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in a pseudo-intracellular space, which forms a branched meshwork structure inside the cell

([@bib10]). If one assumes a simple flux of

multivesicular bodies in and out of the lysosome, the observed influx and efflux of autophagosome lumens with vesicular structures indicates that the

majority of intraluminal vesicles/vesicles and vesiculogenomes are formed inside the

lysosomes. Formation of these vesicles inside the lysosomes could be restricted to the perinuclear space. This might help to explain the observed

enrichment of multivesicular bodies at the perinuclear space ([@bib36]). Our analysis of the vesicular structures in dynamic optical sections throughout cell death supports this hypothesis.

The ability of autophagosomes to deliver their content to the lysosomes depends on the fusion of autophagosomes with lysosomes. The only other

example of fusion of autophagosome with any cellular component is the fusion of autophagosomes with ER membranes, which is also an important process for the degradation of ER cargo ([@bib16; @bib41]). A study on the dynamic behavior of autophagosomes during mitosis showed that lysosomal activity is most strongly reduced after mitosis but remains relatively constant through the G1 phase of the next cell cycle. Fusion of autophagosomes with the ER could result in their redistribution into the perinuclear space ([@bib8]). This statement is in agreement with the results of our live cell imaging study ([Fig. S2](#app2){ref-type="sec"}). During mitosis there was a pronounced increase in the number of autophagic structures in the perinuclear region of the cell as well as a decrease in the number of autophagosomes in the peripheral cytoplasm. This suggested that autophagosomes fuse with the ER before mitosis. During the G1 phase of the next cell cycle

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Waves Complete - Complete in this bundle: VST, RTAS, TDM plugins, from 6 of our stellar processors: FFSim, RetroFilter, 29 2 PDF Bundle Digital Audio Workstation (DAW) - The EAW plugin is a component containing all of the most important .Apoptosis induced by 2-chloro-2'-deoxyadenosine in human glioma cells is mediated through the Akt pathway. 2-chloro-2'-deoxyadenosine (CdA) is a key drug in the treatment of human malignancies, including glioblastoma. The goal of this study was to investigate whether CdA induces apoptosis in human glioma cells and if Akt plays a role in this

process. CdA induced apoptosis in the human glioma cell line U-251MG at low concentrations (6.25 to 12.5 microM), and the most sensitive concentration was 7.5 microM. The apoptotic process was characterized by internucleosomal DNA fragmentation and a decrease in mitochondrial transmembrane potential. As apoptosis induction was correlated with the activation of caspases-3 and-9, flow cytometry experiments detected a significant increase in apoptotic cells after 5 microM CdA treatment, and Western blot assays demonstrated a time-dependent increase in the level of pro-apoptotic Bax protein and a decrease in the anti-apoptotic Bcl-2 protein. In addition, Akt activation was detected by Western blot analysis in U-251MG cells after 5 microM CdA treatment. Furthermore, inhibition of Akt by transfection with the anti-sense Akt oligonucleotide resulted in a decrease of the CdA-induced apoptosis. These results demonstrate that CdA exerts an effect on mitochondrial apoptosis in glioma cells, and suggest that Akt plays an important role in CdA-induced apoptosis.

Multifunctional n-HA-coated Ag-NPs: Chitosan and gelatin as biocompatible stabilizers. In this work, n-hydroxyapatite (n-HA) was used as a capping agent for silver

nanoparticles (n-HA-Ag). At first, n-HA and chitosan (CS) were separately prepared by sol-gel method. Then, a water soluble and biocompatible