

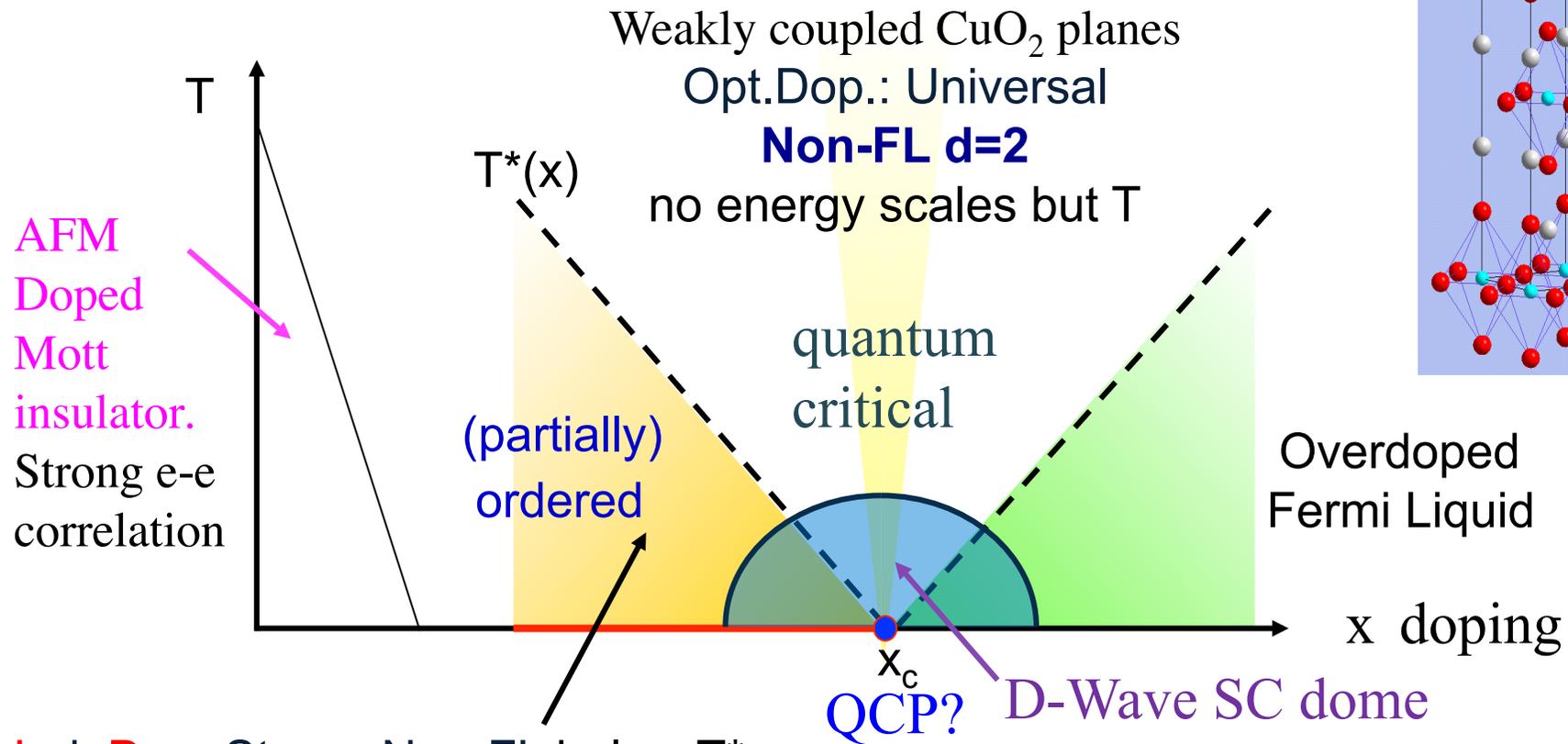
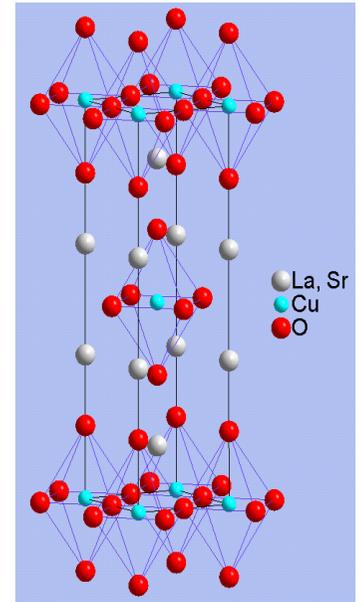
Ferronematic order in underdoped Cuprates

Collaborators: G. Seibold, M. Capati, M. Grilli and J. Lorenzana.

Hidden Ferronematic order in underdoped cuprates.
arXiv:1204.2119v1 [cond-mat.supr-com]

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Generic phase diagram of Cuprates, e.g. $\text{La}_{2-x}\text{Sr}_x\text{CuO}_4$



Und. Dop. Strong Non-FL below T^* .
 Pseudogap associated with a variety of spin and charge ordered structures:
 stripe-like (dynamical with smectic order) (LSCO, YBCO)
 checkerboard or droplets (Bi2212)

Non-FL in $d > 1$? Competing Order?
 The critical fluctuations of the competing order can be the glue for d-wave pairing and the cause of the Non-FL behavior.
 Hidden QCP (many proposals) →

QCP for the formation of ICDW.

Strong correlations reduces double occupancy

Why ICDW?

Phase Separation is easy

Generic of all models for CuO_2 planes

Add Coulomb forces \rightarrow

Frustrated PS on local basis

From the high doped correlated Fermi Liquid (Rome):

Local (Hubbard) repulsion reduces the homogenizing kinetic energy term, favoring PS in the presence of phonon mediated (*Holstein*) attraction.

Frustrate with Coulomb



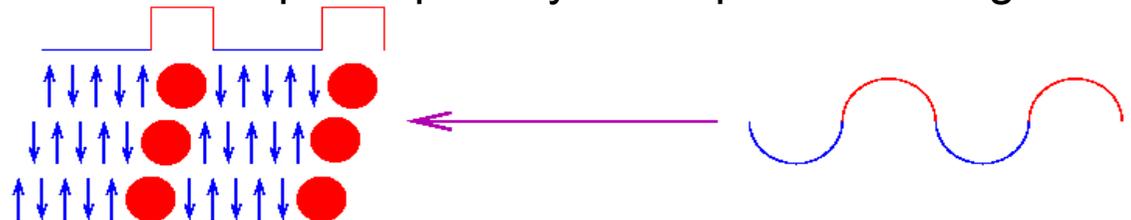
ICDW instability with finite q_c modulating wave vector $[(0,1), (1,0)$ direction], QCP ($x_c=0.19$), end-point of a critical line $T_{\text{CDW}}(x) \approx T^*$

STRIPES

From low doping (Emery, Kivelson):
Expulsion of holes from AF background;
Zaanen, Tranquada, ... antiphase stripes.

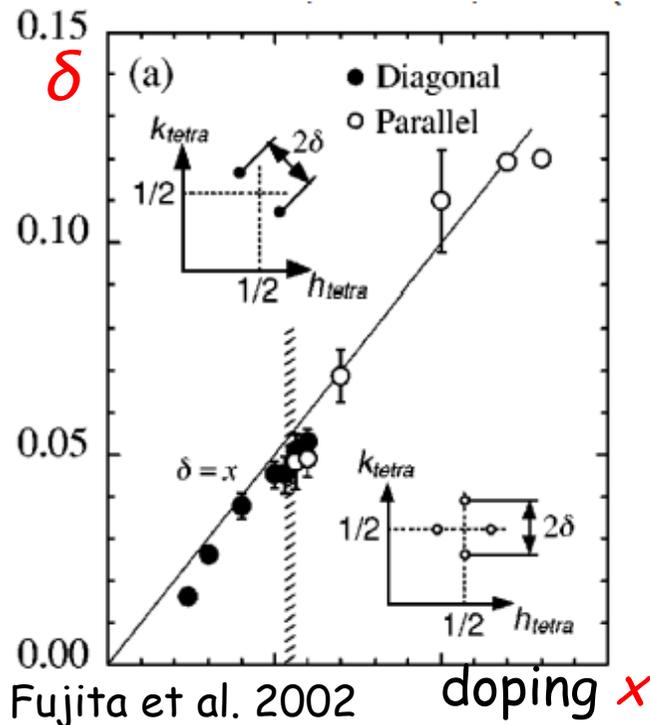
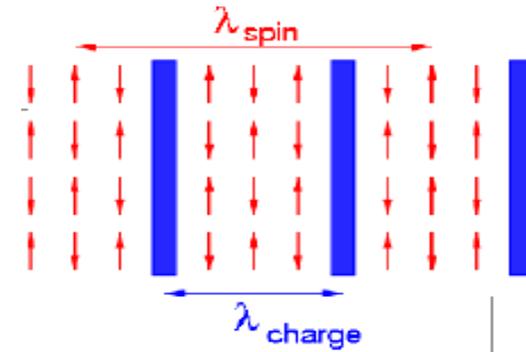
Indeed new resonant x-ray scattering experiments in YBCO [Ghiringhelli et al; Chang et al 2012] identify a 2D-ICDW as the order competing with superconductivity at intermediate doping. "The incipient CDW phase transition ...is preempted by the superconducting transition,..."

ICDW smoothly evolving into anharmonic stripes

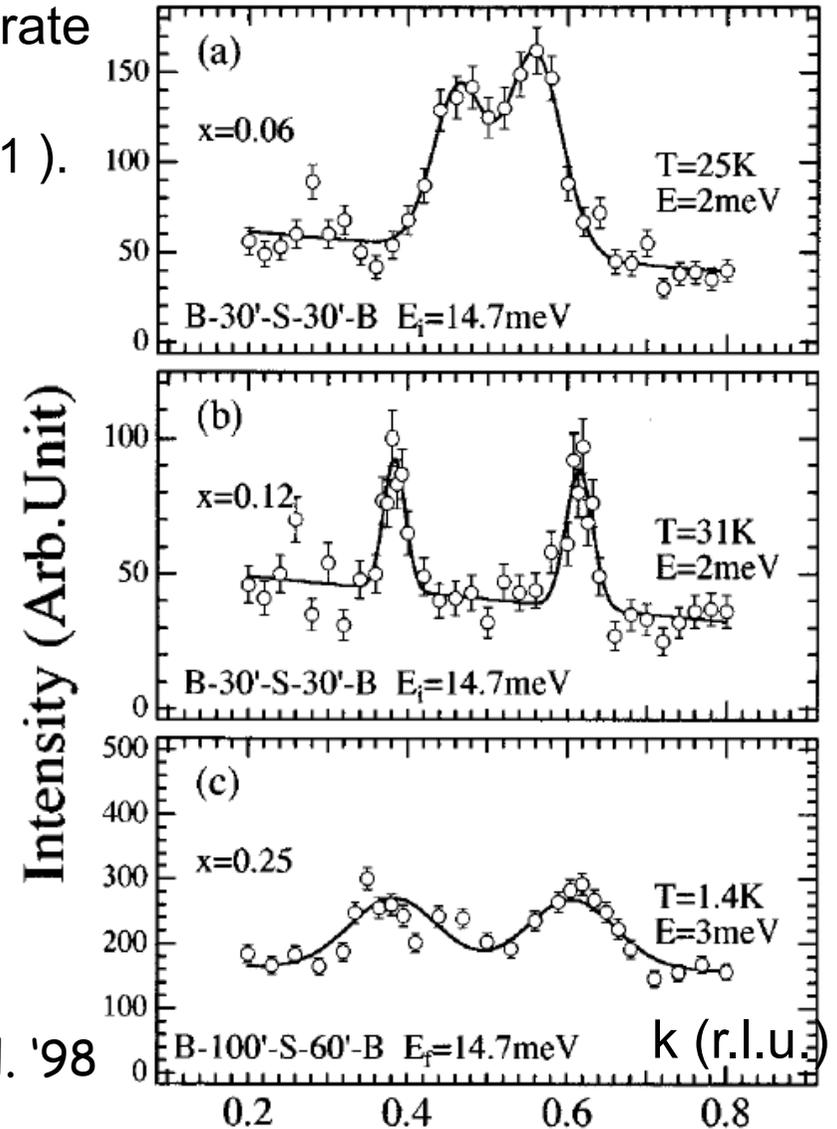


However **long range stripe order** observed in codoped LSCO only (e.g. $\text{La}_{1.48}\text{Nd}_{0.4}\text{Sr}_{0.12}\text{CuO}_4$)
 Elastic neutron scattering: Splitting of magnetic and superlattice peaks denotes periodic **spin** and **charge** modulation ($q_s = q_c/2$)

Generally, YBCO, LSCO, Bi2201: - Incommensurate spin scattering ($q_s \parallel$ Cu-O bond in YBCO;
 $q_s = (\pi/a)\delta$ diagonal for $x < 0.06$ in LSCO, Bi2201).
 -No signature of charge modulation.
 -Breaking of rotational symmetry



Yamada et al. '98



Therefore from low doping

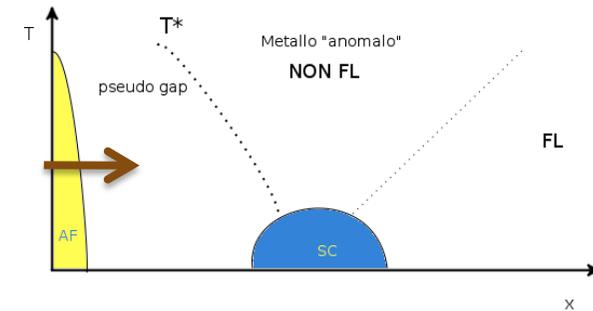
LSCO, Bi2201, YBCO:

-Incommensurate spin modulation, smectic like, response.

-No Charge modulation

-Evidence of four field rotational symmetry breaking

Nematic charge order also in $\text{Bi}_2\text{CaCu}_2\text{O}_{8+y}$, (Lawler et al. 2010 [STM])



By doping the AF background we will answer the questions:

Can **incommensurate spin correlations arise** without charge modulation as instead required by the stripe model?

Is **nematicity** a sign of fluctuating spin and charge density wave order, as it would be given by dynamic melted stripes, or has it an independent origin?

Doped holes form **Vortex-Antivortex (V-A) segments, nematic seeds (random in space) for incommensurate spin smectic like structure,** a new phase, dubbed by us “ferronematic”.

(G. Seibold et al. . **arXiv:12042119**)

See also Berciú and John 2004, Timm and Bennemann 2000.

Dilute limit

1-band extended Hubbard model (U, t, t' , $U/t=8$, $t'/t = \dots$)
 Variational calculations based on Gutzwiller approxim.
 Doped **charged holes** in the anti-ferro background
 form Vortex-Antivortex (**V-A**) pairs which, then,
 tend to arrange in **1D V-A segments**.

Analogy with dipolar fluids

In 3D, the maximal dipole-dipole attraction is
 when two dipole spheres touch each other 
nose-to-tail, favoring an endless alignment (as for stripes).



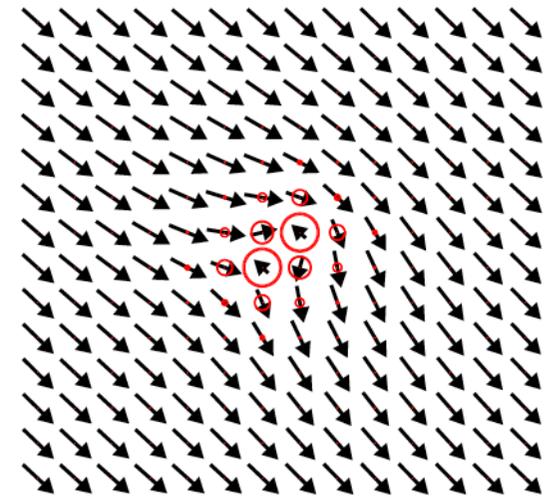
In 2D, the dipole-dipole interaction does not distinguish
 between the *nose-to-tail* and the **antiparallel side-by-side**
quadrupolar alignment



$$V_{dip-dip} = \frac{p_1 p_2}{l^2} - 2 \frac{(p_1 l)(p_2 l)}{l^4} = -|p_1||p_2|/l^2$$

V-A pair:

staggered spin/**charge**
 structure; $t'/t = -0.1$

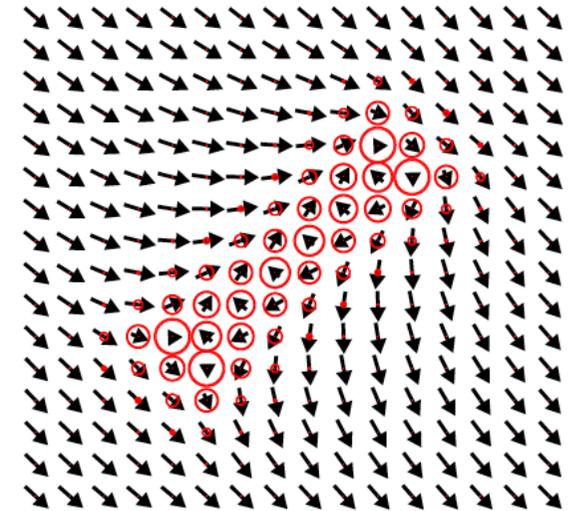


However in our variational approach short range interaction is important and the V-A pairs gain energy for **nose-to-tail alignment** (as in dipolar fluid) due to the local distribution of holes in magnetically ordered environment.

Direction is determined by t'/t .

For $-0.3 < t'/t < -0.2$ crossover from **diagonal to vertical-horizontal** configuration.

16x16 lattice sites,
 $t'/t = -0.2$
4 V-A pairs
 staggered spin/**charge**



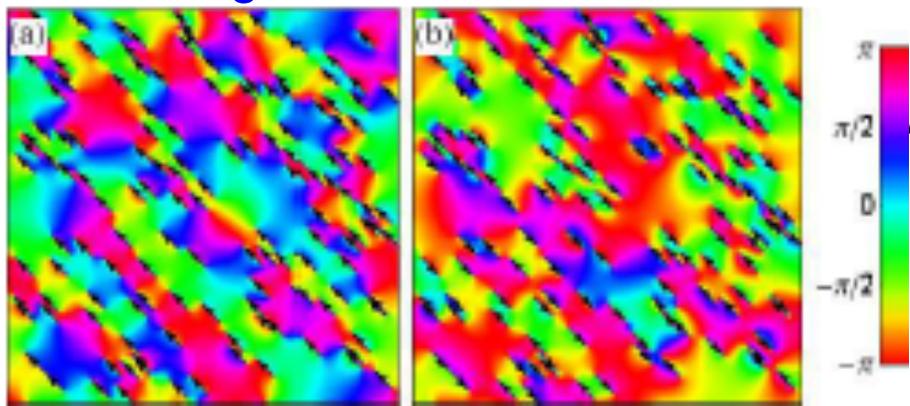
Each segment of N_{pairs} and length l has $N_{ch} (=2N_{pairs})$ charges immersed in a compensating charged background: Long range Coulomb interaction (Coulomb charge energy is increasing as $N_{ch} \ln N_{ch}$) limits **chain's length l to few lattice constants.**

For N_{seg} , the magnetic short range interaction favors, as for two V-A pairs, a ferronematic alignment.

*Distribution of N_{seg} segments each of length l and charge N_{ch} :
effect of orientation*

A classical XY-model is used (variational GA is too limited in sizes). Segments are chains of vacancies, alternately centers of vortex and antivortex. Parameters obtained by comparing the phase change across a single segment. Minimization of classical energy.

Oriented segments random orientation



Spin phase distribution is obtained at given doping $x=N_{ch}N_{seg}/L^2=0.03$ on a lattice of 160x160 sites,, segment sites $l+1=8$, filling factor $\nu=N_{ch}/(l+1)=0.7$.

In fig.a, with macroscopic polarization, phase modulation (stripe-like in the spin sector) appears, but with random space distribution of oriented segments.

Fig.b, instead, disaggregates in large areas of equal phase.

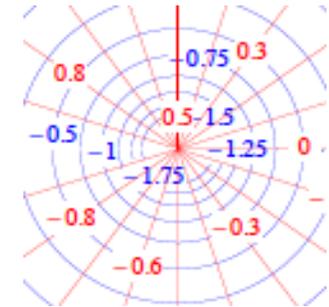
The influence of a collection of equally oriented V-A segments on spins at each point r is obtained by evaluating the total phase change Φ at point r

$$S^x(\mathbf{r}) = S_0 \exp(i\mathbf{Q} \cdot \mathbf{r}) \cos \Phi(\mathbf{r}), \quad S^y = S_0 \exp(i\mathbf{Q} \cdot \mathbf{r}) \sin \Phi(\mathbf{r}), \quad \mathbf{Q} = (\pi, \pi),$$

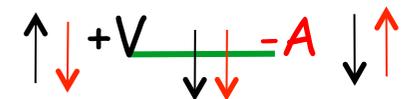
$\text{grad}\Phi$ gives the *incommensuration* of spin response.

-In a single vortex, Φ is the *velocity potential* and *coincides with* the radial angle φ .

The (spin) velocity v , tangential to the *streamlines*, is $v = \text{grad}\varphi = 1/r$.



-Pair of Vortex and Antivortex at distance l



The velocities *add* to the value $v = 2/l/2$ between V and A and tend to *cancel outside*. v is *orthogonal* to the dipole l

The phase disturbance, averaged on a square (L^2), extends on a circle of radius $l/2$:

$$\langle \text{grad}\Phi \rangle \approx \pi(l/2)^2 / L^2 \mathbf{v} = \pi l \mathbf{x}(\text{versz}) / L^2.$$

-For a segment of V-A pairs, the dipole is the length l .

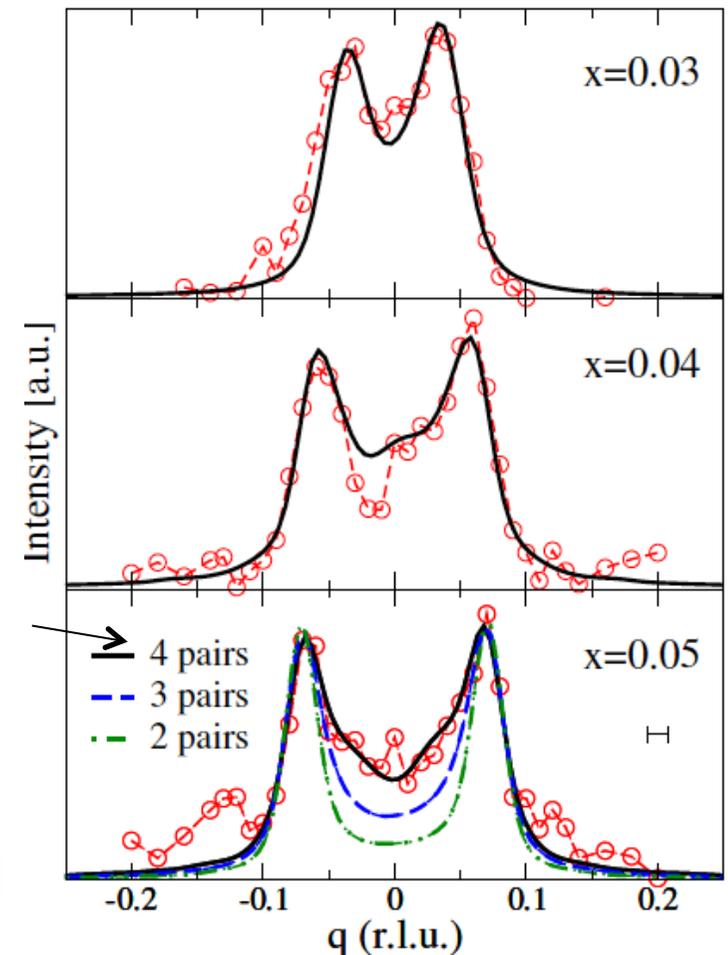
-Distribution of N_{seg} macroscopically polarized segments, random in space, length l , doping $x=N_{seg}N_{ch}/L^2$, filling factor $\nu=N_{ch}/(l+1)$ ($l+1$ sites):

$$\langle grad\Phi \rangle \approx N_{seg} \langle grad\Phi \rangle_{segm} = N_{seg} \pi l x (\text{versz}) / L^2 = q$$

$q = \pi \frac{x}{\nu} \frac{l}{l+1} \equiv \pi P$ is the incommensurate modulation vector of spin response (as in the stripes); q determines peak position

q perpendicular to the dipole segments and to their macroscopic polarization $P = N_{seg} l / L^2$, the "ferronematic" order parameter, (incommensurability in YBCO? Hinkov et al 2009)

Fits of the experiments (Wakimoto et al. 2000) on spin structure factor for doping $x=0.3, 0.4, 0.5$, $l+1=8$, *no smectic order*. Average over 20~30 configurations of 160x160 lattice



Conclusion

Inhomogeneous State as a bridge between FL and doped AFM separated by a **hidden QCP, which**, coming from high doping, **arises as a correlated-Fermi-liquid instability** :

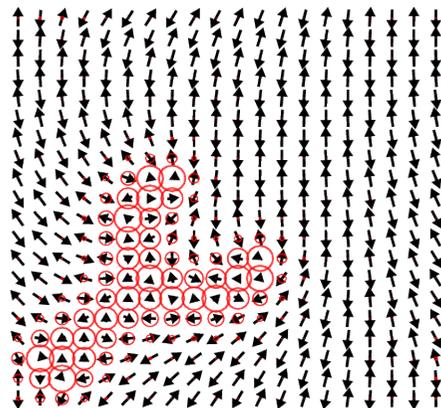
At low doping **glass of nematic V-A chain segments** gives rise to spin **smectic like correlations** with the polarization of the segments as an order parameter of a phase, dubbed **ferronematic** by us.

Open problems: -how **the two topologically different phases (stripes and ferronematic segments)** are interconnected?

- $T \neq 0$? Nematic and polarization transition? Simultaneous or which first?

Increasing doping and T , branching is favored: more complex structures (checkerboard, bubbles,...) and a gas of topological excitations (ends, junctions, edge dislocations) should appear out of the nematic glass.

junctions:
20x20 lattice
14 holes,
 $U/t=8$,
 $t'/t=-0.1$



junction in
dipolar fluid

