

Quantum Suppression of Alignment in Spinning Nanoparticles

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- **Davis-Greenstein Alignment by Magnetic Dissipation**
- **Suppression of Dissipation in Cold Nanoparticles**
- **Alignment vs. Grain Size**
- **Predicted Polarization vs. Frequency**

Davis-Greenstein Alignment by Magnetic Dissipation

- Grain spinning in static $\mathbf{B}_0$
 θ = angle between $\mathbf{J}$ and $\mathbf{B}_0$
- Grain material tries to magnetize
- In grain frame:
 - Rotating component $\mathbf{B}_0 \sin \theta$
 - Magnetization $\mathbf{M}$ lags
- torque $\mathbf{M} \times \mathbf{B}_0$ on grain
Reduces component of $\mathbf{J} \perp \mathbf{B}_0$
Leaves component of $\mathbf{J} \parallel \mathbf{B}_0$ unchanged
→ alignment of $\mathbf{J}$ with $\mathbf{B}_0$
- If $\text{Im}(\chi) = K\omega$, then
Davis-Greenstein alignment time

$$\tau_{\text{DG}} = \frac{2\rho a^2}{5KB_0^2} \propto \frac{a^2}{K}$$

vs. disalignment time (time to collide with own mass of gas)

$$\tau_{\text{M}} \propto a$$

Hence:

$$\frac{\tau_{\text{DG}}}{\tau_{\text{M}}} \propto \frac{a}{K}$$

Naive DG theory:
for standard estimate of

$$K \approx \frac{1 \times 10^{-13} \text{ s}}{T_{\text{gr}}/18 \text{ K}}$$

small grains should be aligned.

Alignment is result of *dissipation*:

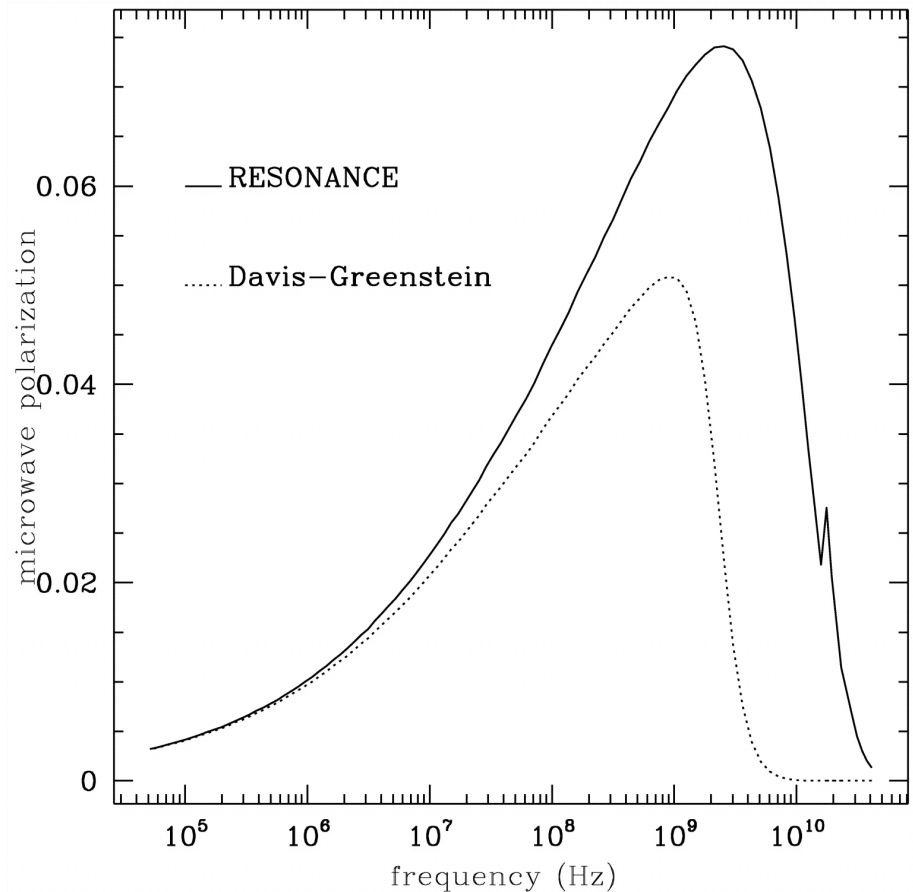
J^2 is *reduced*:

rotational KE is converted to **heat**

Dissipation in Nanoparticles

Lazarian & Draine (2000):

- *Need to dissipate $\hbar\omega_{\text{rot}}$ as heat.*
- *When nanoparticle is in very low vibrational level, there are no other vibrational states within $\Delta E = \hbar\omega_{\text{rot}}$:*
- *Quantization of low-lying vibrational modes leads to suppression of paramagnetic dissipation in small nanoparticles.*



Prediction: $P \lesssim 4\%$ at $\nu > 10$ GHz

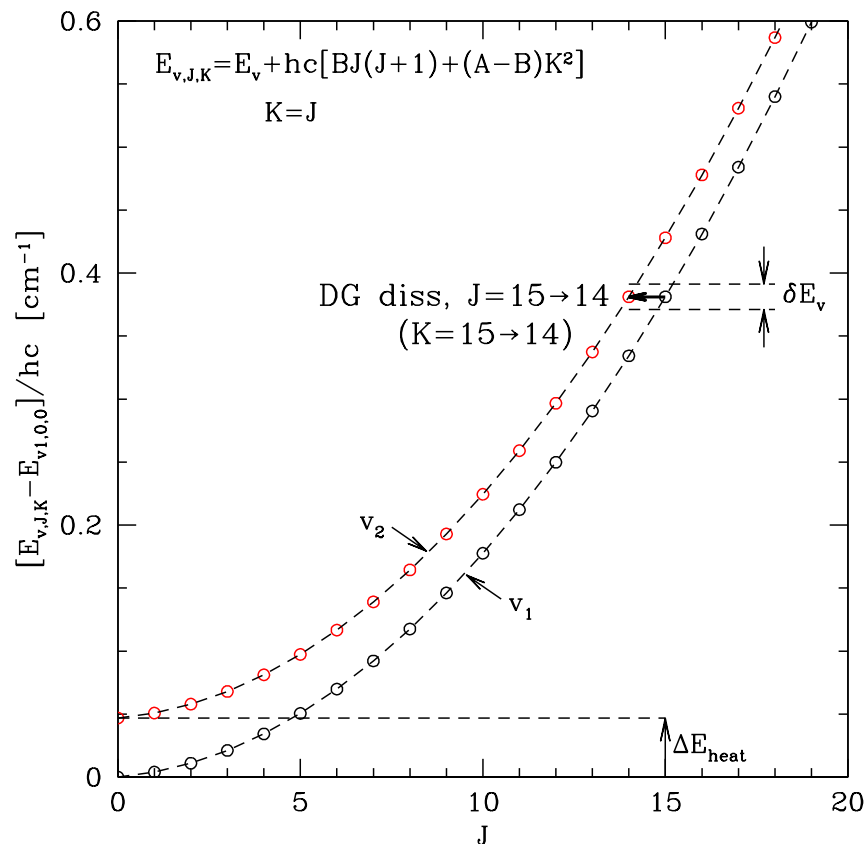
Fig. 1a from Lazarian & Draine (2000)

Draine & Hensley (2016): quantum suppression of dissipation is more severe than estimated by Lazarian & Draine (2000).

Dissipation in Nanoparticles

“heat” = vibrational KE in grain

Dissipation of rotational KE requires that there be a vibrational transition that can “accept” the energy.



The two energy levels must coincide to within energy level uncertainty δE_v

Energy level uncertainty δE_v from

- spontaneous emission
- collisions

For $T_{\text{vib}} \lesssim 100$ K, δE_v is very small:

$$\frac{\delta E_v}{hc} \lesssim 10^{-12} \text{ cm}^{-1}$$

What is the probability that another energy level will exist within energy δE_v of the first level?

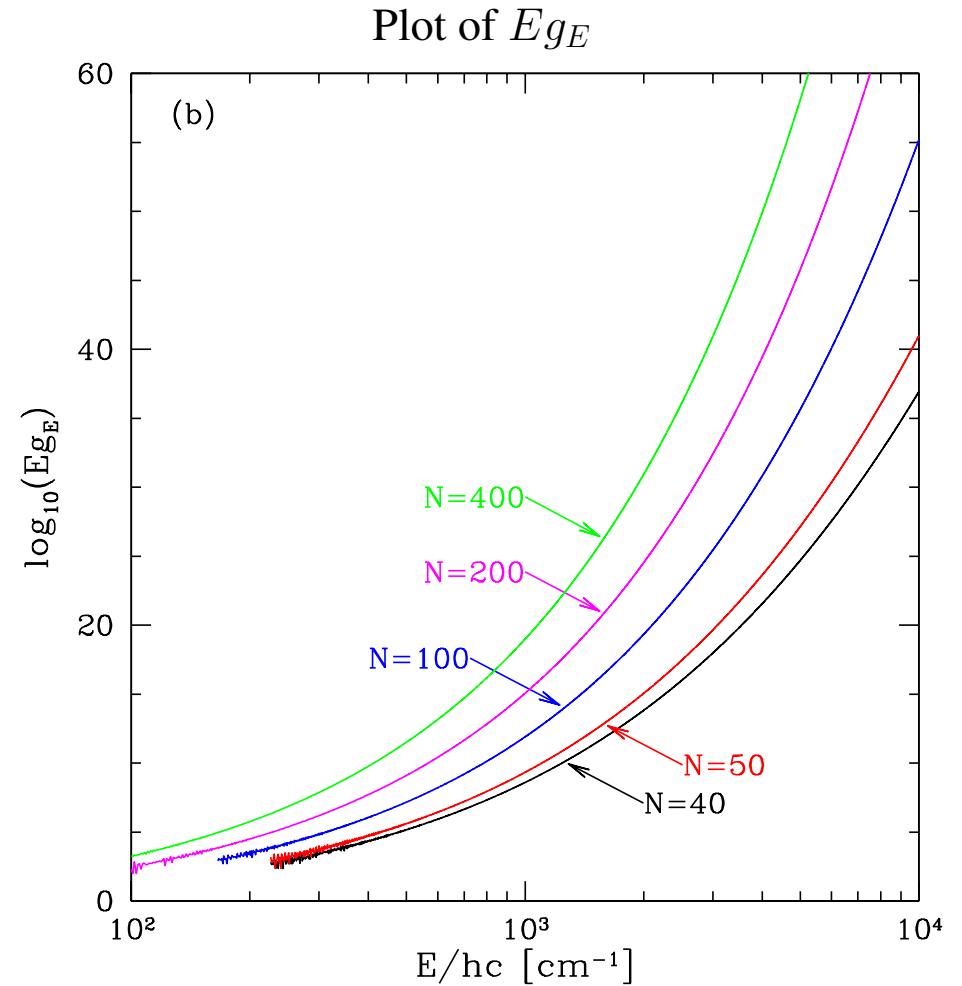
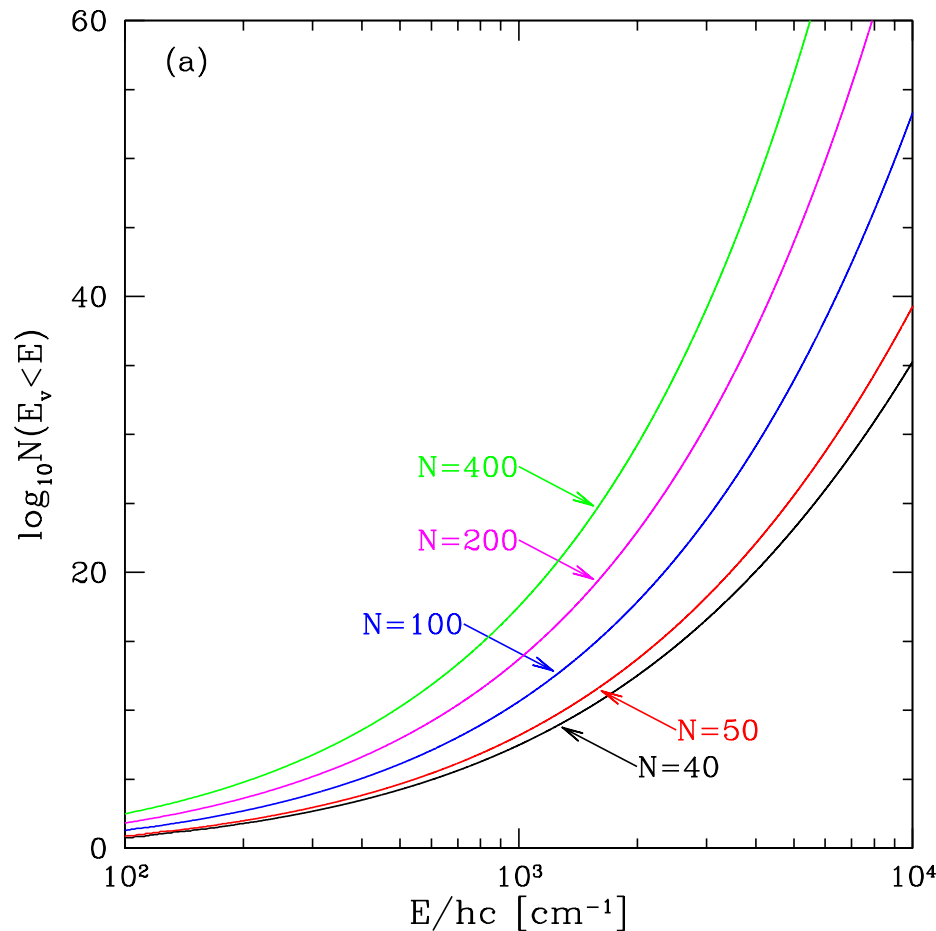
Vibrational Density of States in Nanoparticles

Nanoparticle with N atoms: $3N - 6$ vib. modes.

Each mode: harmonic oscillator with fundamental ω_j .

Directly calculate $N_v(E) \equiv$ number of *distinct* vibrational states with *total* vibrational energy $< E$.

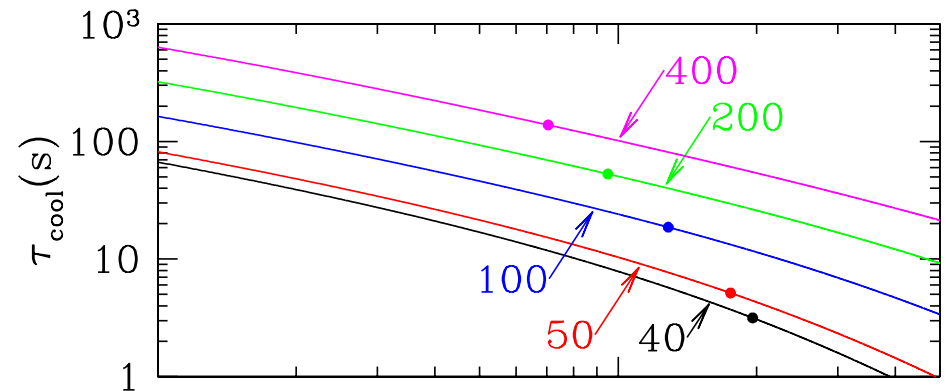
Density of states $g_E \equiv dN_v/dE$



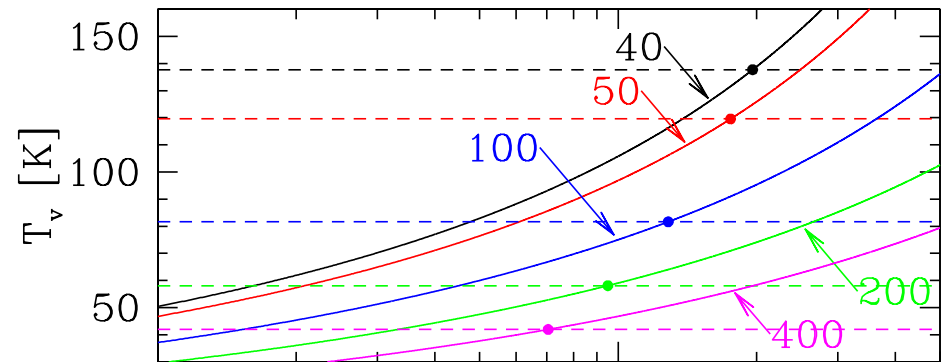
Examples assuming fundamental mode spectra for amorphous silicates

Quantum Suppression Factor ψ_q

Cooling time

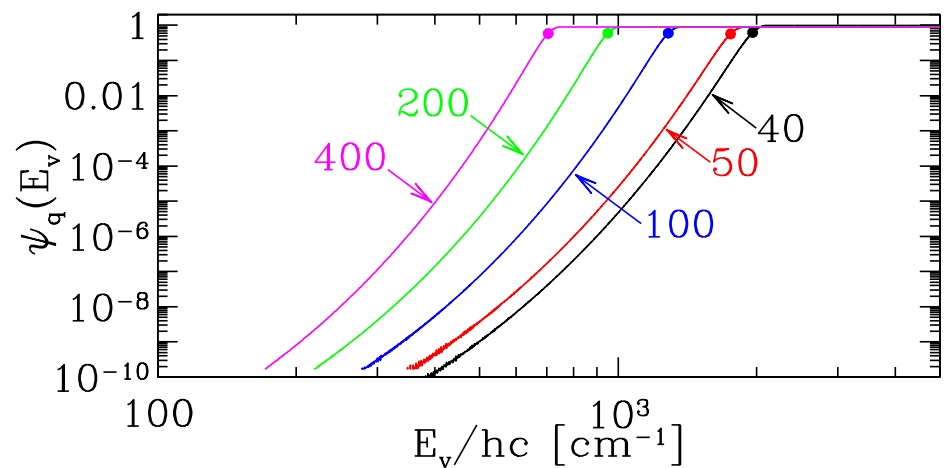


Vibrational Temperature



Probability of one or more levels
in interval δE

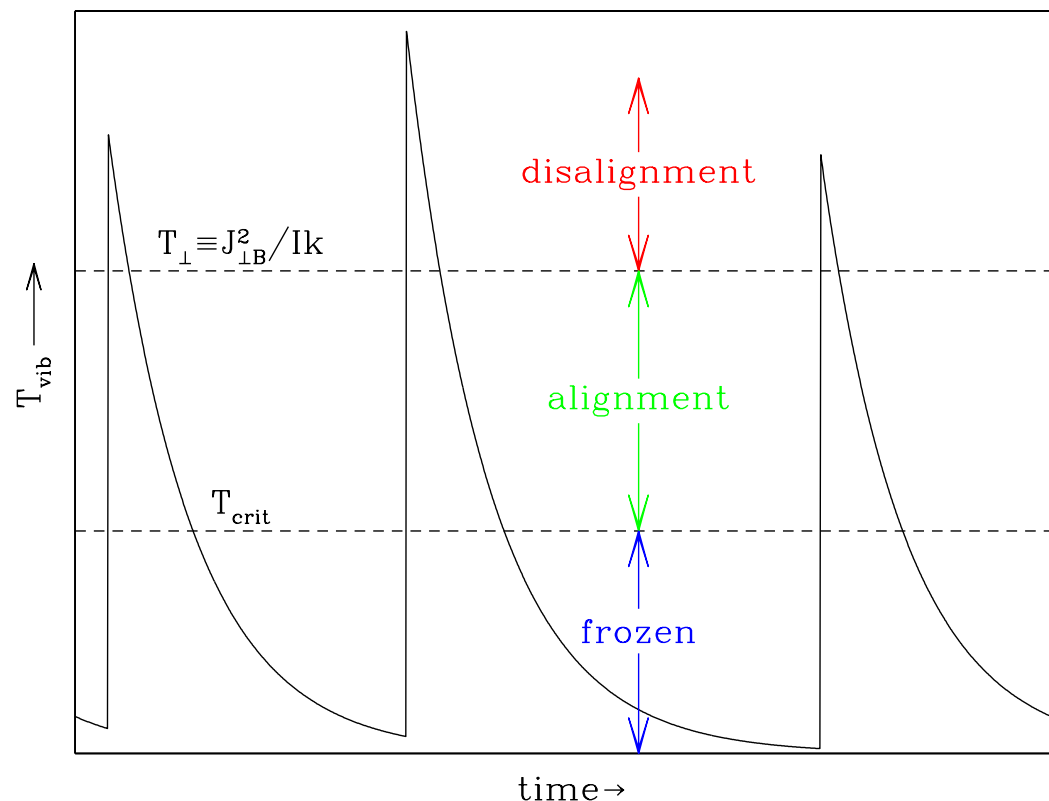
$$\psi_q(E) = 1 - \exp(-\delta E \times g_E)$$



$E_v \equiv$ vibrational energy

Time-Averaged Quantum Suppression Factor

- Stochastic heating by starlight: Fluctuating $T_{\text{vib}}(t)$.
- When grain is “hot”, magnetic **fluctuations** dominate magnetic dissipation: magnetic torques act to *increase* $\mathbf{J} \perp \mathbf{B}_0$
→ *disalignment*
- Intermediate T_{vib} : magnetic **dissipation**: magnetic torques act to *decrease* $\mathbf{J} \perp \mathbf{B}_0$
→ *alignment*
- Low temperatures: dissipation suppressed: alignment is “frozen”



Steady-State Rotational Excitation and Alignment

Excitation and damping of grain rotation by

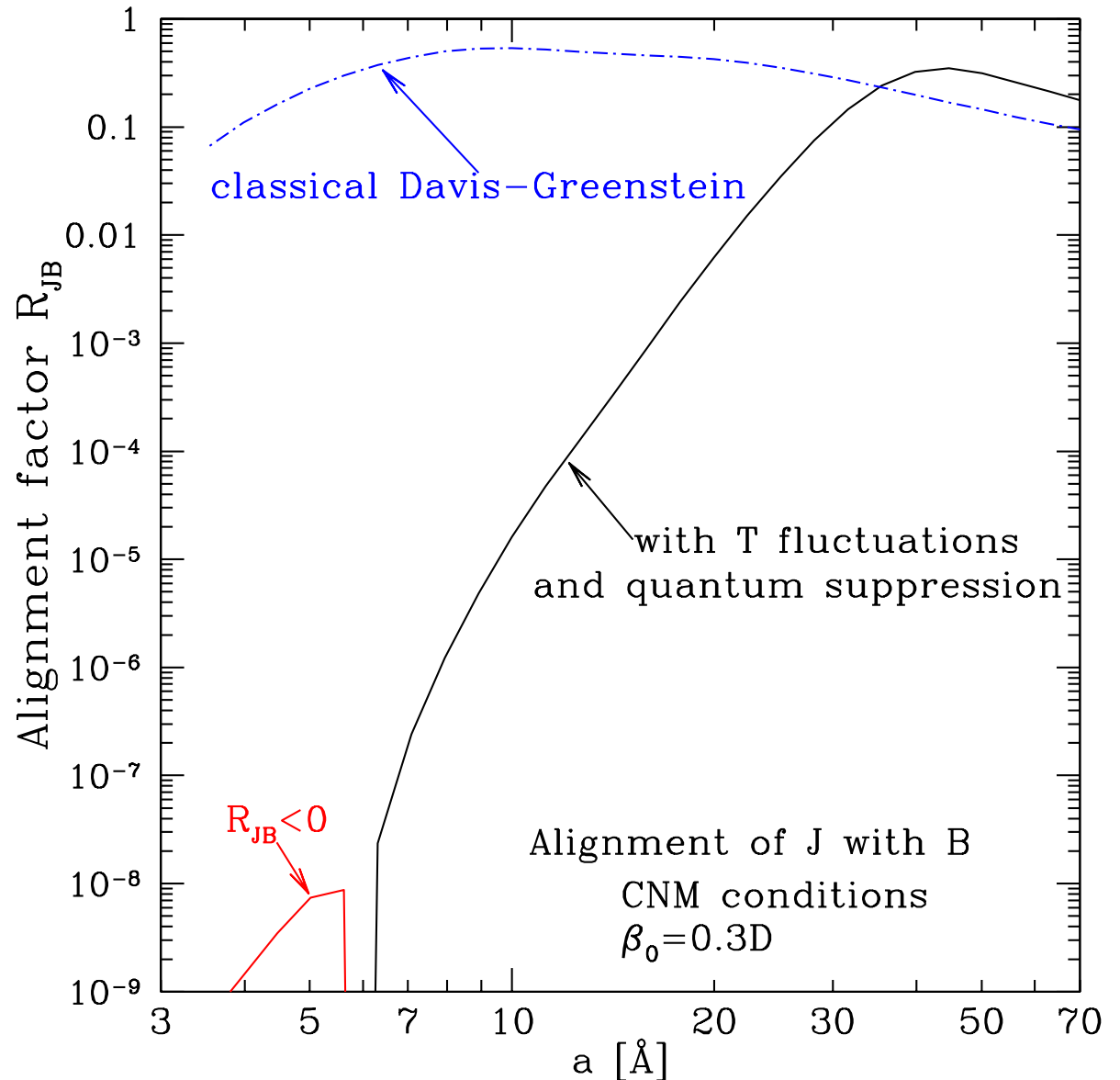
- collisions
- “plasma drag”
- absorption of starlight
- emission of IR photons
- rotational emission
- paramagnetic dissipation

Find rotation $\parallel \mathbf{B}_0$ and $\perp \mathbf{B}_0$:

$$\langle \cos^2 \theta \rangle = \frac{\langle J_{\parallel}^2 \rangle}{(\langle J_{\parallel}^2 \rangle + \langle J_{\perp}^2 \rangle)}$$

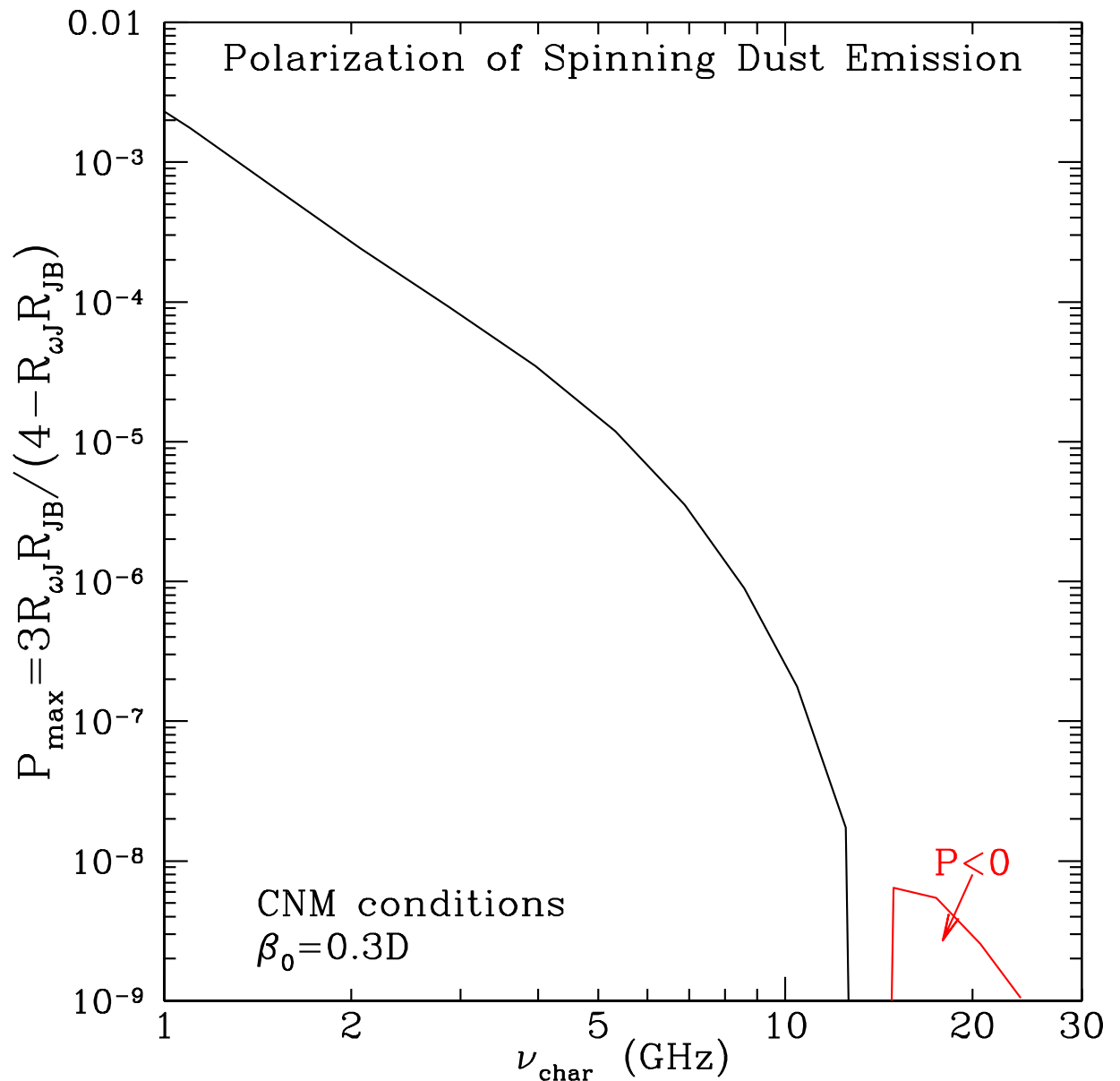
Alignment measure:

$$R_{\text{JB}} \equiv \frac{3}{2} (\langle \cos^2 \theta \rangle - \frac{1}{3})$$

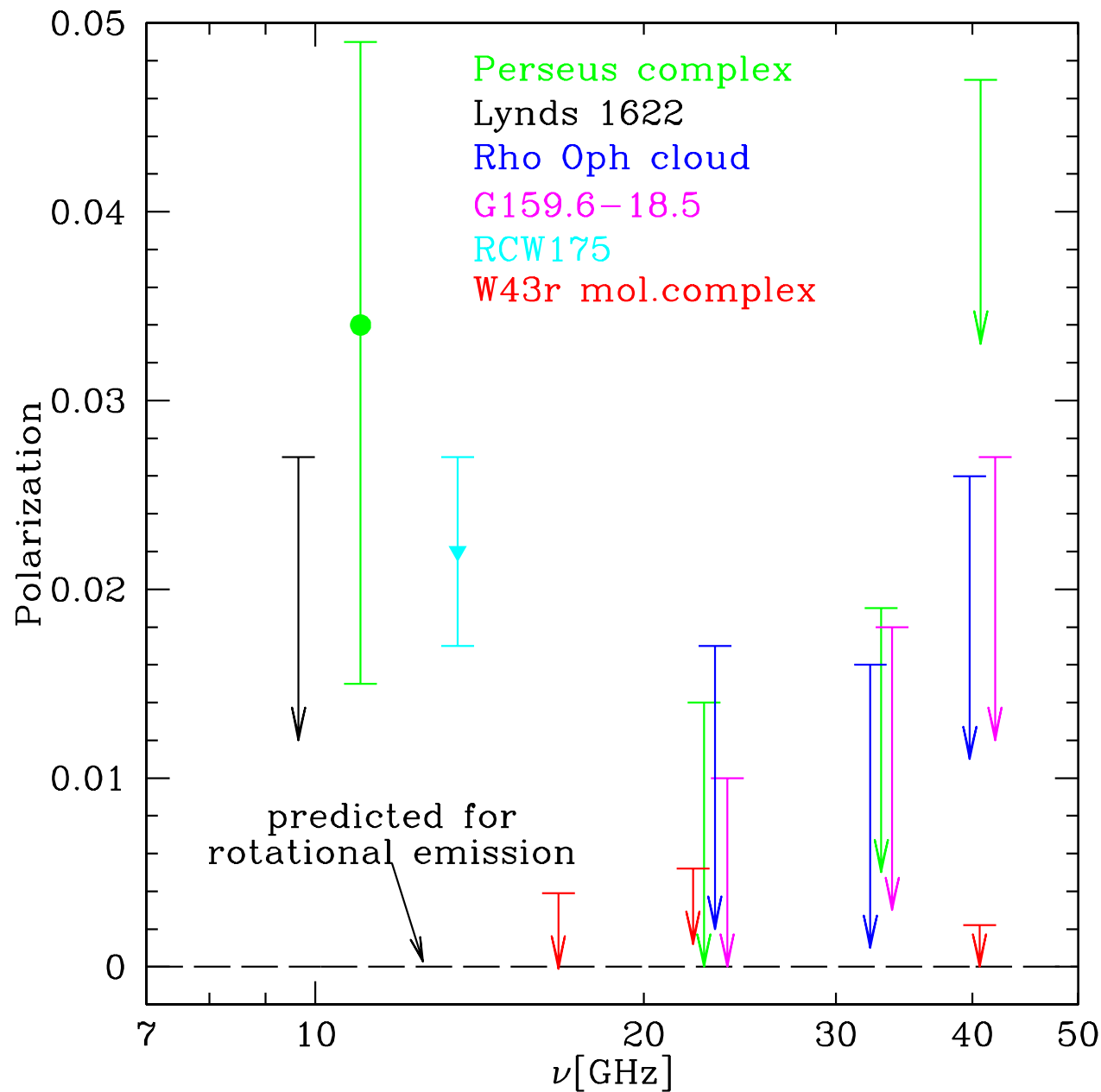


Polarization of Rotational Emission

- Observer's line-of-sight $\perp \mathbf{B}_0$
- Assume each size to spin at characteristic frequency $\langle \omega^4 \rangle^{1/4}$
- Allow for imperfect alignment of ω with $\mathbf{J}$ ($R_{\omega J} \approx 0.9$)



Observations



Perseus mol. complex: Battistelli et al. (2006), Dickinson et al. (2011)

Lynds 1622: Mason et al. (2009)

ρ Oph: Dickinson et al. (2011)

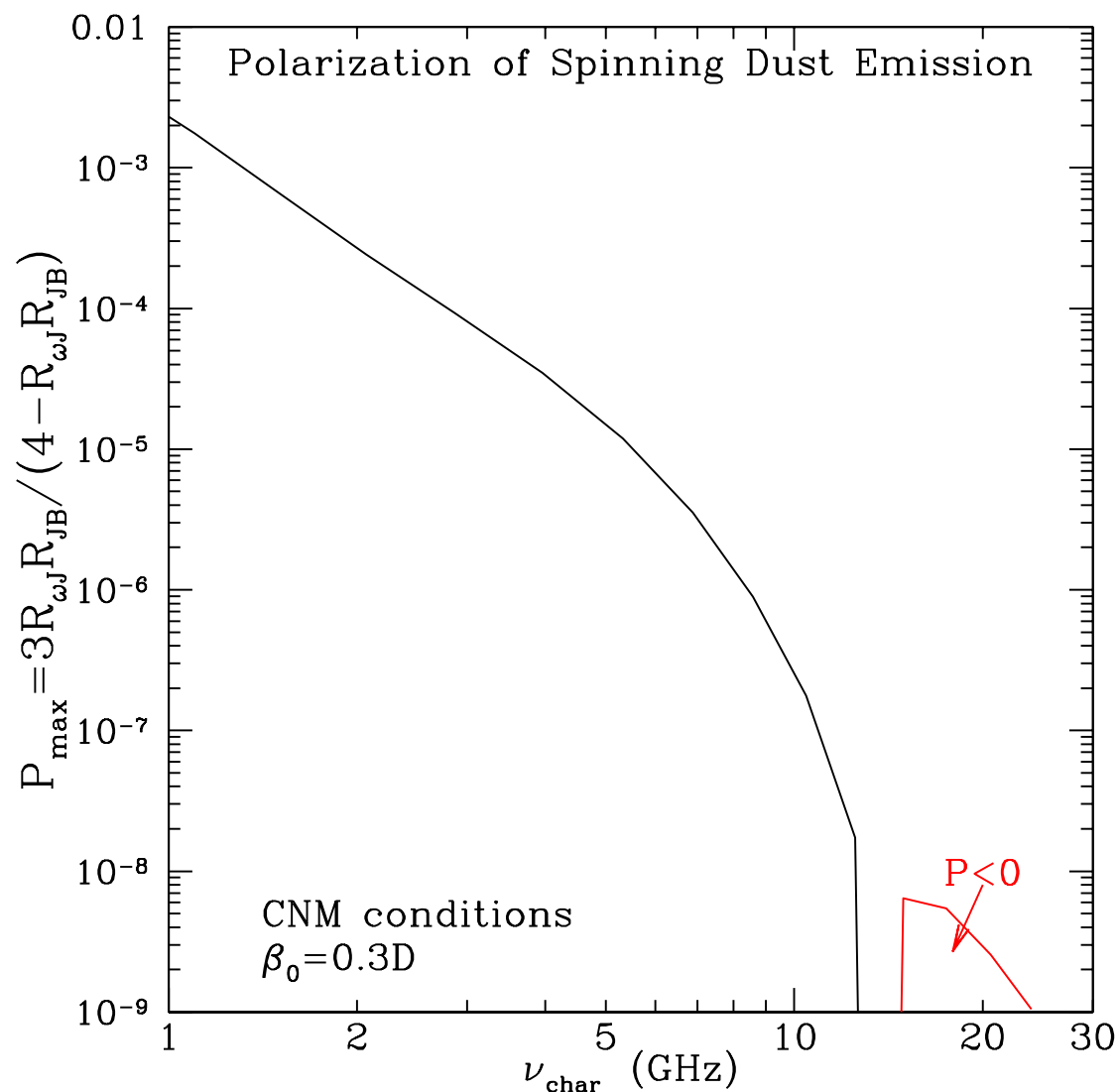
G159.6-18.5: López-Caraballo et al. (2011), Génova-Santos et al. (2016)

RCW175: Battistelli et al. (2015)

W43r mol. complex: Génova-Santos et al. (2016)

Summary

- Quantization of vibrational energy levels $\rightarrow$ suppression of dissipation in very small, cold nanoparticles.
- Suppression of dissipation $\rightarrow$ suppression of alignment of $\mathbf{J}$ with $\mathbf{B}_0$
- Emission from $a \lesssim 10 \text{ \AA}$ nanoparticles will be negligibly polarized.





THANK YOU

References

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