA, and recent studies of the virus assembly process within native cells to demonstrate the power sub-tomogram averaging for in situ structure determination.