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in vivo quantitative FRET small animal imaging: intensity versus lifetime-based FRET

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ABSTRACT Förster Resonance Energy Transfer (FRET) microscopy is used in numerous biophysical and biomedical applications to monitor inter- and intramolecular interactions and conformational changes in the 2–10 nm range. FRET is currently being extended to *in vivo* optical imaging, its main application being in quantifying drug-target engagement or drug release in animal models of cancer using organic dye or nanoparticle-labeled probes. Herein, we compared FRET quantification using intensity-based FRET (sensitized emission FRET analysis using 3-cube IVIS imager) and macroscopic fluorescence lifetime (MFLI) FRET using a time-resolved ICCD system in small animal optical in vivo imaging. The analytical expressions and experimental protocols required to quantify FRET efficiency E and the fraction of donor molecules actually involved in FRET, f_D, are described in detail for both methodologies. Dynamic *in vivo* FRET quantification of transferrin receptor-transferrin binding was acquired in live intact nude mice upon intravenous injection of near infrared-labeled transferrin FRET pair and benchmarked against *in vitro* FRET using hybridized oligonucleotides. Even though both *in vivo* imaging techniques provided similar FRET quantification trends of receptor-ligand engagement, we demonstrate that MFLI FRET has significant advantages. Whereas the sensitized emission FRET approach using IVIS imager required 9 measurements (8 of which are used for calibration) acquired from three mice, MFLI FRET needed only one measurement collected from a single mouse. Hence, MFLI represents the method of choice for FRET measurements in intact, live mice to assess targeted drug delivery in longitudinal preclinical studies, as well as for many other *in vivo* biomedical applications.

WHY IT MATTERS

FRET measurements in live animals open a unique window into drug-target interaction monitoring, by sensing the close proximity between a donor and acceptor-labeled molecular probes. To perform these measurements, a 3-cube fluorescent intensity measurement strategy can be adopted, as is common for in vitro FRET microscopy studies. However, it is challenging to translate this already cumbersome approach to *in vivo* small animal imaging. Here, we compare this standard approach, for which we provide a revised analytical framework, to a conceptually much simpler and more powerful one based on fluorescence lifetime measurements. Our results demonstrate that the technical challenge of *in vivo* fluorescence lifetime macroscopic imaging is well worth surmounting to obtain quantitative, whole-animal information regarding molecular drug-target engagement.

1 INTRODUCTION

Förster Resonance Energy Transfer (FRET) has been extensively used in fluorescence microscopy as a nanometer-range (2-10 nm) proximity assay (1, 2), addressing a distance range that even super-resolution microscopy cannot resolve (< 20-30 nm) in live cells. FRET provides information on the distance between donor (D) labeled and acceptor (A) labeled proteins for each specific donor-acceptor fluorophore pair, independently of the resolution provided by the fluorescence imaging methodology used to acquire FRET measurements (3–6). Thus, FRET can be performed at both visible as well as near-infrared (NIR) wavelengths and can be measured by a wide variety of fluorescence-based imaging methodologies, beyond traditional microscopy approaches (7–9). These characteristics make FRET broadly applicable and one of the most extensively used imaging approaches in living cells as well as in model organisms, including bacteria, yeast, c. elegans, drosophila and mice (6).

There are several different types of FRET assays in fluorescence biological imaging. *Intra*-molecular FRET is used mostly to detect transient and dynamic signaling events using genetically encoded FRET-based biosensors in living cells (10–14).

These biosensor constructs provide a constant 1:1 donor:acceptor stoichiometry in each pixel, allowing for the implementation of ratiometric intensity-based imaging for a fast and qualitative FRET analysis. However, ratiometric FRET is very sensitive to the signal-to-noise ratio, has limited dynamic range and requires a significant number of image processing steps, such as background subtraction, shade/flatfield correction, image alignment and photobleaching correction (15–24). These problems are further compounded in tissues, where measurements are compromised by autofluorescence and the variable attenuation of fluorescence in heterogeneous tissues (25). *Inter*-molecular FRET has been established to monitor protein-protein interactions in live cells using separate donor- and acceptor-labeled proteins (26, 27). However, in inter-molecular FRET the relative abundance of donor and acceptor fluorophores is not controllable and can change over time, limiting the information that can be extracted from ratiometric measurements to the *apparent* or *average* energy transfer efficiency $\langle E \rangle$, which depends on the D to A distance (FRET efficiency for DA pairs, E) but also on the fraction f_{DO} of DO molecules that are not involved in energy transfer.

Fluorescence lifetime microscopy (FLIM) is regarded as the most robust means to collect FRET data since it is largely not influenced by probe concentration, signal intensity, or spectral bleed-through contamination (6). FLIM quantifies FRET occurrence by measuring the reduction of the fluorescence lifetime of the donor when in close proximity to the acceptor. Since the acceptor effectively behaves as a quencher of the donor's fluorescence, this quenching process is accompanied by a reduction in the quantum yield and lifetime of the donor (7). FLIM can measure FRET in each biological sample via the collection of the donor emission channel only. However, FLIM requires complex instrumentation and model-based fitting analysis, which makes it less accessible and straightforward compared to intensity-based FRET, which can be implemented with standard fluorescence microscopes and involves simple algebraic data processing (28–30).

Extending FRET assays to *in vivo* non-invasive macroscopy is one of the last frontiers of FRET imaging. Recently, *in vivo* FRET imaging approaches have been implemented to measure nanoparticle drug delivery and release, drug-target engagement, and dynamic probe uptake or biosensor-based signaling in various pre-clinical animal models (8, 9, 31–36). A major issue preventing full application of FRET into small animal imaging is the need to red-shift FRET into the NIR range to reduce absorption and autofluorescence as well as to increase depth of penetration in thick tissues (37). Development of NIR-labeled donor and acceptor pairs has permitted the implementation of non-invasive longitudinal FRET as well as the multiplexing of FRET pairs with metabolic imaging application in intact living mice (38–41).

Here, we address this challenge by comparing intensity- and lifetime-based NIR intermolecular FRET imaging assays designed to monitor receptor-ligand interactions in live intact mice (38, 41–44). In the context of ligand-receptor systems, FRET between donor-labeled and acceptor-labeled ligands occurs upon their binding to membrane-bound dimerized receptors. Using intensity- and lifetime-based FRET microscopy, we have demonstrated that protein ligands (*e. g.* transferrin: Tf), do bind extracellular domains of membrane-bound receptors (*e. g.* transferrin receptor: TfR) (45, 46). Moreover, *in vivo* MFLI FRET measurements have been successfully validated via *ex vivo* histochemistry, establishing that *in vivo* FRET signal directly reports on receptor-ligand engagement in intact live animals (40, 41, 44, 47, 48).

In the present study, we revisited the standard 3-cube equations for intensity-based FRET and systematically compare its results to lifetime-based FRET measurements analysis. The comparison was first done *in vitro* with double-stranded DNA FRET standard samples. We then extended our comparison to *in vivo* pharmacokinetics of ligand-receptor engagement monitored over a 90 min time period. Altogether, we show that while intensity-based FRET analysis *in vivo* can be attempted, lifetime-based *in vivo* FRET analysis is a much more robust and reliable approach for whole-animal quantitative FRET imaging.

2 MATERIAL AND METHODS

2.1 Macroscopic fluorescence lifetime-FRET (MFLI-FRET) with gated-ICCD

MFLI was performed using a time-resolved widefield illumination and a a time-gated intensified charge-coupled device (ICCD) (39). The system's excitation source was a tunable Ti-Sapphire laser (Mai Tai HP, Spectra-Physics, CA, USA) set to 695 nm. Power at the imaging plane was approximately 2 and 3 mW/cm^2 for dsDNA and *in vivo* MFLI, respectively. A digital micro-mirror device (DLi 4110, Texas Instruments, TX, USA) was used for widefield illumination over the sample plane. During animal imaging, active illumination was applied to ensure the bladder's intensity did not saturate the camera (49). The time-gated ICCD (Picostar HR, LaVision, GmbH, Bielefeld, Germany) was set to acquire gate images with a gate width of W_{ICCD} = 300 ps, separated gate steps δt = 40 ps (details provided elsewhere (39)). Data was acquired over a window of duration shorter than the full laser period (generally G = 160 total gate images per acquisition, i.e. D = 6.4 ns). During fluorescence imaging, a bandpass filter 720 ± 6.5 nm (FF 720/13, Semrock, Rochester, NY) and a longpass filter 715 nm (FF 715/LP25, Semrock, Rochester, NY) were applied to selectively collect fluorescence signal. In addition, for fluorescence imaging, the ICCD's microchannel plate (MCP) voltage was increased to 560 V and the gate integration time adjusted to 300 ms per gate image. Instrument response functions (IRFs) were acquired with equivalent illumination conditions to those used for fluorescence imaging, except for the emission filter, which were removed.

Table 1: Notations used in the text to refer to various background-corrected sample signals and their description. The notation $F(S)^{E_{em}}_{X_{ex}}$, used in ref. (50), represents the signal from species "S" (e.g. donor D) excited by excitation channel X (lower index notation, e.g. D_{ex} for donor excitation laser) and detected in emission channel E (e.g. D_{em} for donor emission channel.

Symbol	Sample	Excitation	Emission	Interpretation
$F^{D_{em}}_{D_{ex}}(DO)$	Donor	Donor	Donor	Donor only sample signal obtained with donor excitation detected in donor emission channel.
$F^{A_{em}}_{D_{ex}}(DO)$	Donor	Donor	Acceptor	Donor only sample with donor excitation, detected in acceptor emission channel.
$F^{A_{em}}_{A_{ex}}(DO)$	Donor	Acceptor	Acceptor	Donor only sample signal obtained with acceptor excitation, detected in acceptor emission channel.
$F^{D_{em}}_{D_{ex}}(AO)$	Acceptor	Donor	Donor	Acceptor only sample signal obtained with donor excitation, detected in donor emission channel.
$F^{A_{em}}_{D_{ex}}(AO)$	Acceptor	Donor	Acceptor	Acceptor only sample signal obtained with donor excitation, detected in acceptor emission channel.
$F^{A_{em}}_{A_{ex}}(AO)$	Acceptor	Acceptor	Acceptor	Acceptor only sample signal obtained with acceptor excitation, detected in acceptor emission channel.
$F^{D_{em}}_{D_{ex}}(DA)$	FRET	Donor	Donor	Donor-acceptor pair sample signal obtained with donor excitation, detected in donor emission channel.
$F^{A_{em}}_{D_{ex}}(DA)$	FRET	Donor	Acceptor	Donor-acceptor pair sample signal obtained with donor excitation, detected in acceptor emission channel.
$F^{A_{em}}_{A_{ex}}(DA)$	FRET	Acceptor	Acceptor	Donor-acceptor pair sample signal obtained with acceptor excitation, detected in acceptor emission channel.

2.2 Intensity-based FRET imaging using IVIS Imager

All samples, including donor-only (DO), acceptor-only (AO) and double-labeled (DA) samples, were imaged simultaneously in the same field-of-view of an IVIS Lumina XRMS Series III imaging system (Perkin Elmer), including heated stage (37 °C) and isoflurane anesthesia connections. Excitation wavelengths were set to 660 ± 10 nm for the donor and 740 ± 10 nm for the acceptor fluorophores. The emission filters were set to 713 ± 20 nm for the donor and to 793 ± 20 nm for the acceptor fluorophores. The intensity used in IVIS was constant throughout all imaging experiments. Three spectral channels were acquired for intensity FRET imaging: 1) donor channel (donor excitation and donor emission), 2) acceptor channel (acceptor excitation and acceptor emission) and 3) FRET channel (donor excitation and acceptor emission). The exposure time was set to 5 seconds per channel. The image size was 256×256 pixels.

2.3 Intensity-based FRET Data Analysis

Intensity-based FRET efficiency measurement relies on quantifying the amount of FRET-induced acceptor fluorescence (also called sensitized emission) in a sample, relative to that measured in the donor emission channel.

In an ideal situation where each donor fluorophore is located at a fixed distance from an acceptor fluorophore, (*i. e.* samples in which 100% of the donor molecules undergo FRET, denoted *DA* to emphasize that each donor forms a pair with an acceptor), a simple ratiometric approach using only signals obtained upon excitation with a donor-specific wavelength can be used to obtain the so-called proximity ratio (PR), or uncorrected ratiometric FRET efficiency, given by Eq. 1 (50):

$$PR = \frac{F^{A_{em}}_{D_{ex}}(DA)}{F^{A_{em}}_{D_{ex}}(DA) + F^{D_{em}}_{D_{ex}}(DA)} \tag{1}$$

where $F^{A_{em}}_{D_{ex}}(DA)$ and $F^{D_{em}}_{D_{ex}}(DA)$ are background-corrected acceptor and donor intensities of the sample undergoing FRET measured upon donor excitation, respectively (see Table 1 for notations). Measuring PR is a semi-quantitative approach for approximately quantifying FRET efficiency in a FRET sample where donor and acceptor are for instance conjugated to the same molecule, but it leaves aside contributions such as donor emission crosstalk (donor signal detected in the acceptor emission channel) and acceptor cross-excitation (direct excitation of the acceptor with donor excitation wavelengths) among other effects. Indeed, generally, the total fluorescence collected in each emission channel is a contribution of acceptor emission from FRET, donor emission leakage, and acceptor emission from direct excitation.

Sensitized emission FRET (SE-FRET) approaches have been designed to correct for these additional effects and require data acquired with separate excitation and emission combinations (the so-called 3-cube approach) (16, 22, 50–54).

A first-order correction consists in subtracting the direct acceptor excitation and the leakage of the donor emission from the measured acceptor signal to obtain a better estimate of the FRET-induced acceptor emission (*i.e.*, the relevant FRET emission signal), using Eq. 2:

$$F^{FRET} = F_{D_{ex}}^{A_{em}}(DA) - d_A F_{A_{ex}}^{A_{em}}(DA) - l_D F_{D_{ex}}^{D_{em}}(DA) \tag{2}$$

where d_A is the direct acceptor excitation correction factor, and l_D is the donor leakage correction factor.

The first correction factor d_A is measured using an acceptor-only (AO) sample excited separately at two excitation wavelengths (donor and acceptor) and detected in the acceptor emission channel. Correction factor d_A is calculated using Eq. A.13 (17, 50, 54) in the Appendix.

The second correction factor l_D is measured using a donor-only (DO) sample excited with a donor excitation wavelength (donor excitation channel) and detected in both emission channels (donor and acceptor). Correction factor l_D is calculated using Eq. A.12 (17, 50, 54) in the Appendix.

The FRET efficiency E can then be computed as (50):

$$E = \frac{F^{FRET}}{F^{FRET} + \gamma F_{D_{ex}}^{D_{em}}(DA)} \tag{3}$$

where γ is the detection-correction factor defined as $\gamma = \phi_A \eta_A^A / \phi_D \eta_D^D$. ϕ_A and ϕ_D are the acceptor and donor quantum yields, respectively and η_A^A (resp. η_D^D) is the acceptor (resp. donor) detection efficiency in the acceptor (resp. donor) channel.

While Eqs. 2 and 3 are adequate in many situations, certain experimental situations result in further signal contamination, when for instance the donor fluorophore can be excited at the acceptor excitation wavelength, or when the acceptor fluorophore can be detected in the donor emission channel. The first effect contributes some unwanted signal to a quantity used to correct the sensitized emission of the acceptor in Eq. 2, while the second requires further correction of the donor channel signal. In those cases, some donor signal needs to be subtracted from $F_{A_{ex}}^{A_{em}}(DA)$ and some acceptor signal from $F_{D_{ex}}^{D_{em}}(DA)$.

These corrections involve two additional correction factors (d_D and l_A), as discussed in the Appendix. To retrieve d_D, two measurements of a donor-only sample are needed: i) excitation at the donor wavelength and recording in the acceptor emission channel and ii) excitation at the acceptor wavelength and recording in the acceptor emission channel. Correction factor d_D is calculated using Eq. A.12 in the Appendix.

The last correction factor l_A requires two measurements of an acceptor-only sample: i) excitation at the donor wavelength and recording in the donor emission channel and ii) excitation at the donor wavelength and recording in the acceptor emission channel. Correction factor l_A is calculated using Eq. A.13 in the Appendix.

It should be noted that correction factors d_A, l_A, d_D and l_D are specific to fluorophores as well as imaging systems, and ideally require constant excitation intensities throughout the series of measurements. At the very least, one must take into account differences in excitation intensity (and integration time) if those need to be adjusted for experimental reasons. Consequently, these correction factors need to be estimated every time the experimental conditions are modified (excitation intensities, integration times, filters, fluorophores or molecular environments).

After all four correction terms have been retrieved, the sensitized emission FRET signal, F^{FRET}, can be calculated according to (see Appendix Eq. A.21 for details):

$$F^{FRET} = \frac{1 + \alpha l_A l_D}{1 - d_A d_D} F_{D_{ex}}^{A_{em}}(DA) - \frac{d_A}{1 - d_A d_D} F_{A_{ex}}^{A_{em}}(DA) - \frac{l_D}{1 - l_A l_D} F_{D_{ex}}^{D_{em}}(DA) \tag{4}$$

where parameter α is defined by Eq. A.19 in the Appendix. If $l_A = d_D = 0$, one recovers Eq. 2.

Similarly, the FRET efficiency can be obtained by a modified version of Eq. 3 (see Appendix Eq. A.22):

$$E = \frac{F^{FRET}}{F^{FRET} + \frac{\gamma}{1 - l_A l_D}\left(F_{D_{ex}}^{D_{em}}(DA) - l_A F_D^A(DA)\right)} \tag{5}$$

an expanded version of which can be found in the Appendix (Eq. A.18). If $l_A = d_D = 0$, one obviously recovers Eq. 3.

The above equations only apply to a pure FRET sample (DA), as mentioned at the beginning of the discussion and indicated by the notations. In general, a real sample will contain a mixture M of species: donor-only (DO), acceptor-only (AO) and FRET (DA), whose respective fractions are fully specified by the total number N_D of donor molecules and total number N_A of

acceptor molecules, and the fraction f_D of donor molecules and fraction f_A of acceptor molecules involved in FRET interaction (with $f_D N_D = f_A N_A$). This mixture of species will be characterized by 3 different types of signals $F_{X_{ex}}^{E_{em}}(\mathrm{M})$, each verifying:

$$F_{X_{ex}}^{E_{em}}(M) = F_{X_{ex}}^{E_{em}}(AO) + F_{X_{ex}}^{E_{em}}(DO) + F_{X_{ex}}^{E_{em}}(DA) \tag{6}$$

As derived in the Appendix, the product $f_D E$ of the fraction of donor f_D involved in FRET and the FRET efficiency E of the FRET sample can then be expressed in terms of the 3 measured quantities $F_{D_{ex}}^{D_{em}}(M)$, $F_{D_{ex}}^{A_{em}}(M)$ and $F_{A_{ex}}^{A_{em}}(M)$ and the coefficients defined by Eqs. A.12-A.13,A.15 & A.19 as (Appendix Eq. A.27):

$$f_D E = \frac{(1+\alpha l_A l_D) F_{D_{ex}}^{A_{em}}(M) - \alpha l_D F_{D_{ex}}^{D_{em}}(M) - d_A F_{A_{ex}}^{A_{em}}(M)}{(1+\alpha l_A (l_D - \gamma) F_{D_{ex}}^{A_{em}}(M) + \alpha(\gamma - l_D) F_{D_{ex}}^{D_{em}}(M) - d_A F_{A_{ex}}^{A_{em}}(M)} \tag{7}$$

which turns out to be the same formula as obtained for a pure DA sample (Eq. 5) with the replacement of E by $f_D E$.

Note in particular that, in the case of a mixture of D, A and DA species, it is not possible to disentangle E from f_D without further information on the sample. Fortunately, this quantity can also be estimated using lifetime measurements as discussed in the next section, allowing a direct comparison of both methods.

2.4 Lifetime-based FRET Data Analysis

In the ideal case, quantification of FRET using fluorescence lifetime imaging (FLI) only requires measuring the fluorescence lifetime of the donor undergoing FRET and that of the isolated donor (no FRET condition). The result of FRET is a reduction (quenching) of the donor fluorescence lifetime.

There are two conventional methods to obtain lifetime-based FRET quantification: 1) multi-exponential fitting and 2) phasor analysis (55). In typical FRET-FLI analysis, two lifetime contributions, τ_{DA} (lifetime of donor undergoing FRET) and τ_{DO} (lifetime of donor not undergoing FRET) are assumed, which can be modeled using a bi-exponential function (Eq. 8):

$$F_T(t) = IRF_T(t) * [A_1 e^{-t/\tau_1} + A_2 e^{-t/\tau_2}] + F_0 \tag{8}$$

where $F_T(t)$ is the T-periodic fluorescence intensity as function of time t after laser excitation (sometimes referred to as a temporal point spread function or TPSF). $IRF_T(t)$ corresponds to the T-periodic instrument response function, which is convolved with the fluorescence decay (symbol $*$, interpreted as a cyclic-convolution over a single period T) (56). A_1 and A_2 correspond to the amplitudes of the quenched and unquenched donor contributions, while $\tau_{DA} = \tau_1$ and $\tau_{DO} = \tau_2$ are the quenched and unquenched lifetimes, respectively.

The relative amplitudes of each component are related to the fraction of the donor in each species (donor-only and FRET pair) by(43):

$$f_{DA} = f_D = \frac{A_1}{A_1 + A_2}, \quad f_{DO} = \frac{A_2}{A_1 + A_2} = 1 - f_{DA} \tag{9}$$

The unquenched donor lifetime ($\tau_{DO} = \tau_2$) can be obtained experimentally (for instance as the longest lifetime component in a 2-exponential fit or by a separate measurement of a donor-only sample acquired in identical conditions as the FRET sample) or from the literature. The *amplitude-weighted* average lifetime of the sample is calculated using Eq. 10, which is sometimes used as a "proxy" to quantify the fraction of FRET-undergoing species at a given location.

$$\langle \tau \rangle = \frac{A_1 \tau_1 + A_2 \tau_2}{A_1 + A_2} \tag{10}$$

The donor-only lifetime ($\tau_{DO} = \tau_2$) and the FRET sample lifetime ($\tau_{DA} = \tau_1$) are related to the FRET efficiency (E) by:

$$E = 1 - \frac{\tau_{DA}}{\tau_{DO}} \tag{11}$$

Combining Eqs. 9-11, we get the following expression for the product $f_D E$:

$$f_D E = 1 - \frac{\langle \tau \rangle}{\tau_{DO}} \tag{12}$$

Because the FRET efficiency of the DO species is equal to zero ($E_{DO} = 0$), Eq. 12 can be rewritten as:

$$f_D E = f_{DO} E_{DO} + f_D E = (1 - f_D) E_{DO} + f_D E = \langle E \rangle \tag{13}$$

which states that the quantity $f_D E$ computed with Eq. 12 is the average FRET efficiency of the sample.

Importantly, acquisition of fluorescence lifetime data for FRET quantification does not require acceptor fluorescence information if the correct filter set is used. In other words, by selectively collecting emission from the donor fluorophore and filtering signal from the acceptor, emission spectral bleedthrough can be made negligible. In addition, the necessary calibration of the system (*i. e.*, IRF) or background correction are much simpler to obtain via this approach.

Eqs. 7 and 12 provide a way to directly compare intensity-based and lifetime-based measurements of the same sample.

2.5 Double-stranded DNA sample preparation

NIR dyes were obtained from ThermoFisher Scientific (Waltham, MA). Oligodeoxynucleotides (oligo-DNAs) were synthesized and labeled by IBA Lifesciences (Göttingen, Germany). The sequences of two complementary oligo-DNA were as in Ref. (50). The following NIR FRET pair was used to conjugate the oligo DNAs: Alexa Fluor 700 (AF700) and Alexa Fluor 750 (AF750). The sequence of the "top" strand was 5'-TAA ATC TAA AGT AAC ATA AGG TAA CAT AAC GGT AAG TCC A-3'. The donor (AF700) was conjugated to dT at position 1 of the top strand, and the acceptor (AF750) was conjugated to dT at three separate positions (17, 22 and 27) on the "bottom" strand for each FRET sample respectively. All purchased fluorescently conjugated oligo-DNAs were provided purified using high-performance liquid chromatography (HPLC) and lyophilized. Unlabeled strands were purified using desalting method and delivered as lyophilized form. Each lyophilized oligo-DNA was first resuspended with Tris-EDTA buffer (pH 8, Sigma-Aldrich, St. Louis, MO) to make a 100 nM stock solution. To perform the hybridization, AF700 oligo-DNA strands were mixed with AF750 oligo-DNA strands at 1:1 molar ratio at 50 nM final concentration for 100 µL reaction volume. For donor-only and acceptor-only samples, unlabeled oligo-DNAs were used as complementary strands at 1:1 molar ratio. The mixture of oligo-DNAs was heated at 95 °C for 5 minutes using dry heating block and cooled at room temperature for 30 minutes to obtain a mixture of double-stranded DNS (dsDNA) and residual unhybridized single-labeled oligo-DNAs.. Identical samples were used for the IVIS and the wide-field MFLI measurements.

2.6 Animal experiments

All animal procedures were conducted with the approval of the Institutional Animal Care and Use Committee at both Albany Medical College and Rensselaer Polytechnic Institute. Animal facilities of both institutions have been accredited by the American Association for Accreditation for Laboratory Animals Care International. Athymic female nude mice were purchased from Charles River (Wilmington, MA). All animals were in healthy condition. Tf probes were prepared by conjugating iron-bound Tf with fluorophores per manufacturer's instruction. AF00 and AF750 were used as donor and acceptor, respectively. The animals were anesthetized with isoflurane before being retro-orbitally injected with 40 µg of AF700-Tf (donor) and/or 80 µg of AF750-Tf (acceptor) conjugates and imaged immediately. The time-resolved data were acquired continuously for 90 minutes ($\approx$ 43 seconds per acquisition). Each intensity FRET measurement involved three mice. The single-donor mouse was injected with AF700-Tf, the single-acceptor mouse with AF750-Tf and the double-labeled FRET mouse was injected with a mixture of AF700-Tf and AF750-Tf at acceptor:donor (A:D) ratio of 2:1 (40 µg of AF700-Tf and 80 µg of AF750-Tf).Each lifetime measurement used only a mouse injected with a mixture of donor and acceptor. During imaging, mice were kept anesthetized using isoflurane, and their body temperature maintained using a warming pad (Rodent Warmer X2, Stoelting, IL, USA) on the imaging plane.

2.7 Immunohistochemistry

Mice were injected with 40 µg Tf-biotin conjugates (Sigma-Aldrich, Inc., MO) or PBS buffer and sacrificed 6 h post-injection. Bladders were collected, fixed in 4 % paraformaldehyde for 24 h and processed for embedding and sectioning (40). Tissue sections were analyzed by immunohistochemistry using ABC Elite and NovaRed peroxidase substrate kit (Vector laboratories) to visualize Tf-biotin. Parallel bladder sections were stained with Hematoxylin and Eosin and imaged using 10x lens microscope for tissue morphology visualization.

2.8 FRET quantification using decay fits of MFLI data

2.8.1 dsDNA samples

$f_D E$ (product of the fraction of donor involved in FRET and the FRET efficiency of the FRET sample) was quantified by fitting the fluorescence decays in each pixel of selected regions of interests (ROIs) to a bi-exponential model τ_M (Eq. 8) and retrieving the amplitude-weighted averaged lifetime $\langle\tau\rangle$ (Eq. 10). IRFs were acquired using a sheet of white paper as sample after removing all emission filters. After convolution, the tail portion of each pixel's decay (99%-2% of the peak value) was fit

using the MATLAB function *fmincon*() for least squares minimization of the cost function. After τ_M was calculated for every decay of interest (including donor-only and double-labeled FRET sample), Eq. (12) was used to calculate $f_D E$.

2.8.2 Dynamic Tf-TfR FRET *in vivo* imaging

The liver and bladder ROIs were delineated via intensity thresholding. Since the mouse did not move laterally along the imaging plane during the 90 minutes of imaging, the same mask was appropriate for all time-points. The donor-only lifetime was retrieved using the averaged mean-lifetime values from the urinary bladder over the first five acquisitions ($\tau_{bladder}$ = 1.03 ns). This method neglects environment-dependent changes of lifetime between mouse organs. Though this assumption is valid herein, it may very well not be for other applications (e.g., pH dependence on lifetime (48)). All other analysis steps and calculation of FRET efficiency were performed similarly as described above for the dsDNA sample.

2.9 FRET quantification using sensitized emission analysis of IVIS data

2.9.1 dsDNA samples

Background subtraction was performed on all excitation/emission channels. Then, the correction factors (d_D, l_D, d_A and l_A) were determined using Eq. A.12-A.13. Additionally, the γ correction factor was determined using the known quantum yields and fluorescence emission spectra (Fig.S1) for the NIR dyes, as well as filter specifications and camera quantum efficiency of the IVIS imaging setup. The calculated value was $\gamma = \phi_{AF750}\eta^{AF750}_{AF750_{em}}/\phi_{AF700}\eta^{AF700}_{AF700_{em}} = 0.41$. Afterwards, $f_D E$ was calculated according to Eq. 7.

2.9.2 Dynamic Tf-TfR FRET *in vivo* imaging

The correction factors (d_D, l_D, d_A and l_A) were determined in a dynamic fashion at each time-point using the intensities of the liver and bladder ROIs of all three mice and Eqs. A.12-A.13 (Fig. 2). Intensity-based $f_D E$ was then calculated as described above for the dsDNAs.

3 RESULTS

3.1 Short double-labeled dsDNA strands as FRET standards

double-labeled double-stranded DNA (dsDNA) molecules provide a simple and convenient way to design molecules with well-defined distances between donor and acceptor fluorophores that can be used as FRET standards (57, 58). In an ideal case, the base pair separation between donor and acceptor dyes determines the FRET efficiency based on the B-DNA model structure. The larger the separation, the lower the FRET efficiency, which depends on the ratio of the distance between the two fluorescent donor and acceptor fluorophore molecules to to their Förster radius R_0 ($R_{AF700/AF750}$ = 7.8 nm). For this study, three dsDNA FRET standard samples were prepared by hybridization of donor- or acceptor-labeled complementary 35 oligonucleotide long single-stranded DNA (ssDNA) molecules characterized by donor-acceptor distances of 17, 22 and 27 nucleotides (Fig. 1A-B) (50). The fluorophores used here (Alexa Fluor 700 and Alexa Fluor 750) are near-infrared (NIR) emitting fluorophores widely adopted for *in vivo* imaging applications (38, 59). The donor fluorophore (AF700) is located at the end of the same ssDNA, while the acceptor (AF750) is located in different positions of the complementary strand, but surrounded by a common nucleotide pattern, in order to ensure a constant environment (and therefore a common Förster radius) for all samples (50). These dsDNA oligos samples were imaged for both FLI- and intensity-based FRET analysis using MFLI and IVIS imagers, respectively.

In contrast with MFLI, which requires the imaging of one or two samples only (FRET, i.e. donor + acceptor and optionally, donor-only), three samples are necessary for intensity FRET analysis: donor-only, acceptor-only and FRET sample (notated as donor-acceptor n, or, "DA_n", where n indicates the number of nucleotides separating donor and acceptor). All samples were imaged with donor, acceptor and FRET excitation/emission channels, with the same field of view and excitation power settings (Table 1). The fluorescence intensity maps of all samples from all channels are shown in Fig. 1C. Correction factors d_A and l_A were obtained from the acceptor-only dsDNA sample (Fig. 1D, Table S1). Additionally, corrections factor d_D and l_D were obtained from the donor-only dsDNA sample (Fig. 1D, Table S1). The correction factor γ, was calculated using the known quantum yield of both dyes, as well as by calculating detection efficiency with known emission spectra and detection wavelength bands used herein ($\gamma = \phi_{AF750}\eta^{AF750}_{AF750_{em}}/\phi_{AF700}\eta^{AF700}_{AF700_{em}} = 0.41$). Using this information, calculation of the FRET efficiency of the sample using Eq. 7 resulted in $f_D E$ values shown in Fig. 1E, F (Table S2).

The fluorescence decays and the intensity, average lifetime, and $f_D E$ maps and values of all samples from all channels are shown in Fig. 1G-J. The amplitude-weighted mean lifetime of a dsDNA donor-only sample consisting of the donor-labeled ssDNA

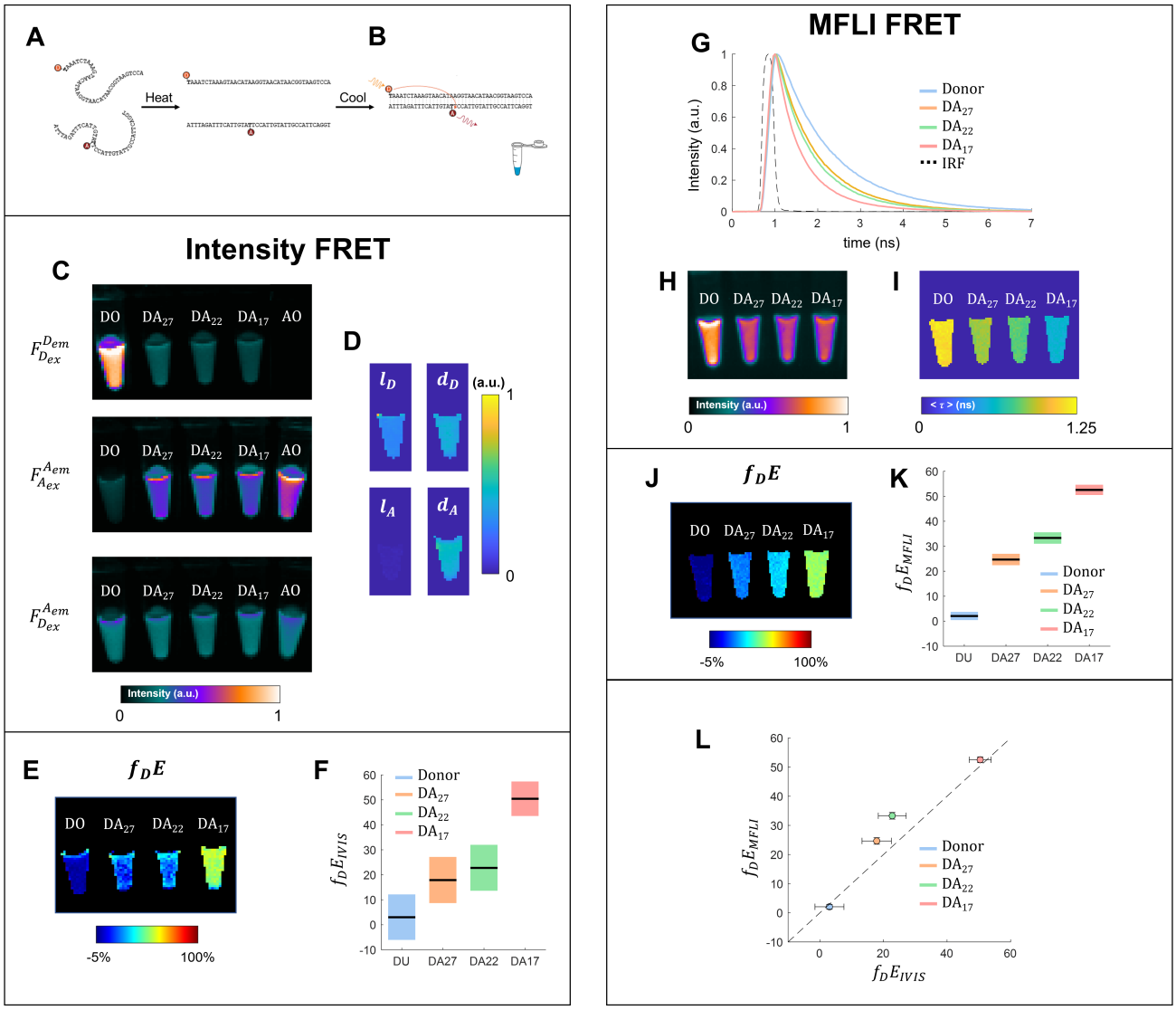

Figure 1: ***in vitro* comparison of intensity- and MFLI-FRET imaging methods**. A-B, Oligonucleotide sequences used for hybridized DNA FRET sample DA_{17}. The donor, Alexa Fluor 700 was conjugated to dT at position 1. The acceptor, Alexa Fluor 750, was conjugated to dT positions 17, 22 and 27 on the complementary strand. Hence, the distance between the donor and the acceptor for "DA 17" after hybridization was 17 base pairs, which corresponds to approximately 5.8 nm. L, Scatter plot of $f_D E$ results (mean ± standard deviation) retrieved through intensity- and FLI- FRET. C, Fluorescence intensity data acquired with the IVIS Lumina XRMS Imaging setup: donor only (DO) dsDNA, acceptor only (AO) and FRET samples (DA_{17}, DA_{22} and DA_{27}) imaged with donor wavelength/donor detection ($F^{D_{em}}_{D_{ex}}$), acceptor wavelength/acceptor detection ($F^{A_{em}}_{A_{ex}}$) and donor wavelength/acceptor detection ($F^{A_{em}}_{D_{ex}}$). Results from DNA FRET standard sample. D, Spectral correction coefficient maps. E, $f_D E$ map retrieved through intensity FRET. F, Boxplot of $f_D E$ values retrieved using intensity FRET. G, Normalized MFLI decays measured from the donor-only and FRET dsDNA samples (whole vial ROI). H, Max-normalized intensity measurements using a gated-ICCD. G, Amplitude-weighted mean lifetime of the donor-only and FRET dsDNA samples. I, $f_D E$ map retrieved through lifetime-based FRET. J, Boxplot of $f_D E$ values retrieved using lifetime-based FRET.

hybridized to an unlabeled complementary ssDNA strand was measured as $\tau_{DO} = 1.19 \pm 0.05$ ns. Using the amplitude-weighted mean lifetimes of the double-labeled dsDNA samples and Eq. 12, led to the different values of $f_D E$ represented in Fig. 1J,K. We hypothesize that the difference between both sets of results is due to residual donor bleedthrough when excited with the acceptor wavelength (see Fig. 1C, $F_{A_{ex}}^{A_{em}}(DO)$). To properly correct for this, the measurement of $F_{A_{ex}}^{D_{em}}(X)$ (X = DO, AO and the 3 FRET samples) would be needed and additional correction factors included in the analysis. This further highlights the complexity of the intensity-based FRET approach for these applications.

The larger standard deviation of the intensity-based FRET results compared to the lifetime-based results noticeable in Fig. 1L is likely due to the lower signal-to-noise ratio (SNR) of the IVIS data. This lower SNR also prevented us from performing a pixel-wise estimation of correction factors. Using pixel-wise correction factors is typically not recommended for this very reason (23). Importantly, these complications are not present for FLI-FRET quantification, and we observed low standard deviation across the vial ROIs (see Fig. 1J,K).

3.2 *in vivo* FRET imaging of transferrin-transferrin receptor binding

Herein, we have demonstrated dynamic monitoring of ligand-receptor engagement *in vivo* using sensitized emission FRET and compare it to MFLI-FRET.

The Tf-TfR system was chosen as a model for *in vivo* FRET imaging of ligand-receptor engagement since transferrin has been used widely as a carrier for drug delivery (60). Tf-TfR binding was monitored by either IVIS imaging according to intensity FRET imaging protocol as described in Material & Methods or using the MFLI imager as described elsewhere (39). The animals were intravenously injected with NIR-Tf fluorescently labeled probes and imaged continuously for 90 min at 30 sec interval steps. As expected, there is a significant increase in fluorescence accumulation in the liver and the urinary bladder as a function of time, while very little is detected in other organs (40, 41, 59). This finding was consistent across the intensity and the lifetime FRET measurements.

3.2.1 Sensitized emission FRET quantification using IVIS data

As shown in Fig. 2, single-labeled donor-only or acceptor-only mice showed increased fluorescence intensity in the urinary bladder and the liver in all excitation/emission channels, respectively. Fluorescence of donor-only mouse was negligible in the acceptor excitation/emission channel, and that of acceptor-only mouse was negligible in the donor excitation/emission channel. However, fluorescence intensity levels are clearly detected in the liver and bladder in the FRET excitation/emission channel (donor excitation and acceptor detection) in donor-only and acceptor-only single-labeled mice, indicative of spectral bleedthrough in both organs.

In the double-labeled FRET mouse, FRET excitation/emission channel also showed an increase in fluorescence intensity in the liver as well as in the urinary bladder (Fig. 2A). In agreement, FRET channel intensity measurements in the double-labeled mouse showed an accumulation of fluorescence probe in the liver and the urinary bladder over time (Fig. 3A). These results are suggestive of FRET occurring upon ligand-receptor interaction in the liver as expected considering its role in iron metabolism and high level of TfR expression. In contrast, the finding of significant intensity levels in the urinary bladder (used as a negative control) in the FRET excitation/emission channel of the double-labeled FRET mouse suggests contaminations of the FRET signal with donor spectral leakage and/or direct excitation of the acceptor. Hence, it was essential to perform sensitized emission FRET analysis in a dynamic fashion to account for spectral bleedthrough at each time-point.

Herein, we used the data from the single-labeled mice for spectral correction of the FRET mouse data (i.e., mouse injected with a mixture of donor and acceptor-labeled Tf). For this spectral correction, information from the donor-only and acceptor-only mice was used to obtain all four correction coefficients for acceptor and donor spectral bleedthrough, respectively (Fig. 2B). All coefficients were calculated and applied at each time-point based on the variation of intensity at each time point (Fig. 2C-F). In addition, we applied γ correction factor as used for dsDNA FRET standard samples during the *in vitro* calibration. Though this factor does not fully correct for optical absorption variation across wavelengths in biological tissue, given the use of NIR fluorophores herein, we assume the variation is negligible. Using these parameters and the FRET-induced mouse intensity in the FRET excitation/emission channel, $f_D E$ was calculated at each time point. Intensity-based dynamic $f_D E$ of Tf-TfR ligand-receptor interaction successfully showed increasing FRET efficiency in the liver and no change of $f_D E$ in the urinary bladder (Fig. 4B). This result directly correlates with the known physiology of Tf, which binds to its receptor in the liver allowing FRET to occur. The expected absence of FRET in the urinary bladder indicates excretion, and inability to bind TfR, of degraded Tf or their degradation products (free fluorophores). However, $f_D E$ values retrieved over the first 20 minutes in the mouse liver are nonsensically negative (Fig. 4B). Indeed, the calculated F^{FRET} (Eq. 4) over this time-interval is negative. We hypothesize that this is due to variation in fluorophore accumulation kinetics post-injection across the donor-only and acceptor-only mice compared to the FRET mouse (or, from mouse-to-mouse). Unfortunately, this complexity is intrinsically

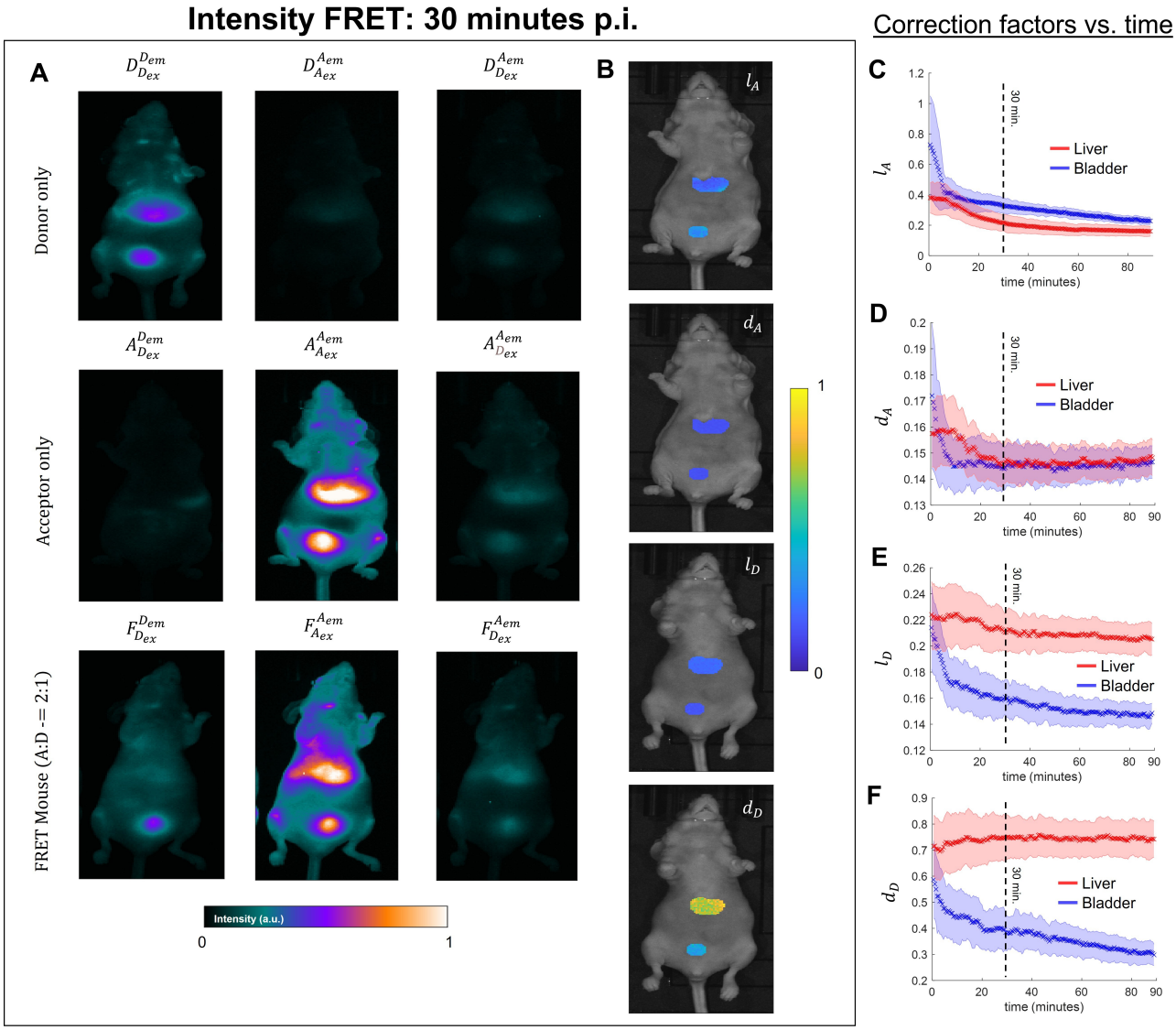

Figure 2: **Dynamic SE-FRET spectral correction *in vivo* using IVIS**. A, Image data of mice injected with NIR-Tf 30 minutes after injection. Top row, middle row and bottom row correspond to mice injected with donor only, acceptor only and FRET Tf, respectively. B, spectral correction coefficient maps for this time-point at the liver and bladder ROI (top right of each image provides factor). C-F, Average and standard deviation of spectral correction coefficients across each organ ROI over the duration of imaging.

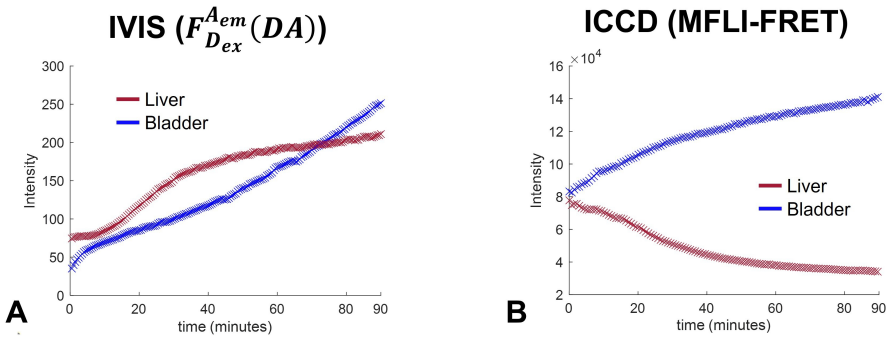

Figure 3: **Example of intensity dynamics in longitudinal *in vivo* optical imaging**. A, Dynamic FRET intensity plot from donor/acceptor excitation/emission channel using IVIS. B, Intensity (summed over entire fluorescence decay) using MFLI. All y-values mark the averaged intensity value across each organ ROI at each time-point.

unavoidable when using the three-cube intensity FRET imaging protocol alone across three independent mouse subjects.

3.2.2 Lifetime-based FRET quantification using MFLI data

In comparison, lifetime-based FRET analysis was much more straightforward. MFLI data from the donor emission only (695 nm excitation, 721 ± 6 nm detection) from a double-labeled mouse was sufficient to evaluate ligand-receptor interaction without any additional calibration samples (or mice) for correction.As shown in Fig. 3B, there was an increasing accumulation of fluorescence probe in the urinary bladder during the imaging session. However, the liver fluorescence decreased over the span of imaging due to FRET quenching, as expected (Fig. 3B). In this TfR-Tf based FRET system, the donor lifetime was calculated from the urinary bladder ($\tau_{bladder}$ = 1.03 ns, see Material & Methods).Application of MFLI FRET to other biomedical assays that do not provide internal negative FRET control such as the bladder, can instead use the long lifetime component obtained from bi-exponential fitting of tissues or organs in which FRET takes place. An alternative solution may involve obtaining the donor-only lifetime from a single-labeled mouse, as indicated in Table S3, or a donor-labeled probe injected into the double-labeled mouse in a nearby location, as an internal negative FRET control, although both approaches add a degree of uncertainty due to the use of different animals.

Lifetime FRET analysis of the donor fluorescence decay curves with bi-exponential fitting yielded the amplitude-weighted average lifetime from which f_DE values were calculated using Eq. 12. f_DE in the liver is increasing over time and is significantly higher than that of the urinary bladder and elsewhere due to high expression of TfR (Fig. 4C-D). In addition, hepatic sinusoids also have relatively large vascular fenestration (100-175 nm). This structure allows passive accumulation of Tf in the sinusoid. The local accumulation with a high presence of TfR in the liver resulted in increasing Tf internalization over time, which was represented by increasing f_DE. However, f_DE within the urinary bladder was less than 10%, which was significantly lower than that observed at later time points in the liver (Fig. 4C-D). f_DE in the urinary bladder did not increase throughout the imaging session despite the increase of the fluorescence intensity. This finding suggests that there was negligible FRET in the urinary bladder. At least two possibilities could explain this observation, including: 1) there was no internalization of Tf into the urinary bladder, or 2) that the fluorescence of the donor in the urinary bladder was a result of Tf degradation – leading to residual fluorophore accumulation. In agreement with the second possibility, intensity levels are shown to increase in bladders, as the FRET signal remains at very low levels over time, as shown in Fig. 3B and Fig. 4D, respectively. Moreover, in a separate control experiment, immunohistochemical analysis of bladder tissues from mice intravenously injected either with biotin-labeled Tf or PBS (negative control) showed no Tf staining in the lining of all bladders (Fig. S2). Overall, these results indicate that bladder tissues do not accumulate TfR-bound Tf upon intravenous injection. In any case, with exception of the negative values retrieved over the first 20 minutes through intensity FRET for the liver, the f_DE quantification obtained through both approaches are in good agreement with each other, considering that the measurements were performed using different mice (and in the case of the intensity-based FRET analysis, multiple mice were used for obtaining the different correction parameters). Remarkably, analysis of the two kinetic curves shown in Fig. 4B-D using a simple exponential model (data not shown) yielded similar time constants τ_{kin} in both experiments (intensity-based analysis: τ_{kin} = 19.7 ± 0.3 min, lifetime-based analysis: τ_{kin} = 31.2 ± 1.6 min), comparable to those observed in previous experiments analyzed using a different approach (43).

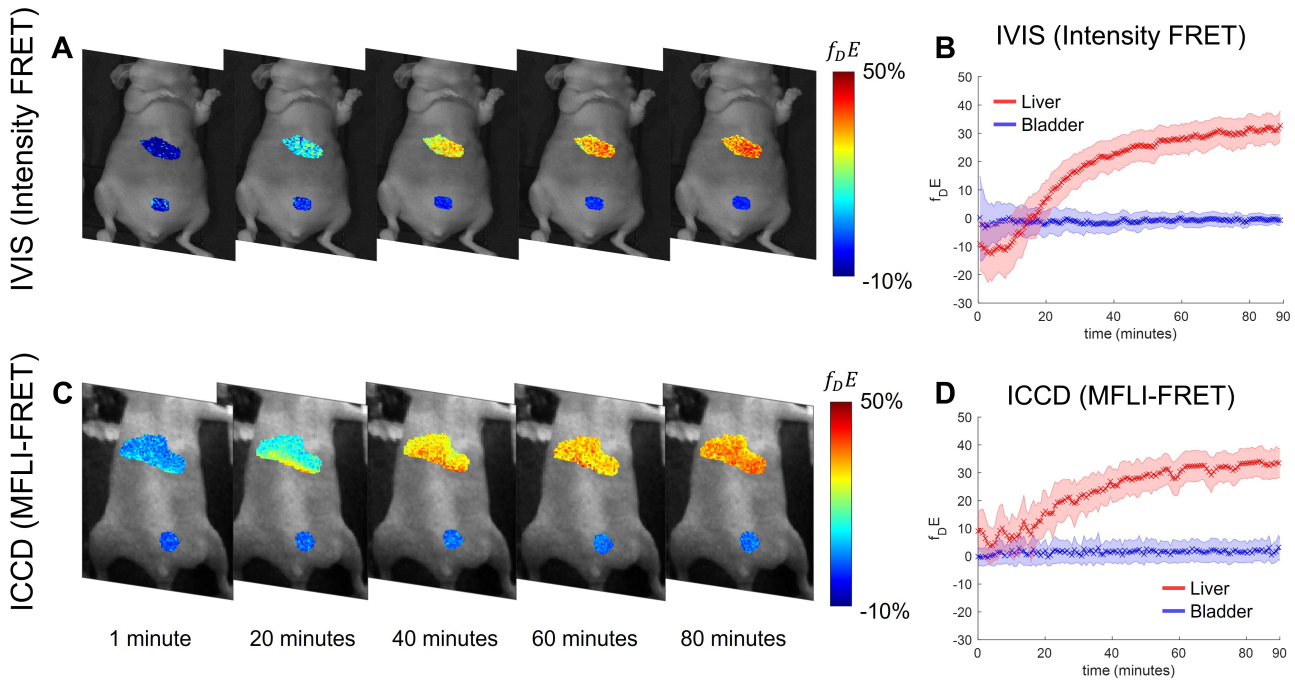

Figure 4: ***in vivo* comparison of intensity-and MFLI-FRET dynamic imaging**. A, Intensity-based dynamic *in vivo* f_DE in the liver and the urinary bladder of a mouse injected with Tf A:D 80:40 µg at selected time points (total time points = 180). B, f_DE retrieved at all time-points using intensity FRET (solid line denotes average, shaded area marks the standard deviation). C, Lifetime-based dynamic *in vivo* f_DE in the liver and the urinary bladder of a mouse injected with Tf A:D 80:40 µg at selected time points (total time points = 135). D, f_DE retrieved at all time-points using MFLI-FRET. For B and D, solid line marks the average value and shaded area denotes the standard deviation across all pixels within each organ ROI.

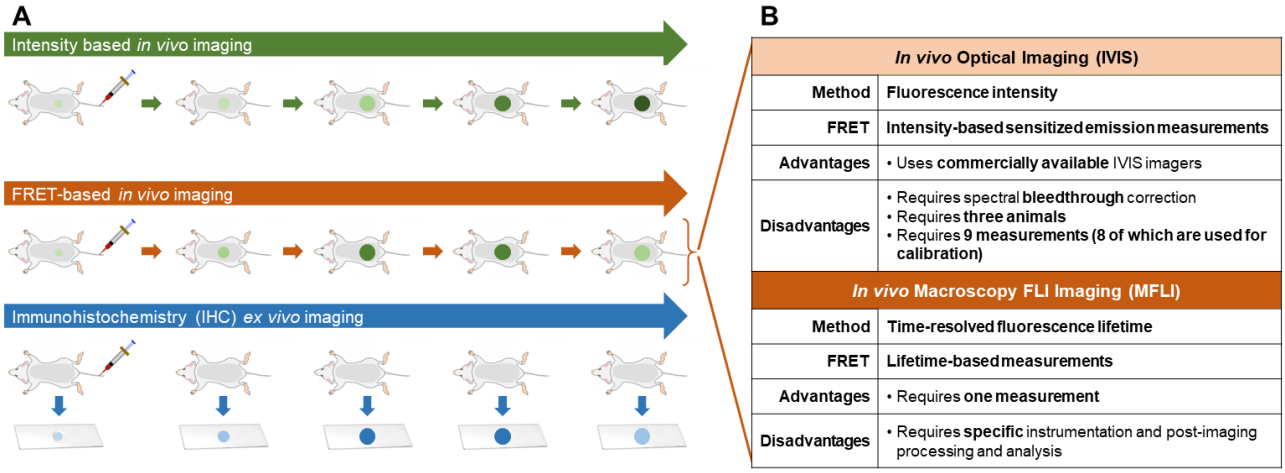

Figure 5: **Comparison of different approaches for longitudinal preclinical molecular interaction monitoring**. A, Conventional preclinical longitudinal studies such as IHC (bottom, blue color scheme) require invasive analysis at each time point, thus increasing the number of animals used and introducing inter-animal variability. In contrast, preclinical longitudinal molecular imaging (top, middle; green and red color schemes) enables the use of the same animals throughout the whole experiment, reducing the number of animals and minimizing inter-animal variability. Circles with different contrast indicate location of probe accumulation and recorded signal intensities. B, However, intensity-based imaging cannot discriminate bound- vs. non-bound-probes, measuring only the passive accumulation of probes in the organ of interest. In contrast, FRET in vivo imaging can measure dynamic receptor-ligand engagement using either intensity-based sensitized emission or lifetime-based in vivo imaging approaches.

4 DISCUSSION

Noninvasive molecular imaging approaches have been used for assessment of drug distribution and delivery in vivo with great success (61, 62). Noninvasive imaging enables longitudinal assessment of preclinical drug candidates without the need to sacrifice animals at every time point of interest. Moreover, using the same animal across multiple time points minimizes inter-animal variation (Fig. 5A). The localization of imaging contrast agents provides insight into the distribution of pharmaceutical or biopharmaceutical compounds administered to the animals. Hence, molecular targeted imaging has been used for in vivo studies of pharmacokinetics and drug distribution using nuclear imaging (PET and SPECT) (63, 64). The output of these modalities is intensity information, which is used to represent the localization of the drug. Unfortunately, this information cannot be used to distinguish between co-localization in the same spatial region and the accurate direct measurement of target binding or cellular delivery. This limitation often requires an invasive ex vivo approach to fully reveal binding of the administered compound to its respective target. The method of choice is histopathology – including immunohistochemistry or immuno- (fluorescence) staining (Fig. 5A). Though, analysis of ex vivo samples lacks whole-body drug distribution context, which should include other important organs besides the pathological ones. Additionally, ex vivo investigations require sacrifice of the animal for each time point considered, leading to increased biological variations.

We have previously shown that dynamic TfR-Tf receptor-ligand engagement can be studied in vivo using MFLI FRET imaging (40, 41, 59). Transferrin (Tf) is an iron-carrying protein which can bind its homodimeric transferrin receptor (TfR) at the surface of all cells in the organism. TfR is a homodimeric membrane-bound glycoprotein characterized by an inter-dimer distance less than 10 nm, which allows the monitoring of Tf binding using FRET (59). Tf-TfR binding has been monitored both in vitro and in vivo using FLI FRET imaging and validated by immunohistochemistry (40, 41, 43). As NIR-Tf probes are introduced into the body via intravenous injection, they will primarily label the liver, which acts as a major location for iron homeostasis regulation and displays a high level of TfR expression. Free dye and/or small labeled degradation products of NIR-Tf probes end up accumulating non-specifically in the urinary bladder, due to its role as an excretion organ (43).

As demonstrated here, this type of study can be performed using both intensity- and lifetime-based approaches (Fig. 5). However, the intensity-based approach requires spectral correction that are cumbersome to implement experimentally, as additional calibrating samples are necessary. The intrinsic complexity of the corresponding 3-cube method commonly employed

in vitro is further compounded by the fact that different animals need to be employed, raising questions about the reproducibility of this approach. Altogether, 9 independent measurements involving three mice were required (Table S3; Fig. 5B). While the data obtained with the two mice injected with donor-only and acceptor-only probes can in principle be reused for correction of new measurements with new mice injected with both probes, this requires that no change in acquisition parameters (and setup) takes place from one experiment to another, which might be difficult to ensure. In practice, it could very well be necessary to repeat these calibration measurements from time to time, increasing the cost and complexity of these measurements. Moreover, for the correction factors defined in Eqs. A.12 and A.13 to be valid, it is critical that the dye environment in the different mice used for their estimation, as well as the properties of the surrounding tissues, are similar (as implicitly assumed in the derivation). This may prove extremely difficult to ensure due to mouse-to-mouse variability, in particular when perturbations such as tumor xenografts are involved, since xenografts grown from the same cell line often possess variable size, cell density and microenvironments across different animals (40, 65).

In vivo FRET measurement protocols have traditionally relied on reporting a relative increase in acceptor intensity of FRETing sample compared to non-FRETing sample (*i.e.*, proximity ratio). However, imaging throughout the body of small animals upon probe injection, results in significant variation of fluorescence intensity as well as confounding emission leakage. Therefore, as in microscopy, intensity-based FRET in vivo macroscopy imaging should use the sensitized emission method, in which confounding emission leakages are properly corrected. However, sensitized emission FRET approach has not been adopted in small animal optical imaging, probably due to its complexity. Nevertheless, intensity-based sensitized emission FRET approach can be applied using widely available small animal imaging instruments such as the IVIS platform (PerkinElmer), making this imaging methodology accessible to many researchers. Sensitized emission FRET in vivo small animal imaging would allow the visualization of spatial drug distribution in a dynamic manner, enabling the understanding of the cellular mechanisms under pathophysiological context and providing valuable information for precision pharmacokinetics.

Lifetime-based FRET quantification provides robust and quantitative measures of target receptor engagement in vivo in a direct and non-invasive fashion. Even in the case of unique tumor microenvironments, lifetime-based FRET can be analyzed in each mouse independently. Hence, in its macroscopic implementation, MFLI-FRET is uniquely positioned to extract information regarding protein-protein interaction across entire small animals with high sensitivity (Fig. 5B). Importantly, MFLI FRET has been expanded to measure antibody-target engagement using NIR-labeled trastuzumab, an anti-HER2 clinically relevant antibody in HER2 breast tumor xenograft models (41, 48). Therefore, NIR MFLI FRET imaging is a quantitative and non-invasive tool for the optimization of targeted drug delivery systems based on receptor-ligand or antibody-target engagement in tumors in vivo. MFLI-FRET should find broad applicability in in vivo molecular imaging and could be extended to applications as diverse as image guided-surgery or optical tomography as well as other antibody-target systems, including other HER or immunotherapy receptors. Considering the recent development of next-generation time-resolved SPAD cameras, which are simpler to use and more affordable than the gated-ICCD technology used in this study and have recently been validated in MFLI-FRET imaging of tumor xenografts in mice models of human breast and ovarian cancer (48), MFLI-FRET appears uniquely well-positioned to impact the field of molecular imaging.

5 DATA AND SOFTWARE AVAILABILITY

All data and results are available on a public cloud repository (66) in order to ensure reproducibility.

6 ACKNOWLEDGMENTS

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7 AUTHORS CONTRIBUTION

NS performed MFLI experiments. AR conducted IVIS imaging. JTS analyzed the data. XM derived the equations. XI and MB designed and supervised the work. JTS, XM & MB wrote the manuscript. All authors reviewed the final manuscript.

8 DECLARATION OF INTERESTS

The authors declare no competing interests.

A APPENDIX: DERIVATION OF INTENSITY FRET EQUATIONS USED IN THE TEXT

A.1 Definitions and notations

We use the notations of Ref. (50) with minor modifications. In particular, we drop the 'ex' and 'em' indices in the quantities $F^{E_{em}}_{X_{ex}}$ used in the main text, replacing them by F^E_X in order to simplify the notation.

As a reminder, data acquisition involves one of two types of excitation channels X (laser line in Ref. (50)), bandpass filter in the IVIS device), corresponding to the donor ($X = D$) or acceptor ($X = A$) excitation wavelengths, and two emission channels E, characterized by bandpass filters specific for the donor emission ($E = D$) or acceptor emission ($E = A$). 4 possible combinations of excitation and emission channels are therefore possible in principle: $(X, E) \in \{(D, D), (D, A), (A, A), (A, D)\}$. The last combination is rarely used in practice, as it is uncommon to observe emission in a wavelength band (D) shorter than the excitation band (A). While it could have been relevant to use it in the measurements described in the main text, no such data was collected, and therefore the formalism described next will ignore it.

For a given molecular species S, the signal collected using an (X, E) excitation/emission pair is denoted $F^E_X(S)$, which we will assume to be corrected for background (data acquired in the same sample in the same conditions but with excitation source blocked).

Three different molecular species S are relevant in this study: $S \in \{DO, AO, DA\}$, where DO designates a donor-only species (molecule labeled with a donor fluorophore only), AO designates an acceptor-only species (molecule labeled with an acceptor fluorophore only) and DA designates a double-labeled (donor and acceptor) molecule. In principle, the DA species could be comprised of different sub-categories $\{DA_i\}_{i=1...n}$, characterized by different stoichiometries and/or different attachment sites and/or D-A distances. Formally, the same could be true of the DO and AO species, as fluorophore quantum yield could depend on the attachment site. In that case, we need to consider $\{DO_i\}_{i=1...d}$ and $\{AO_i\}_{i=1...a}$. We will here assume a single configuration for each species.

Following previous notations (50), we further distinguish the physical process Z at the origin of the recorded signal using the notation ${}^Z F^E_X(S)$. There are 3 processes of interest in this type of experiment:

(i) $Z = D$: direct excitation of the donor, followed by donor emission,

(ii) $Z = A$: direct excitation of the acceptor, followed by acceptor emission,

(iii) $Z = D{\rightarrow}A$: direct excitation of the donor, followed by non-radiative transfer to the acceptor, and acceptor emission.

We therefore have the following identities:

$$\begin{cases} F^E_X(DO) = {}^D F^E_X(DO) \\ F^E_X(AO) = {}^A F^E_X(DO) \\ F^E_X(DA) = {}^D F^E_X(DA) + {}^{D\rightarrow A} F^E_X(DA) + {}^A F^E_X(DA) \end{cases} \tag{A.1}$$

The first two simply state that no matter what excitation channel and emission channel are considered, only one single process needs to be considered when a single fluorophore species is present. The last identity expresses the fact that, in the case of the double-labeled species, three types of process can contribute to the signal: direct donor excitation/emission, donor excitation followed by FRET and acceptor emission, or direct acceptor excitation/emission.

Finally, for a sample comprised of a mixture of N_D donor molecules and N_A acceptor molecules, a fraction f_D (resp. f_A) of which are part of a D-A pair, we have for the total number N of fluorophores in the sample, and the respective numbers N_S of species S:

$$\begin{cases} N = N_A + N_D \\ N_{DO} = (1 - f_D) N_D \\ N_{AO} = (1 - f_A) N_A \\ N_{DA} = f_D N_D = f_A N_A \end{cases} \tag{A.2}$$

A.2 Fundamental equations

The equations used in ref. (50) were defined for single-molecules and thus require a simple multiplication by one of the N_S factors and reintroducing the terms neglected in that work:

$$F^D_D(DO) = N_{DO} I_D \sigma^D_{D_{exc}} \phi_D \eta^D_{D_{em}} \tag{A.3}$$

$$F_D^A\,(DO) = N_{DO} I_D \sigma_{D_{exc}}^D \phi_D \eta_{A_{em}}^D \tag{A.4}$$

$$F_A^A\,(DO) = N_{DO} I_A \sigma_{A_{exc}}^D \phi_D \eta_{A_{em}}^D \tag{A.5}$$

$$F_D^D\,(AO) = N_{AO} I_D \sigma_{D_{exc}}^A \phi_A \eta_{D_{em}}^A \tag{A.6}$$

$$F_D^A\,(AO) = N_{AO} I_D \sigma_{D_{exc}}^A \phi_A \eta_{A_{em}}^A \tag{A.7}$$

$$F_A^A\,(AO) = N_{AO} I_A \sigma_{A_{exc}}^A \phi_A \eta_{A_{em}}^A \tag{A.8}$$

$$F_D^D\,(DA) = N_{DA} I_D \left[\sigma_{D_{exc}}^D \phi_D\,(1-E)\,\eta_{D_{em}}^D + \sigma_{D_{exc}}^A \phi_A \eta_{D_{em}}^A + \sigma_{D_{exc}}^D E \phi_A \eta_{D_{em}}^A\right] \tag{A.9}$$

$$F_D^A\,(DA) = N_{DA} I_D \left[\sigma_{D_{exc}}^D \phi_D\,(1-E)\,\eta_{A_{em}}^D + \sigma_{D_{exc}}^A \phi_A \eta_{A_{em}}^A + \sigma_{D_{exc}}^D E \phi_A \eta_{A_{em}}^A\right] \tag{A.10}$$

$$F_A^A\,(DA) = N_{DA} I_A \left[\sigma_{A_{exc}}^D \phi_D\,(1-E)\,\eta_{A_{em}}^A + \sigma_{A_{exc}}^A \phi_A \eta_{A_{em}}^A + \sigma_{A_{exc}}^D E \phi_A \eta_{A_{em}}^A\right] \tag{A.11}$$

where Eqs. A.5-A.6 were assumed to be equal to zero in ref. (50) as were the last two terms of Eq. A.9 and the first and last term of Eq. A.11.

In the above equations:

- I_X is the X-excitation intensity (expressed in events per unit area, as detectors such as cameras do not measure photon energy, and instead only count the number of photon absorption events), which factors in integration time;
- $\sigma_{X_{exc}}^F$ is the absorption cross-section of fluorophore F (= D or A) at wavelength X_{exc} (or the average absorption cross-section in the X excitation wavelength band);
- ϕ_F is the quantum yield of fluorophore F (= D or A);
- $\eta_{E_{em}}^F$ is the detection efficiency of fluorophore F (= D or A) in emission channel E (= D or A);
- E is the FRET efficiency of the DA pair.

Note that we ignore all $F_A^D(X)$ terms in this analysis (species excited with the acceptor wavelength and detected in the donor emission channel), as their contribution should be negligible in the present case, but some experimental situations might require their consideration to obtain fully corrected quantities.

Finally, these expression neglect any higher order photophysical effects such as re-excitation, saturation, etc., which could potentially play a role in some specific experimental situations but are deemed negligible here.

Based on these general equations, we can now look at the two "reference" samples measured in this study, namely donor-only (DO, $N = N_{DO}$, $N_A = N_{DA} = 0$) and acceptor-only (AO, $N = N_{AO}$, $N_D = N_{DA} = 0$). Using Eqs. A.3-A.5 we obtain:

$$\begin{aligned} l_D &= \frac{F_D^A(DO)}{F_D^D(DO)} = \frac{\eta_{A_{em}}^D}{\eta_{D_{em}}^D} \\ d_D &= \frac{F_A^A(DO)}{F_D^A(DO)} = \frac{I_A \sigma_{A_{exc}}^D}{I_D \sigma_{D_{exc}}^D} \end{aligned} \tag{A.12}$$

and using Eqs. A.6-A.8:

$$\begin{aligned} l_A &= \frac{F_D^D(AO)}{F_D^A(AO)} = \frac{\eta_{D_{em}}^A}{\eta_{A_{em}}^A} \\ d_A &= \frac{F_D^A(AO)}{F_A^A(AO)} = \frac{I_D \sigma_{D_{exc}}^A}{I_A \sigma_{A_{exc}}^A} \end{aligned} \tag{A.13}$$

Coefficients l_D and d_A correspond to coefficients l and d in Ref. (50) and represent the donor leakage coefficient and the acceptor direct-excitation coefficient respectively. The two new coefficients d_D and l_A are counterparts of these coefficients, and are negligible if the donor absorption cross-section at the acceptor excitation wavelength, $\sigma_{A_{exc}}^D$, is negligible, and the detection efficiency of the acceptor in the donor emission channel, $\eta_{D_{em}}^A$, is negligible. These coefficients can be estimated

from the *DO* and *AO* signals measured with the different excitation-emission (X, E) combinations, provided these quantities all correspond to the same integration time, and more generally, same detection parameters, as well as constant excitation intensity for a given (X, E).

A.3 Pure *DA* sample case

The next step consists in extracting from Eqs. A.9-A.11, which are valid for "pure" *DA* species, an expression for E in terms of the measurable quantities $F_D^D(DA)$, $F_D^A(DA)$ and $F_A^A(DA)$. In order to simplify notations, we will define D, A and F as:

$$\begin{cases} D = I_D \sigma^D_{D_{exc}} \phi_D \eta^D_{D_{em}} \\ A = I_A \sigma^D_{A_{exc}} \phi_D \eta^D_{A_{em}} \\ F = I_D \sigma^A_{D_{exc}} \phi_A \eta^A_{A_{em}} E = \gamma E D \end{cases} \tag{A.14}$$

where the γ factor is defined by (50):

$$\gamma = \frac{\phi_A \eta^A_{A_{em}}}{\phi_D \eta^D_{D_{em}}} \tag{A.15}$$

With these notations, Eqs. A.9-A.11 can be rewritten:

$$\begin{cases} F_D^D(DA) = N\left[(1-E)D + d_A l_A A + l_A F\right] \\ F_D^A(DA) = N\left[(1-E) l_D D + d_A A + F\right] \\ F_A^A(DA) = N\left[(1-E) l_D d_D D + A + d_D F\right] \end{cases} \tag{A.16}$$

Replacing F by $\gamma E D$ results in 3 equations for the 3 unknowns D, A and E, the latter one being the only one of interest. Simple algebra yields:

$$\begin{cases} F_D^A(DA) - d_A F_A^A(DA) = N(1-d_A d_D)\left[l_D + (\gamma - l_D)E\right] D \\ F_D^D(DA) - l_A F_D^A(DA) = N(1-l_A l_D)(1-E) D \end{cases} \tag{A.17}$$

Taking the ratio of these two expression eliminates N and D, yielding the following result:

$$E = \frac{(1+\alpha l_A l_D) F_D^A(DA) - \alpha l_D F_D^D(DA) - d_A F_A^A(DA)}{(1+\alpha l_A (l_D - \gamma)) F_D^A(DA) - \alpha (l_D - \gamma) F_D^D(DA) - d_A F_A^A(DA)} \tag{A.18}$$

where we have introduced α defined by:

$$\alpha = \frac{1 - d_D d_A}{1 - l_D l_A} \tag{A.19}$$

This formula is identical to Eqs. (10)-(11) of Ref. (50) when $d_D = l_A = 0$ ($\alpha = 1$).

Note that Eq. A.19 can be expressed in terms of the sensitized emission FRET term $F^{FRET} = NF$ and some additional terms. To express NF as a function of $F_D^D(DA)$, $F_D^A(DA)$ and $F_A^A(DA)$, we look for (u, v) such that $F_D^A(DA) - u F_D^D(DA) - v F_A^A(DA)$ contains no D and A terms, based on the definitions of Eq. A.16. We obtain:

$$\begin{cases} u = \frac{1-d_A d_D}{1-l_A l_D d_A d_D} l_D \\ v = \frac{1-l_A l_D}{1-l_A l_D d_A d_D} d_A \end{cases} \tag{A.20}$$

From this, we obtain the following expression for $F^{FRET} = NF$:

$$F^{FRET} = \frac{1+\alpha l_A l_D}{1-d_A d_D} F^{A_{em}}_{D_{ex}}(DA) - \frac{d_A}{1-d_A d_D} F^{A_{em}}_{A_{ex}}(DA) - \frac{l_D}{1-l_A l_D} F^{D_{em}}_{D_{ex}}(DA) \tag{A.21}$$

It is then straightforward to verify that E in Eq. A.18 can be rewritten:

$$E = \frac{F^{FRET}}{F^{FRET} + \frac{\gamma}{1-l_A l_D}\left(F^{D_{em}}_{D_{ex}}(DA) - l_A F_D^A(DA)\right)} \tag{A.22}$$

based on the definitions of Eq. A.16.

A.4 General case: *D*, *A* and *DA* mixture

When the sample is a mixture M as defined by Eq. A.2, the 3 measured signals are given by the sums:

$$\begin{cases} F_D^D(M) = F_D^D(DO) + F_D^D(AO) + F_D^D(DA) \\ F_D^A(M) = F_D^A(DO) + F_D^A(AO) + F_D^A(DA) \\ F_A^A(M) = F_A^A(DO) + F_A^A(AO) + F_A^A(DA) \end{cases} \tag{A.23}$$

where the terms in the right hand side of Eq. A.23 are given by their expressions in Eqs. A.3-A.11. Using the definitions of Eq. A.14, we obtain:

$$\begin{cases} F_D^D(M) = N_D(1-f_D)D + N_A(1-f_A)d_A l_A A + N_D f_D[(1-E)D + d_A l_A A + l_A F] \\ F_D^A(M) = N_D(1-f_D)l_D D + N_A(1-f_A)d_A l_A A + N_D f_D[(1-E)l_D D + d_A A + F] \\ F_A^A(M) = N_D(1-f_D)d_D l_D D + N_A(1-f_A)A + N_D f_D[(1-E)l_D d_D D + A + d_D F] \end{cases} \tag{A.24}$$

Using the identity $F = \gamma E D$ and the fact that $f_D N_D = f_A N_A$ (Eq. A.2), this can be rewritten:

$$\begin{cases} F_D^D(M) = N_D(1 - f_D E + l_A \gamma f_D E)D + N_A d_A l_A A \\ F_D^A(M) = N_D((1-f_D E)l_D + \gamma f_D E)D + N_A d_A A \\ F_A^A(M) = N_D((1-f_D E)d_D l_D + d_D \gamma f_D E)D + N_A A \end{cases} \tag{A.25}$$

To eliminate $N_A A$, the same combinations as in Eq. A.17 give:

$$\begin{cases} F_D^A(DA) - d_A F_A^A(DA) = N_D(1-d_A d_D)[l_D + (\gamma - l_D)f_D E]D \\ F_D^D(DA) - l_A F_D^A(DA) = N_D(1-l_A l_D)(1-f_D E)D \end{cases} \tag{A.26}$$

which are identical to Eq. A.17, save for the replacement of N by N_D and E by $f_D E$. The final result for the product $f_D E$ is therefore identical to Eq. A.18, except for the quantities involved, which are now the intensities recorded for the mixture rather than the pure *DA* sample:

$$f_D E = \frac{(1+\alpha l_A l_D)F_D^A(M) - \alpha l_D F_D^D(M) - d_A F_A^A(M)}{(1+\alpha l_A(l_D-\gamma))F_D^A(M) - \alpha(l_D-\gamma)F_D^D(M) - d_A F_A^A(M)} \tag{A.27}$$

Note that this result is slightly different from the one proposed by Zal & Gascoigne (51), although the actual numerical difference will in general be negligible for $l_A << 1$ and $d_D << 1$, which is generally the case.

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SUPPLEMENTAL TABLES & FIGURES

Table S1: Intensity FRET correction parameters obtained herein using the dsDNA standards (Fig. 1).

l_D	d_D	l_A	d_A
0.28 ± 0.06	0.39 ± 0.03	$1.5\times10^{-2}\pm 4.0\times10^{-3}$	0.43 ± 0.05

Table S2: $f_D E$ quantification obtained herein using both intensity and FLI-FRET for the dsDNA standards (Fig. 1).

Technique	DO	DA_{27}	DA_{22}	DA_{17}
Intensity FRET	3.0 ± 9.1 %	18.0 ± 9.2 %	22.8 ± 9.2 %	50.4 ± 6.8 %
FLI-FRET	2.1 ± 1.6 %	24.7 ± 2.3 %	33.3 ± 2.3 %	52.5 ± 2.0 %

Table S3: Measurements required for *in vivo* FRET analysis. Intensity FRET quantification required multiple animals, as well as multiple acquisition channels. MFLI-FRET only requires a single animal sample and single acquisition channel. (*) Indicates measurements that need to be repeated assuming equivalent experimental conditions. (**) When an internal negative FRET control is not available a donor-only labeled animal may be needed for in vivo lifetime FRET analysis.

Intensity FRET			Lifetime FRET		
No.	Sample	Channel	No.	Sample	Channel
1	Donor Injection	Donor	1	FRET injection*	Donor
2	Donor Injection	Acceptor	2	Donor injection**	Donor
3	Donor Injection	FRET			
4	Acceptor Injection	Donor			
5	Acceptor Injection	Acceptor			
6	Acceptor Injection	FRET			
7	FRET Injection*	Donor			
8	FRET Injection*	Acceptor			
9	FRET Injection*	FRET			

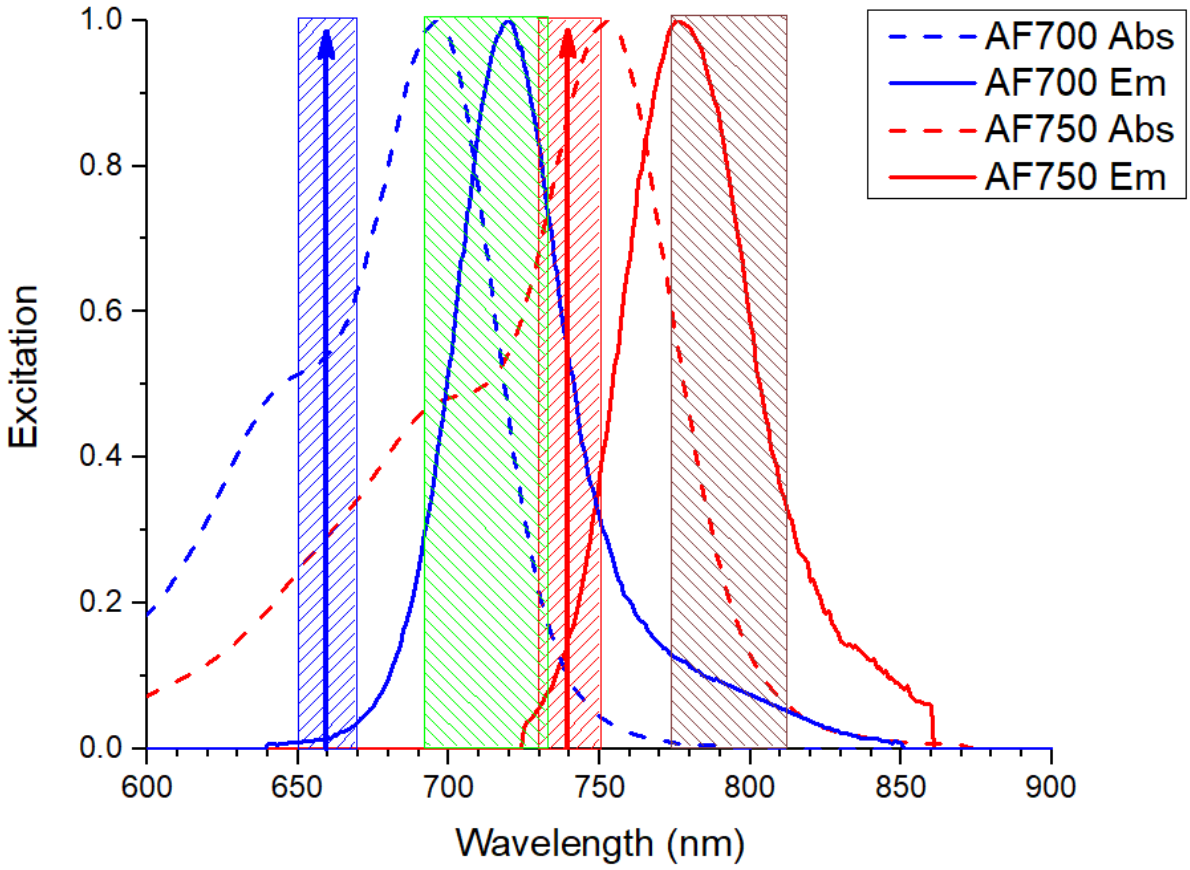

Figure S1: Excitation (dashed curves) and emission spectra (plain curves) of AF700 and AF750, as well as the excitation and emission filter characteristics used (hashed bands, the vertical arrows indicating the center of the excitation filters).

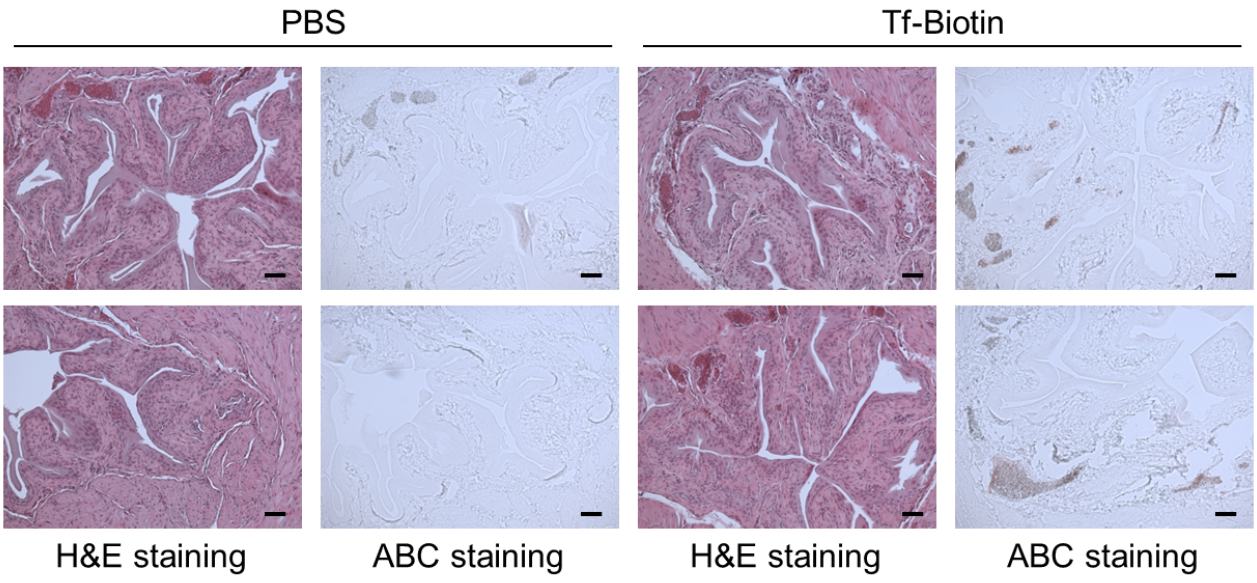

Figure S2: Mice were injected with biotin-Tf or PBS and sacrificed 6 h post-injection. Bladders were collected and processed for immunohistochemistry using ABC Elite and NovaRed kit (ABC staining) to visualize Tf-biotin. Parallel bladder sections were stained with Hematoxylin and Eosin (H & E staining) and imaged using 10x lens microscope for tissue morphology visualization. Arrows indicate staining of transferrin in blood vessels. Scale bar: 100 μm.