# 0.1 The biological process

## 0.2 The simulation framework

#### 0.2.1 The simulation domain

A UV beam is shot vertically thorough the nucleus and considered to affect all chromatin in its path similarly. The simulation in therefore placed in a 2-dimensional domain. This is in-line with the experimental signal values, reported as values of a maximal z-projection [REF].

### 0.2.2 The polymer model

We use a cross-linked Gaussian chain [REF] of N monomers connected by harmonic springs to represent a coarse-grained model of the chromatin. Springs connecting adjacent monomers in the linear chain are assigned a minimal length  $L_0$ . Cross-linking are added randomly between pairs of monomers by a harmonic spring with minimal length zero. Cross-linking measure is given by the percentage  $\alpha$  of non-nearest-neighbor monomers connected of the N monomers of the chain. In each realization of the simulation, a random set of non-nearest neighbor monomers is chosen for cross-linking, according to the value set for  $\alpha$ . The cross-linked polymer is simulated up to relaxation time, at which time the UV beam is shot.

#### 0.2.3 UV irradiation

At the end of relaxation steps, UV beam focal point is set to the polymer's center-of-mass. The damage region (DR) is represented by a 2-dimensional fixed circular region of area  $A_0$  centered at laser's focal point. For each UV dose u, damages caused by UV are homogeneously distributed in  $A_0$  [REF] among the polymer's monomers. Damaged monomers are chosen randomly, such that the average number of damages in  $A_0$  increases as  $k_t u$ , with  $k_t$  in units of bp/msec.

#### Affect of UV irradiation on the polymer, the repair stage

To simulate crowding of repair proteins around damaged sites, a circular exclusion region is centered at the location of each damaged monomer. Ex-

clusion region is represented by an elastic spring pushing force of radius  $r_p$ , originating from each damaged monomer. The elastic force applied on any monomer within the exclusion range is thus oriented outwards. In addition to the exclusion region, all cross-links from and to damaged monomers are removed.

The system will evolve into a new steady configuration which represents the chromatin 15 minutes post UV-C. At which point, the region of interest (ROI) is defined as the circle containing 95% of the damaged monomers and centered at their center-of-mass. The ROI remains a fixed region used to track the number of damaged and undamaged monomers within it. Measurements are done off-line, such that the ROI is always centered at the center-of-mass of the monomers known to be damaged.

### 0.2.4 Post repair stage

As damaged monomers are repaired the exclusion region is removed from damaged monomers. Cross-links are reintroduced gradually according to the spatial distance between monomers. The amount of cross-links re-introduces is such that the initial cross-linking percentage  $\alpha$  is restored.