

# T3R Antibunching - Slow Decay

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HowTo (Experimental); SymPhoTime; How to perform Hanbury Brown Twiss Antibunching experiments; Diamond NV centers; MicroTime 200; Microtime Specials; Antibunching Measurements; Polarisation resolved measurements with the MT200 (for internal use only); total correlation

### Q.: How can we avoid decay at very long time delay (up to 5us)?

#### In this show-case the external preconditions are:

- detectors: perkin elmer SPCMAQR14
- 700 nm SP filter in front of one of the apds
- sample: diamond nanocrystals (100 nm), we are measuring the antibunching of the NV centers emission
- laser wavelength: 532 nm, cw
- T3R Measurement with PicoHarp Software

# A.: The decay can be caused by blinking of the nanodiamonds.

### A decay caused by correlation statistics

On top of an eventual blinking you will have a decay caused by correlation statistics.

This happens because you correlate the first arriving photon against a photon at the longer time,

which will be an increasingly unlikely event because the earliest stop photons always win and the experiment restarts.

I attached a simulation.

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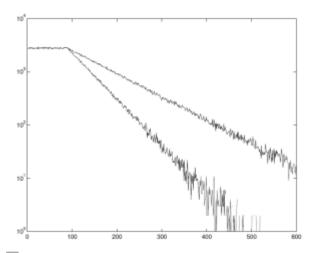


Fig. 1: Logfile logf.png

==== You will I need the raw data ==== The timescale is in ns. The plateau is the dead-time. The decay slope depends on the count rate. You can zoom into your data and you will see roughly the same behavior - linear decay after the dead-time, which than of course is overshadowed by the background. The way to avoid this is is to calculate the total correlation. This is not a start-stop correlation but the correlation of every photon against every photon. It can be done measuring in T2 mode and then correlating using the SymPhoTime Software (http://www.picoquant.com/products/sw\_mt/sw\_mt.htm). For a trial you will I need the raw data - a pt2 file containing all the photon data - not just the correlated curve. You can record this using the button I highlighted in the attached screenshot.

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