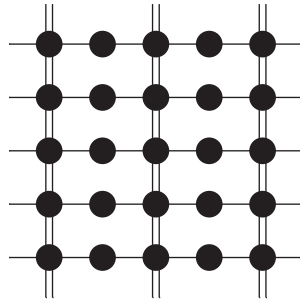
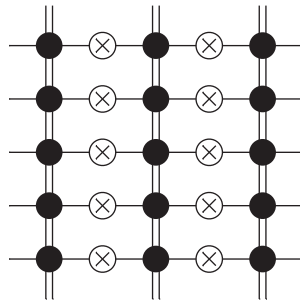


The second procedure's step is to approximate the Hamiltonian by “sliding” half of the bond interactions in the \hat{y} direction to the other half as shown in the figure below. In practice: we neglect half of the vertical bond interactions, while the other half becomes *twice as strong*.



The third procedure's step is to sum over the spins which are interacting *only in the horizontal direction* as shown in the following figure.



Together the three procedure's steps form our initial RG step.

Question 1: Write down the effect of an RG step to the couplings: $\{K'_x, K'_y\} = RG_1[\{K_x, K_y\}]$.

From the answer to the first question it should be clear that it doesn't make sense to iterate this single RG step.

Question 2: Can you explain qualitatively why?

Now consider another RG step which we call $\{K'_x, K'_y\} = RG_2[\{K_x, K_y\}]$ which is of the same form but has x and y switched (so it is rotated by $\pi/2$).

Question 3: Consider the nested steps $\{K'_x, K'_y\} = RG_2[RG_1[\{K_x, K_y\}]]$. What are the fixed points for K_x and K_y ? (Do it numerically.) Using either fixed point estimate the exponent α as done in class with scaling methods. Using Josephson's hyperscaling identity estimate the exponent ν too.

Optional question 4: Are these estimates better or worse than those obtained with the method shown in class? Can you give some qualitative argument why?