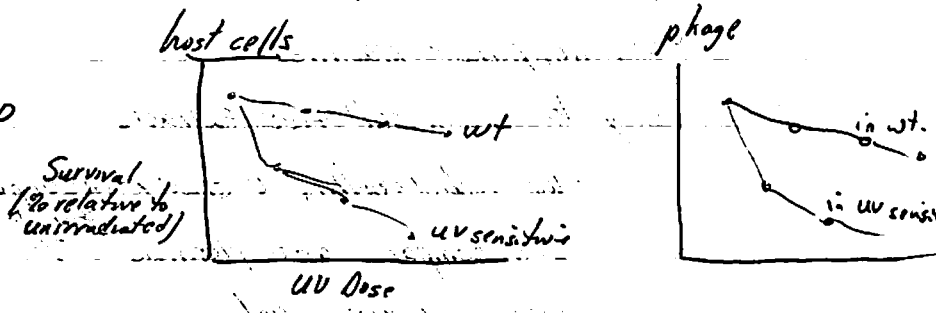


# Ann Groneman - Use of Viruses & Plasmids to Study DNA Repair

Why - because you can manipulate DNA

## Host cell reactivation

© Ellison et al 1960



- wild type strain repairs bacteriophage DNA

## Phage vs. Plasmids

© Phage have own mechanism to insert DNA

© Irradiate  $\phi$ X174 DNA

DS - much host cell reactivation } suggests that most host cell reactivation  
 SS - no host cell reactivation } due to excision repair

© Irrad. plasmid DNA

(a) recA mutations = ~~host cell recA~~ same hcr as wild type  
 -  $\therefore$  recA dependent repair NOT important here

(b) insert into human cells (use CAT selection)

- XPA + O show lower CAT activity than "normal"

- take plasmid w/ many  $^{14}C$ 's
- allow turnover
- assay plasmid for C/A

Why

- ① can damage the DNA w/o affecting repair system
- ② plasmids/viral genomes are small
- ③ can manipulate plasmids
- ④ easy to use in cell-free extracts R. Wood
  - use radiolabelling to study incorporation
  - can compare diff. plasmids <sup>on joint</sup> w/ UV light
  - run on agarose gel
  - more radioactivity in UV irradiated plasmids
  - can compare plasmids with specific lesions