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melanoma and lung and breast cancer cells from patients with xeroderma pigmentosum \[\[@R41]-\[@R44]\], and a striking example for a tumor cell line derived from a patient with Fanconi's anemia (the HS578T breast cancer cell line) \[\[@R45]\]. Also, re-analysis of the above mentioned studies suggested that there might be other examples where differences between the NER phenotypes of various cancer cells from patients with similar DNA repair defects would enable cross-comparison of their NER efficiencies. Finally, the observations obtained for cell lines from patients with DNA repair defect of XPA-BRCA1 heterozygotes \[\[@R46]\] and XP-DDB2 patients \[\[@R47]\] were consistent with our original hypothesis that cancer cell lines might display a wider range of NER phenotypes than their non-transformed counterparts, likely reflecting stochastic changes during immortalization of normal human somatic cells or the accumulation of secondary mutations in cancer cell lines c6a93da74d

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