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# Time gated Fourier transform spectroscopy with burst excitation for time-resolved spectral maps from the nano- to millisecond range†

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**We demonstrate burst-mode Time Gated Fourier Transform Spectroscopy (bmTG-FTS), a technique for simultaneously capturing and disentangling emission signals from short- (ns) and long-lived ( $\mu\text{s}$ –ms) states. We showcase the possibilities of the technique by preparing time gated temporal-spectral maps from a dual-emissive DNA-stabilized silver nanocluster (DNA-AgNC).**

Among the most essential parameters for the characterization of novel luminescent materials are the spectral features, which include emission maximum and shape, and luminescence decay time.<sup>1–9</sup> Emission spectra and decay curves are often measured under different experimental configurations. Even when recording decay curves, different equipment is often used depending on the decay time. Nanosecond decays are usually obtained with high repetition rate pulsed lasers and time-correlated single photon counting (TCSPC), while slower decays are traditionally acquired with low repetition rate pulsed Xe flashlamps.<sup>10,11</sup> We recently demonstrated how a burst-mode excitation scheme in combination with TCSPC could be useful for simultaneously measuring the temporal response of the short- and long-lived component of a DNA-AgNC with spectrally overlapping dual emission.<sup>12–15</sup> While this allowed for measuring temporal dynamics from the sub-ns to ms timescale in a single measurement, it did not yield any information on the spectral features.<sup>12</sup> In another recent publication, we showed how the insertion of an interferometer in the emission path of a standard TCSPC setup allowed for collecting and disentangling the spectra of short- and long-lived components.<sup>16</sup> In this case, however, the temporal dynamics of the long-lived component was unresolved.

In this contribution, we have effectively combined our two previously introduced methods into one,<sup>12,16</sup> which we term

burst-mode time gated Fourier transform spectroscopy (bmTG-FTS). With bmTG-FTS, a single measurement is conducted that allows for collecting spectral-temporal maps of disentangled short-lived fluorescence and long-lived luminescence contributions on the nano- to millisecond timescale. Thus, bmTG-FTS simultaneously provides some of the most important spectral and temporal parameters for the characterization of novel luminescent materials. In this paper, we will describe the working principle of bmTG-FTS and demonstrate its applicability with an exemplary DNA-AgNC that shows dual emission on the nano- to millisecond timescale.<sup>17</sup>

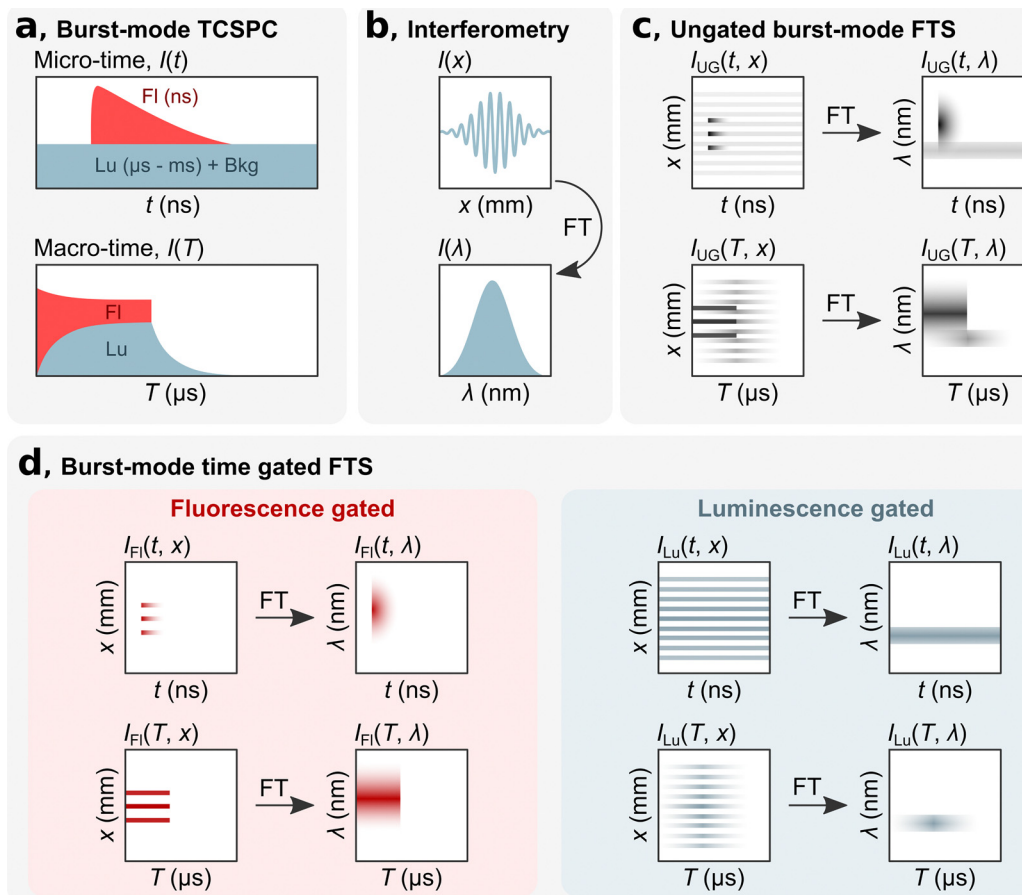
bmTG-FTS is based on three key elements: TCSPC hardware, an interferometer, and a high repetition laser (MHz) that can be switched on and off on a microsecond time scale. Using TCSPC, each detected photon is assigned a ‘micro-time’ and a ‘macro-time’ (Fig. 1a). The micro-time ( $t$ ) is the time between a detection event and the next sync pulse and has a sub-ns time resolution. This time domain is commonly used for determining the decay time of short-lived fluorescence (ns) that decays within two consecutive excitation pulses. In such measurements, long-lived luminescence ( $\mu\text{s}$ –ms) appears as a virtually flat background along with the detector dark counts and after pulsing events. The macro-time ( $T$ ) is the time between the start of the experiment to a detection event (note that what is recorded is the time of the next sync pulse). It has a time resolution equal to the reciprocal of the laser’s repetition rate (*e.g.*, 100 ns for a repetition rate of 10 MHz, or faster if an internal clock is used). This time domain can be used for measuring slower processes, like the rise and decay of a long-lived state ( $\mu\text{s}$ –ms).<sup>12</sup> By implementing a burst excitation scheme, it is possible to simultaneously measure the short- and long-lived response of a luminescent species. With the combined micro-macro information, it is possible to numerically time gate the photons according to their micro-times and calculate the number of photons originating from either the short- or long-lived state/species, as depicted in Fig. 1a. More details on the burst excitation scheme and gating of photons according to their micro-times can be found in Liisberg *et al.*<sup>12</sup>

The spectral aspect of bmTG-FTS is obtained with a common-path birefringent interferometer (Translating-Wedge-based

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**Fig. 1** Principle of bmTG-FTS. (a) In burst-mode TCSPC measurements, micro- and macro-times are recorded. Burst-mode excitation and time gating allow for temporally disentangling the short-lived fluorescence (FI) from the long-lived luminescence (Lu). (b) In interferometry, an interferogram is recorded by displacement over a known distance and the corresponding emission spectrum is obtained by FT. (c) In ungated bmFTS, burst-mode TCSPC is combined with interferometry allowing for following the spectral and temporal evolution of short- and long-lived components on both timescales. However, when used ungated, spectral signals might be overlapping. (d) In bmTG-FTS, the signals are gated according to the micro-times, which makes it possible to completely disentangle the fluorescence from the luminescence spectrally and temporally.

Identical pulses encoding System; TWINS). The working principle of the TWINS interferometer has been described in detail previously.<sup>18</sup> Briefly, the TWINS interferometer is positioned in the emission path, and luminescence is passed through it. In the TWINS interferometer, the two polarization components of the emission light travel on the same path but will inherit a delay due to the birefringent medium. An interferogram is produced by recording the luminescence signal as a function of their time delay ( $I(x)$ , Fig. 1b). The time delay is controlled by moving a birefringent wedge pair a distance,  $x$ . Fourier transforming (FT) the recorded interferogram and performing a series of calibrations yields the corresponding emission spectrum ( $I(\lambda)$ ).<sup>18</sup>

By combining TCSPC with the TWINS interferometer, it is possible to record 2D temporal-spectral maps, which has previously been demonstrated in the micro-time domain.<sup>16,19,20</sup> In the micro-time domain, it is possible to temporally disentangle spectrally overlapping signals with ease if their decay times are sufficiently distinct (e.g., a short-lived state of 1 ns and a long-lived state of 1  $\mu$ s). This allows for determining the decay time

of the short-lived state, but the temporal evolution of the long-lived state is not resolved due to the limited size of the time-to-amplitude (TAC) window.<sup>16</sup>

By utilizing a burst excitation scheme, it is possible to have a single technique that can simultaneously capture both spectral and temporal features on a short (ns) and long ( $\mu$ s–ms) timescale. To exemplify this, we consider here the hypothetical case of a dual emissive species with a short- and long-lived component. In the micro-time domain, the results are similar as to experiments performed without bursts.<sup>16</sup> An ungated micro time-resolved interferometric map (mTRIM,  $I_{UG}(t, x)$ ) will reveal a short- and long-lived component. FT of the mTRIM yields a micro time-resolved emission map (mTREM,  $I_{UG}(t, \lambda)$ ) revealing two spectrally distinct bands (Fig. 1c), one decaying and synchronized with the excitation pulse and one seemingly unsynchronized and virtually flat over the time window. The addition of the burst excitation scheme allows for using the macro-time information for investigating the temporal dynamics of the long-lived component. Similar to the micro-time domain, an ungated macro TRIM (MTRIM,  $I_{UG}(T, x)$ ) is prepared, which by





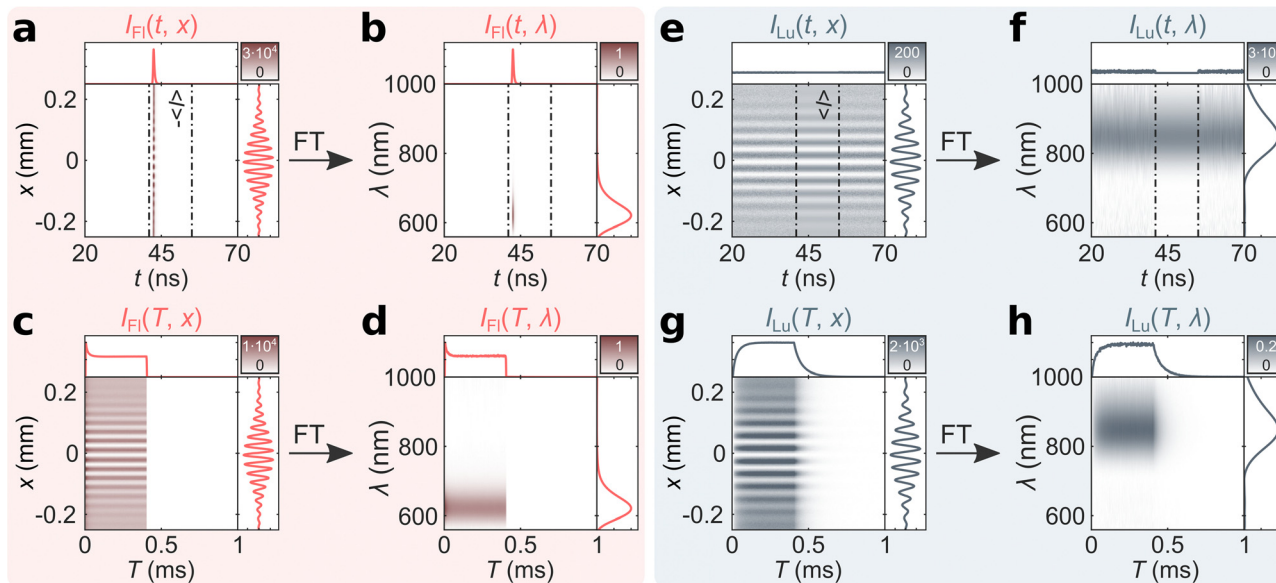


Fig. 3 Fluorescence and luminescence gated bmTG-FTS data of  $(\text{DNA})_2[\text{Ag}_{18}]^{12+}$ . Fluorescence gated (a) mTRIM, (b) mTREM, (c) MTRIM, and (d) MTREM. Luminescence gated (e) mTRIM, (f) mTREM, (g) MTRIM, and (h) MTREM.  $\langle I \rangle$  represents the mean luminescence intensity outside the fluorescence time gate.<sup>16</sup>

of the fluorescence and luminescence during the burst cycle. No significant changes could be observed with respect to the emission maxima before and after reaching the steady state condition (see Fig. S7, ESI†).

In summary, we introduced bmTG-FTS, a new technique for simultaneously capturing and spectrally disentangling the signals of both short- (ns) and long-lived ( $\mu\text{s}$ –ms) luminescence signals from the sub-nanosecond to millisecond timescale in a single measurement. We demonstrated the technique on a dual-emissive DNA-AgNC, from which we were able to extract essential photophysical parameters including spectral features and luminescence decay times in a single measurement.

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## Conflicts of interest

There are no conflicts to declare.

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