

Application of zone refining to the development of NaI(Tl) detectors for SABRE North

Sabre North collaboration

The SABRE Project: Sodium Iodide with Active Background REjection

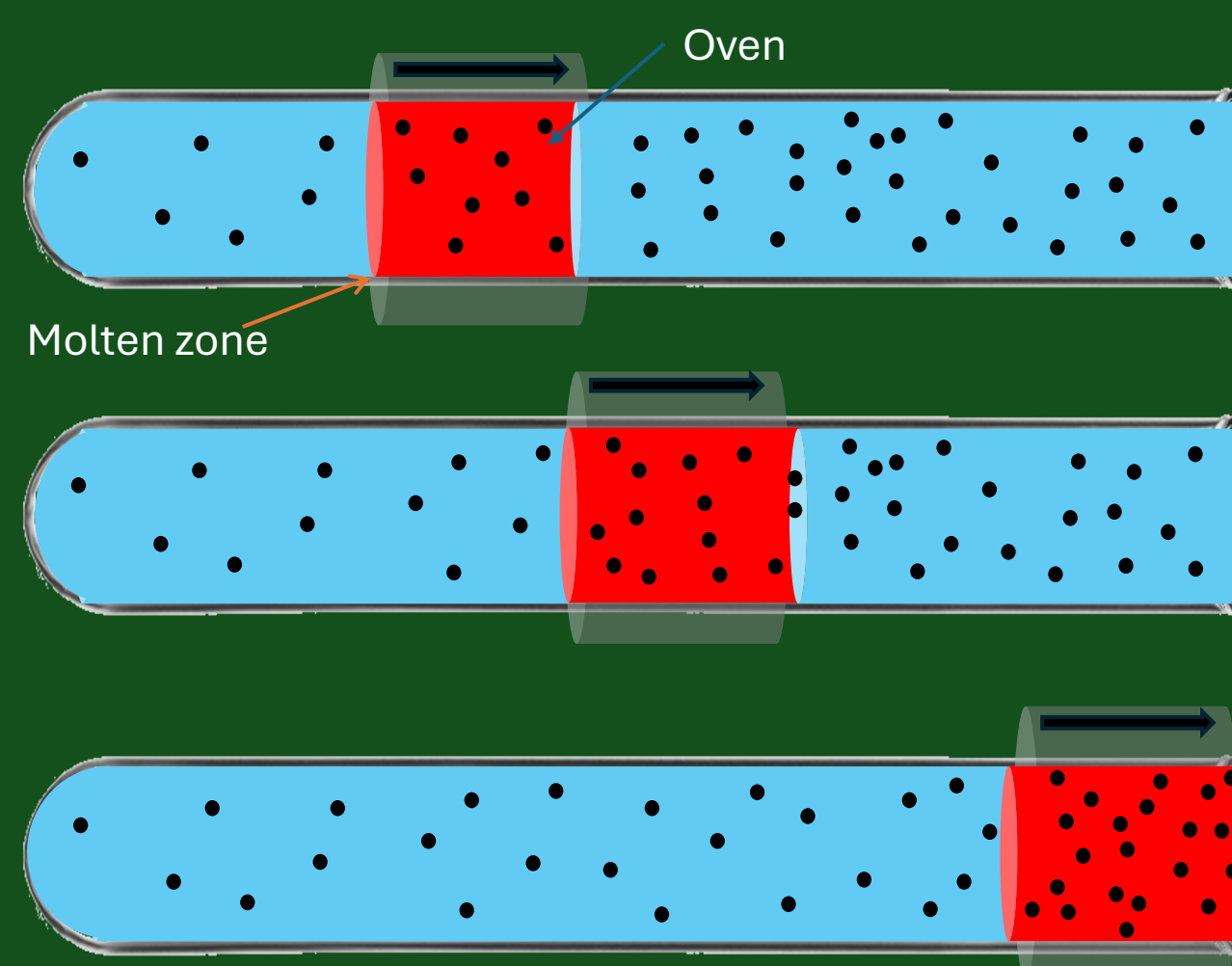
GOAL: Search for Dark Matter through the annual modulation of the experimental rate.

STRATEGY:

- Development of ultra-high purity NaI(Tl) crystals (~0.5 counts/keV/kg/day)
 - High purity NaI powder - further improved with Zone Refining technique
 - Clean crystal growth method
 - Reducing total detector background will increase the relative amplitude of a DM modulation signal $S_m/(S_0+B)$, 1-2% of the rate in DAMA/LIBRA.
- Low energy threshold
 - High QE PMTs directly coupled to the crystal
- Passive shielding + active veto
 - Unprecedented background rejection and sensitivity with a NaI(Tl) experiment
 - With active veto order of 90% K background rejection (only SABRE South)
- Two identical detectors in northern (LNGS) and southern hemispheres (SUPL)
 - seasonal backgrounds have opposite phase in northern and southern hemispheres
 - dark matter signal has same phase

Zone Refining (ZR) Technique

A narrow region of a crystal is melted, and this molten zone is moved through the crystal. The molten region melts impure solid at its forward edge and leaves a wake of purer material solidified behind it as it moves through the ingot.



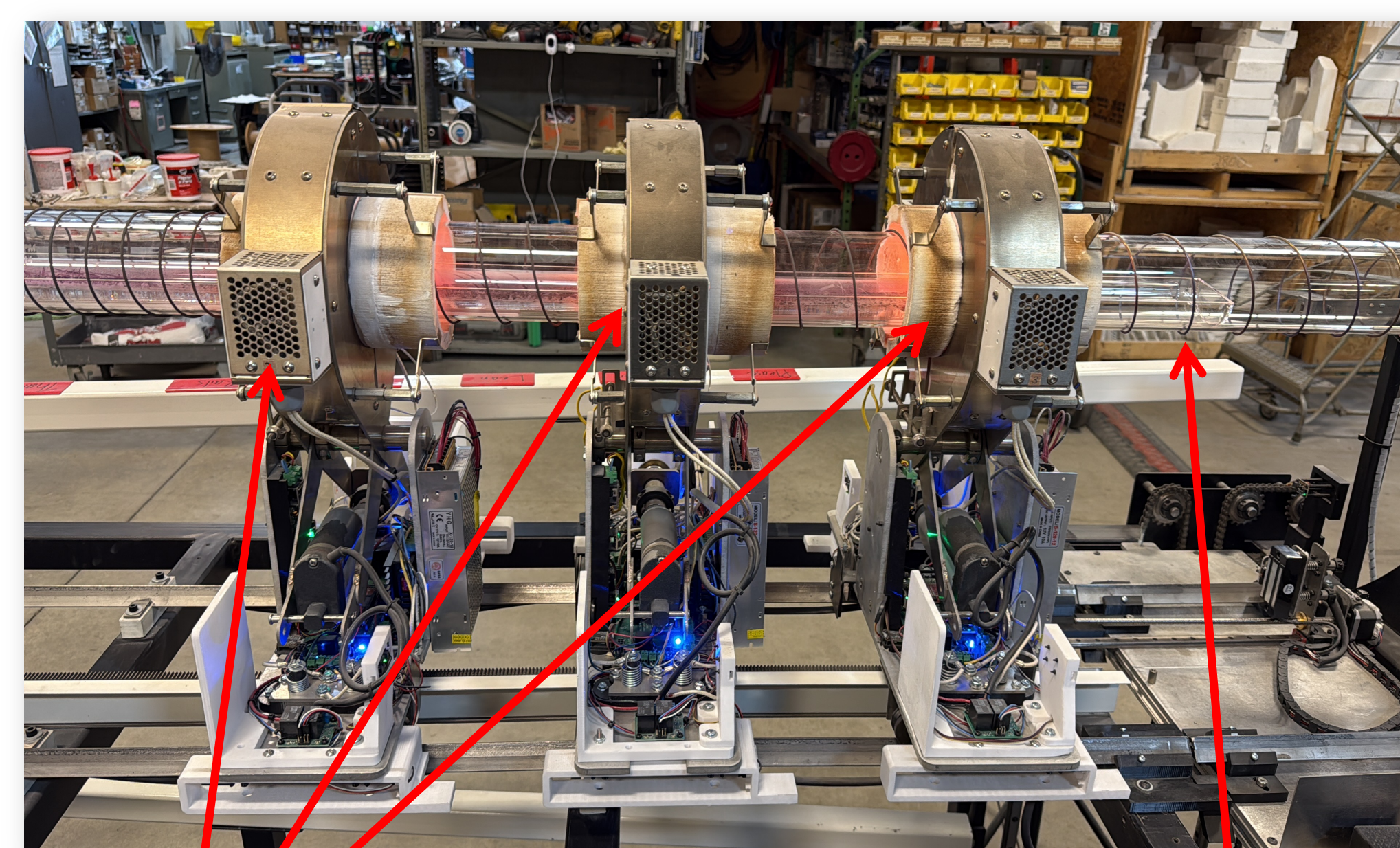
After solidification in the interface between liquid and solid a fraction k (coefficient of segregation) of impurities contained in the liquid remain in the solid

$$C_s = kC_L$$

where C_s (C_L) is the concentration of pollutant per volume unit in the solid (liquid). If $k < 1$ ($k > 1$) the pollutant remains preferably in the molten (frozen) zone.

The procedure can be repeated many times to increase the purity of the ingot. After cutting the tail we remain with a purified ingot.

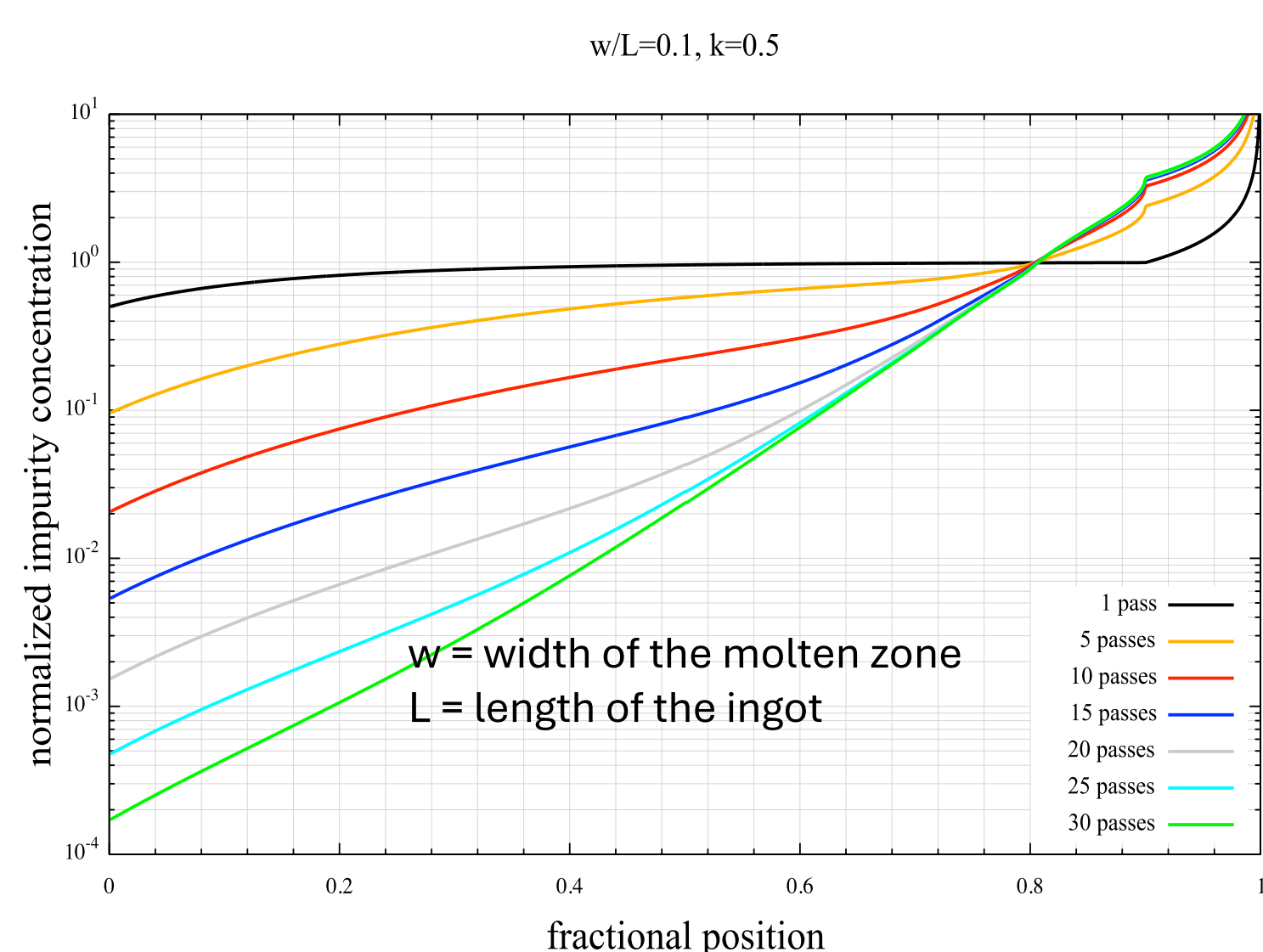
Zone refiner at MELLEN company (Concord, NH)



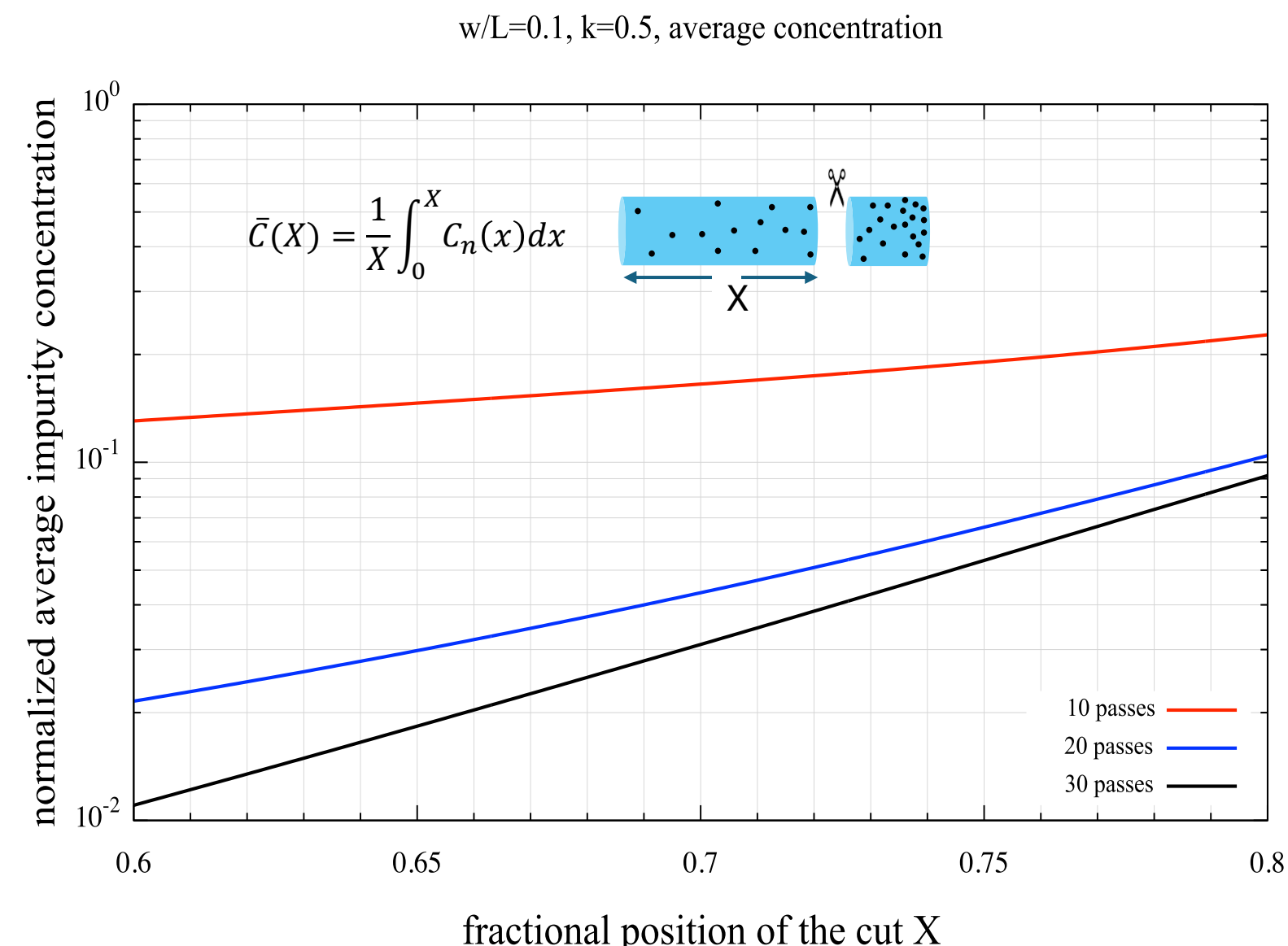
Ovens

Ampoule

Distribution of impurities after n passes

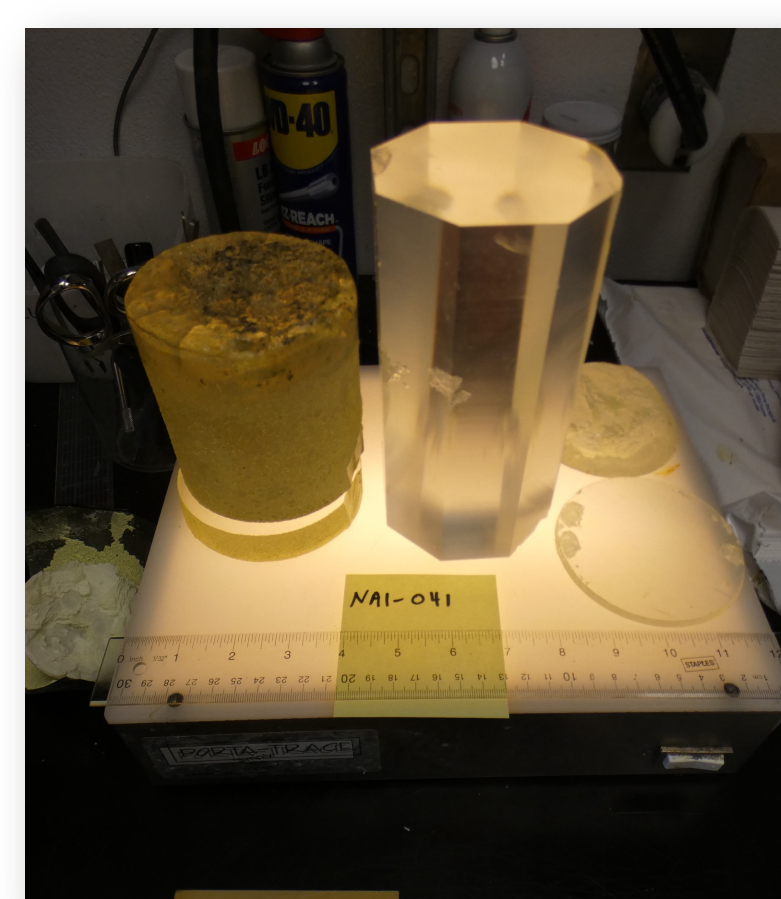


Average impurity concentrations as function of the cut



ZR Experimental Procedure

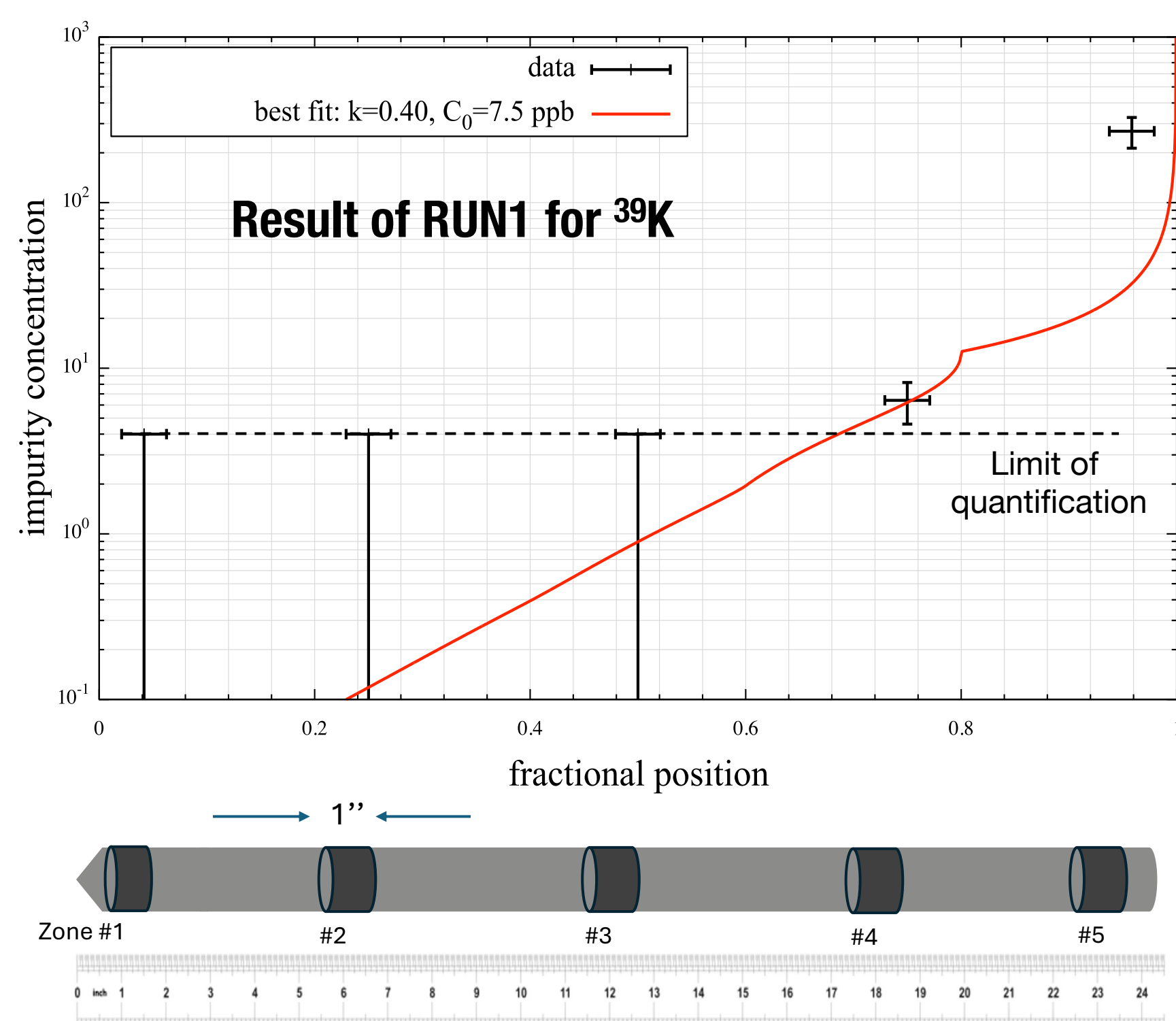
- Astrograde (former sigma-Aldrich) NaI powder used to grow the crystal
- Ampoule preparation @ RMD (Boston) with carbon coating or $SiCl_4$ vapour treatment to prevent sticking of molten powder to ampoule
- Zone refining done @ Mellen Company (Concord)
- Ampoule length $L=122cm$ (48"), molten zone width $w=12cm$
- 26 passes @ 1.5" per hour
- The purified part of the ingot is crumbled and recast in a single crystal through the *Bridgman method* i.e., by melting the NaI in a vertical cylindrical crucible which is extracted very slowly (1cm/day) from the oven.
- Two runs have been considered for this analysis (RUN1 and RUN2)



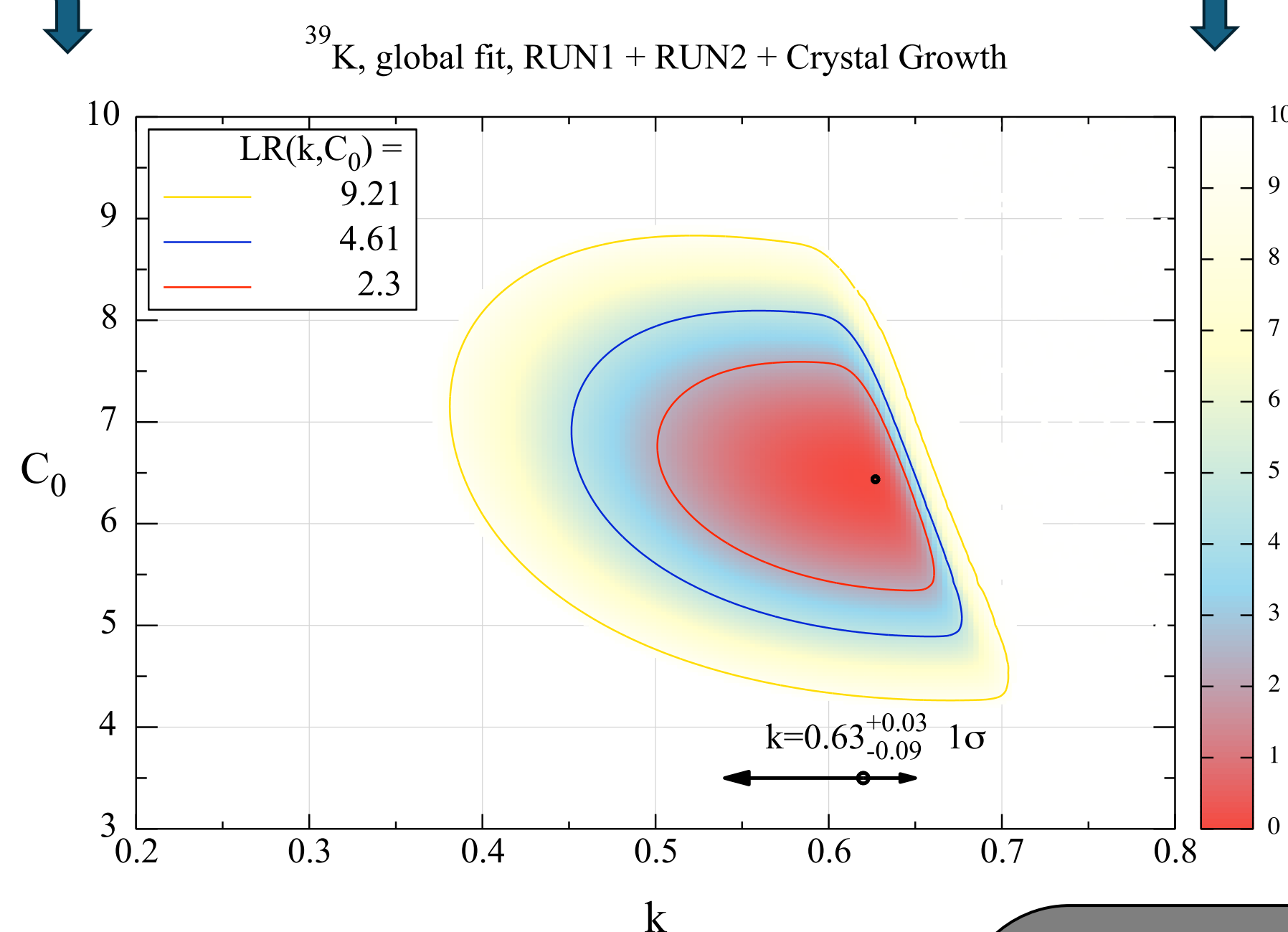
ZR Sample Analysis

- Five samples in RUN1 and four in RUN2 have been extracted from the refined ingot
- Various contaminants (in particular, ^{39}K and ^{208}Pb) have been measured using inductively coupled plasma mass spectrometry (ICP-MS) at Seastar
- Also, 3 samples of the final crystal have been also analyzed
- For each sample, a 20% assumed systematic error was added in quadrature to statistical error
- A maximum likelihood analysis has been performed to extract the initial impurity concentration C_0 and the segregation coefficient k .

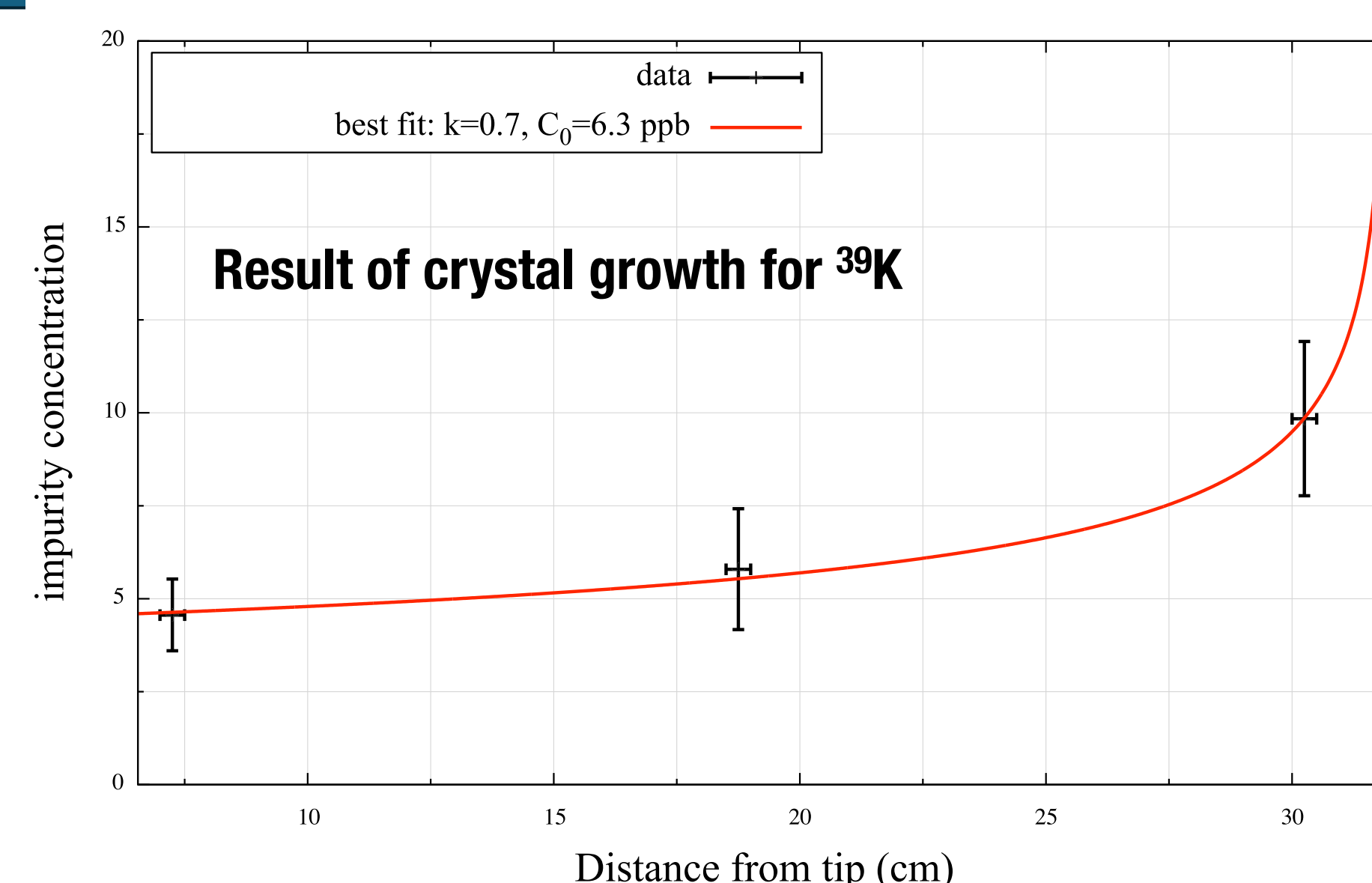
^{39}K , $W/L=0.2$, $N_{pass}=26$, RUN 1



Combined Analysis (^{39}K)



^{39}K , crystal growth



Conclusions

Zone-refining has shown to be an efficient method to purify NaI crystals

- Potassium is reduced by factor 5 (but further improvement is achievable). ^{40}K is the main radioactive background for Dark Matter detection
- Unfortunately, the technique cannot reduce lead contamination since the segregation coefficient for Pb is $k \approx 1$

➤ More details can be found in the paper [arXiv:2601.15998](https://arxiv.org/abs/2601.15998) (submitted to JINST)



	^{138}Ba	^{44}Ca	^{39}K	^{24}Mg	^{208}Pb	^{88}Sr (RUN2)
$f_{pur, best}$	0.06	1.21	0.24	1.	1.1	1.23
$k \in 1\sigma$ range	$0.03 \div 0.14$	$1.20 \div 1.22$	$0.13 \div 0.31$	$0.94 \div 1.04$	$1.08 \div 1.13$	$1.22 \div 1.24$
$\bar{C}$, best (ppb)	0.018	45.1	1.55	1.97	1.98	0.61
$(k, C_0) \in 1\sigma$ allow.	$0.008 \div 0.036$	$38.6 \div 51.7$	$0.64 \div 1.80$	$1.76 \div 2.0$	$1.78 \div 2.18$	$0.61 \div 0.62$

Coefficient of purification $f_{pur} = C/C_0$ and coefficient of segregation k for 6 selected contaminants for 80% cut of the initial part the ingot