

Microscopic Understanding of the Effects of Impurities in Low RRR SRF Cavities

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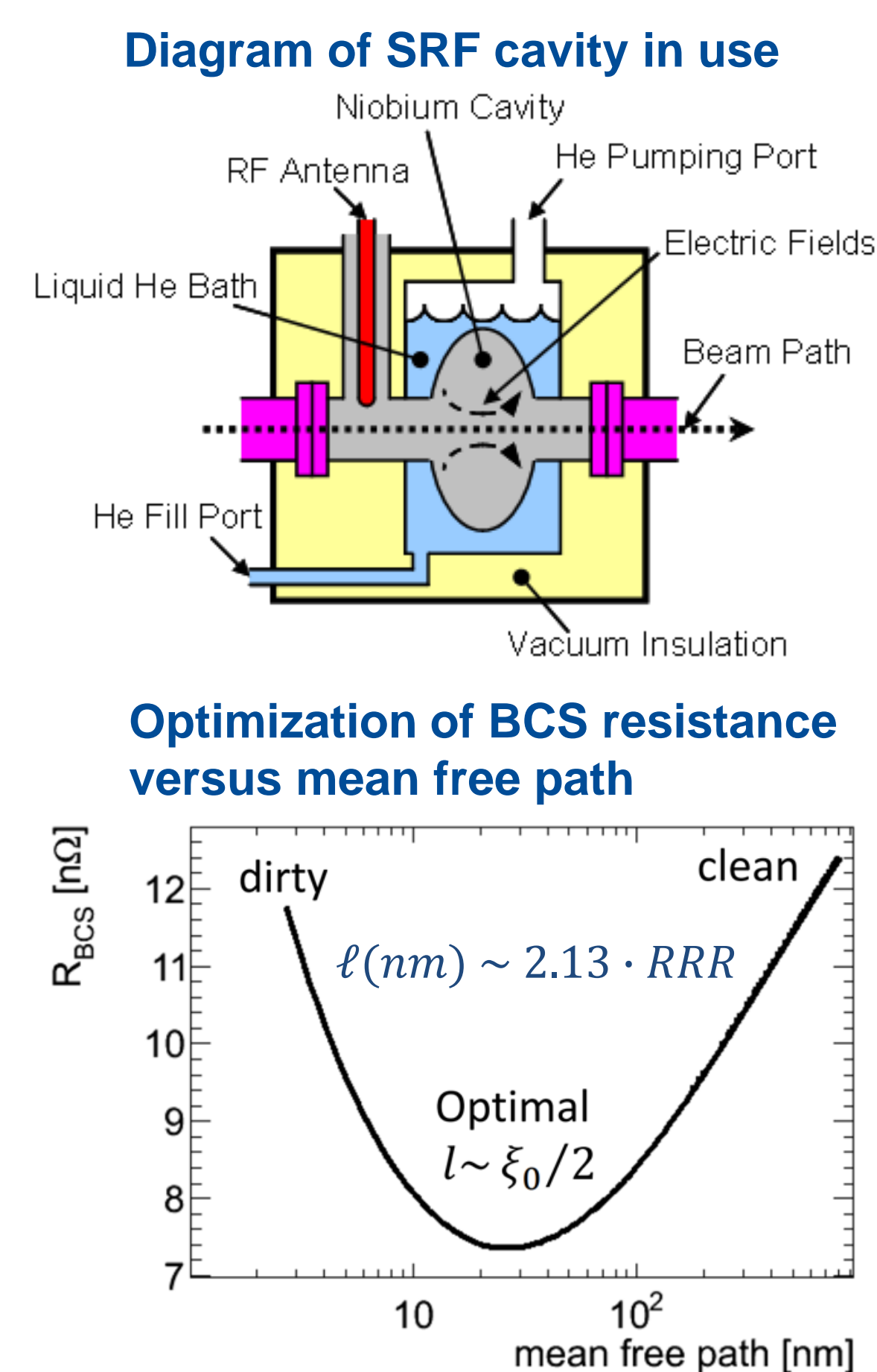
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SRF = Superconducting Radio-Frequency

- SRF cavities are resonant structures made from niobium that generate the electric field along the beamline
- Goal of SRF studies is to design surface profile to increase quality factor (Q_0) and accelerating gradient

Motivation

- Many SRF studies follow a “clean bulk dirty surface” technique to optimize the BCS resistance by adding extrinsic impurities
- What role do intrinsic impurities serve?
 - Might perform similar functions as extrinsic impurities which have been shown to improve performance
- Understanding of intrinsic impurities will enable **future** high Q_0 /high gradient **surface treatments**



RRR = Residual Resistance Ratio

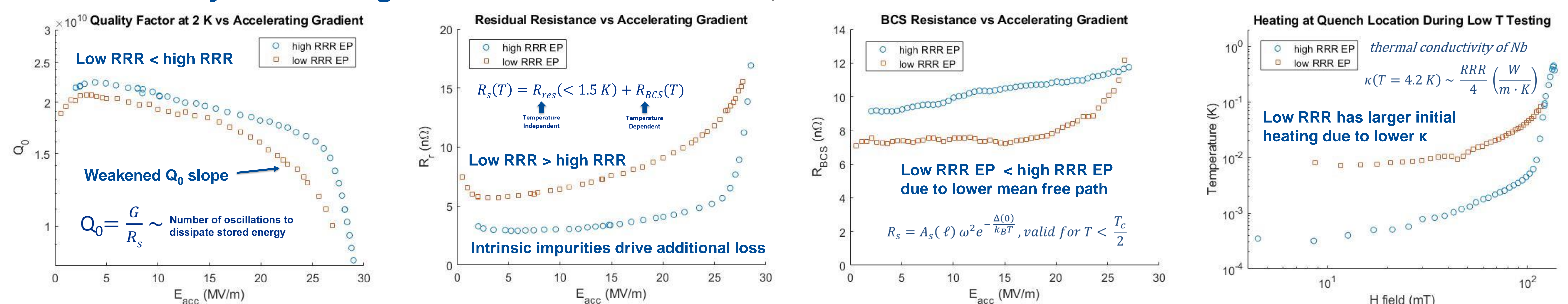
$$RRR = \frac{\rho(300\text{ K})}{\rho(10\text{ K})}$$

- RRR is lowered by impurities in the Nb
- High RRR for SRF is ~300

Experiment

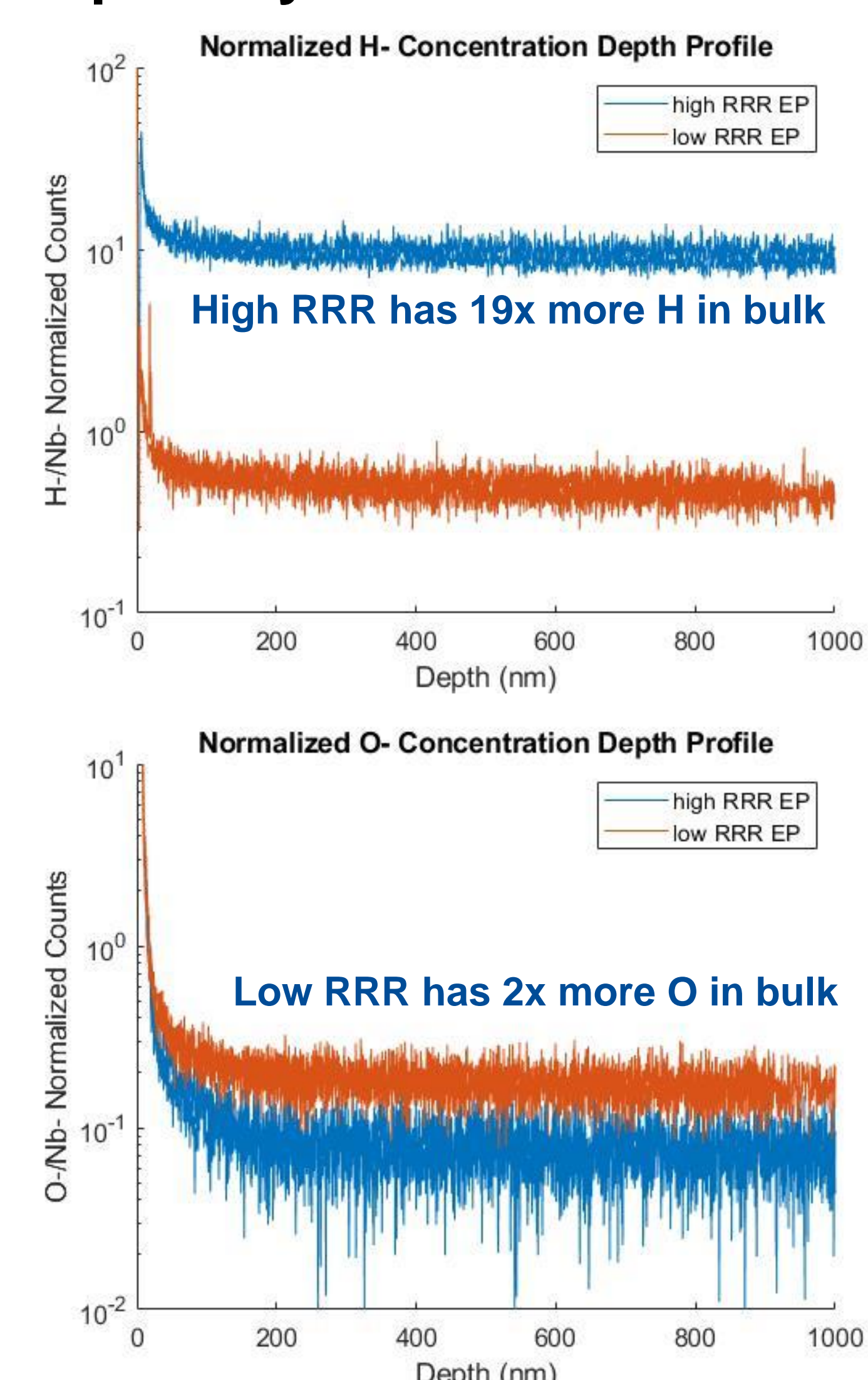
- RF testing on 1.3 GHz single-cell TESLA-shaped low RRR cavity in electropolished (EP) condition
- Quality factor vs accelerating gradient
- Temperature mapping
- Sample study on low RRR material
- RRR measurement = 35.11** → mfp ~ 75 nm
- Secondary ion mass spectrometry (SIMS)
- Electron backscatter diffraction (EBSD)

Cavity RF Testing Results: worse performance in general but low BCS resistance

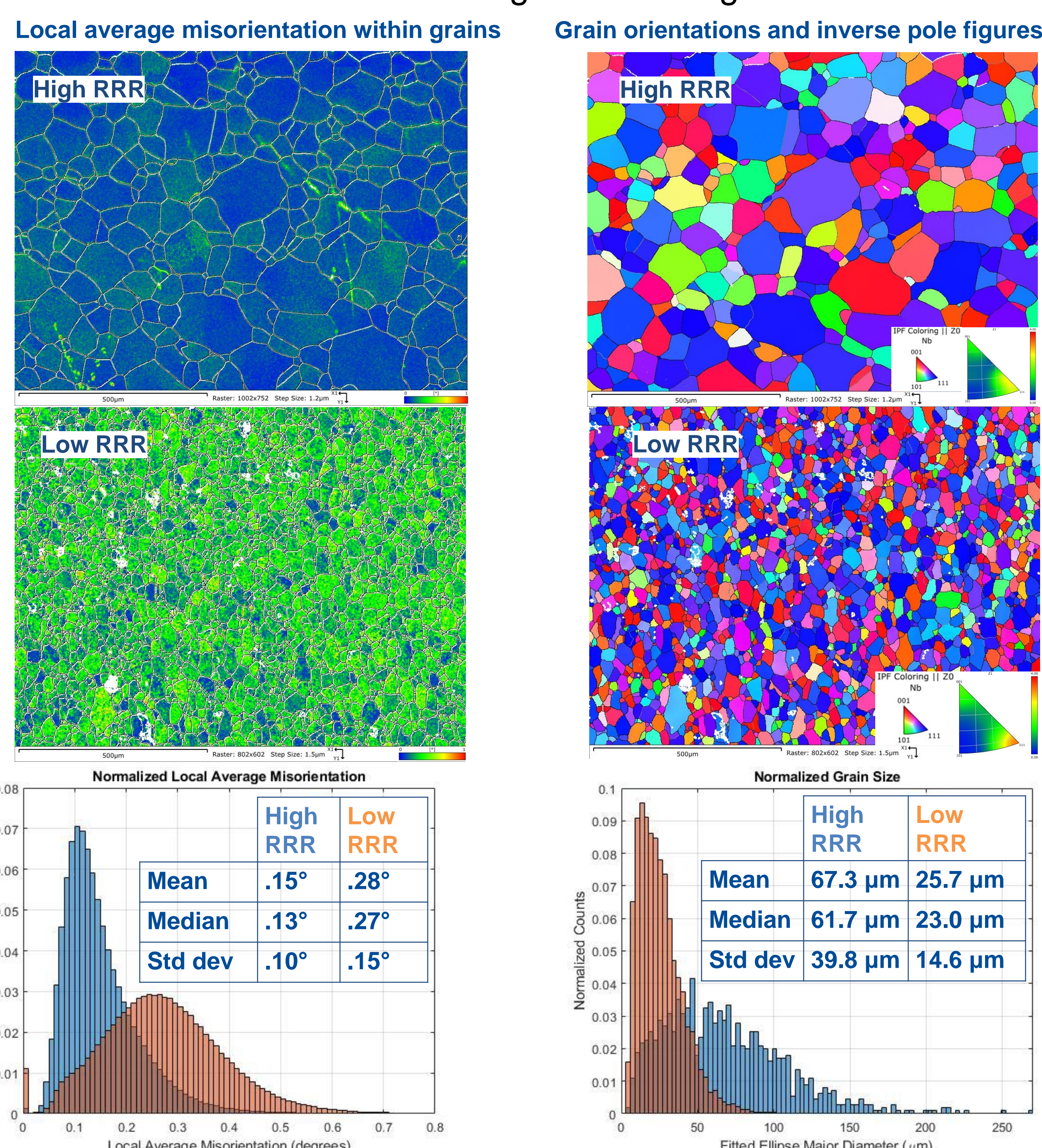


Impurity Profiles

- RRR is a bulk property so we only care about deep in bulk
- Impurity concentration is not primary reason for low RRR**



Grain Structure: small grains with high misorientation



Conclusions

- Low RRR cavity has:
 - Mitigated HFQS
 - High residual resistance
 - Low BCS resistance
 - Increased heating at quench location
- Low RRR material has:
 - Less hydrogen in bulk
 - No “key” impurity in bulk
 - High misorientation within grains**
 - Small grains**
- The low RRR is caused by misorientation of Nb lattice itself, not from impurities
- Does high misorientation mitigate grain growth?
- Are high misorientation/small grains synonymous with low mean free path?

Next Steps

- Measure strain in lattice via X-ray diffraction
- Repeat sample measurements after high temperature anneal to promote grain growth

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