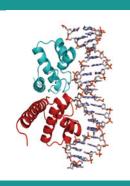
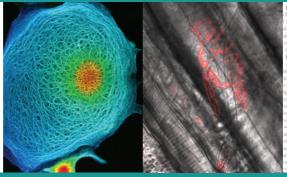
Biophysics at All Scales



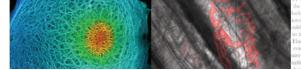


It is interesting to note that if one had a specific marker for the GTP cap. Nevertheless, one could measure how the GTP bulin density decreases as one moves away from the tip of the icrocubule. Fitting that density to an exponential curve, on ould be able to measure the characteristic length 2 and compan to the prediction of our model shown in Fig. 11. Fluctuations in the length of the microtubule cap at the + end evident over time as shown in Fig. 12 for two free tubulin imer concentrations, d=2 and d=5. The sharp decreases in ngth occur when a single GTP-tubulin dimer takes a long time hydrolyse to GDP dimer. For example, between the arrows in ig. 12 the hydrolysis time is 30 s compared to the average GTI drolysis time of 3 s. The decreased length depicts the new ngth of the cap which is from the next closest GTP-tubulit mer to the base of the crown. Moreover we can evaluate the ugth of the cap and this averages approximately 60 dimers for =2 and 300 dimers for d=5. Another way of evaluating the ze of the cap is to count the number of GTP-tubulin dimer er the 13 protofila- ments. These data are shown in Fig. 1: rther supporting the variations in cap length seen in Fig. 14

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