

Case with more significant corrections

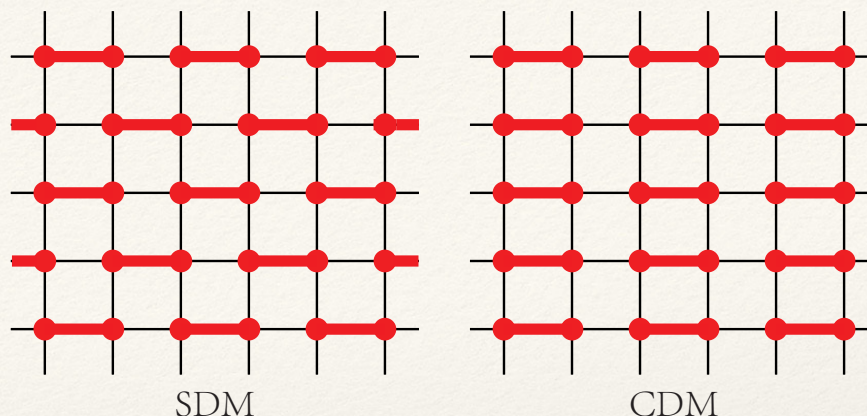
- common at quantum critical points

$S=1/2$ Heisenberg model with

- columnar dimers (CDM)
- staggered dimers (SDM)

The SDM has been controversial

- $O(3)$ or new universality class
- strange scaling behaviors



PHYSICAL REVIEW LETTERS **121**, 117202 (2018)

Anomalous Quantum-Critical Scaling Corrections in Two-Dimensional Antiferromagnets

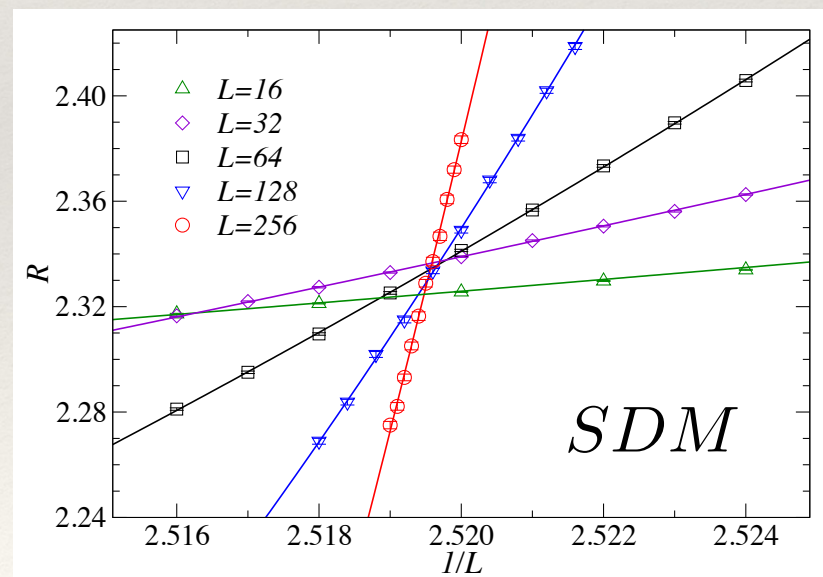
Nysen Ma,^{1,2,3} Phillip Weinberg,³ Hui Shao,^{4,3} Wenan Guo,^{5,4} Dao-Xin Yao,^{1,*} and Anders W. Sandvik^{3,2,†}

Analyze critical behavior with two scaling corrections taken into account

$$O(g, L) = f[(g - g_c)L^{1/\nu}, \lambda_1 L^{-\omega_1}, \lambda_2 L^{-\omega_2}, \dots]$$

Taylor expand, analyze crossing points for different dimensionless quantities

Compare CDM and SDM behaviors

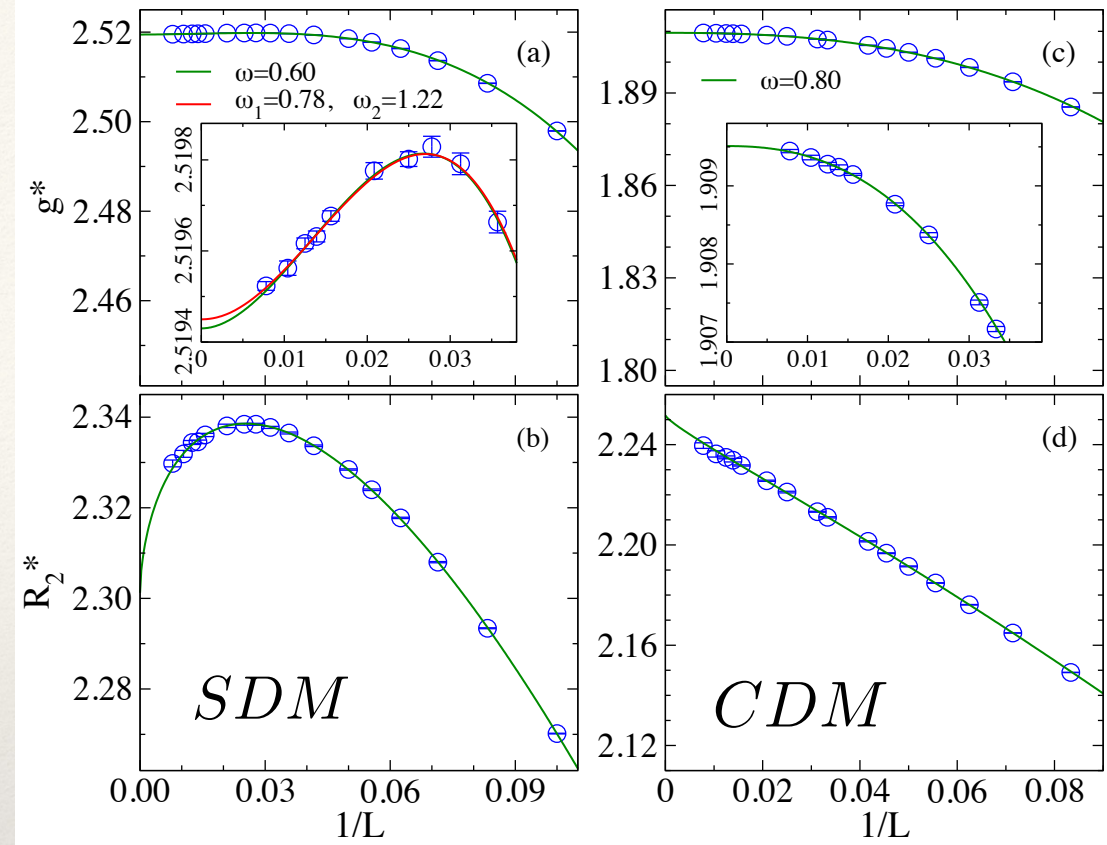
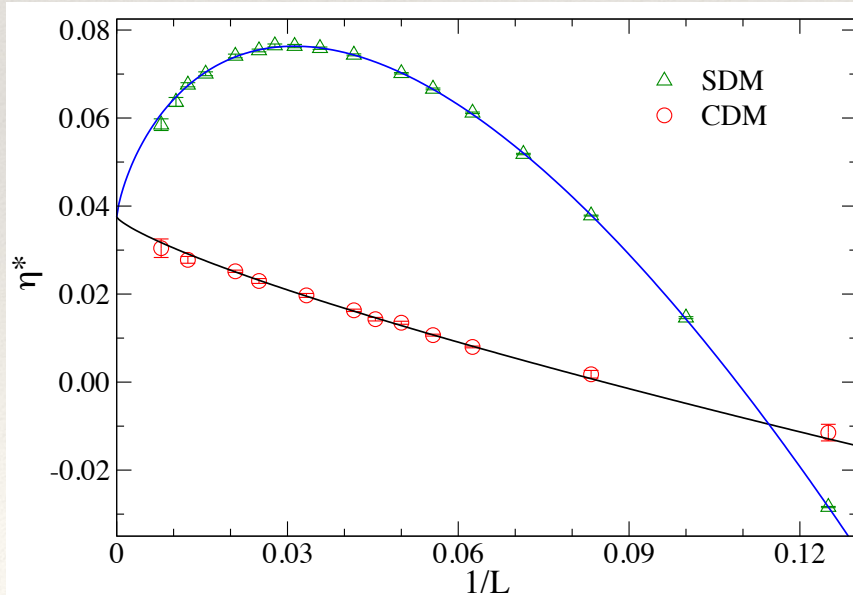


Leading-order cross-point shifts

$$g^*(L) = g_c + aL^{-\omega_1 - 1/\nu},$$

$$O^*(L) = O_c + bL^{-\omega_1},$$

- Works for CDM, $\omega_1 \approx 0.78$
- Two corrections needed for SDM $\omega_1 \approx 0.78, \omega_2 \approx 1.25$
- Fits within theory where the SDM field theory needs a new term (Fritz et al, PRB 2012)



Order parameter at the critical point

$$\langle m^2 \rangle_c \propto L^{-(1+\eta)} (1 + b_1 L^{-\omega_1} + b_2 L^{-\omega_2} + \dots)$$

$$\eta^*(L) = \ln[\langle m^2(L) \rangle_c / \langle m^2(2L) \rangle_c] / \ln(2) - 1$$

$$\eta^*(L) = \eta + c_1 L^{-\omega_1} + c_2 L^{-\omega_2} + \dots$$

Field-theory aspects of the anomalous scaling [Fritz et al., PRB 2011]

The SDM leads to a cubic term in the field-theory action

$$\mathcal{S} = \frac{1}{2} \int d^D r [m_0 \varphi_\alpha^2 + (\vec{\nabla} \varphi_\alpha)^2] + \frac{u_0}{4!} \int d^D r (\varphi_\alpha^2)^2 + i\gamma_0 \int d^D r \vec{\varphi} \cdot (\partial_x \vec{\varphi} \times \partial_y \vec{\varphi})$$

Is the cubic term relevant or irrelevant at the O(3) critical point?

$$C(\vec{r}) = \langle \mathcal{O}(\vec{r}) \mathcal{O}(0) \rangle \propto \frac{1}{|\vec{r}|^{2\Delta_{\mathcal{O}}}} \quad \mathcal{O}(\vec{r}) = \vec{\varphi}(\vec{r}) \cdot (\partial_x \vec{\varphi}(\vec{r}) \times \partial_y \vec{\varphi}(\vec{r}))$$

MC of the classical Heisenberg model to extract scaling dimension

$$H = -J \sum_{\langle ij \rangle} \vec{S}_i \cdot \vec{S}_j$$

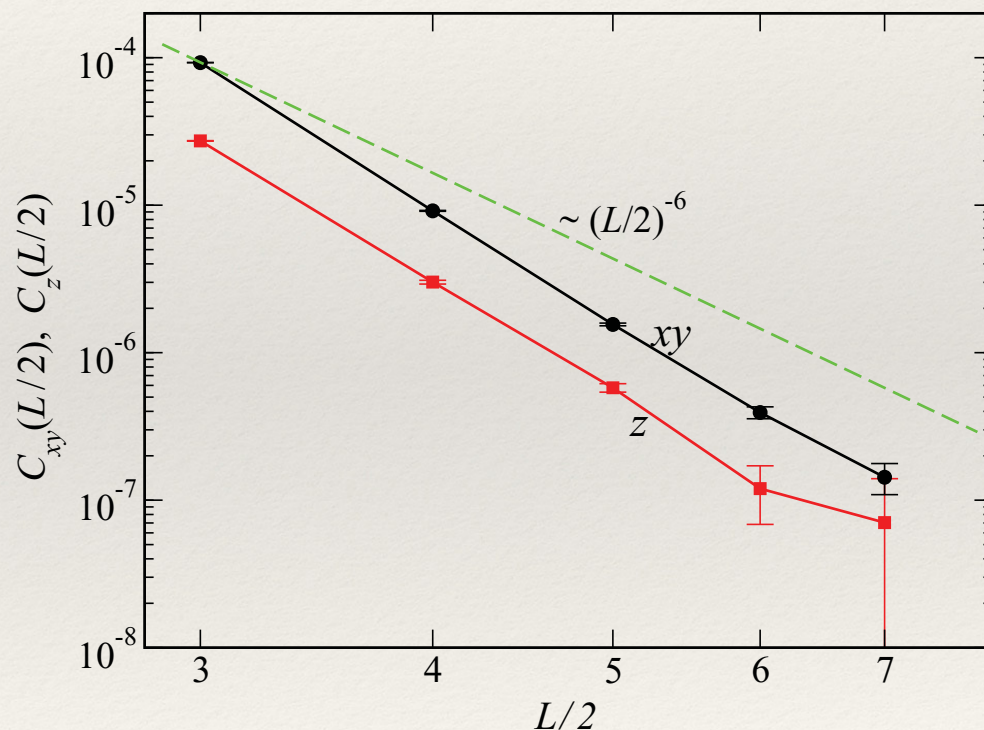
$$\mathcal{O}_{i,\text{lattice}} = \vec{S}_i \cdot (\vec{S}_{i+e_x} \times \vec{S}_{i+e_y})$$

They found $\Delta_o \sim 3.2 \dots 3.5$

- corresponds to $\omega \sim 0.2 - 0.5$
- would be leading correction

Inconsistent with SDM QMC

- $\omega \sim 1.3$
- second correction



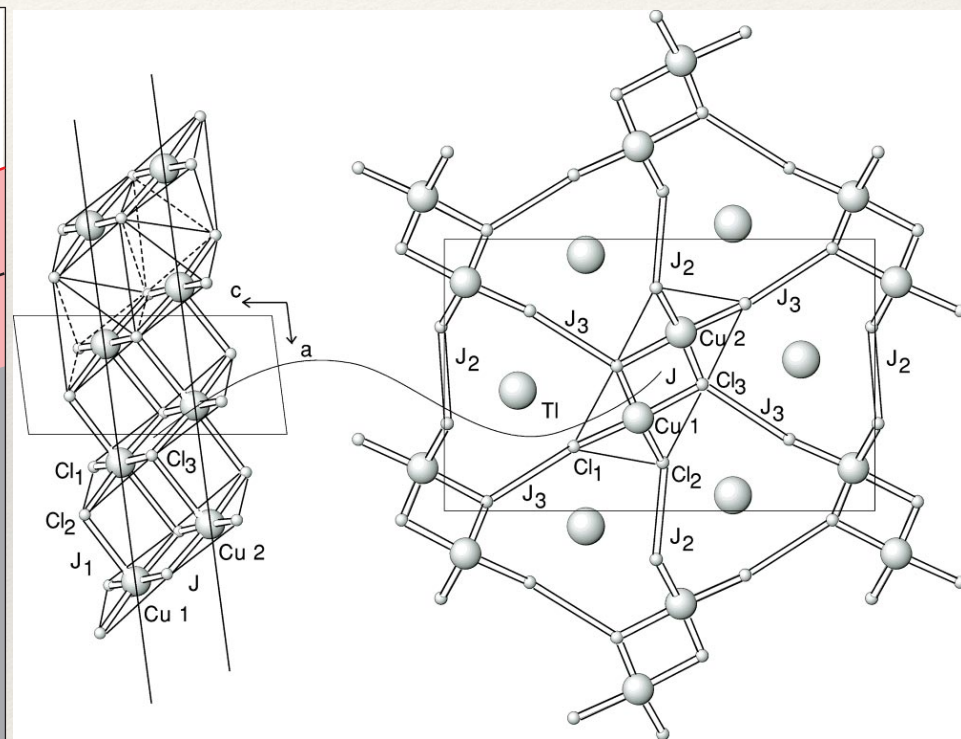
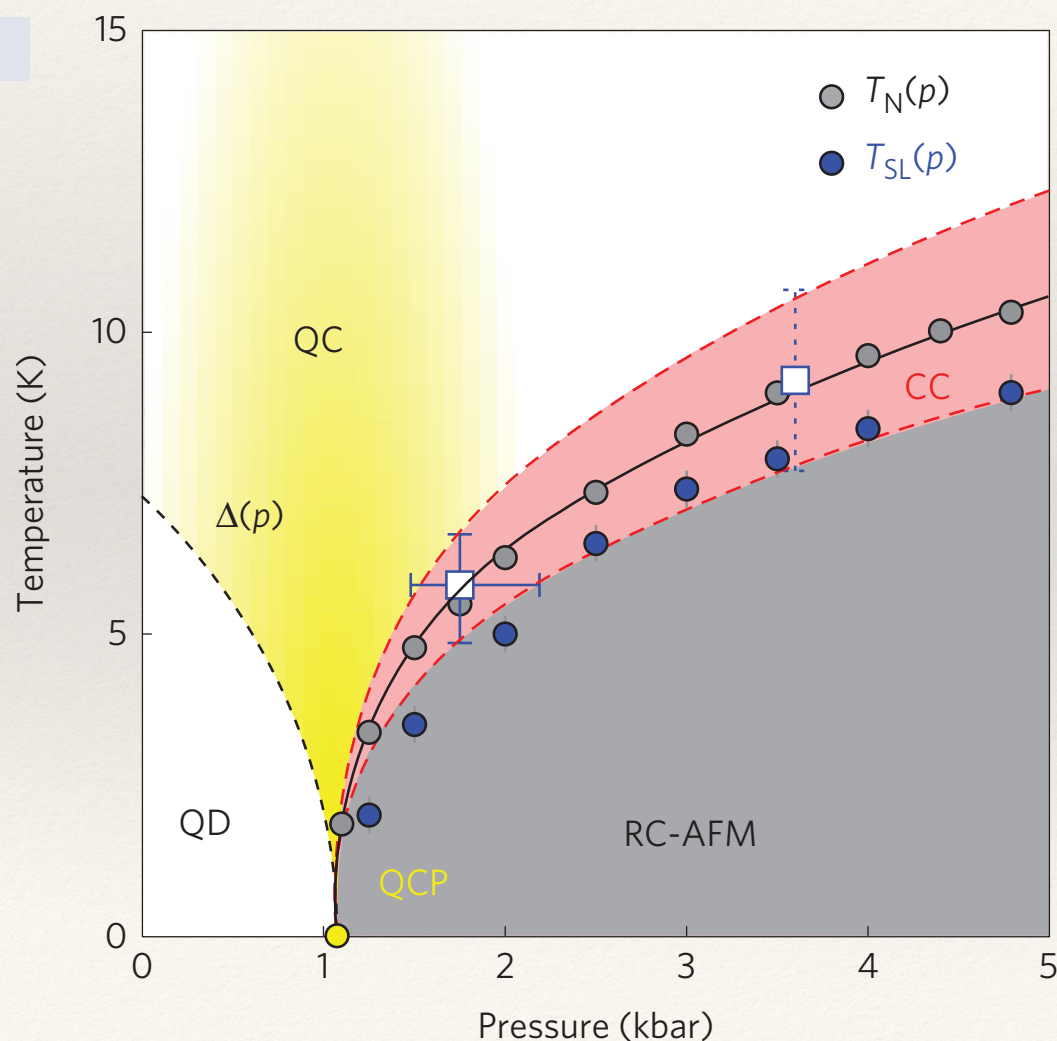
Conformal bootstrap method: There is an operator with $\omega \sim 1.4$

- improved classical MC would be useful

Quantum and classical criticality in a dimerized quantum antiferromagnet

P. Merchant¹, B. Normand², K. W. Krämer³, M. Boehm⁴, D. F. McMorrow¹ and Ch. Rüegg^{1,5,6*}

3D Network of dimers
- couplings can be changed by pressure



From: M Matsumoto, B Normand, TM Rice, M Sigrist, PRB (2004)

Universality of the Neel temperature in 3D dimerized systems?

[S. Jin, AWS, PRB2012]

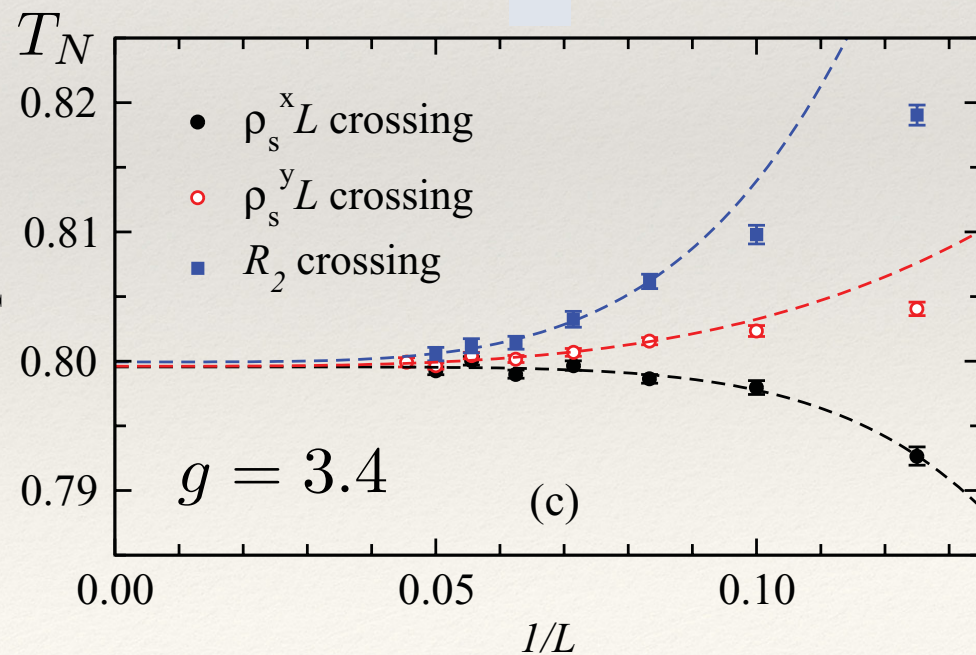
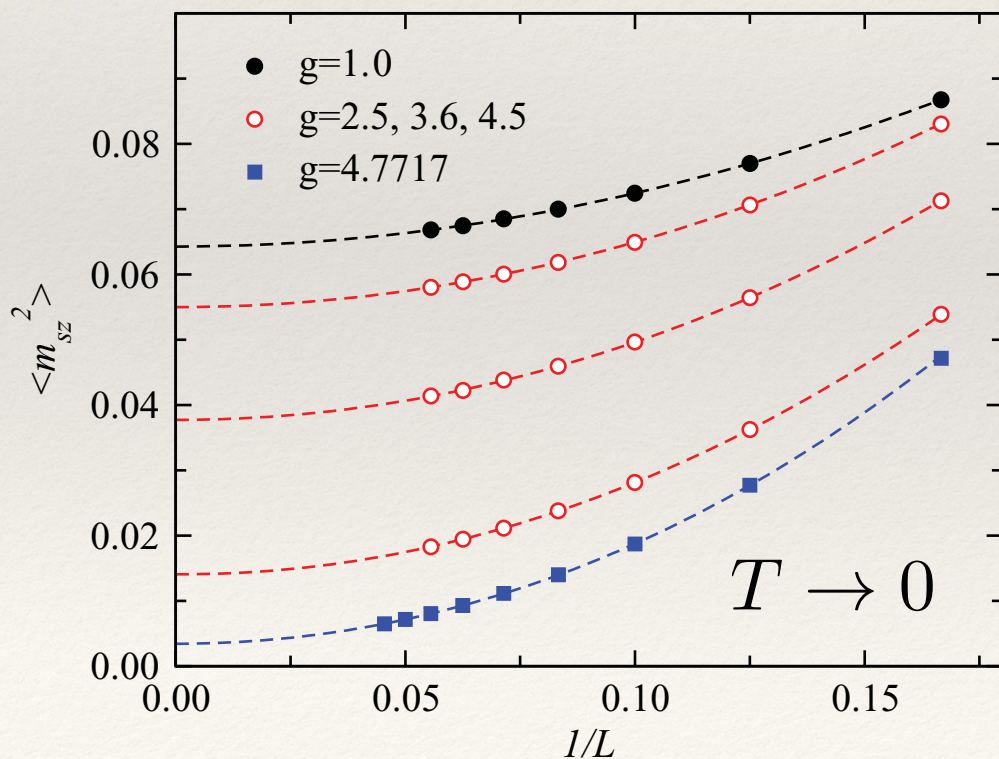
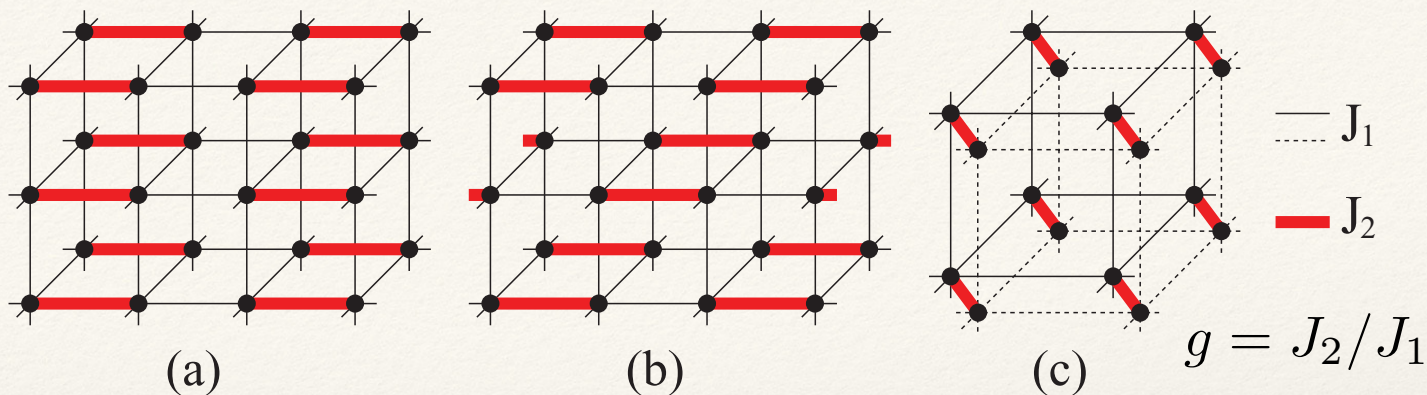
Determine the Neel ordering temperature

T_N and the $T=0$ ordered moment

m_s for 3 different dimerization

patterns

Example: Columnar dimers

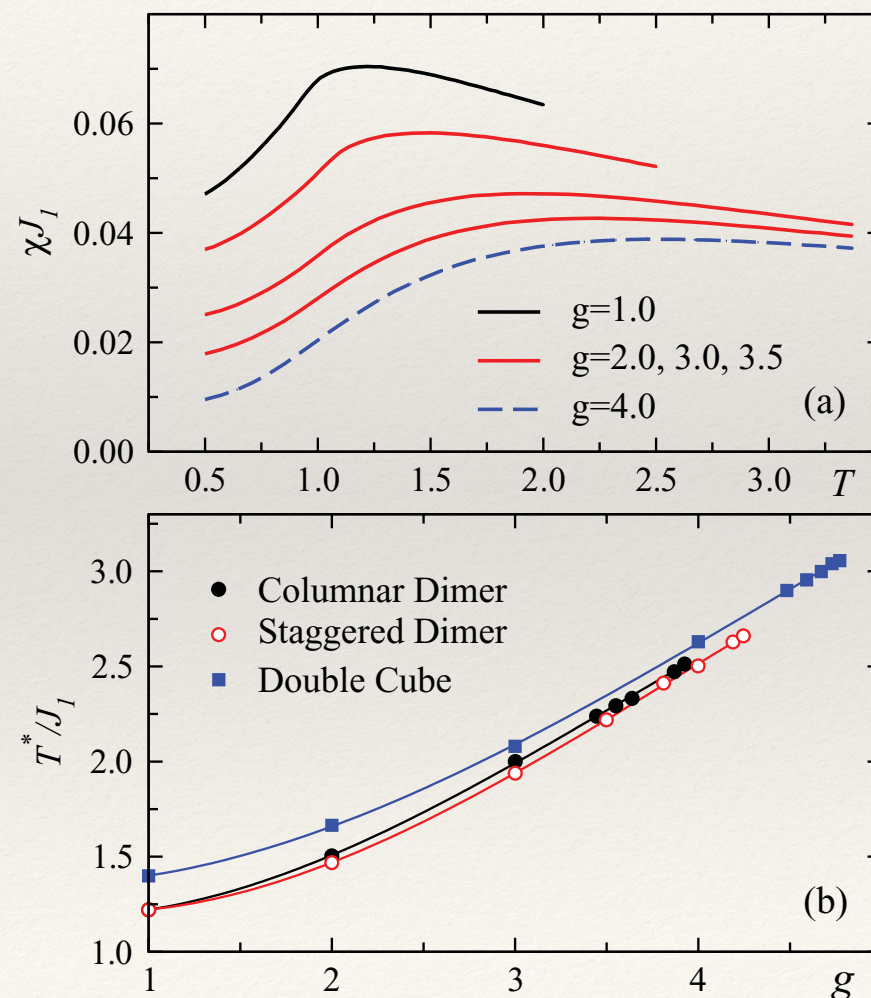
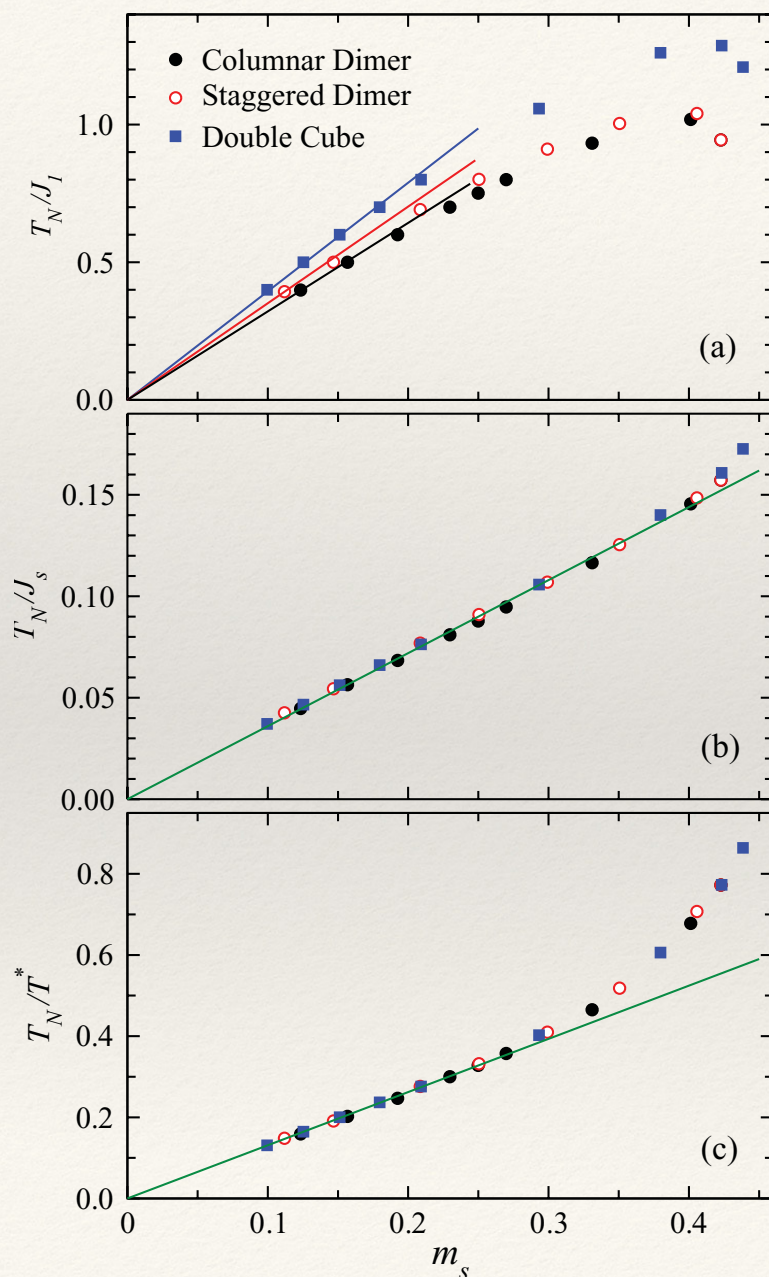


Couplings vs pressure not known experimentally

- plot T_N vs m_s to avoid this issue and study universality
- but how to normalize T_N ?

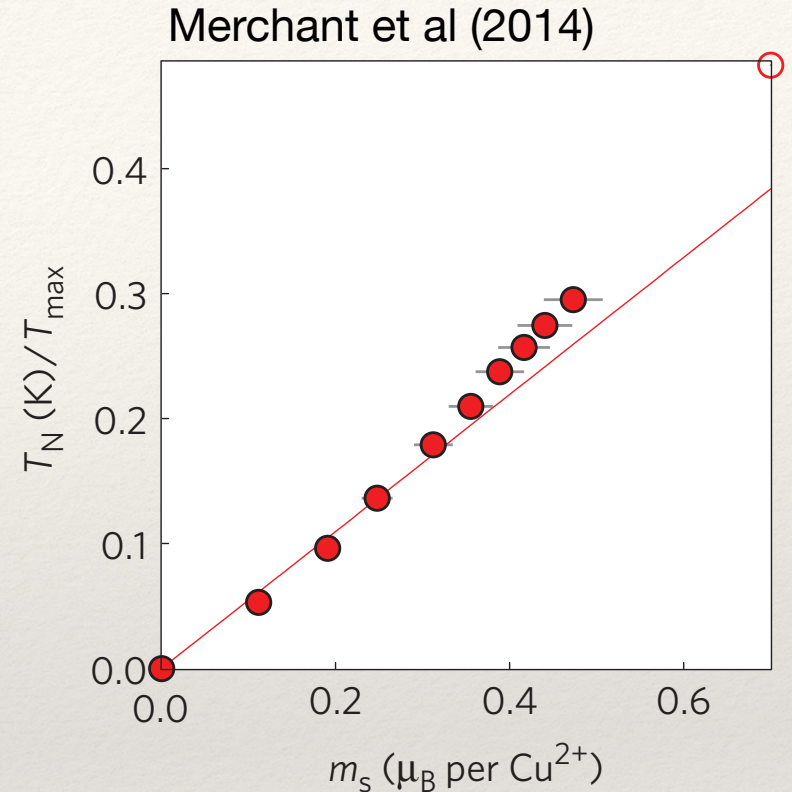
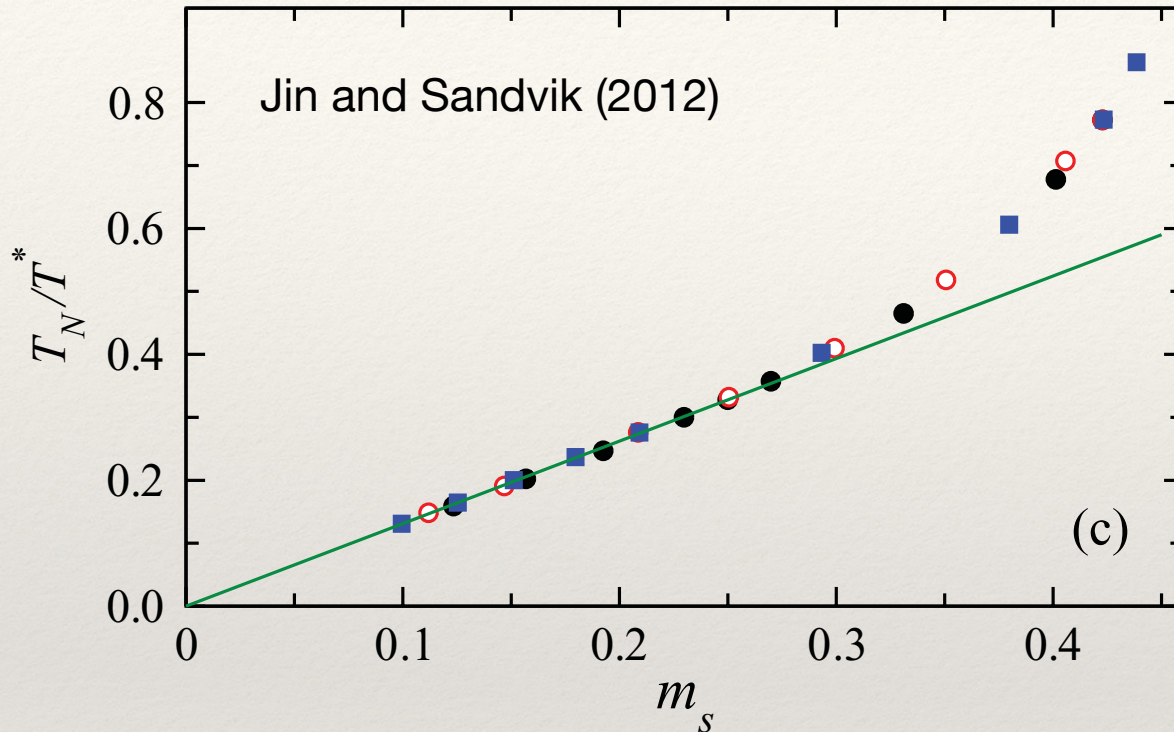
Three normalizations

- weaker coupling J_1
- sum J_s of couplings per spin
- peak T^* of magnetic susceptibility



T* normalization is accessible experimentally

- some experimental susc. results available
- neutron data analyzed with this normalization



Same features observed in models and experiment

- experimental slope about 25% lower if g-factor =2 assumed (what exactly is the g-factor?)

More recent works to study log corrections, dynamics,....

Qin, Normand, Sandvik, Meng, PRB 2015, PRL 2017

Why the linear form $T_N = am_s$ ($a=\text{constant}$)?

The ordered state can be qualitatively described by mean field theory:

$$H = \sum_{ij} J_{ij} \vec{S}_i \cdot \vec{S}_j$$

$$H_0 = \left(\sum_i J_{i0} \right) \langle \vec{m}_s \rangle \cdot \vec{S}_0$$

The order parameter m_s is reduced from its maximum value by two effects

- quantum fluctuations
- thermal fluctuations

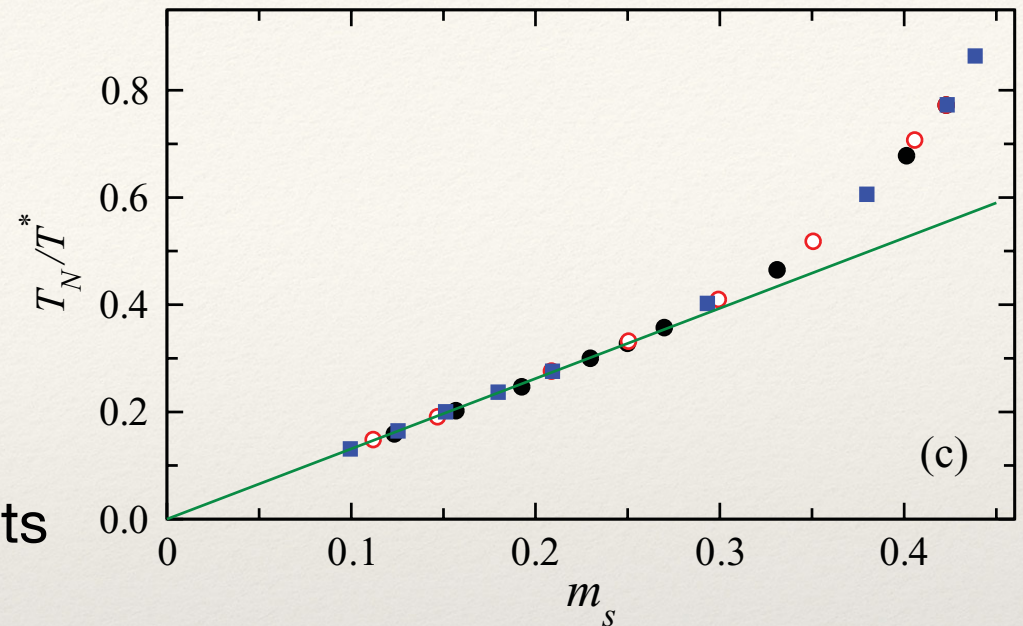
Assume that these fluctuations decouple:

$$\langle m_s \rangle = m_0(T=0) f(T)$$

Then we have an effective mean-field coupling $J_{\text{eff}} \propto m_0$

In mean-field theory: $T_c \propto J_{\text{eff}} \rightarrow T_c \propto m_0$

The violations of the linear form indicate the temperature where quantum and thermal fluctuations cannot be decoupled



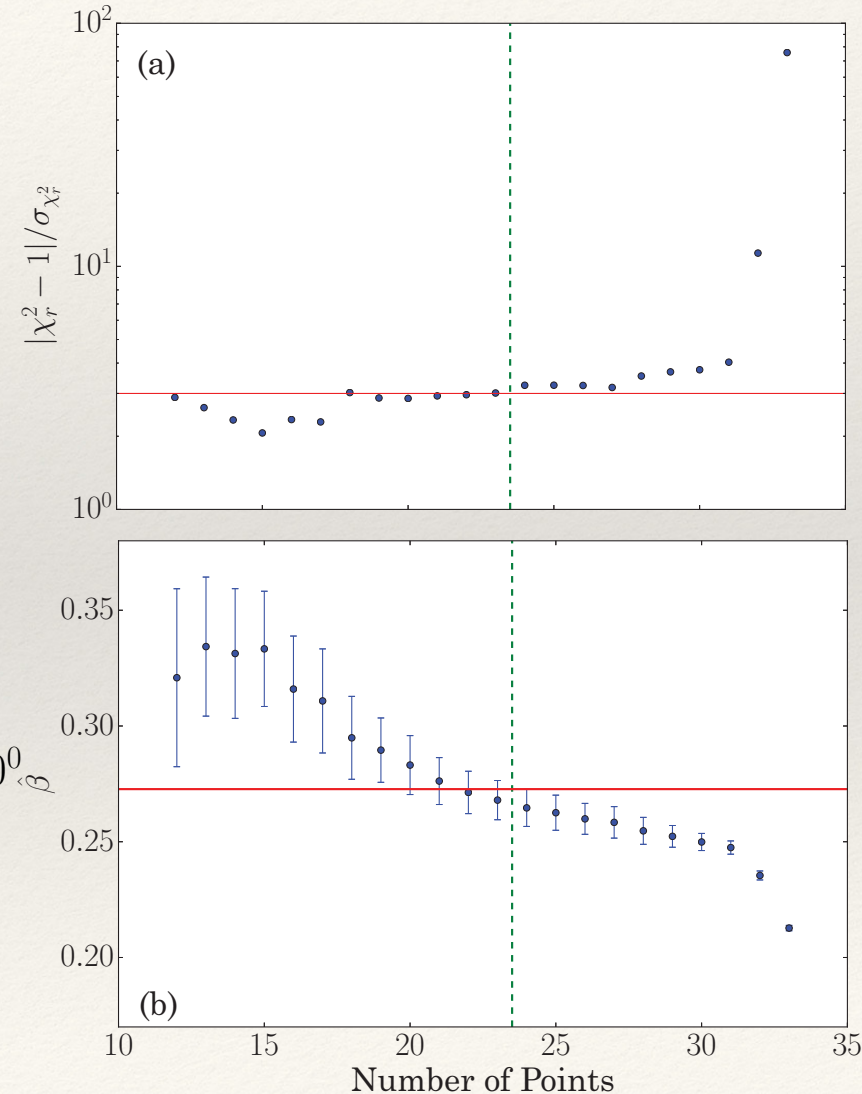
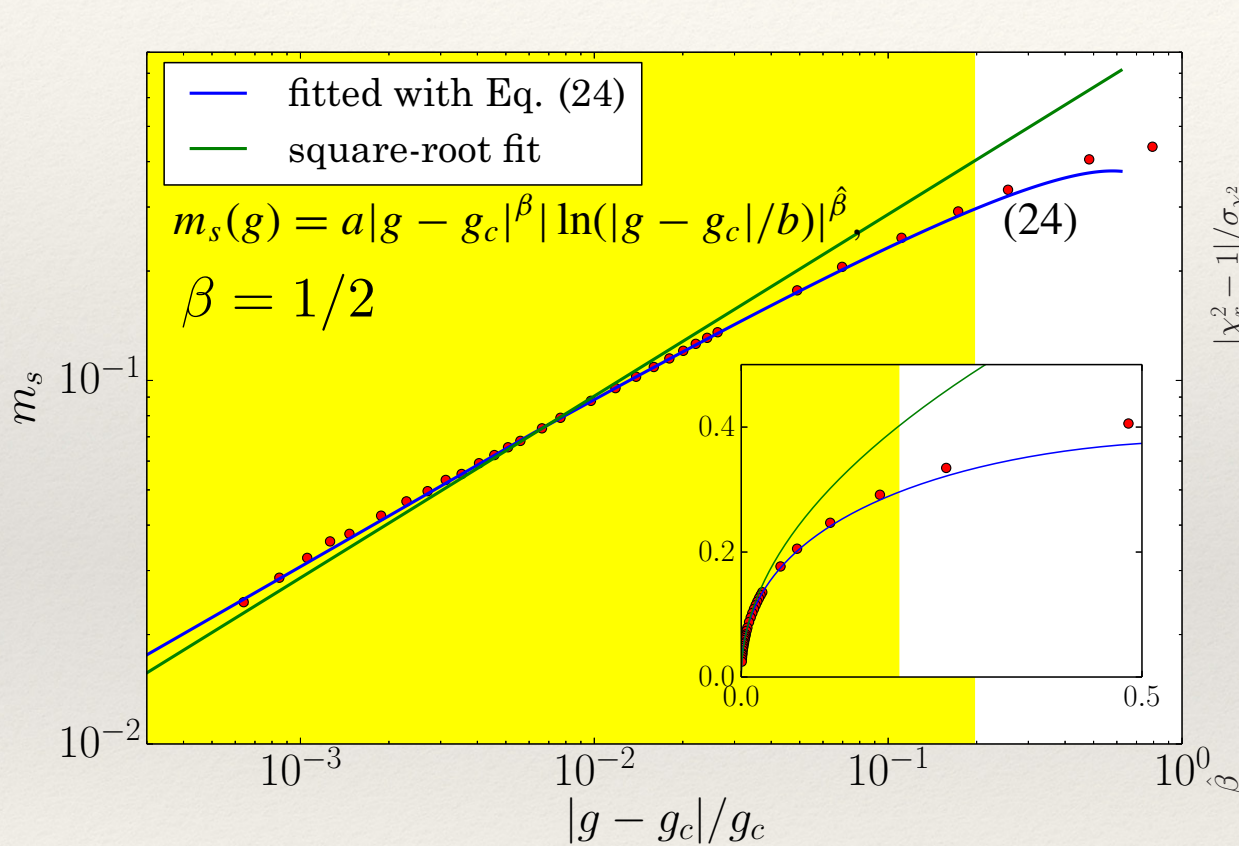
Logarithmic correction at the upper critical dimensionality

Mean-field theory is exact above the upper critical dimensionality d_u

- trivial critical exponents
- exactly at $d=d_u$ there are logarithmic corrections to the power laws

Test of expected log correction in the double-cube model ($d_u=4=3+1$)

[Y. Q. Qin, B. Normand, A. W. Sandvik, Z.Y. Meng, PRB 2015]



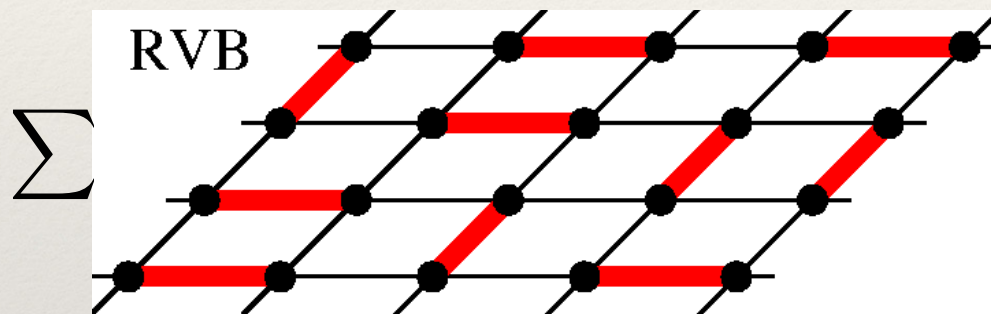
Conclusion: Log corrections can be detected if the leading exponent is known/fixed

More complex non-magnetic states; systems with 1 spin per unit cell

$$\mathbf{H} = J \sum_{\langle i,j \rangle} \mathbf{S}_i \cdot \mathbf{S}_j + g \times \dots$$

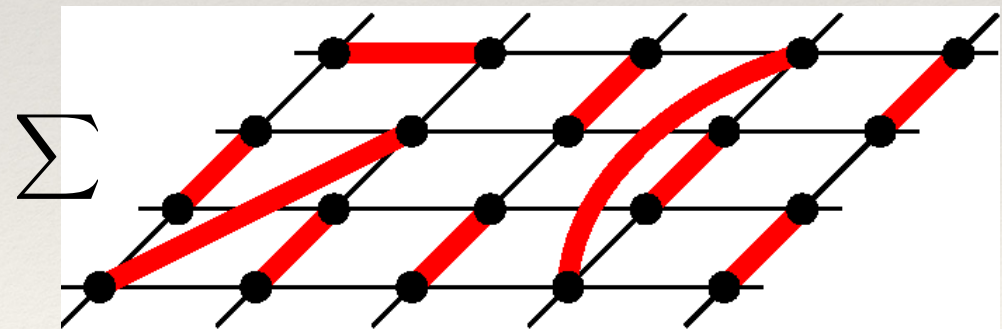
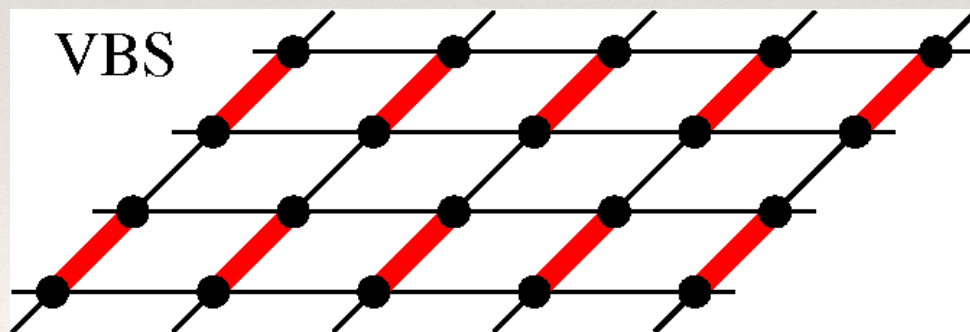
- highly non-trivial non-magnetic ground states are possible, e.g.,
 - ➔ resonating valence-bond (RVB) spin liquid
 - ➔ valence-bond solid (VBS)

Non-magnetic states often have natural descriptions with **valence bonds**



$$\begin{array}{c} \bullet \\ \diagdown \quad \diagup \\ i \quad j \end{array} = (\uparrow_i \downarrow_j - \downarrow_i \uparrow_j) / \sqrt{2}$$

The basis including bonds of all lengths is **overcomplete** in the singlet sector



- non-magnetic states dominated by short bonds