

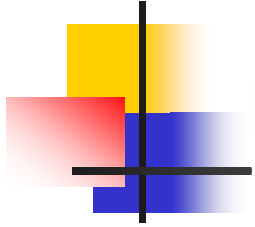
Radiobiology in Brachytherapy

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Overview

- Need
- History
- Key factors & their effects
- RB models
- Combination & utilities
- Future
- Conclusion



Need

- For changing the fractionation schedules
- Change of dose rate systems (LDR/HDR)
- Gap correction
- Combining EBRT with Brachytherapy
- Choosing the right isotope
- Comparison of data between centres



- Experimental- Ram testis

- Stranquist Curves 1944

- NSD –Ellis 1969

- Orton TDF Brachytherapy 1974

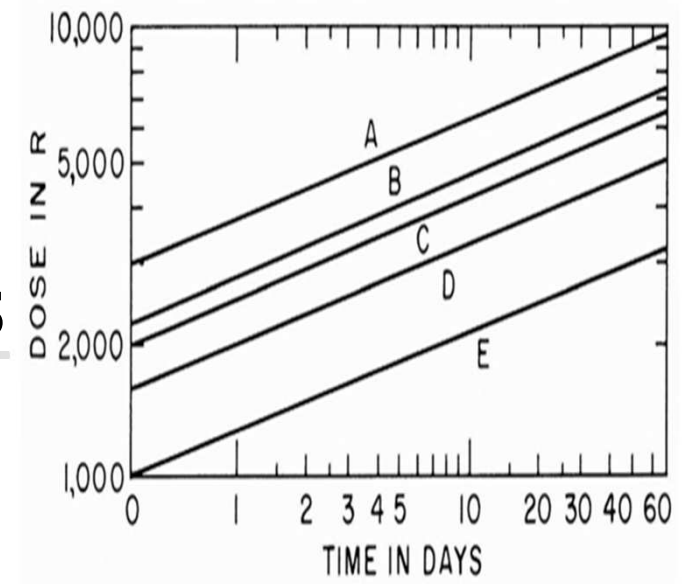
- Elkind kind of repair

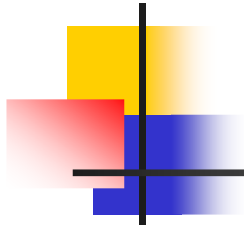
- Inter Fr time, Dose/Fr, Dose Rate

- Cumulative Response Dose(CRE)

- Kirk et al

- Tumor Significant Dose (TSD)





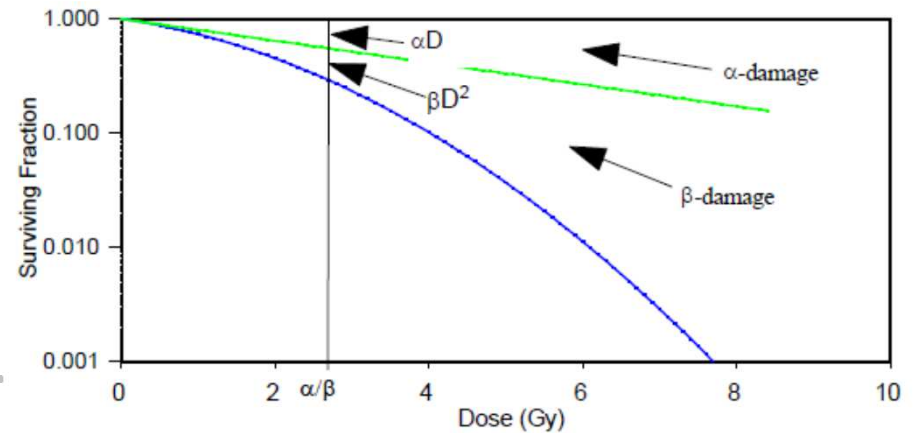
- It does not take all complex biological process
- Doubts on validity of NSD relation with tissue type
- Doubts on validity of NSD relation with different effect in same tissue type
- Range of number of fractions. The formula is provided
- Concern on the time factor taken.



Biological Models

- Linear Quadratic Model
 - Lea & Catcheside
 - Various modifications
- BED : Biological Equivalent dose
- EUD : Equivalent Uniform Dose

L Q model



- It quantitatively predicts the dose/fraction dependence. The principal determinants are α & β
- α – Linear portion in Cell Survival Curve.
 - Occurs along a single ionizing tract
 - Tumor cells (rapid proliferation rates & short duration for repair)
- β - Quadratic portion in Cell Survival Curve
 - Occurs along a two different ionising tracts
 - Normal cells (coordinated repair & hence requiring double hit)
- α/β ratio
 - Dose where there is proportionate of cell death due to linear & quadratic portions



Interpretation of α/β

- High α/β ratio

For a particular dose of radiation either the

- α DNA injury is higher

or

- β DNA injury is lower

- Low α/β ratio

For a particular dose of radiation either the

- α DNA injury is lower

or

- β DNA injury is Higher



Pros & Cons

Low α/β means

↓ in dose/fraction

less injury to normal tissue

High α/β means

↑ in dose/fraction

More injury to normal tissue

- Limitations

Fractionated Rx delivered @ regular interval period (once in 24hrs) & 5Fr/Wk.

Gap in Rx in pt **NOT** considered



Biological Effective Dose (BED)

- Concept used to compare the effectiveness of cell killing by different fractionation regimen by using LQ Model
- Use
 - Intercomparison of various RT schedules
 - Intercomparison of different types of radiotherapy
- Formula
 - = Total Physical dose [D] x Relative effectiveness [RE]



Factors considered in BED

- Physical
 - Dose
 - Dose/fr
 - Inter fraction interval .
- Radiobiology
 - α/β
 - Repair rate
 - T pot
 - Repopulation & Redistribution .
 - Overall treatment time



BED contd . . .

- BED differs for different normal tissue & also for different tumor biology
- BED is represented as numerical value of dose with suffix indicating the α/β value.
- Eg: 100 BED₃, 65 BED₁₀
- Relative Effective factor
= Phy Factor + RB Factor

Repopulation



- RE for repopulation when taken into consideration uses subtractive repopulation correction factor w.r.t repopulation rate and Rx time.
- $BED = D \times RE - RCF$ (repopulation correction factor)
- $RCF = K (T - T_{\text{delay}})$
 T_{delay} is delay time after beginning of treatment before the repopulation rate becomes significant.

Eg: 28 days for HNSCC



Equivalence

- Each treatment – specific type of biologic effect in N & T tissue
- 2 treatment if they produce the same effect then it is called Equivalent.
- Equivalence does not mean that both treatments produce the same amount of biological damage in all the irradiated structures; rather it means that both produce the same differential pattern of damage.
- In BT the source geometry is important;
 - Small difference between the 2 treatments
 - Results in change to the physical dose distribution
 - Which itself will render equivalence impossible.



Equivalent Uniform dose

- Formula suggested by Niemerko
- Modified for IMRT
- Available in many TPS
- Different weightage
 - Target
 - Normal structure (Parallel or serial)

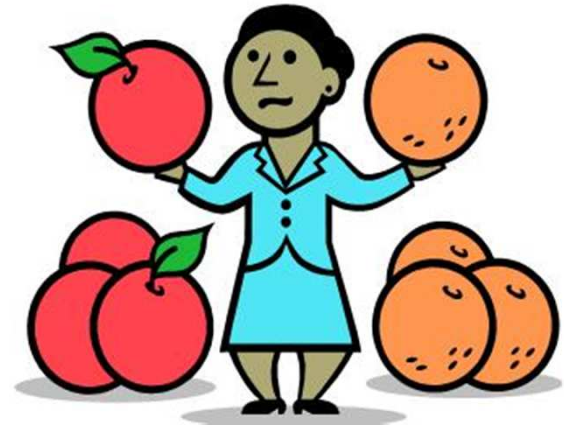


Combination & Utilities

- EBRT vs Brachytherapy
- LDR vs HDR
- Volume & anatomical site
- Tumor shrinkage
- Reirradiation
- EBRT+ BT combination



Differences



EBRT

- Large Volume
- Homogeneous
- -5% to +7% acceptable
- Small dose, protracted time (weeks)
- Full repair

Brachytherapy

- Isodose encircling a small target volume
- Very heterogenous
- High dose, short treatment (hours to days)
- Short interval (HDR), Continuously (LDR)

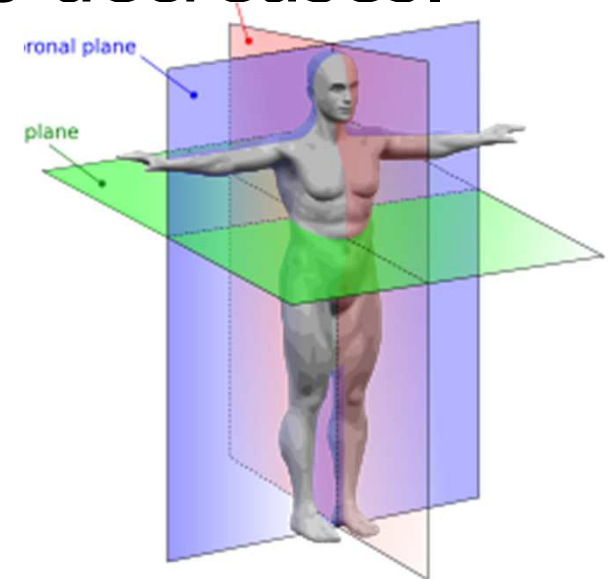
LDR vs HDR



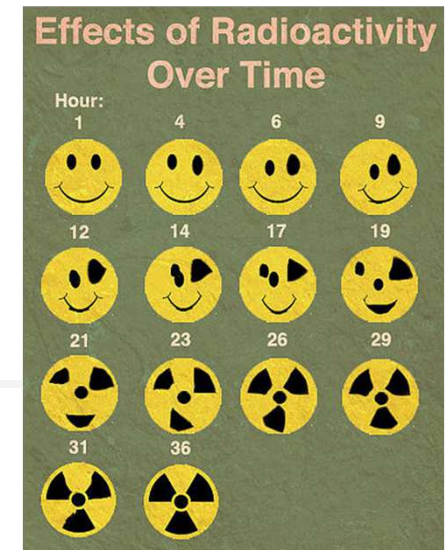
- Several Trials comparing LDR Vs HDR
 - Historical data
- Most cases similar results
- HDR beneficial with equivalent normal tissue tolerance & the tumoricidal doses
- Severe Complications
 - 3.44% (>7 Gy)& 1.28% (<7 Gy)

Volume, Anatomical site

- The Dose reqd increases with size of tumor
- As Dose increases tolerance of late responding normal tissues decreases.



Reirradiation



- High doses can be delivered to previously irradiated area
- Can be tolerated if delivered to a limited volume. recovery seems to be less in some like CNS
- No clarity regarding minimal interval between two irradiations.

Combined with EBRT



- Because BEDs are additive the determination of biological effects associated with combined modality treatments is straightforward.
- Total BED
$$= \text{BED}_{(\text{EBRT})} + \text{BED}_{(\text{Brachy})} - \text{RCF}$$
RCF (repopulation correction factor) is reqd only for tumor calculations and should be calculated using the overall treatment time of the combined treatments.
- In Brachytherapy calculation allowance for dose gradient effect. Should also be considered.

Future



- Genetherapy
 - Arrest Accelerated repopulation
- Commercially TPS incorporating biological models
 - Bioplan
 - Orbit
 - LQ Survivor





Conclusion

- Quantification of CLDR & FHDR
- Quantification of dose rate effect
- Quantification Permanent implants
- Quantification of PDR Brachytherapy
- Treatment intercomparison
- Designing new Fr & Rx schedules
- Calculating dose equivalence used with different isodose



