

# **SIMS MEASUREMENT of OXYGEN in FZ-SILICON**

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In order to detect O at levels relevant to diffusion into FZ Si, it was necessary to employ  $\text{Si}^{18}\text{O}_2$  as a source; [Mikkelsen, 1986]

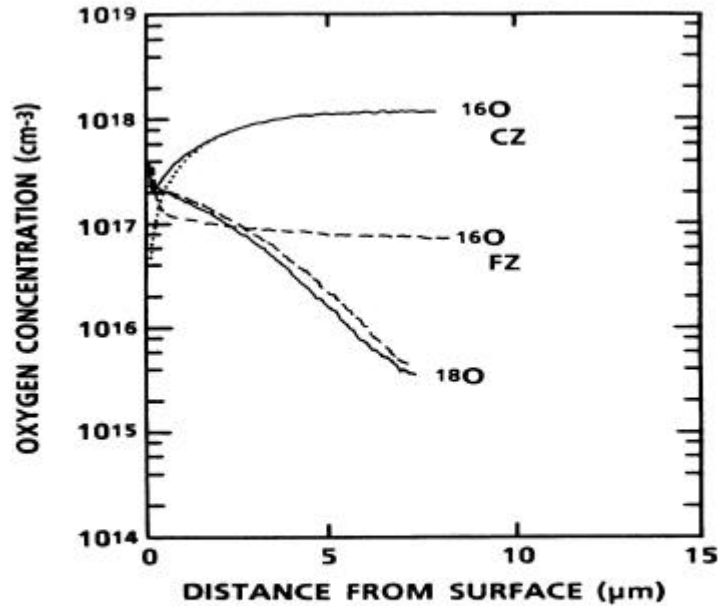


Fig. 1. SIMS profiles of CZ and FZ crystals indiffused under 1 atm  $\text{H}_2^{18}\text{O}$  oxidations at 1000C for 30 min. Dotted curve is the expected  $^{16}\text{O}$  out-diffusion profile for the CZ crystal.

Our first results in ROSE using  $^{16}\text{O}$ ; note decreasing concentration at surface

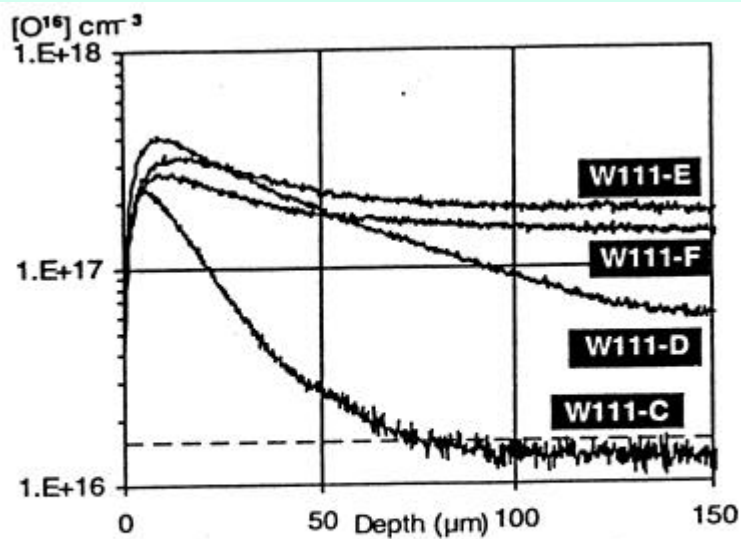
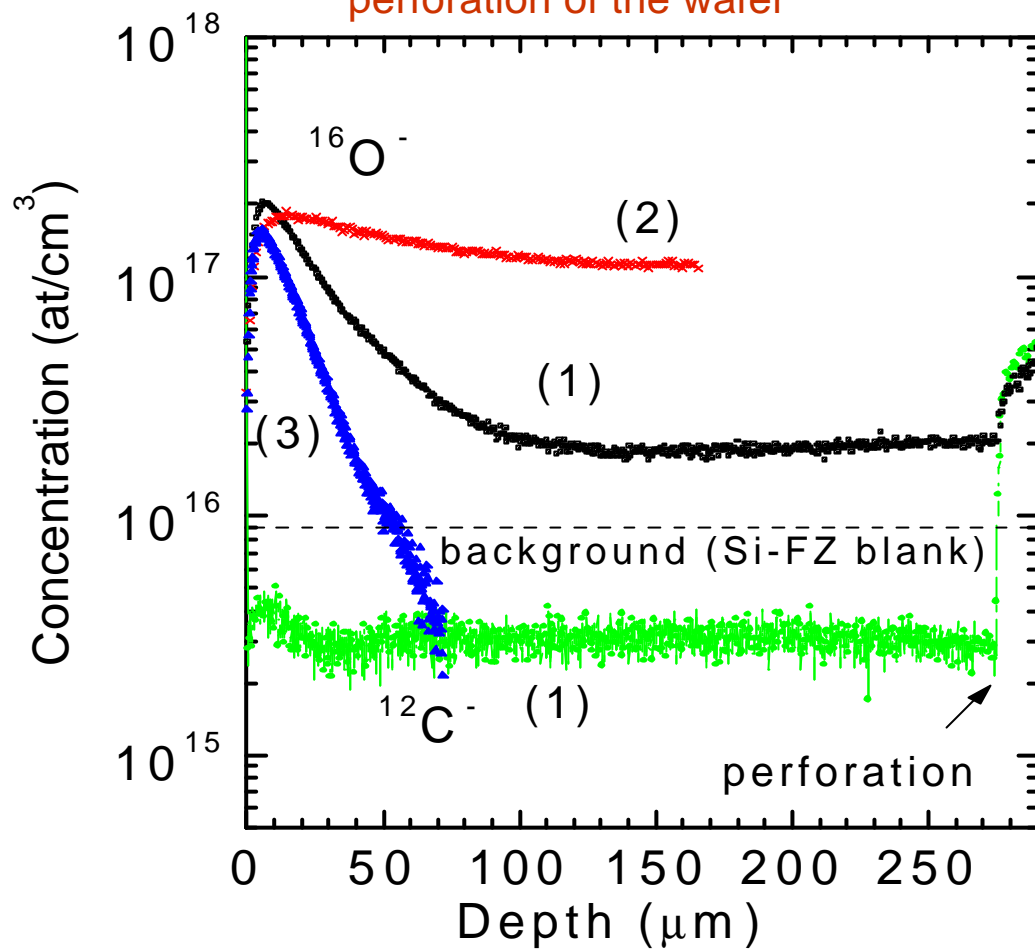


Fig. 2. Oxygen concentrations obtained with different processing times.

To check correctness of the profiles, one analysis was done until perforation of the wafer

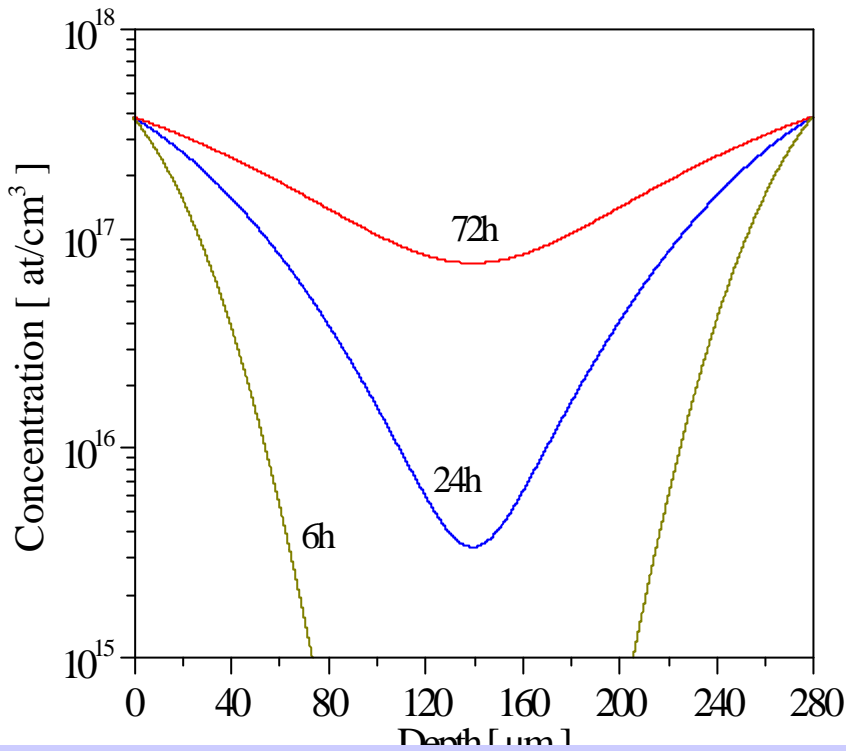


Oxygen depth profiles for FZ-Si wafers annealed at 1150 °C for:  
(1) – 16 hrs, (2) – 72 hrs, (3) – 6 hrs.

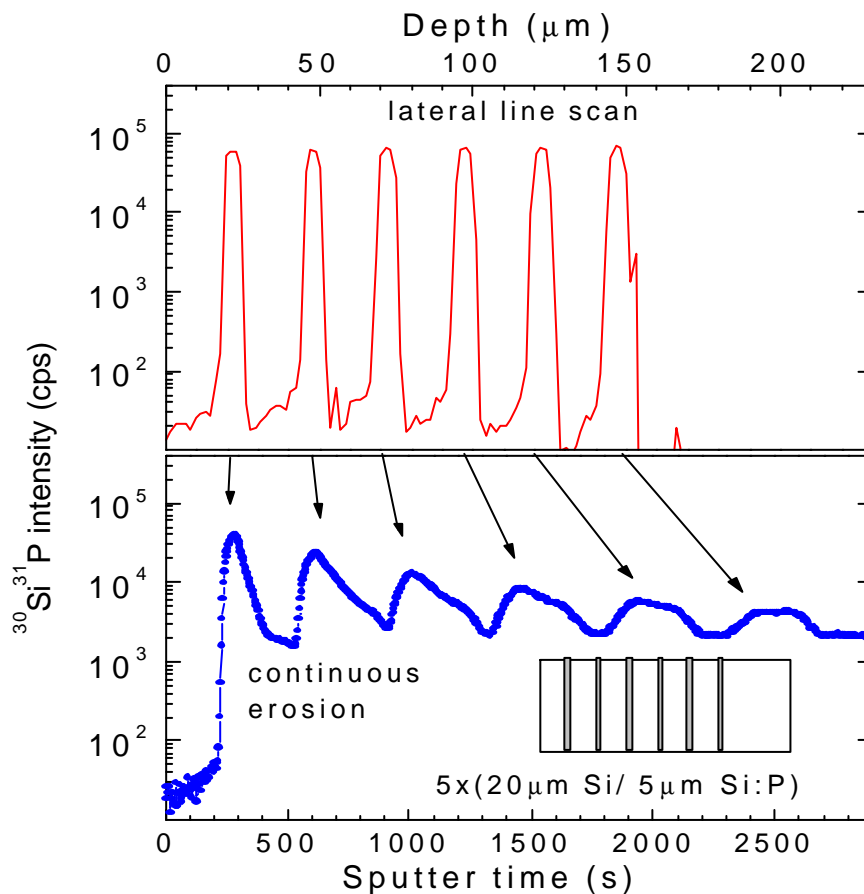
immediate observations :

- i – the profile 1 is not symmetrical with respect to the mid-thickness as it should be since diffusion proceeds from both sides of the (oxidized) 280 μm thick wafer.
- ii – the O signal levels off at  $\sim 2 \times 10^{16}$  i.e. well above the *bkg*: hence, direct background correction not feasible.
- iii – the mean sputter rate differs markedly from that established at shallow depths from either stylus profilometry or the location of the implant maximum.

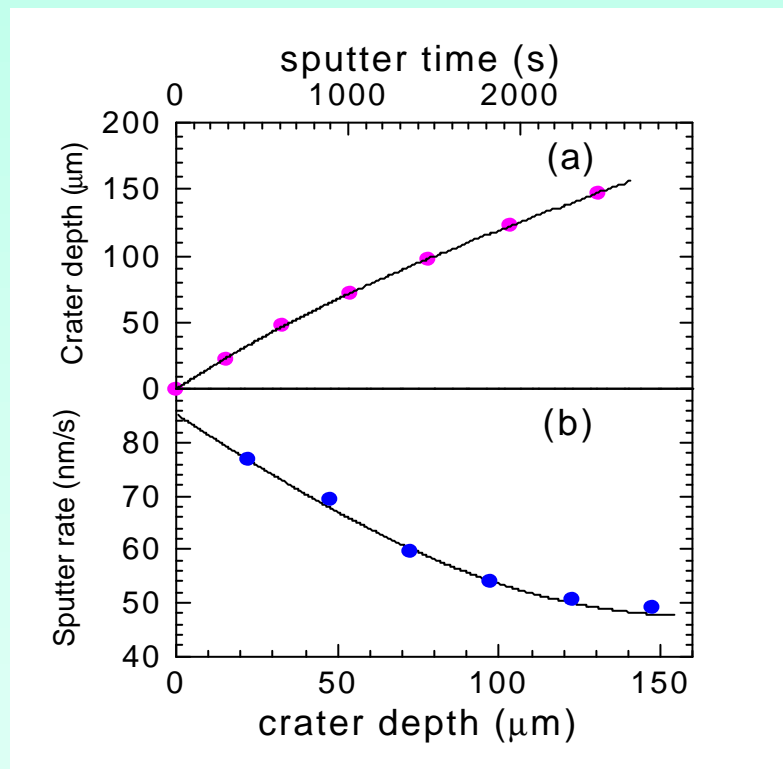
## Calculated curves for diffusion into a slab



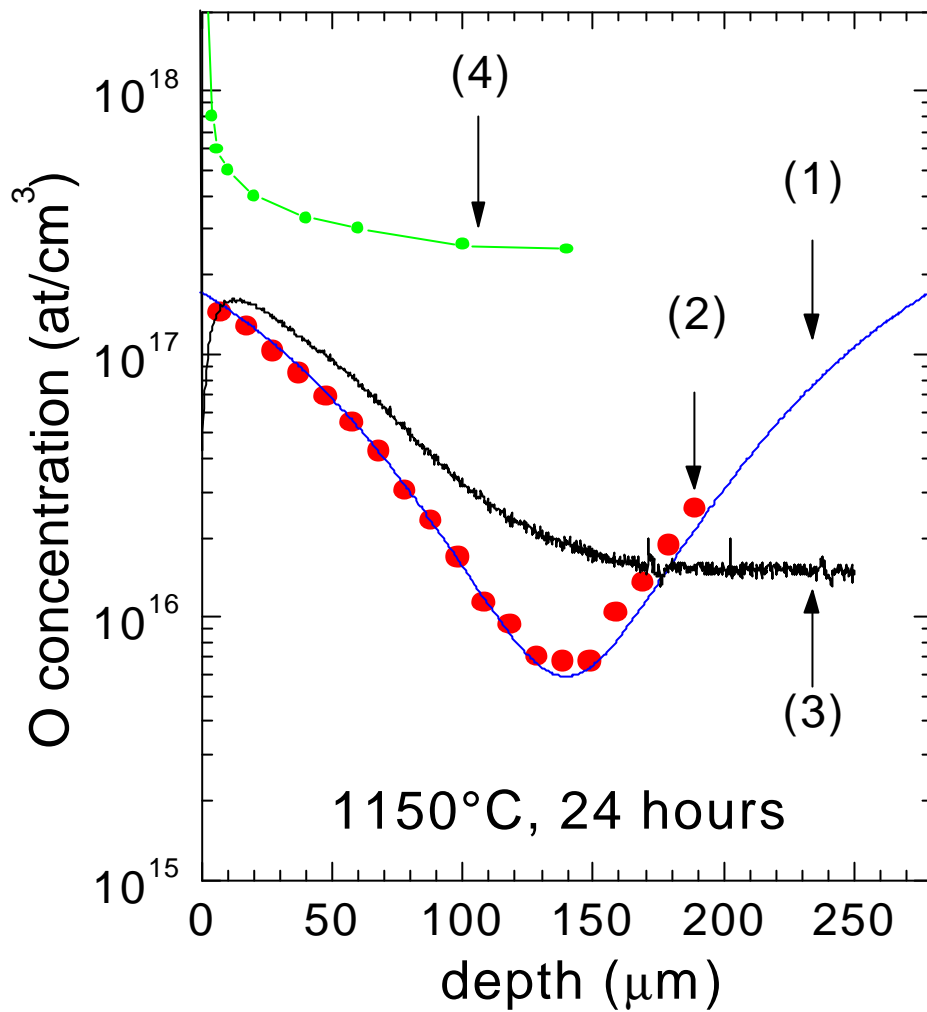
To verify the degree of reliability - a test epi-structure was manufactured for step-by-step measurement on beveled surface (top) or by standard profiling (bottom)



The loss of depth resolution is evident, also the removal rate is not constant; a polynomial fit was derived to enable corrections



$$Y = -0.398 + 0.085t - 1.71t^2 + 3.408t^3 - 2.632t^4$$



Comparison of different results concerning deep O diffusion into FZ Si: (1) calculated, for  $D=2e-10$ ,  $C_0=1.7e17$ , (2) line scan with *bkg* correction,; continuous profiling: (3) – this work, (4) - ref. [6]

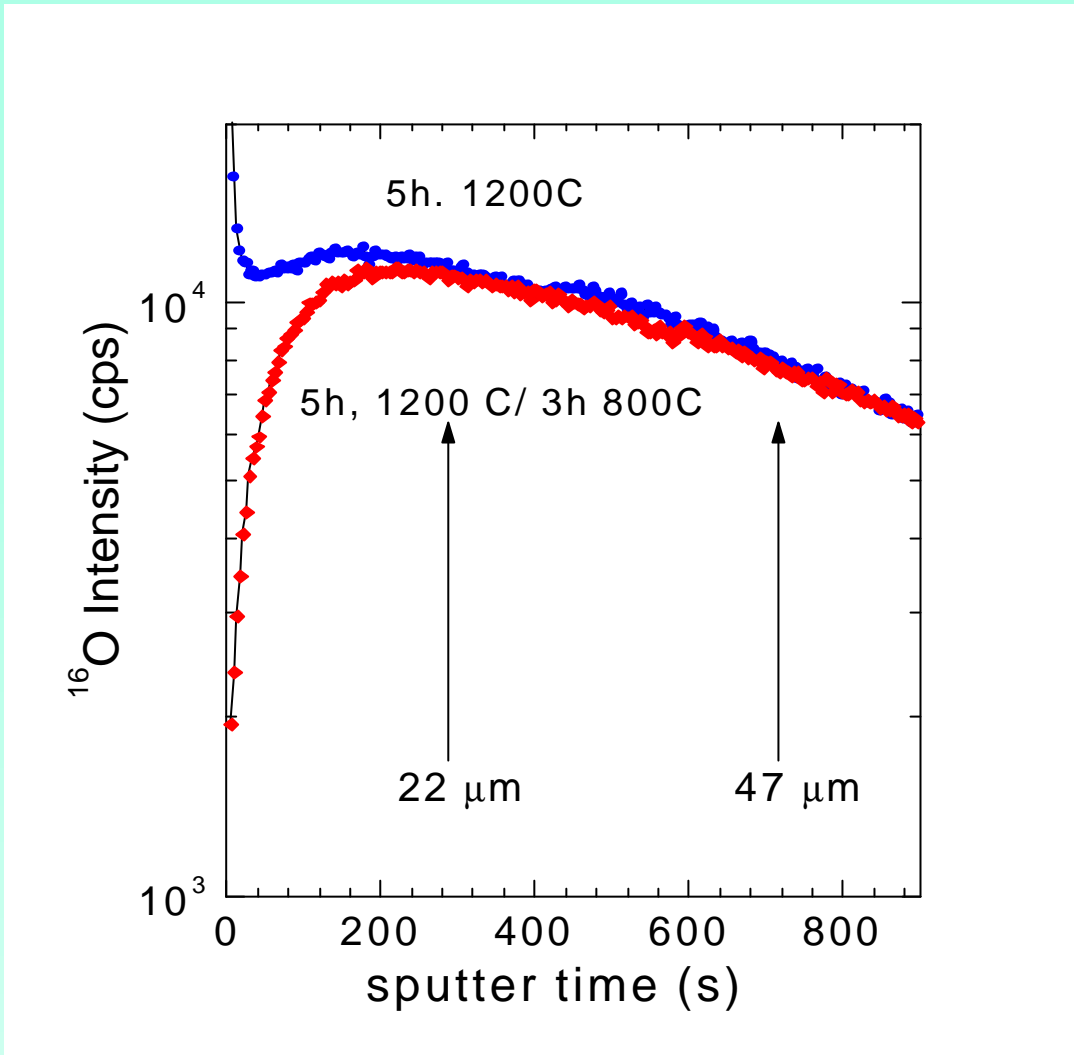


Figure showing the origin of the observed „depletion” at the surface: it takes place upon prolonged cooling and segregation back to the oxide

## CONCLUSIONS

When a dynamic range of at least two orders of magnitude is required over a depth exceeding 50-70  $\mu\text{m}$ , the beveling of the sample followed by line-scan analysis is necessary.

For shallower [O] distributions, a continuous analysis can be applied; however, the variation of the sputter rate with depth should be taken into account.

When investigating deep diffusion together with the process of oxygen segregation at the  $\text{SiO}_2$  – Si interface, both methods are recommended.