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Structural exploration and Förster theory modeling for the interpretation of gas-phase FRET measurements: Chromophore-grafted Amyloid- β peptides

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The distance-dependence of excitation energy transfer, e.g. being described by Förster theory (FRET), allows the use of optical techniques for the direct observation of structural properties. Recently, this technique has been successfully applied in the gas phase. The detailed interpretation of the experimental FRET results, however, relies on the comparison with structural modeling. We therefore present a complete first-principles modeling approach that explores the gas-phase structure of chromophore-grafted peptides and achieves accurate predictions of FRET efficiencies. We apply the approach to Amyloid- β 12-28 fragments, known to be involved in amyloid plaque formation connected to Alzheimer's disease. We sample structures of the peptides that are grafted with 5-carboxyrhodamine 575 (Rh575) and QSY-7 chromophores by means of replica-exchange molecular dynamics simulations upon an Amber-type forcefield parametrization as a function of the charge state. The generated ensembles provide chromophore-distance and -orientation distributions which are used with the spectral parameters of the Rh575/QSY-7 chromophores to model FRET-efficiencies for the systems. The theoretical values agree with the experimental average "action"-FRET efficiencies and motivate to use the herein reported parametrization, sampling and FRET-modeling technique in future studies on the structural properties and aggregation-behavior of related systems.

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I. INTRODUCTION

The investigation of protein structure and aggregation behavior is integral to understand protein function and dysfunction and in particular protein misfolding-related diseases¹. For example, the first stages of how the amyloid plaques of Alzheimer patients are formed and what causes this aggregation to happen is not yet fully understood. To tackle this quest, global structure-determination methodologies have to evolve that allow monitoring structural parameters of the precursor molecules and to observe misfolding at the molecular level. The gas phase provides a well-suited environment to implement such global strategy as the conditions for the experiments can be precisely defined and the overall structure of complex molecular edifices can be preserved using soft ionization techniques^{2,3}.

Förster resonance energy transfer (FRET) is a phenomenon that allows the probing of structural parameters by means of optical techniques. In the picture of Förster Theory developed in the late forties⁴, the efficiency for a resonant energy transfer between the chromophores depends on their mutual distance as well as orientation of their transition dipole moments. FRET is usually measured by recording a decrease in the fluorescence of the donor chromophore, with the concurrent emergence of the acceptor's fluorescence or by changes in fluorescence decay times. The distance dependence is described by Förster's approximation, (equation 1)

where the so-called critical Förster distance R_0 defines the distance at which 50% of the donor excitation is transferred to the acceptor moiety. It is routinely applied as a model for donor-acceptor distances on a length scale from 1 to 10 nm and when dipole-dipole (or convergent multipole-multipole) approximation for the electronic coupling can be employed appropriately for the donor-acceptor interaction⁵.

$$E_{FRET} = \frac{1}{1 + (r/R_0)^6} \quad (1)$$

The energy transfer is dominated by the $\frac{1}{r^6}$ distance-dependence that imprinted the term "spectroscopic ruler"⁶ on the technique. Thus, the interpretation of FRET results relies on the possibility to graft specific chromophores at well-defined sites on a molecule. Taking into account the flexibility of the chromophores and their linkers, the measured intensities provide information on the mutual distance of these specific sites⁷⁻¹⁰ e.g. to monitor structural changes in biomolecules^{6,11}.

The use of multiple dye pairs allows for triangulation of biomolecules, which provides three-dimensional structural information^{9,12-16}. Consequently, combining this phenomenon with the concept of single molecule detection¹⁷⁻¹⁹ and time-resolved experiments²⁰ allows for obtaining a unique glimpse into molecular structure as well as inter- and intra-molecular motions. While there exist numerous applications of FRET in the condensed phase²¹⁻²⁴, gas-phase FRET measurements²⁵⁻³⁴ suffer from difficulties in recording the fluorescence of a small number of ions in a mass spectrometer. The development of techniques capable of probing FRET in the gas

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phase with higher sensitivity is thus of high interest. This FRET detection problem has been recently overcome by combining photofragmentation action-spectroscopy with FRET (“action”-FRET)³⁵, opening up the possibility to integrate gas-phase FRET into a global approach for structure determination of proteins and protein complexes^{36–38}.

Extracting exact distances from an experimental calibration of FRET efficiencies alone is difficult because additional structural parameters like the orientation of the chromophore’s transition dipole moments affect the excitation energy transfer efficiency significantly. In fact, experimental single-molecule FRET (smFRET) studies on DNA strands allowed to specify those orientations and showed a clear orientation-modulation^{39,40} indicating that a negligence of the explicit orientational distribution in “fixed conformations” limits the predictive power. Even in systems where the dye orientation is not strongly restricted, a significant influence on the overall FRET-efficiency was suggested⁴¹.

Therefore, modeling efforts that access the structure of the chromophore-grafting sites and their distribution or temporal evolution are most attractive. These serve as the basis for the calculation of excitation energy transfer efficiencies and thus assign a detailed molecular picture to the experiments. The accuracy of FRET-efficiency prediction is limited by two major factors. Firstly, the Förster formula is straightforward in its use, but it relies on the validity of its underlying assumption of dipole-dipole interaction. Excitation energy transfer rates based calculated with approaches beyond Förster Theory⁵ can overcome this issue. For a recent review on the calculation of electronic coupling in excitation energy transfer see ref.⁴². In particular, nonadiabatic dynamics^{43,44} or semiclassical treatments^{45,46} of the excited state dynamics could deliver accurately the excitation energy transfer efficiencies (though at higher computational costs). Secondly, the prediction’s accuracy depends critically on the reliability of the structural exploration - being the underlying foundation for the subsequent numerical modeling. Notably, Hoefmüller *et al.* recently combined single molecule FRET experiments with the mutual dye orientation statistics obtained from Molecular Dynamics (MD) simulations. From the simulated time-dependent mutual orientations and distances, FRET efficiencies were calculated and the full statistics of individual photon absorption, energy transfer, and photon emission events were obtained from subsequent Monte Carlo (MC) simulations of the FRET kinetics⁴⁷. The technique is dedicated to time-coherent single-molecule studies in solution and its algorithms and implementation are part of a computational toolkit for the simulation of smFRET on the basis of molecular dynamics (MD) trajectory ensembles⁴⁸. Dedicated to gas-phase FRET, Talbot *et al.* have combined gas-phase fluorescence measurements and MD-based numerical modeling of the critical Förster distance²⁷. The qualitative insight from the calculations allowed them to interpret molecu-

lar structures on basis of their modeled conformations. However, conventional MD may not overcome barriers between the guessed initial configuration and the global minimum and may therefore fail to achieve sampling of a representative structural basis. The resulting errors in the modeling of FRET-efficiencies can be qualitative in nature also using accurate methods for excitation energy transfer calculations and can be overcome only with techniques that enhance the exploration of conformational space.

Such an enhanced structural sampling of gas-phase chromophore-grafted peptides together with numerical modeling of FRET-efficiencies based on distances and orientations at the same time has not yet been reported. In this work, we devise a strategy that succeeds in the accurate numerical first-principles modeling of FRET efficiencies based on a thorough exploration and enhanced sampling of gas-phase structures. In brief, we exploit Replica-Exchange Molecular Dynamics⁴⁹ (REMD) for the sampling procedure that offer a convenient framework to gas-phase conformations and statistical analyses⁵⁰. Taking into account the distance distributions and chromophore-orientation distributions at a given temperature, we calculate averaged FRET-efficiencies according to Förster-Theory and finally compare the results with experimentally measured “action”-FRET efficiencies published elsewhere⁵¹. The paper is structured as follows: we first describe the model peptide and chromophores used throughout this paper, then we explain our modeling setup detailing the procedure to obtain structural ensembles by means of REMD. Finally, we report the numerical modeling of FRET efficiencies and compare them to their experimental counterparts.

II. RESULTS AND DISCUSSION

A. System and experimental gas-phase FRET

In order to establish our strategy for converging “action-FRET” experiments and REMD ensemble based FRET modeling, we chose the Amyloid- β 12-28 fragment with the sequence V¹²HHQKLVFFAEDVGSNK²⁸ for which we recently published experimental data⁵². It has been shown that the naturally occurring mutations in the hydrophobic cluster (residues 17-22) are an important factor of Amyloid- β aggregation and which are probably indicating an enhanced risk of developing Alzheimer’s disease^{53–55}. We used the FRET-donor-acceptor pair that we have already proven suitable for this purpose³⁵. The acceptor chromophore is QSY7-C5 maleimide, a dark quencher absorbing and fragmenting efficiently at 545 nm in the gas phase^{35,56}. The donor chromophore is 5-carboxyrhodamine 575 (Rh575) C5 maleimide, absorbing at 505 nm in the gas phase. Rh575 was also used as the donor chromophore by Talbot *et al.*²⁷, and both the fluorescence excitation

FIG. 1. Chemical formulas of the QSY7-C₅ maleimide and Rhodamine 575-C₅ maleimide chromophores grafted to a Ac-cysteine-NH₂ peptide used for the parametrization of the nonstandard aminoacid residues (denoted as RDM and QSY residues in the following).

and emission spectra have been previously studied in detail⁵⁷. Both chromophores exhibit a C5 linker chain and a terminal maleimide group that allows thiol-specific grafting under mild conditions. The grafting does not affect significantly the absorption and emission properties of the chromophores. As the target peptide must contain 2 thiol groups for the grafting, our analyte molecule exhibits capped N- and C-terminal cysteine residues (Ace-CV¹²HHQKLVFFAEDVGSNK²⁸C-NH₂). The chemical formulas of the chromophores grafted onto the cysteine residues are depicted in figure 1. The QSY7-C₅ maleimide and Rh575-C₅-maleimide grafted cysteine residues will be denoted throughout the paper as QSY and RDM, respectively. Complementing the herein established theoretical technique, an experimental-theoretical study of the gas-phase structure of chromophore-grafted Amyloid- β peptides is published elsewhere⁵¹.

B. Computational setup and structural sampling

Replica Exchange Molecular Dynamics (REMD) simulations⁴⁹ were conducted to obtain structural ensembles for the chromophore grafted Amyloid- β peptides. The peptide was described on basis of the the Amber99⁵⁸ force field, that, different to more recent versions, was parametrized against gas-phase ab-initio calculations. It was completed with the Generalized Amber Force Field (GAFF⁵⁹) parametrizing the chromophore grafted cysteine residues QSY and RDM. GAFF is designed to “interoperate” with the Amber force field and is suitable for a broad variety of organic molecules. The dye residues were constructed by first finding a low-energy gas-phase conformation of Ace-QSY-NH₂ and Ace-RDM-NH₂ molecules. For this purpose we ran Molecular Dy-

namics (MD) with simulated annealing based on forces obtained with the semiempirical AM1 Hamiltonian⁶⁰ using a local code. To obtain the GAFF parameter sets we exploited AM1-bcc⁶¹ calculations that reproduce HF/6-31G* RESP charges. We used the software acpype⁶² in conjunction with Antechamber of the AmberTools version 14 suite of programs⁶³ for this process. From the parameter sets, we created nonstandard internal residues by removing either the Ace or the NH₂ cap and applying manually a small offset to the partial charges of the connecting atoms in order to re-establish the correct total charge of the residues. The residue libraries in Amber format can be obtained from the supplemental material. The interface between the GAFF and Amber99 residues was achieved by assigning Amber99 amino acid bond-, angle- and dihedral-parameters of the interfacing peptide bond.

All forcefield MD simulations and optimizations were carried out with the GROMACS suite of software in its version 4.6.5^{64–66}. Topologies of the chromophore-grafted Amyloid beta peptides in 3+, 4+, 5+ and 6+ charge states were constructed using the Leap program of the Ambertools distribution and converted to GROMACS topologies with acpype. The Ace-CV¹²HHQKLVFFAEDVGSNK²⁸C-NH₂ peptides were created in a full extended conformation and the cysteine residues mutated to QSY and RDM in both possible sequences. Additional to the fixed charges from the chromophores (amounting to +2), one, two or three protons were distributed over the LYS (in priority) and HIS residues (LYS 16, LYS 28, HIS 13, HIS 14) and by using the respective parameter set of protonated or neutral residues. For the 3+ and 5+ system, two different topologies resulting from the different possible locations of the proton were obtained. ASP and GLU residues were treated in their neutral form (ASH and GSH Amber residue types) since for relatively small peptides zwitterionic structures are generally less stable in the gas phase. For the MD simulations the systems were treated non-periodically without solvent molecules or implicit solvation and with no cutoff for nonbonded interactions. For optimizations, the L-BFGS algorithm was employed until a potential energy convergence of 1 kJ/mol was achieved.

MD trajectories were propagated using the velocity-verlet integrator with a timestep of 0.001 ps. Thermalization was achieved via coupling with an external heat bath in frame of the velocity-rescaling algorithm⁶⁷ that incorporates a stochastic term and thus provides the correct canonical ensemble. Replica exchange molecular dynamics simulations as implemented in GROMACS were conducted by propagating 20 trajectories in parallel. The temperatures of the replicas span a range of 220 K to 850 K (220, 236, 254, 272, 292, 314, 337, 362, 389, 417, 448, 481, 517, 555, 596, 640, 687, 737, 792, 850). After a 0.1 ns equilibration of the replica at their temperatures, sampling of the structures with REMD was begun. A REMD simulation was conducted with a length of 9.9 ns (9.9·10⁶ steps) totaling to 10 ns of MD trajec-

tory in all temperatures. Exchanges between adjacent replica were attempted every 200 integration steps. The chosen temperatures gave rise to exchange probabilities of 0.01 to 0.1 that ensured the mobility of the replicas through the whole temperature range. After a completed REMD sampling, the 220 K trajectory was quenched by extracting and optimizing every 1000th step to the next local minimum. A new calculation cycle was started with the found lowest-energy (LE) structure as starting point. The convergence of the procedure was monitored by the energy of the optimized LE structure and the potential energy distribution function (PEDF) of the 220 K and 292 K ensemble. Selecting the quenched lowest-energy structure from the 220K trajectory as starting point for a new run turned out to achieve convergence of the overall sampling procedure more quickly than using the 292K trajectory. When a convergence (energy of optimized LE structure and stationarity of the PEDF) was achieved, the 292 K trajectory was used to represent the ensemble of structures under ambient conditions. In total, about 25 μ s of MD trajectory were propagated.

In order to validate our force field parametrization, we tested it against electronic structure calculations with the PM7 Hamiltonian⁶⁸. The found lowest-energy structures from the 220 K REMD cycles were reoptimized without constraints using the MOPAC2012 code⁶⁹. The root mean square deviations (RMSD) of all atoms (including hydrogens) of the corresponding optimized structures are reported in table I. These indicate an overall correspondence of the structures with RMSD value between 0.4 and 0.8 Å (the difference between the semiempirical versus force-field optimized structures is visualized for the QSY-A β -RDM sequence in figure SA1 in the supplemental material) and confirm the expected accuracy of the chromophore-distance prediction of about 1 Å.

The semiempirical energetics can also be used to quantify the importance of the different possible tautomers (different charge locations) for the 3+ and 5+ systems. The relative PM7 energies for the lowest-energy structures of QSY-A β -RDM as well as RDM-A β -QSY in its 3+ and 5+ charge states are given in table I. Here the label LYN denotes which of the lysine residues was changed to the neutral form, while HIP denotes the position of the protonated histidine residue in Amber notation. The tautomers differ by at least 0.29 eV, so that only the lowest-energy tautomer ensemble was selected for further processing.

C. Calculation of FRET-efficiencies

The critical Förster distance for a chromophore pair is calculated as

$$R_0 = \left(\frac{9 \ln(10)}{128 \pi^5 N_A} \cdot \frac{\kappa^2 \Phi_D}{n^4} \cdot J \right)^{\frac{1}{6}}, \quad (2)$$

where Φ_D is the quantum yield of the donor in the absence of the acceptor, J the spectral overlap integral, N_A

TABLE I. Semiempirical calculations: RMSD values of atomic coordinates for force-field compared to PM7-optimized structures, relative PM7 energies ΔE of different PM7-optimized tautomers for 3+ and 5+ charge states

System	RMSD [Å]	ΔE [eV]
QSY-A β -RDM		
3+ (LYN 16)	0.41	0.37
3+ (LYN 28)	0.65	0.00
4+	0.78	
5+ (HIP 13)	0.75	0.83
5+ (HIP 14)	0.65	0.00
6+	0.59	
RDM-A β -QSY		
3+ (LYN 16)	0.53	0.00
3+ (LYN 28)	0.59	0.29
4+	0.58	
5+ (HIP 13)	0.79	0.38
5+ (HIP 14)	0.48	0.00
6+	0.59	

Avogadro's number, n the index of refraction being 1 in vacuum, and κ^2 is the orientation factor^{17,70,71}. We use a fluorescence quantum yield of 0.82 as known from Rh575 in ethanol⁷².

We first evaluated the sensitivity of FRET-efficiencies in the expected range of chromophore separations as a function of the distance between the optical units of the chromophores and the orientational factor κ^2 utilizing a fixed set of spectral parameters of the two chromophores (parameters are given in table II). The optically active parts of the QSY7 (left part in figure 1) and Rh575 (right part in figure 1) are marked in blue. The geometrical center of these units has been taken to evaluate the chromophore separation. To use the nuclear coordinates to define the center of transition dipole moments is a common strategy (see e.g. Refs.^{47,48}).

The donor and acceptor spectra were approximated assuming a Gaussian lineshape around absorption and emission maximum $\tilde{\nu}_{max}$ with the half-width of 1000 cm^{-1} . Equation 3 yields the absorption extinction coefficient ϵ_A in $\text{L/mol}^{-1}\text{cm}^{-1}$ from the oscillator strength f_e^{QSY} of the acceptor that we have obtained from a TDDFT calculation (see supplemental material), transition energies given in cm^{-1} . The calculated spectra were used to calculate the spectral overlap integral (equation 4 with $F_D(\lambda)$ being the normalized fluorescence intensity).

$$\epsilon_A(\tilde{\nu}) = \frac{2.175091 \cdot 10^8}{\sigma} \cdot f_e^{QSY} \cdot e^{-2.772 \left(\frac{\tilde{\nu} - \tilde{\nu}_{max}}{\sigma} \right)^2} \quad (3)$$

$$J(\lambda) = \int_0^\infty F_D(\lambda) \epsilon_A(\lambda) \lambda^4 d\lambda \quad (4)$$

The numerical results for these chromophore-parameters are visualized in figure 2 (compare also table

TABLE II. Spectral properties of the QSY7 and RDM chromophores used for FRET-modeling

variable	value	taken from
λ_{em}^{RDM}	505 nm	gas-phase action spectrum ³⁵
λ_{abs}^{QSY}	539 nm	gas-phase action spectrum ³⁵
Φ_D	0.82	Ref. ⁷²
f_e^{QSY}	0.85	this work (TDDFT calculation)
R_0^{calc}	40.3 Å	this work

FIG. 2. Calculated FRET efficiency of the QSY7-Rh575 chromophore pair as a function of the chromophore separation and the orientational factor κ^2 (equation 1,2,3,4 with the spectral parameters of QSY7 and RDM575 in table II). Color coding of E_{FRET} values according to the legend on the right. The average value of an isotropic orientational distribution $\langle \kappa^2 \rangle_{iso} = 2/3$ is indicated as a white line.

II). The calculated critical Förster distance R_0 for this chromophore pair amounts 40.3 Å (equation 2). The influence of a slightly different fluorescence quantum yield ($\Phi_D=0.9$) on the final FRET-efficiency values is only marginal ($R_0[\Phi_D=0.9]=40.9$ Å) as is a slightly different oscillator strength ($R_0[f_e=0.95]=41.0$ Å). Within a chromophore separation of 80 Å the FRET efficiency has dropped to nearly zero even when orientation would favor excitation energy transfer (high κ^2). On the other hand, the occurrence of high FRET-efficiencies can be substantially decreased by a perpendicular chromophore orientation leading to κ^2 approaching zero. Consequently, if a simple distinction between *near* and *far* is to be extended by a more accurate quantitative measure, an inclusion of the orientational effects is mandatory.

In order to predict the average FRET-efficiencies for the chromophore-grafted Amyloid- β peptides, we have postprocessed the calculated structural samples in the 3+, 4+, 5+ and 6+ charge states and have computed explicitly their FRET-efficiencies. In the following, we compare two different strategies with respect to the structural basis for the FRET-efficiency calculation: a) single low-energy conformations and b) the inclusion of the full canonical ensembles. The results for the modeled FRET-efficiencies in comparison with the offset-corrected exper-

FIG. 3. Explicitly calculated average FRET efficiencies $\langle E_{FRET} \rangle$ as a function of charge state: In red the experimental offset-corrected FRET-efficiency is given. $\langle E_{FRET} \rangle$ is calculated using a) the chromophore-distance of the lowest energy optimized structure r_{LE} and $\kappa^2 = 2/3$ from an isotropic orientation distribution b) explicit distribution of chromophore-distances as well as orientational factors $[r, \kappa^2]^{292K}$. Experimental values are taken from ref.⁵¹.

imental efficiencies are displayed in figure 3.

Figure 3a depicts the FRET efficiencies obtained for the lowest-energy structure (LE) with an assumed isotropic chromophore-orientation distribution ($\kappa^2 = 2/3$). The agreement of the efficiencies in figure 3a is qualitative, showing that already the single lowest-energy structure is indicative for the overall trend and may be used to classify chromophores as *near* or *far*. Here, thermal effects are not accounted for as the lowest-energy structure corresponds to a temperature of 0 K. This adds to the negligence of chromophore-orientational effects and explains the mediocre agreement.

There are two main strategies to improve the agreement. On the one hand, chromophore-orientations can be explicitly accounted for rather than assuming their isotropic distribution, and, on the other hand, a statistically representative thermal ensemble can be used as a structural basis. We included both effects by using explicit orientational and distance distributions in the 292 K ensemble and report the resulting E_{FRET} values in figure 3b. Evidently, this strategy - which we discuss in detail in the following - reproduces the experimental values in good agreement.

The results from analyzing the thermal ensembles are displayed in figure 4 together with a visualization of a small ensemble of structures. The sampled distance distribution functions as a function of charge state are displayed in figure 4a. The two sequences QSY-A β -RDM versus RDM-A β -QSY in the 3+ charge state exhibit different chromophore separations indicated by the two distinct features of the distribution function. In the 6+ charge state, three different conformations of the two sequences appear to be sampled at 292 K. In conclusion, a single distance from only one single structure might be a poor representation of the structured and nonsymmetric distance distributions present at 292 K.

FIG. 4. Normalized probability distribution functions of a) chromophore-distances and b) FRET-efficiencies E_{FRET} in the samples of Rh575/QSY7-labeled Amyloid- β peptide fragments (smoothed by Gaussians with the half-widths of 1 Å and 0.01 for r and E_{FRET}) in the charge states 3+, 4+, 5+ and +6 (from top to bottom). On top, a superposition of 5 structures (blue and red: Rh575 and QSY7 chromophores, green: Amyloid- β 12-28) from the 4+ ensemble visualizes the thermal effect.

The orientation factor κ^2 can be calculated from a structure according to equation 5 with the N-atoms of the chromophores defining the transition dipole moment vectors μ of the optically active units. Upon the sampled canonical ensembles of structures at 292 K we have calculated the κ^2 distributions shown in figure SA3 in the supplemental material.

$$\kappa^2 = \mu_D \cdot \mu_A - 3(\mu_D \cdot \mathbf{r}) \cdot (\mu_A \cdot \mathbf{r}) \quad (5)$$

From the distance distributions and the κ^2 distributions we calculated FRET-efficiency distributions in the 292 K ensembles as shown in figure 4b. These assume a very different functional form compared to the distance distributions. While for the 3+ and 6+, the distributions are centred around a value near to 1 and near to 0, they are spread over a large interval for the 4+ and 5+ charge state. This behavior is originating from the broad shape of the orientational factor κ^2 distribution function with values ranging between 0 and 4.

From the κ^2 distribution functions, the potential impact of chromophore orientations on the FRET-efficiencies can be deduced. For this reason, we quantified the deviation between the observed and the theoretical isotropic κ^2 distribution functions. The integral absolute deviation between the two functions (see supplemental

FIG. 5. Correlation between chromophore distance and FRET efficiency for Rh575/QSY7-labeled Amyloid- β peptide fragments in the charge range from +3 to +6. Blue color map shows the normalized probability distribution for the occurrence of a specific E_{FRET}, r pair in the samples (dark blue regions indicate a high probability). Discrete E_{FRET} values were calculated for all sampled distance and chromophore orientations of all charge states and both possible sequences RDM-A β -QSY and QSY-A β -RDM and smoothed by two-variate Gaussian functions. The ensemble-average E_{FRET}, r pairs for each charge state are given as green dots. In red, the analytic FRET-efficiency curve is given for a fixed orientation factor of $\kappa^2 = 2/3$.

material, figure SA3) amounts to 0.7 for the 3+ compared to about 0.35 for 4+, 5+ and 6+. This high deviation for the 3+ charge clearly indicates a nonisotropic distribution. In the molecular picture, the narrow κ^2 distribution is associated with discrete conformations being fixed by intramolecular interactions. Notice, that the particular conformation adopted is not only imposed by the overall charge state, but also depends on the charge location (that there exists an interplay between charge distribution, conformation and FRET-efficiencies can be depicted from the different 3+ κ^2 distributions in figure SA4 of the supplemental material). In the 4+, 5+ and 6+ charge systems a quasi-free rotation of the optically active units can be concluded from the κ^2 distributions - where fixing interactions are absent. Overall, the observed decrease in E_{FRET} with growing charge state is in agreement with the expectation that the Coulomb-repulsion between the charges is the main driving force for the structural changes upon protonation.

For making the overall effect of thermal distributions on FRET-efficiencies more comprehensive, we visualize the E_{FRET} versus distance correlation in figure 5. This 2-dimensional plot maps the calculated FRET-efficiencies onto the space of sampled distances. The ensemble-average values $\langle E_{FRET} \rangle$ depicted by the green dots correspond to the values reported in 3b. This 2D map demonstrates that there exists no one-to-one mapping of a single distance with a single FRET-efficiency. An

unambiguous assignment of a single conformation of a molecule to an experimental E_{FRET} value is therefore only possible when distance and κ^2 distributions are narrow and possess only one maximum. This limits the isotropic orientational distribution assumption ($\kappa^2=2/3$) often made. Concerning the ensemble-average of chromophore separations at 292 K, however, the ($\kappa^2=2/3$) assumption seems reasonable for the studied system. In conclusion, the knowledge of the explicit thermal structural ensembles is mandatory to perform a structural assignment to a FRET efficiency and to judge assumptions of orientational distributions. For this reason a global experimental-theoretical strategy is the most promising route towards the determination of biomolecular structures in the gas phase.

III. CONCLUSIONS AND OUTLOOK

In the presented work, we have performed a complete first-principles modeling of FRET-efficiencies of the chromophore-grafted Amyloid- β (12-28) fragment in the charge states 3+, 4+, 5+ and 6+. The modeling is based on an extensive exploration of the conformational space of twelve different systems (charge states, tautomers) by Replica-Exchange Molecular Dynamics (REMD) totaling to about 25 μ s of MD trajectory. We compared the FRET-efficiencies calculated using Förster Theory with measured values from the gas-phase “action”-FRET experiments (Daly *et al.*⁵¹). By analyzing the agreement between experiment and theory employing different structural representations, we found that an inclusion of distance distributions of the canonical ensemble outperforms calculations where only single low-energy conformations are considered. The former agrees very well with the experimental data. The average FRET-efficiencies of the systems studied here mainly depend on the average chromophore separations and orientational effects do not play an important role. For the 3+ charge state however, we have identified fixed chromophore orientations with a reduced rotational freedom which leads to clear non-isotropic κ^2 distribution. The fact that the extent of conformational fixing is depending on the particular system, suggests an explicit inclusion of chromophore orientation distributions into FRET modeling as best practice. Including thermal and conformational effects, we obtain excellent correspondence of theory and experiment, thereby assigning structural parameters such as the interchromophore distance distributions to the experimental signal. An “elongation” of the peptide and the resulting drop in FRET efficiency with increasing charge can therefore be interpreted. The achieved agreement encourages us to use the structural model, parametrization and sampling scheme for further investigations. The simulation protocol can be easily adapted to other peptides charge states and protonation patterns. It has to be noticed, however, that the bottleneck for the numerical accuracy of the procedure is a complete exploration

of the molecule’s conformational space. In view of the increasing conformational freedom of bigger systems, the presented strategy might be extended with biasing and steering algorithms⁷³.

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- ¹C. M. Dobson, *Nature* **426**, 884 (2003).
- ²B. T. Ruotolo and C. V. Robinson, *Curr. Opin. Chem. Biol.* **10**, 402 (2006).
- ³R. H. van den Heuvel and A. J. R. Heck, *Curr. Opin. Chem. Biol.* **8**, 519 (2004).
- ⁴T. Förster, *Annalen der Physik* **2**, 55 (1948).
- ⁵G. D. Scholes, *Annu. Rev. Phys. Chem.* **54**, 57 (2003).
- ⁶L. Stryer and R. P. Gaughland, *Proc. Natl. Acad. Sci. U. S. A.* **58**, 719 (1967).
- ⁷S. Kalinin, S. Felekyan, A. Valeri, and C. A. M. Seidel, *J. Phys. Chem. B.* **112**, 8361 (2008).
- ⁸H. Neubauer, N. Gaiko, S. Berger, J. Schaffer, C. Eggeling, J. Tuma, L. Verdier, C. A. M. Seidel, C. Griesinger, and A. Volkmer, *J. Am. Chem. Soc.* **129**, 12746 (2007).
- ⁹A. K. Woźniak, S. Nottrott, E. Kühn-Hölsken, G. F. Schröder, H. Grubmüller, R. Lührmann, C. A. Seidel, and F. Oesterheld, *RNA* **11**, 1545 (2005).
- ¹⁰S. Sindbert, S. Kalinin, H. Nguyen, A. Kienzler, L. Clima, W. Bannwarth, B. Appel, S. Müller, and C. A. M. Seidel, *J. Am. Chem. Soc.* **133**, 2463 (2011).
- ¹¹B. Wieb van der Meer, G. Coker III, and S. Y. S. Chen, *Resonance Energy Transfer: Theory and Data*, 1 edition (John Wiley & Sons, 1994).
- ¹²A. K. Woźniak, G. F. Schröder, H. Grubmüller, C. A. M. Seidel, and F. Oesterheld, *Proc. Natl. Acad. Sci. U. S. A.* **105**, 18337 (2008).
- ¹³A. T. Brunger, P. Strop, M. Vrljic, S. Chu, and K. R. Weninger, *J. Struct. Biol.* **173**, 497 (2011).
- ¹⁴M. G. Düser, Y. Bi, N. Zarrabi, S. D. Dunn, and M. Börsch, *J. Biol. Chem.* **283**, 33602 (2008).
- ¹⁵G. F. Schröder and H. Grubmüller, *Comput. Phys. Commun.* **158**, 150 (2004).
- ¹⁶S. Uphoff, S. J. Holden, L. L. Reste, J. Periz, S. van de Linde, M. Heilemann, and A. N. Kapanidis, *Nat. Methods* **7**, 831 (2010).
- ¹⁷E. Sisamakias, A. Valeri, S. Kalinin, P. J. Rothwell, and C. A. Seidel, in *Single Molecule Tools, Part B: Super-Resolution, Particle Tracking, Multiparameter, and Force Based Methods*, Methods in Enzymology, Vol. 475, edited by N. G. Walter (Academic Press, 2010) pp. 455 – 514.
- ¹⁸T. Ha, T. Enderle, D. F. Ogletree, D. S. Chemla, P. R. Selvin, and S. Weiss, *Proc. Natl. Acad. Sci. U. S. A.* **93**, 6264 (1996).
- ¹⁹E. B. Shera, N. K. Seitzinger, L. M. Davis, R. A. Keller, and S. A. Soper, *Chem. Phys. Lett.* **174**, 553 (1990).
- ²⁰A. J. Berghlund, A. C. Doherty, and H. Mabuchi, *Phys. Rev. Lett.* **89**, 068101 (2002).
- ²¹S. Weiss, *Science* **283**, 1676 (1999).
- ²²P. R. Selvin, *Nat. Struct. Biol.* **7**, 730 (2000).

- ²³B. Schuler, E. A. Lipman, and W. A. Eaton, *Nature* **419**, 743 (2002).
- ²⁴B. N. G. Giepmans, A. S. R., M. H. Ellisman, and R. Y. Tsien, *Science* **312**, 217 (2006).
- ²⁵A. S. Danell and J. H. Parks, *Int. J. Mass Spectrom.* **229**, 35 (2003).
- ²⁶M. Dashtiev, V. Azov, V. Frankevich, L. Scharfenberg, and R. Zenobi, *J. Am. Soc. Mass Spectrom.* **16**, 1481 (2005).
- ²⁷F. O. Talbot, A. Rullo, H. Yao, and R. A. Jockusch, *J. Am. Chem. Soc.* **132**, 16156 (2010).
- ²⁸V. Frankevich, X. W. Guan, M. Dashtiev, and R. Zenobi, *Eur. J. Mass Spectrom.* **11**, 475 (2005).
- ²⁹A. T. Iavarone, J. Meinen, S. Schulze, and J. H. Parks, *Int. J. Mass Spectrom.* **253**, 172 (2006).
- ³⁰A. T. Iavarone, A. Patriksson, D. van der Spoel, and J. H. Parks, *J. Am. Chem. Soc.* **129**, 6726 (2007).
- ³¹X. G. Shi, D. Duft, and J. H. Parks, *J. Phys. Chem. B* **112**, 12801 (2008).
- ³²V. Frankevich, V. Chagovets, F. Widjaja, K. Barylyuk, Z. Y. Yang, and R. Zenobi, *Phys. Chem. Chem. Phys.* **16**, 8911 (2014).
- ³³N. G. Hendricks, N. M. Lareau, M. Stow, J. A. McLean, and R. R. Julian, *J. Am. Chem. Soc.* **136**, 13363 (2014).
- ³⁴V. Frankevich, V. Chagovets, F. Widjaja, K. Barylyuk, Z. Yang, and R. Zenobi, *Phys. Chem. Chem. Phys.* **16**, 8911 (2014).
- ³⁵S. Daly, F. Poussigues, A.-L. Simon, L. MacAleese, F. Bertorelle, F. Chiro, R. Antoine, and P. Dugourd, *Anal. Chem.* **86**, 8798 (2014).
- ³⁶A. Miranker, C. V. Robinson, S. E. Radford, R. T. Aplin, and C. M. Dobson, *Science* **262**, 896 (1993).
- ³⁷A. J. R. Heck, *Nat. Methods* **5**, 927 (2008).
- ³⁸G. Vonhelden, T. Wyttenbach, and M. T. Bowers, *Science* **267**, 1483 (1995).
- ³⁹A. Iqbal, S. Arslan, B. Okumus, T. J. Wilson, G. Giraud, D. G. Norman, and T. Ha, *Science* **105**, 2 (2008).
- ⁴⁰F. D. Lewis, L. Zhang, and X. Zuo, *J. Am. Chem. Soc.* **127**, 10002 (2005).
- ⁴¹D. B. VanBeek, M. C. Zwier, J. M. Shorb, and B. P. Krueger, *Biophys. J.* **92**, 4168 (2007).
- ⁴²Z.-Q. You and C.-P. Hsu, *Int. J. Quant. Chem.* **104**, 102 (2014).
- ⁴³R. Mitrić, J. Petersen, and V. Bonačić-Koutecký, "Multistate nonadiabatic dynamics on the fly in complex systems and its control by laser fields," in *Conical Intersections* (2011) Chap. 13, pp. 497–568.
- ⁴⁴Y. Okuno and S. Mashiko, *Int. J. Quant. Chem.* **94**, 36 (2003).
- ⁴⁵J. Megow, B. Röder, A. Kulesza, V. Bonačić-Koutecký, and V. May, *ChemPhysChem* **12**, 645 (2011).
- ⁴⁶J. Megow, T. Renger, and V. May, *ChemPhysChem* **15**, 478 (2014).
- ⁴⁷M. Hoeffling, N. Lima, D. Haenni, C. A. M. Seidel, B. Schuler, and H. Grubmüller, *PLoS ONE* **6**, e19791 (2011).
- ⁴⁸M. Hoeffling and H. Grubmüller, *Comp. Phys. Commun.* **184**, 841 (2013).
- ⁴⁹Y. Sugita and Y. Okamoto, *Chem. Phys. Lett.* **314**, 141 (1999).
- ⁵⁰F. Chiro, F. Calvo, F. Albrieux, J. Lemoine, Y. O. Tsybin, and P. Dugourd, *J. Am. Soc. Mass Spectrom.* **23**, 386 (2012).
- ⁵¹S. Daly, A. Kulesza, F. Poussigues, A.-L. Simon, C. M. Choi, G. Knight, F. Chiro, L. MacAleese, R. Antoine, and P. Dugourd, *Chem. Sci.*, DOI:10.1039/C5SC01436H (2015).
- ⁵²T. N. Le, J. C. Pouilly, F. Lecomte, N. Nieuwjaer, B. Manil, C. Desfrancois, F. Chiro, J. Lemoine, P. Dugourd, G. van der Rest, and G. Grégoire, *J. Am. Soc. Mass Spectrom.* **24**, 1937 (2013).
- ⁵³N. S. de Groot, F. X. Aviles, J. Vendrell, and S. Ventura, *FEBS J.* **273**, 658 (2006).
- ⁵⁴A. Baumketner, M. G. Krone, and J.-E. Shea, *Proc. Natl. Acad. Sci. U. S. A.* **105**, 6027 (2008).
- ⁵⁵K. Murakami, K. Irie, A. Morimoto, H. Ohigashi, M. Shindo, M. Nagao, T. Shimizu, and T. Shirasawa, *J. Biol. Chem.* **278**, 46179 (2003).
- ⁵⁶Q. Enjalbert, M. Girod, R. Simon, J. Jeudy, F. Chiro, A. Salvador, R. Antoine, P. Dugourd, and J. Lemoine, *J. Anal. Bioanal. Chem.* **405**, 2321 (2013).
- ⁵⁷M. W. Forbes and R. A. Jockusch, *J. Am. Soc. Mass Spectrom.* **22**, 93 (2011).
- ⁵⁸J. Wang, P. Cieplak, and P. A. Kollman, *J. Comp. Chem.* **21**, 1049 (2000).
- ⁵⁹J. Wang, R. M. Wolf, J. W. Caldwell, P. A. Kollman, and D. A. Case, *J. Comp. Chem.* **25**, 1157 (2004).
- ⁶⁰M. J. S. Dewar, E. G. Zoebisch, E. F. Healy, and J. J. P. Stewart, *J. Am. Chem. Soc.* **107**, 3902 (1985).
- ⁶¹A. Jakalian, B. L. Bush, D. B. Jack, and C. I. Bayly, *J. Comp. Chem.* **21**, 132 (2000).
- ⁶²A. Sousa da Silva and W. Vranken, *BMC Research Notes* **5**, 367 (2012).
- ⁶³J. Wang, W. Wang, P. A. Kollman, and D. A. Case, *J. Mol. Graph. Model.* **25**, 247 (2006).
- ⁶⁴B. Hess, C. Kutzner, D. van der Spoel, and E. Lindahl, *J. Chem. Theory Comput.* **4**, 435 (2008).
- ⁶⁵D. Van Der Spoel, E. Lindahl, B. Hess, G. Groenhof, A. E. Mark, and H. J. C. Berendsen, *J. Comput. Chem.* **26**, 1701 (2005).
- ⁶⁶H. Berendsen, *Comp. Phys. Commun.* **91**, 43 (1995).
- ⁶⁷G. Bussi, D. Donadio, and M. Parrinello, *J. Chem. Phys.* **126**, 014101+ (2007).
- ⁶⁸J. J. P. Stewart, *J. Mol. Model.* **19**, 1 (2013).
- ⁶⁹J. J. P. Stewart, "Mopac2012, stewart computational chemistry, colorado springs, co, usa," <http://OpenMOPAC.net> (2012).
- ⁷⁰B. Schuler, in *Protein Folding Protocols*, Methods in Molecular Biology, Vol. 350, edited by Y. Bai and R. Nussinov (Humana Press, 2006) pp. 115–138.
- ⁷¹S. E. Braslavsky, E. Fron, H. B. Rodriguez, E. S. Roman, G. D. Scholes, G. Schweitzer, B. Valeur, and J. Wirz, *Photochem. Photobiol. Sci.* **7**, 1444 (2008).
- ⁷²F. L. Arbeloa, T. L. Arbeloa, E. G. Lage, I. L. Arbeloa, and F. C. Deschryver, *J. Photochem. Photobiol., A* **56**, 313 (1991).
- ⁷³F. Calvo, F. Chiro, F. Albrieux, J. Lemoine, Y. O. Tsybin, P. Pernot, and P. Dugourd, *J. Am. Soc. Mass Spectrom.* **23**, 1279 (2012).
- See supplemental material at [URL will be inserted by AIP] for an overlay of structures optimized with our force-field parametrization and the semiempirical PM7 level of theory; a detailed analysis of the influence of thermal effects for the distance and orientational distributions on the predicted FRET-efficiency; numerical values of the calculated FRET-efficiencies; inter-chromophore distance and orientational factor distributions of chromophore-grafted Amyloid- β 12-28; orientational factor distributions for a different tautomer (charge location) of the 3+ chromophore-grafted Amyloid- β 12-28; details of DFT/TDDFT calculations to obtain the QSY7 absorption oscillator strength and non-standard GAFF/AMBER99 residue libraries in Amber format for Ac-QSY, Ac-RDM, QSY-NH₂ and RDM-NH₂ residues.

Supplemental Material for:

Structural exploration and Förster theory modeling for the interpretation of gas-phase FRET measurements: Chromophore-grafted Amyloid- β peptides

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Assessment of parametrization

Figure SA1: Overlay of optimized structures of QSY-A β -RDM. The force-field optimized structure is given in yellow, while the PM7 optimized structure is given color-coded for the atom type.

Influence of thermal and orientational effects for FRET modeling

Figure SA2: Explicitly calculated average FRET efficiencies $\langle E_{\text{FRET}} \rangle$ as a function of charge state: In red the experimental offset-corrected FRET-efficiency is given. $\langle E_{\text{FRET}} \rangle$ is calculated using a) the chromophore-distance of the lowest energy optimized structure r_{LE} and $\kappa^2=2/3$ b) the chromophore-distance of the lowest energy structure r_{LE} and the actual value of κ^2 for the lowest-energy structure, c) the average from the thermal distribution of chromophore-distances $\langle [r]^{292\text{K}} \rangle$ and $\kappa^2=2/3$ d) explicit distribution of chromophore-distances as well as orientational factors $[r, \kappa^2]^{292\text{K}}$. For values see table SA1. Experimental values are taken from S. Daly et al., to be published (2015).

Table SA1: Calculated average FRET efficiencies $\langle E_{\text{FRET}} \rangle$ as a function of charge state using a) the chromophore-distance of the lowest energy optimized structure r_{LE} and $\kappa^2=2/3$, b) the chromophore-distance of the lowest energy structure r_{LE} and the actual value of κ^2 for the lowest-energy structure, c) the average from the thermal distribution of chromophore-distances $\langle [r]^{292\text{K}} \rangle$ and $\kappa^2=2/3$ d) explicit distribution of chromophore-distances as well as orientational factors $[r, \kappa^2]^{292\text{K}}$.

	a)	b)	c)	d)
3+	1.00	0.96	1.00	0.96
4+	0.51	0.25	0.37	0.29
5+	0.27	0.14	0.16	0.17
6+	0.15	0.00	0.10	0.06

Figure SA2 shows the influence of distance distributions and orientational distributions on the FRET-efficiency predictions. SA2a) and SA2b) correspond to figure 3a and 3b in the main text (taking the lowest-energy optimized structure and isotropic orientational distribution / explicit thermal distance and orientational distributions) Figure SA2b) shows the FRET-efficiency values from the lowest-energy optimized structures but with the explicit κ^2 value for the structure. Although the results for the 4+ and 5+ appear to better fit the experiment than the calculation in figure 3a/SA2a, a significant error cancellation might be present (QSY-A β -RDM and RDM-A β -QSY FRET efficiencies differ by up to 0.2) The deviation of the 6+ charge state from the experiment is characteristic for a situation in which such cancellation does not take place. Therefore, we regard calculating FRET efficiencies based on a single e.g. LE structure but accounting its explicit chromophore-orientation inferior to using $\kappa^2=2/3$. In SA2c) a thermal ensemble of distances corresponding to 292 K is used for the calculation. Here, the width and shape of the distance distribution function is accounted for in its influence on the FRET-efficiencies. Chromophore-orientations have been assumed to be isotropically distributed. This treatment proves to be rather accurate for the systems studied here, as a very good quantitative agreement with the experiment is achieved. Comparing figure SA2c with SA3d (figure 3b in the main text) indicates only a slight improvement of the numerical results when also explicit orientational distributions are included. Although the 3+ charge state displays fixed conformations with narrow κ^2 distribution, it does not greatly affect the averaged FRET-efficiencies. The low chromophore distance is responsible for FRET-efficiencies that are very near to one, irrespective of their orientation. It has to be pointed out that for a different system, conformational fixing could coincide with separations in the sensitive region (e.g. if conformational fixing would be existent in the 5+ charge state of our system) which would lead to a breakdown of the good agreement with the strategy of figure SA3c.

Analysis of distance distribution and chromophore orientation distribution

Figure SA3: Normalized probability distribution functions for chromophore-chromophore distance r (left) and orientational factor κ^2 (right) in the structure samples (292 K) of chromophore-grafted Amyloid- β fragments in the charge states 3+ to 6+ (smoothing by Gaussian functions with half-widths of 1 Å and 0.05). A 50/50 mixture of the two sequences QSY-A β -RDM and RDM-A β -QSY is assumed. The red curve in the κ^2 depicts ideal isotropic distribution of the transition dipole vectors. Differences in the sampled κ^2 distribution functions from isotropic limit are marked in gray.

Analysis of orientational factors as a function of charge distribution

Figure SA4: Normalized probability distribution functions for orientational factors κ^2 in the converged thermal ensembles for chromophore-grafted Amyloid- β in the charge state 3+ compared to isotropic distribution (green). The black curve $P(\kappa^2)_A$ is shown for the lowest-energy proton location where QSY-A β -RDM (LYN 28) and RDM-A β -QSY (LYN 16) are present (see main text and table 2) while $P(\kappa^2)_B$ denotes the configurations QSY-A β -RDM (LYN 16) and RDM-A β -QSY (LYN 28), which are higher in energy by 0.37 and 0.29 eV, respectively, and therefore not expected to be significantly populated at 293 K. Notice that $P(\kappa^2)_A$ is identical to the one shown in figure SA3 (upper part).

Calculation of QSY7 absorption oscillator strength

The absorption oscillator strength was estimated from a TDDFT calculation of the S1 state of a QSY7 dye. We obtained a low-energy conformation of the molecule by running simulated annealing coupled to molecular dynamics where we used the semiempirical AM1 Hamiltonian [1] for the forces. This structure was optimized at the B3LYP/6-31G** level of density functional theory [2-5] and applying Grimme's D3BJ dispersion correction [6]. The S1 $\leftarrow$ S0 transition properties were calculated with the CAM-B3LYP functional [7] and using the 6-311++G** basis set. All calculations were performed using Gaussian09 [8].

Nonstandard residue parameter files in Amber format

AC-QSY-Nterm.lib – library for the N-terminal QSY residue

AC-RDM-Nterm.lib – library for the N-terminal RDM residue

SupplementalMaterial.CHROMOPHORES-AMBERLIB.frcmod – additional parameters from the Antechamber procedure and used Amber-GAFF interface

QSY-NH2-Cterm.lib – library for the C-terminal QSY residue

RDM-NH2-Cterm.lib – library for the C-terminal RDM residue

References

- [1] M. J. S. Dewar, E. G. Zoebisch, E. F. Healy, J. J. P. Stewart, *J. Am. Chem. Soc.* **107** (1985), 3902–3909.
- [2] A.D. Becke, *J.Chem.Phys.* **98** (1993) 5648-5652
- [3] C. Lee, W. Yang, R.G. Parr, *Phys. Rev. B* **37** (1988) 785-789
- [4] S.H. Vosko, L. Wilk, M. Nusair, *Can. J. Phys.* **58** (1980) 1200-1211
- [5] P.J. Stephens, F.J. Devlin, C.F. Chabalowski, M.J. Frisch, *J.Phys.Chem.* **98** (1994) 11623-11627
- [6] S. Grimme, S. Ehrlich and L. Goerigk, *J. Comp. Chem.* **32** (2011) 1456-65.
- [7] T. Yanai, D. Tew, and N. Handy, *Chem. Phys. Lett.*, **393** (2004) 51-57.
- [8] Gaussian 09, Revision **D.01**, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski, and D. J. Fox, Gaussian, Inc., Wallingford CT, 2009.