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Lecture 11: Assessing nanotoxicity at the single cell level

- 11.1 Need for single cell measures of nanotoxicity
 - 11.1.1 There is more than one way for a cell to die...
 - 11.1.2 "Necrosis" vs. "Apoptosis"
 - 11.1.3 There are other forms of "toxicity"
 - 11.1.4 Some other challenges in measuring toxicity of nanomaterials
- 11.2 Necrosis vs. Apoptosis mechanisms
 - 11.2.1 Necrosis is unplanned "cell injury"
 - 11.2.2 Apoptosis is planned "programmed cell death"
 - 11.2.3 Why it is important to distinguish between necrosis and apoptosis?
- 11.3 Single-cell assays for necrosis and apoptosis
 - 11.3.1 Dye exclusion assays for necrosis
 - 11.3.2 TUNEL assays for late apoptosis
 - 11.3.3 Annexin V assays for early apoptosis
 - 11.3.4 COMET assays for DNA damage and repair
 - 11.3.5 Light scatter assays
 - 11.3.6 Dihydroethidium assays for oxidative stress
- 11.4 Nanotoxicity in vivo some additional challenges
 - 11.4.1 Single-cell nanotoxicity, plus biodistribution measuring challenges....
 - 11.4.2 Accumulations and agglomerations of nanoparticles can change toxicity locally to tissues and organs
 - 11.4.3 Filtration issues of nanoparticles size matters toxicity to kidney, liver and lung
 - 11.4.4 Functional sensitivity of heart and brain to nanotoxicity largely unknown

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