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Standalone Arapuca Analysis

The X-Arapuca without Quartz window (XN) will see the sum of the three spectra, with the assumption that the three wavelengths are shifted with similar quantum efficiency.

$$\begin{aligned}\frac{dXN}{dt}(\text{scint}@128nm + 150nm + 178nm) \\ = K \left(\frac{\tau_{TA}}{\tau_{128}} e^{-\frac{t}{\tau_{TA}}} + \frac{\tau_{TA}}{\tau_{AX}} \frac{(e^{-t/\tau_{TA}} - e^{-t/\tau_{TX}})}{(\tau_{TA} - \tau_{TX})} \right)\end{aligned}$$

The X-Arapuca with the Quartz window (XQ) will only be sensitive to the third spectrum (the one from XeXe*)

$$\frac{dXQ}{dt}(\text{scint}@178nm) = (1 - \varepsilon) K \frac{\tau_{150}}{\tau_{XX} + \tau_{150}} \frac{\tau_{TA}}{\tau_{AX}} \frac{(e^{-t/\tau_{TA}} - e^{-t/\tau_{TX}})}{(\tau_{TA} - \tau_{TX})}$$

The XN and XQ spectra at the different Xe concentrations can be fitted 'simultaneously' with the $\frac{dXN}{dt}$ and $\frac{dXQ}{dt}$ functions to extract the common value of τ_{TA} and τ_{TX} .

A linear fit of $\frac{1}{\tau_{TA}}$ and $\frac{1}{\tau_{TX}}$ as a function of the Xenon concentration could allow to estimate estimate of τ_{AX} and τ_{XX} :

$$\frac{1}{\tau_{TA}} = \frac{1}{\tau_{128}} + \frac{1}{\tau_{N2}} + \frac{1}{\tau_{AX}} = (a + b \text{ } Xe[\text{ppm}]) \mu s^{-1}$$

$$\frac{1}{\tau_{128}} + \frac{1}{\tau_{N2}} = a \text{ } \mu s^{-1}$$

$$\frac{1}{\tau_{N2}} = (a - \frac{1}{1.6}) \text{ } \mu s^{-1}$$

$$\tau_{N2} = 1/(a - \frac{1}{1.6}) \text{ } \mu s$$

$$\frac{1}{\tau_{TX}} = \frac{1}{\tau_{150}} + \frac{1}{\tau_{XX}} = c + d \text{ } Xe[\text{ppm}] \mu s^{-1}$$

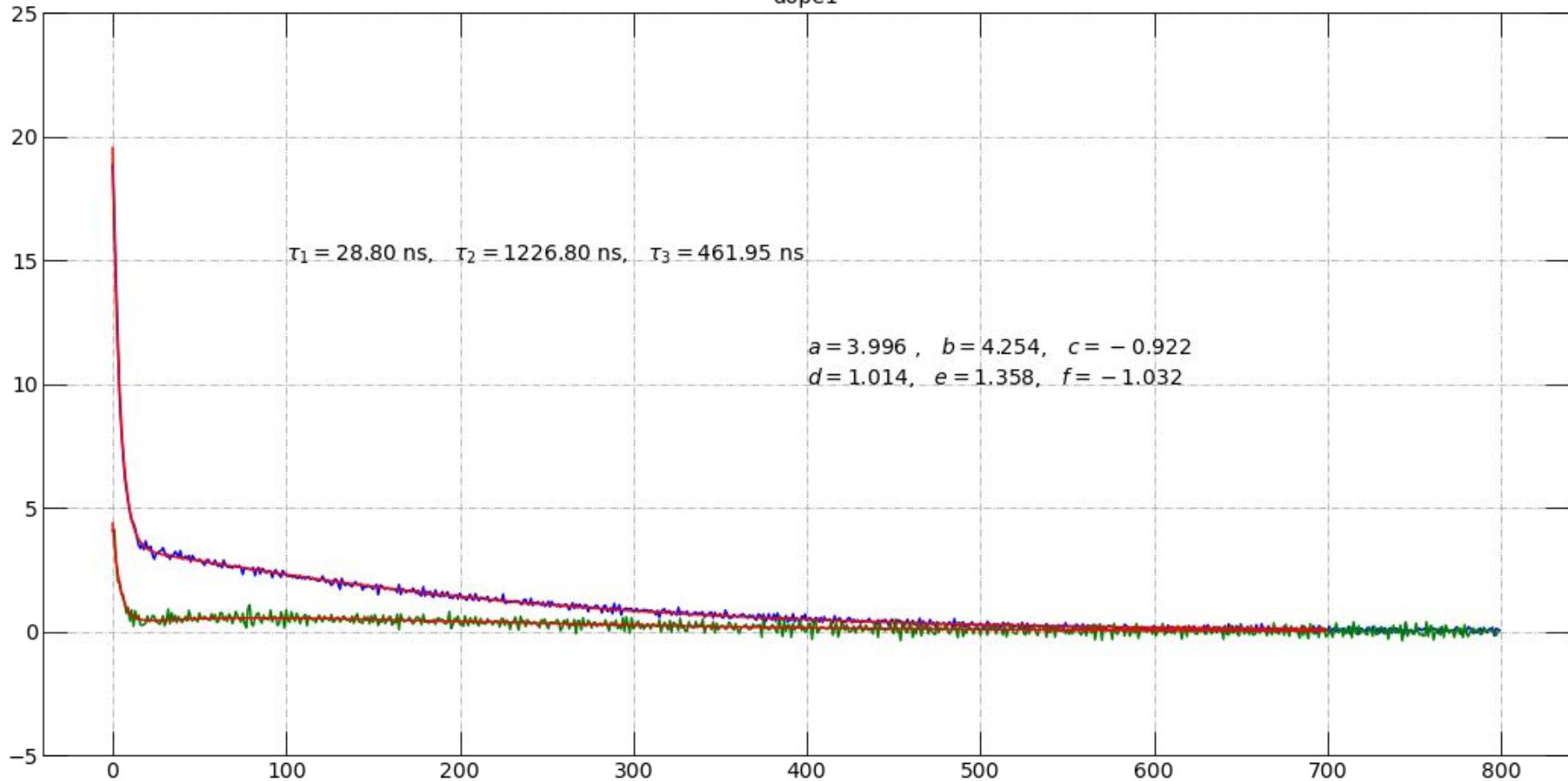
$$\frac{1}{\tau_{AX}} = b \text{ } Xe[\text{ppm}] \mu s^{-1}$$

$$\tau_{AX} = \frac{1/b}{Xe[\text{ppm}]} \mu s$$

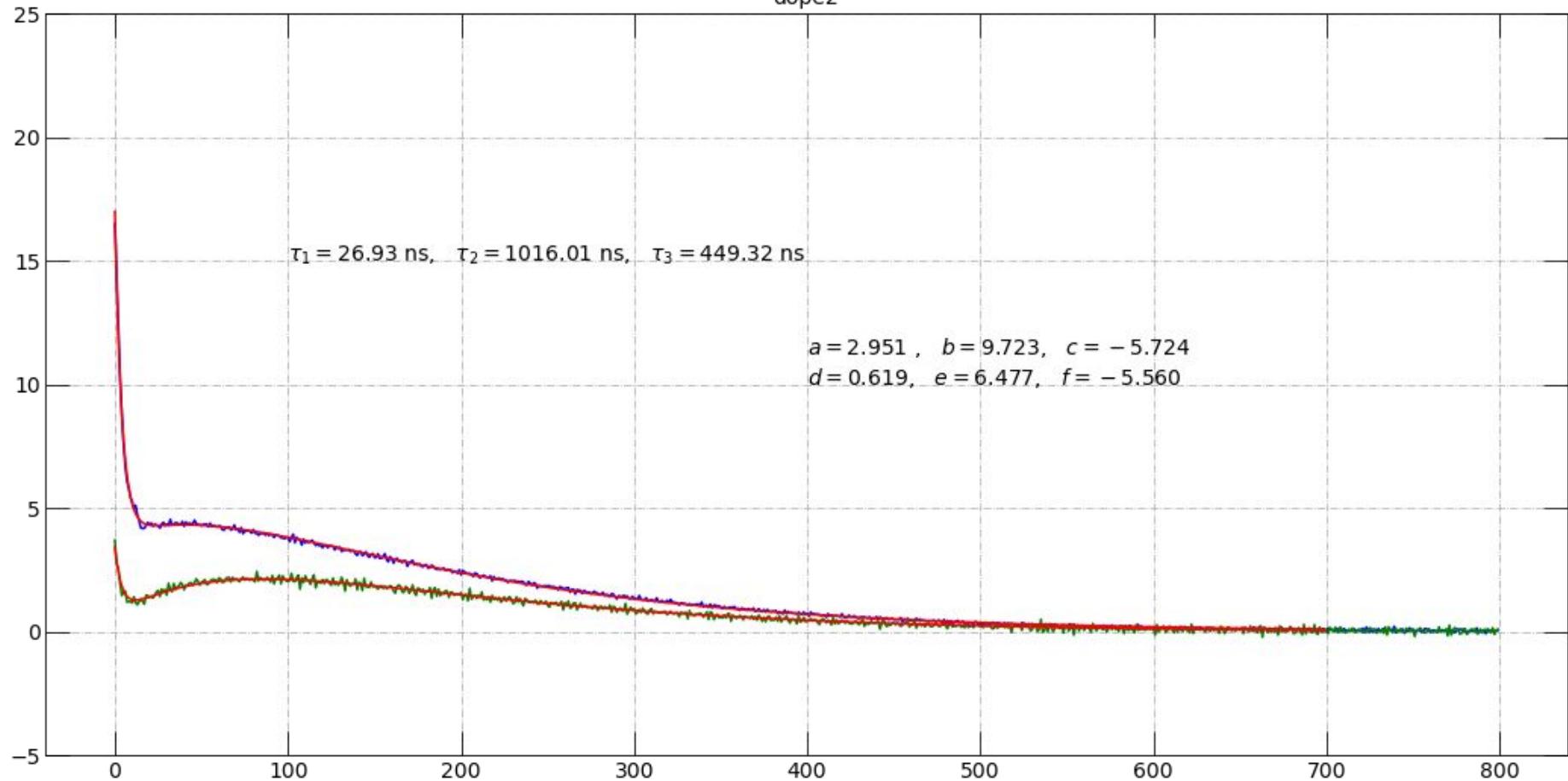
$$\tau_{150} = \frac{1}{c} \mu s$$

$$\tau_{XX} = \frac{\frac{1}{d}}{Xe[\text{ppm}]} \mu s$$

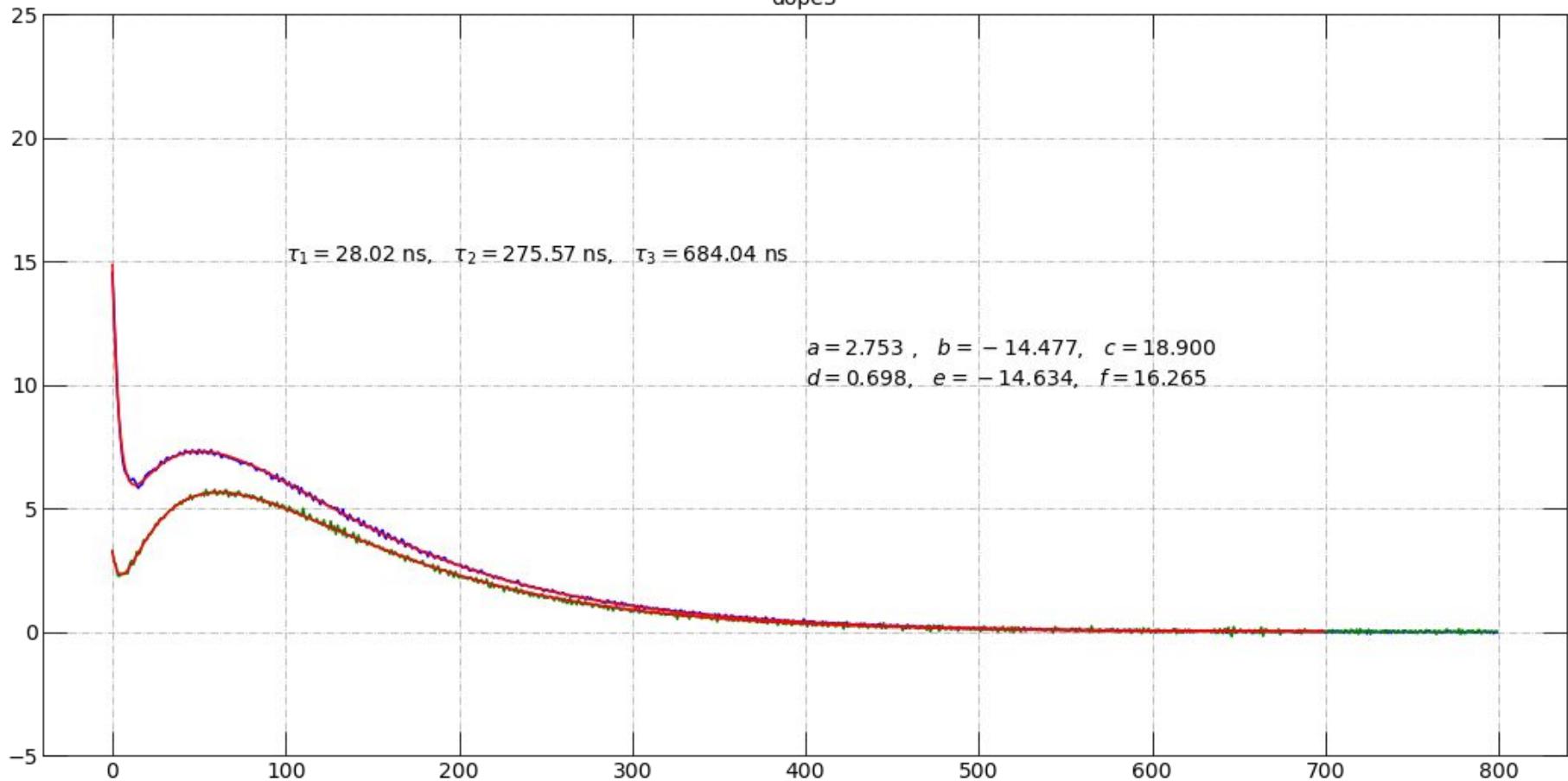
dope1



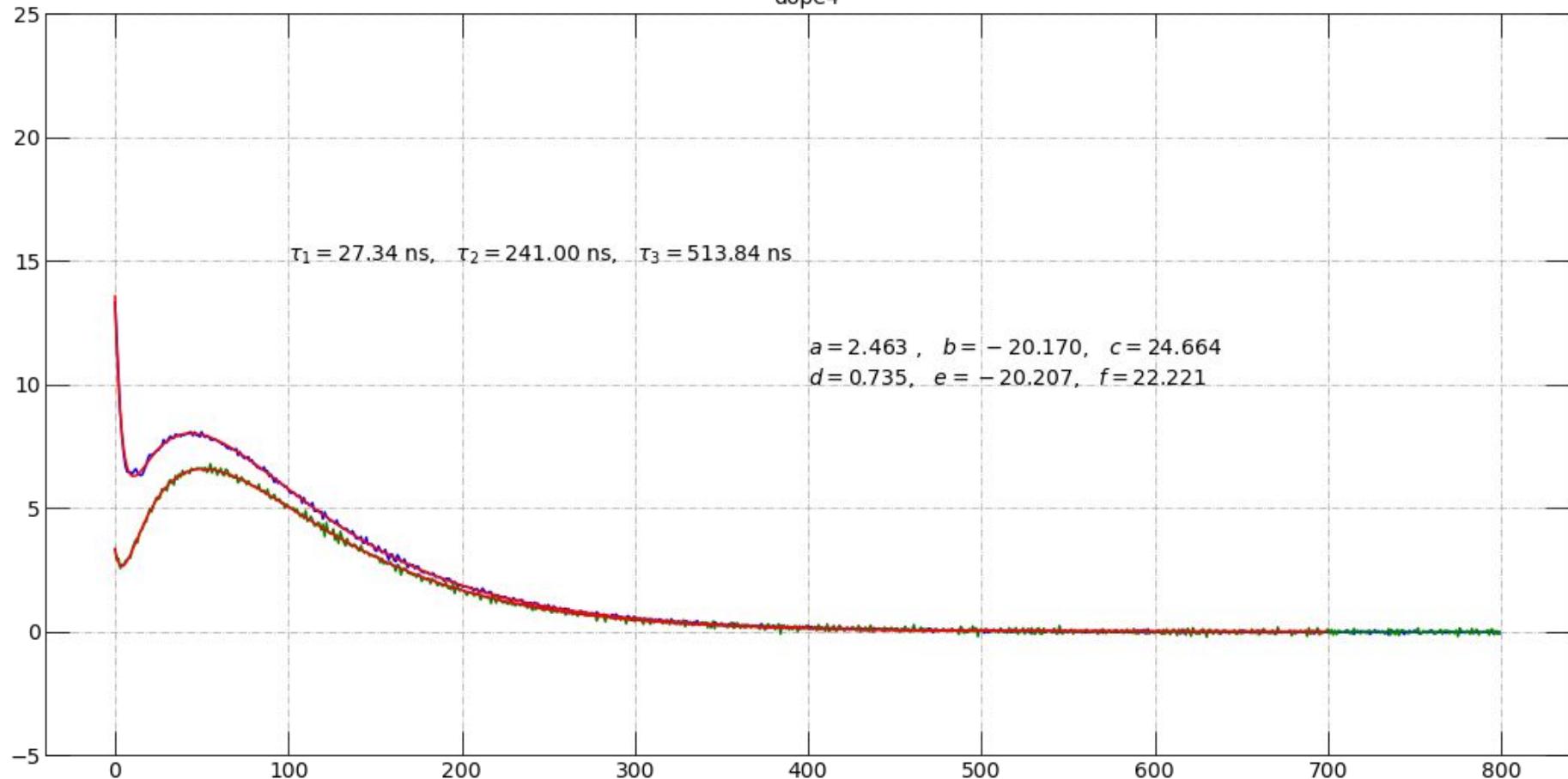
dope2



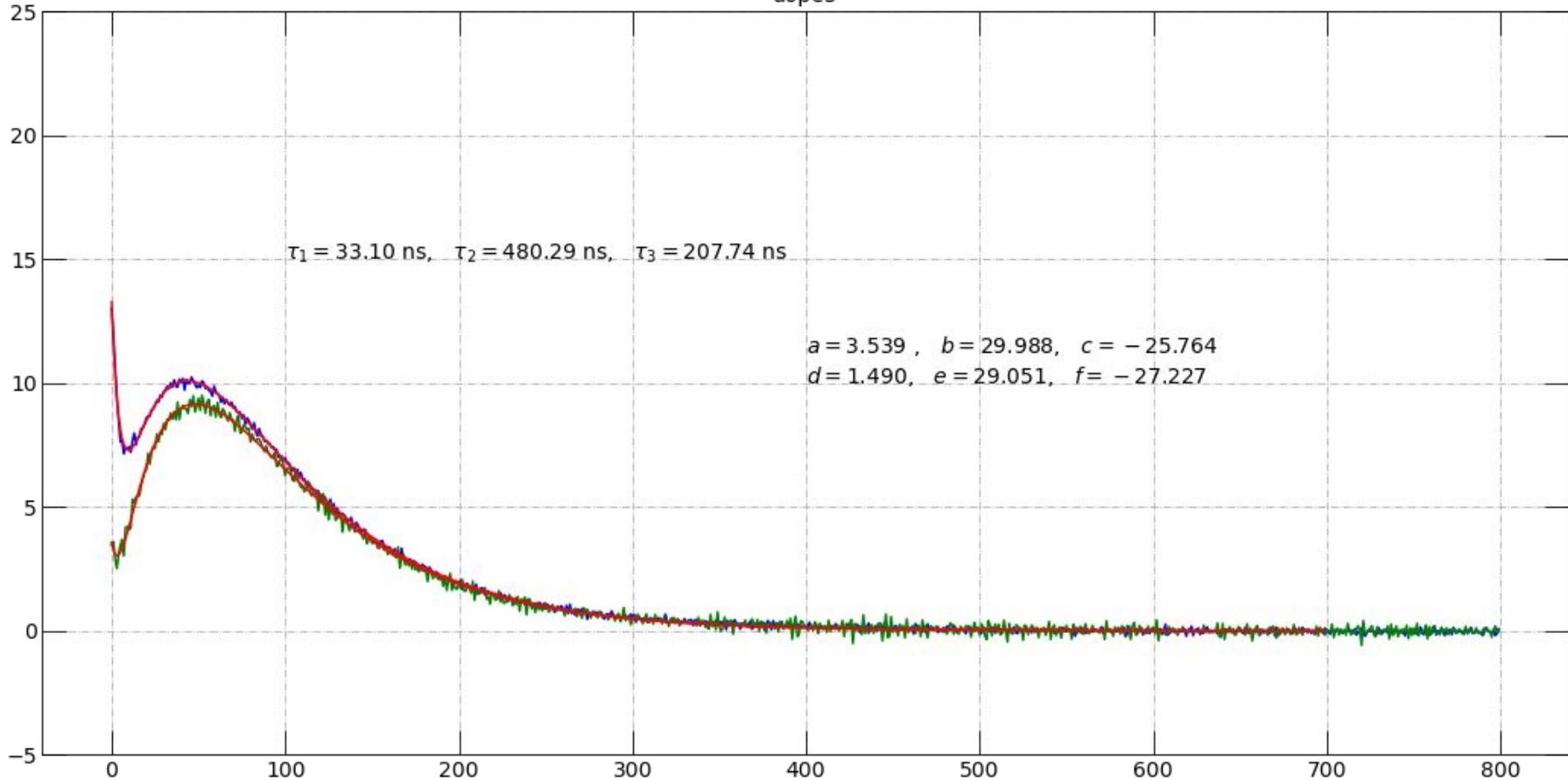
dope3



dope4



dope5



—●— 1/tau2
—●— 1/tau3

— y = 1.5317 + 0.18691x R = 0.99108

— y = 0.68279 + 0.073093x R = 0.99344

