THE ORIENTATIONAL FREEDOM OF MOLECULAR PROBES

THE ORIENTATION FACTOR IN INTRAMOLECULAR ENERGY TRANSFER

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ABSTRACT The measurement of the efficiency of Förster long-range resonance energy transfer between donor (D) and acceptor (A) luminophores attached to the same macromolecular substrate can be used to estimate the D-A separation, R. If the D and A transition dipoles sample all orientations with respect to the substrate (the isotropic condition) in a time short compared with the transfer time (the dynamic averaging condition), the average orientation factor <\chi^2> is 2/3. If the isotropic condition is not satisfied but the dynamic averaging condition is, upper and lower bounds for <\chi^2>, and thus R, may be obtained from observed D and A depolarizations, and these limits may be further narrowed if the transfer depolarization is also known. This paper offers experimental protocols for obtaining this reorientational information and presents contour plots of <\chi^2>_max and <\chi^2>_min as functions of generally observable depolarizations. This permits an uncertainty to be assigned to the determined value of R. The details of the D and A reorientational process need not be known, but the orientational distributions are assumed to have at least approximate axial symmetry with respect to a stationary substrate. Average depolarization factors are derived for various orientational distribution functions that demonstrate the effects of various mechanisms for reorientation of the luminophores. It is shown that in general the static averaging regime does not lend itself to determinations of R.

INTRODUCTION

Orientational molecular probes are old tools of the physical chemist. They permitted Perrin to estimate the lifetimes of excited molecules in solution by the use of steady-state polarization spectroscopy some 50 years ago (Perrin, 1926), and more recently have become useful to the biochemist for probing the dynamics and supramolecular structure of anisotropic systems such as membranes (Badley et al., 1973; Kawano et al., 1977). While these and similar uses of orientational probes are based on the interaction between a single transition dipole and the electric vector of a photon, it is the interaction between the two transition dipoles that plays the important role in determining the energy transfer rate between pairs of luminophores.

In most biochemical applications an orientational probe does not have a unique direction in the framework of a quasi-stationary substrate, but has limited freedom of motion with respect to it. This motion may be rapid or slow compared with the emission lifetime, and the range of allowed orientations may be narrowly defined or close to isotropic.

The work presented here develops a model that makes it possible to estimate the effect of orientational freedom of optical probes on experimental parameters such as fluorescence...
depolarization and Förster energy transfer rates. The last-named application is of primary concern and makes possible more accurate determinations of intramolecular distances, but a discussion of fluorescence depolarization in terms of models for limited orientational freedom based on several axially symmetrical distribution functions is included. The latter is primarily of heuristic value, because experimental methods for determining such distributions are not available at present, except possibly in oriented systems.

In the 30 years or so since Förster developed an exact quantum mechanical theory of resonance energy transfer for the "very weak" dipole-dipole coupling limit (Förster, 1948, 1951, 1965), an extensive and still rapidly growing literature has described the utilization of the phenomenon to determine intramolecular separations (e.g., recently Wu et al., 1976; Langlois et al., 1976; Shepherd et al., 1976; Wright and Takahashi, 1977; Papadakis and Hammes, 1977; Zukin et al., 1977) and conformational dynamics (Haas et al., 1975, 1977; Ohmine et al., 1977) of mainly biological macromolecules. Further impetus to such studies was undoubtedly provided by the finding that the predicted inverse sixth-power distance dependence of the transfer rate holds closely for several series of oligomers having donor (D) and acceptor (A) moieties bound covalently at their ends, the separation of which depends on the number of intermediate monomer units (Stryer and Haugland, 1967; Conrad and Brand, 1968; Gabor, 1968).

However, as recently emphasized (Eisinger and Dale, 1974; Dale and Eisinger, 1974, 1975), the use of the energy transfer technique as a "spectroscopic ruler" (Stryer and Haugland, 1967) has, with but few possible exceptions (Luk, 1971; Baugh er et al., 1974; Maróti and Szalay, 1976), suffered from a lack of knowledge of the relative orientations of the donor and acceptor transition moments, D and A, upon which the rate of transfer is also strongly dependent—to the extent that, in the most unfavorable case, no energy transfer occurs no matter what the separation\(^1\) (Chang and Filipescu, 1972). Usually, a value for the dipole orientation factor \(\kappa^2\), corresponding to a dynamic average or sometimes also a completely inappropriate static average (see below and Appendix A) over all orientations of both D and A is taken to be a good approximation, the tacit or stated assumption being that, even if orientational averaging is limited, extreme values, in particular those near zero, are unlikely. As pointed out earlier (Eisinger and Dale, 1974) and as demonstrated analytically in Appendix B, this argument has no statistical validity; indeed the opposite is true.

The fact that, in many cases, partial averaging of orientations undoubtedly occurs on a time-scale shorter than the transfer time (dynamic averaging), as indicated by some degree of depolarization of D and/or A emission, does not usually justify the use of the isotropic dynamic average value of \(2/3\), as can readily be seen from examples given previously (Dale and Eisinger, 1975). Fortunately, the assumption of isotropic dynamic orientational averaging for D and A need not be relied upon, since polarization spectroscopy can be used to obtain information on both the orientational freedom of the donor and acceptor luminophores and on their relative orientations. While this kind of information does not normally yield a unique dynamic average value for the orientation factor, it always delineates realistic upper and lower bounds. As will be seen, the range of possible values can, under favorable conditions, be quite narrow. However, even if it methodology provides a meaning of \(\kappa^2\) and hence in the D-A separation of D and A.

### ENERGY TRANSFER

The rate of (one-way) energy transfer between D and A moieties of suitable spectral overlap integral, the quotient of the intervention medium.

The orientation factor \(\kappa^2\) gives a measure of the separation between the dipole orientations. It is given by

\[
\kappa^2 = \cos \theta_f
\]

where \(\theta_f\) is the angle between the transition moments (D and A) and

\[
\cos \theta_f = \cos \theta_\rho \cos \phi + \sin \theta_\rho \sin \phi \cos \theta_z
\]

where \(\theta_\rho\) and \(\theta_\phi\) are the angles between the phosphorescence and fluorescence axes, respectively.

For a particular D-A pair, the transfer efficiency \(T\) is given by

\[
T = \frac{1}{4} \kappa^4 \cos \theta_f
\]

\[\text{FIGURE 1: Visualization of the transition moments (D and A) and...}

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1. At very close separations, transfer mechanisms other than the very weak coupling dipole-dipole (Förster) transfer considered here predominate (see, e.g., Eisinger et al., 1969). The remarks above and throughout the rest of this text apply only to Förster transfer.
named application is of primary of intramolecular distances, but a for limited orientational freedom included. The latter is primarily mining such distributions are not
quantum mechanical theory of e coupling limit (Förster, 1948, 1951) has described the utilization of e.g., recently Wu et al., 1976; E etal., 1977; Papadakis and nastics (Haas et al., 1975, 1977; Further impetus to such studies d inverse sixth-power distance of oligomers having donor (D) separation of which depends on land, 1967; Conrad and Brand, 974; Dale and Eisinger, 1974, e spectroscopic ruler" (Stryer and k, 1971; Baugher et al., 1974; if the relative orientations of the ich the rate of transfer is also rable case, no energy transfer 1972). Usually, a value for the average or sometimes also a dix A) over all orientations of stated assumption being that, particular those near zero, are demonstrated analytically in iperse is true.
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narrow. However, even if it is not, as is the case for a fixed D-A configuration, this methodology provides a meaningful estimate of the intrinsic uncertainty in the average value of $\kappa^2$ and hence in the D-A separation, without restrictive assumptions about the orientations of D and A.

ENERGY TRANSFER THEORY
The rate of (one-way) energy transfer in the very weak dipole-dipole coupling limit between D and A moieties of suitable spectroscopic properties is given (Förster, 1948, 1951, 1965) by

$$k_T = (1/\tau_D)(R_0/R)^6,$$  \hspace{1cm} (1)

where $\tau_D$ is the (singly exponential) decay time of D emission in the absence of A, $R$ is the D-A separation and $R_0$ is the characteristic (Förster) separation.

$$R_0 = C\kappa^2,$$  \hspace{1cm} (2)

where $C$ is a constant for the system under investigation, made up of universal constants, a spectral overlap integral, the quantum yield of D in absence of A, and the refractive index of the intervening medium.

The orientation factor $\kappa^2$ gives the dependence of the interaction between two electric dipoles on their orientations. It can be defined by

$$\kappa^2 = (\cos \theta_T - 3 \cos \theta_D \cos \theta_A)^2,$$  \hspace{1cm} (3)

where $\theta_T$ is the angle between the D and A moments, given by

$$\cos \theta_T = \sin \theta_D \sin \theta_A \cos \phi + \cos \theta_D \cos \theta_A$$  \hspace{1cm} (4)

in which $\theta_D, \theta_A$ are the angles between the separation vector R, and D and A, respectively, and $\phi$ is the azimuth between the planes (D, R) and (A, R) (Fig. 1).

For a particular D-A pair, $i$, characterized by an orientation factor $\kappa^2$, the energy transfer efficiency $T_i$ is given by

\begin{figure}[ht]
\centering
\includegraphics[width=0.5\textwidth]{diagram.png}
\caption{Visualization of the angles used to define the relative orientations of the donor and acceptor transition moments (D and A) and the separation vector (R).}
\end{figure}

DALE ET AL. Orientational Freedom and $\kappa^2$
\[ T_i = \frac{k_i}{(\tau_{D,i} + k_T)} = \frac{k_i}{(C^{-1} R^b + \kappa^i)}. \] (5)

As has been emphasized elsewhere (Dale and Eisinger, 1976; Eisinger, 1976), only when each D-A pair in the experimental ensemble has the same fixed orientation, or, equivalently, the same extent of dynamic averaging (fast compared with the fluorescence and transfer rates), is it vital to substitute a single or average value for \( \kappa^i \) in Eq. 5, although for low transfer efficiencies this will be a good approximation in the static averaging regime too. This is true independently of whether the transfer efficiency is obtained by steady-state measurements of relative intensities, as has usually been the case, by time-resolved measurements of the decrease in donor lifetime (Wu and Stryer, 1972), or by analysis of the rise and decay of acceptor emission excited by light absorbed by the donor (Schiller, 1975), a method apparently not exploited experimentally to date. Definition of an appropriate unique or dynamic average value of the orientation factor in any given system is usually impossible without independent crystallographic information, which, with few exceptions (Luk, 1971; Baughner et al., 1974; Maróti and Szalay, 1976), is not currently available and even then is of uncertain validity for the solution conditions generally employed in the energy transfer experiment.

An inappropriate isotropic static average value of 0.476 (Galanin, 1955; Maximov and Rozman, 1962; Steinberg, 1968) has also been quoted quite extensively. Its inapplicability to the problem of intramolecular energy transfer is indicated in Eq. 5, which demonstrates that no average value independent of the D-A separation can be defined, as has been discussed in some detail elsewhere (Dale and Eisinger, 1976). It seems unlikely that isotropic averaging in the static limit will provide an appropriate model in any experimental single D-A pair system, but the effect of this is examined in Appendix A, included less for its practical applicability than to illustrate how much energy transfer efficiencies can be affected by the orientational averaging regime.

**Fluorescence Depolarization**

It has been known for more than 50 years that in general luminescence is partially polarized. This is true even for an assembly of randomly oriented luminophores in solution excited by polarized or, indeed, unpolarized light, provided only that complete thermal orientational relaxation does not occur in the interval between excitation and emission. The explanation of this phenomenon of photoselection (Albrecht, 1961) is that molecules whose transition moments are aligned closer to the direction of the electric vector of the exciting light (and therefore more nearly perpendicular to its propagation direction) are preferentially selected for excitation even when the exciting light is unpolarized. In many cases, certainly in most cases of interest in energy transfer studies, the observed polarization properties of the system can be well described in terms of specific transition moments associated with absorption and emission. The extent of depolarization of the emission transition moment from such a photoselected assembly reflects the change in orientation between the initial absorption and final emission transitions.

The most useful measure of the extent of polarization is the emission anisotropy \( r \) introduced relatively recently (Jablonski, 1960), but used implicitly much earlier (Perrin, 1936). This quantity may be defined by

\[ r = \frac{I_v - I_h}{I_v + I_h} \]

where \( I_v \), \( I_h \) are the vertically and horizontally at right angles to an arbitrarily polar directions being in the horizontal plane polarized intensities and corresponds to a

\[ d \]

where \( d \) represents depolarization factor

\[ d = \theta \]

\( \theta \) being the angle by which the transition is lowered from this limiting value, and acc

\[ \langle d \rangle \]

is appropriate.

Thus, in energy transfer between D-A, instance, the observed EA for the trans

\[ r = 0.4 \]

The azimuthal averaging that justifies because in a solution containing an e distribution of planes containing \( \theta \) ab

\[ 1 \text{In principle these vectors are coincident in a low E } \]

1. In principle these vectors are coincident in a low E, excitation or transfer efficiency, do not lend themselves to the analysis degeneracy of transitions contribute to the or planar degeneracy, such as may account for solutions, can be treated in the same way, at least to represent coincident, i.e. nondegenerate, ab
Eisinger, 1976), only when each orientation, or, equivalently, the fluorescence and transfer rates), is q, 5, although for low transfer averaging regime too. This is true by steady-state measurements of time-resolved measurements of the analysis of the rise and decay of or (Schiller, 1975), a method of an appropriate unique or system is usually impossible with few exceptions (Luk, 1971; nally available and even then is of employed in the energy transfer (Galanin, 1955; Maksimov and extensively. Its inapplicability to Eq. 5, which demonstrates that defined, as has been discussed in likely that isotropic averaging in imental single D-A pair system, ness for its practical applicability be affected by the orientational

\[ r = \frac{(I_v - I_h)}{I}, \]

where \( I_v \), \( I_h \) are the vertically and horizontally polarized components of the emission observed at right angles to an arbitrarily polarized excitation beam, the excitation and emission directions being in the horizontal plane. \( I \) represents the sum of any three orthogonally polarized intensities and corresponds to the total emission intensity. For V-polarized excitation, which is usually employed since it gives rise to the maximum EA,

\[ I = I_v + 2I_h, \]

since the symmetry of the system dictates that the third (unobserved) polarized component of the emission intensity is equal to the (observed) H-polarized component. Under these conditions the EA attains its limiting value of 0.4 if the absorption and emission moments coincide in the luminophore framework, and no extrinsic reorientation due, for instance, to rotation occurs. If a series of depolarizing events intervenes between absorption and emission and each reorientation is azimuthally isotropic with respect to the preceding one, the EA is lowered from this limiting value, and according to Soleilhet's theorem (Soleilhet, 1929)

\[ r = 0.4 \prod_i d_i, \]

where \( d_i \) represents depolarization factors (Soleilhet, 1929; Perrin, 1936) defined by

\[ d_i = \frac{1}{2} \cos^2 \theta_i - \frac{1}{2}, \]

\( \theta_i \) being the angle by which the transition moment is changed in the \( i \)th depolarization step. More commonly, a distribution of \( \theta \) values is of interest, and the use of an average depolarization factor

\[ <d> = \frac{1}{2} <\cos^2 \theta> - \frac{1}{2} \]

is appropriate.

Thus, in energy transfer between D and A fixed with respect to an immobile substrate, for instance, the observed EA for the transferred excitation energy \( (r_T) \) is

\[ r_T = 0.4 \left( \frac{1}{2} \cos^2 \theta_T - \frac{1}{2} \right) = 0.4d_T. \]

The azimuthal averaging that justifies the use of Soleilhet's theorem is assured in this case, because in a solution containing an ensemble of D-A pairs there exists an axially isotropic distribution of planes containing \( \theta_T \) about D.

\[ ^1 \text{In principle these vectors are coincident in a lowest-lying nondegenerate transition. As discussed elsewhere (Dale and Eisinger, 1975), cases of excitation or transfer into higher transitions, the moments of which are not parallel to that of emission, do not lend themselves to the analysis of the energy transfer problem proposed here. Axially symmetric degeneracy of transitions contribute to the overall depolarization process, according to Eqs. 8 and 9, while slight planar degeneracy, such as may account for limiting EA values less than 0.4 even in dilute rigid or highly viscous solutions, can be treated in the same way, at least to first order. In the following, therefore, both D and A will be taken to represent coincident, i.e., nondegenerate, absorption and emission transition moment vectors.} \]

DALE ET AL. Orientational Freedom and \( \kappa^2 \)
POLARIZED INTRAMOLECULAR EXCITATION ENERGY TRANSFER

For many applications of interest to the biochemist or polymer chemist, a realistic model for an ensemble of identical macromolecules in solution, each endowed with an equivalent D-A pair, is one in which both D and A have some degree of reorientational freedom with respect to a rigid macromolecular framework that defines their fixed separation. The macromolecule itself may be free to rotate in the supporting solvent, but the effect of slow substrate rotation, considered artificial in the present analysis, can be accounted for experimentally (Dale and Eisinger, 1975). This and the effect of segmental flexibility, which may alter the D-A separation (Cantor and Pechukas, 1971; Grinvald et al., 1972; Haas et al., 1975, 1977; Ohmine et al., 1977), have been discussed elsewhere (Dale and Eisinger, 1975, 1976) and are excluded from further consideration here.

In the following, this model in the dynamic averaging limit is analyzed to investigate the effects of limited orientational freedom of D and A on the orientation factor in Förster energy transfer and on the transfer depolarization. It is shown how estimates of upper and lower bounds for the average value of the orientation factor may be obtained by utilizing the information available from polarization measurements.

Transfer Depolarization

Fig. 2 illustrates the model employed for calculating the depolarization resulting from energy transfer between two luminophores with limited orientational freedom. The axially symmetric orientational distributions of the donor and acceptor transition moments are indicated by cones with axes D' and A', respectively. Fig. 2a shows schematically the three depolarizing events, those due to reorientation of D and A and that brought about by transfer between them. The overall depolarization between D and A depends only on the relative orientation of these two vectors, specifically on \( \cos \theta_{ij} = (\mathbf{D}_i \cdot \mathbf{A}_j) \) and not at all on the intermediate orientations D and A. In the dynamic reorientational limit, all possible orientations corresponding to D and A are explored many times during the transfer period so that the transfer rate does not depend on either the initial acceptor orientation A, into which the transfer of excitation occurs or the final donor orientation D, from which emission occurs. The average transfer depolarization is therefore simply that of all D, A pairs, each pair contributing with the same weight to the average, so that

\[
<d_T> = \frac{1}{2} \cos^2 \theta_{ij} - \frac{1}{2}.
\]

The azimuthal averaging implied in Eq. 12 is ensured by the fact that in a solution of identical D, A-labeled macromolecules, every vector D, fixed at the moment of excitation in the laboratory coordinate system, has a uniform distribution of planes containing \( \theta_{ij} \) about it.

Under the circumstances detailed above, the transfer depolarization process depicted in Fig. 2b, involving reorientation of D, to the axial orientation D' followed by transfer to the acceptor in its axial orientation A' and reorientation to A, is entirely equivalent to that in Fig. 2a and may formally replace it. Since D, and A are axially symmetrically distributed about D' and A', respectively, and the azimuthal orientation of A' about D' (being a particular case of A, about D) is random, Soleillet's theorem (Eq. 8) applies, and the average transfer

\[
<\text{depolarization factor}> = \text{the pro three depolarizing events depict}
\]

where \( <d_T^b>, <d_T^a> \) will be respectively, and \( d_T^a \) as the ax (mean) orientations, that is, where \( \theta_{ij} \) is the angle between I the following section.

As will be shown below, the depolarizations of D and A ex possible to determine \( d_T^a \), and b orientational distributions. Thi range of values, the signs of <
ENERGY TRANSFER

The chemist, a realistic model for an enzyme with an equivalent D-A orientation freedom with respect to separation. The macromolecule effect of slow substrate rotation, observed for experimentally (Dale and Haas, 1972; Haas et al., 1975, 1977; and Eisinger, 1975, 1976) and are not studied here. It is analyzed to investigate the orientation factor in Förster energy transfer estimates of upper and lower limits be obtained by utilizing the depolarization resulting from energy transfer. The axially symmetric moments are indicated by statistical the three depolarizing light about by transfer between monomers on the relative orientation of not all on the intermediate limit, all possible orientations the transfer period so that the orientation $A_i$ into which the from which emission occurs. The of all $D_i, A_j$ pairs, each pair

\begin{equation}
\langle d_i \rangle = \langle d_{i0}^0 \rangle \langle d_i^T \rangle \langle d_i^A \rangle.
\end{equation}

Figure 2 (a) Schematic representation of the three depolarizing steps after the absorption of excitation energy by the donor: donor depolarization, transfer depolarization, and acceptor depolarization. It is shown in the text that the depolarization corresponding to these steps is the same as in the three depolarizing events shown in $b$: axial depolarization of the donor, depolarization due to transfer between the axes of the orientational distributions of $D$ and $A$, and axial depolarization of the acceptor.

depolarization factor is the product of the three depolarization factors, corresponding to the three depolarizing events depicted in Fig. 2 $b$:

\begin{equation}
\langle d_i \rangle = \langle d_{i0}^0 \rangle \langle d_i^T \rangle \langle d_i^A \rangle.
\end{equation}

where $\langle d_{i0}^0 \rangle, \langle d_i^A \rangle$ will be referred to as the axial depolarization factors for $D$ and $A$, respectively, and $\langle d_i^T \rangle$ as the axial transfer depolarization factor associated with their axial (mean) orientations, that is,

\begin{equation}
\langle d_i^T \rangle = \frac{1}{2} \cos^2 \Theta_i - \frac{1}{2}.
\end{equation}

where $\Theta_i$ is the angle between $D_i^T$ and $A_i^a$ (cf. Fig. 3). A formal derivation of Eq. 13 is given in the following section.

As will be shown below, the axial depolarization factors may be obtained from observed depolarizations of $D$ and $A$ excited and observed separately. From these and $\langle d_i^T \rangle$ it is possible to determine $\langle d_i^T \rangle$, and, by use of Eq. 14, $\Theta_i$, the angle between the axes of the $D$ and $A$ orientational distributions. This determination is not always unique, because in a certain range of values, the signs of $\langle d_{i0}^0 \rangle$ and $\langle d_i^A \rangle$ obtained in this way are indeterminate. In
such situations $\Theta_T$ is rendered two-valued, as discussed in more detail later. Nevertheless, unambiguous structural information can often, perhaps in the majority of cases, be obtained.

**The Orientation Factor**

Clearly, some degree of dynamic reorientational averaging of $D$ and $A$ or both about their mean orientations will be reflected not only in a decreased EA (positive or negative) after energy transfer, but also in an orientation factor closer to the isotropic value of 2/3. Furthermore, as is the case for fixed $D$ and $A$ orientations (Dale and Eisinger, 1974, 1975), the magnitudes of these effects are related.

Consider Fig. 3, in which the model for axially symmetric $D$ and $A$ distributions is illustrated in detail. For the arbitrary donor and acceptor orientations defined, the orientation factor determined by substitution of Eq. 4 into Eq. 3 is

$$k^2 = (\sin \theta_D \sin \theta_A \cos \phi - 2 \cos \theta_D \cos \theta_A)^2,$$

(15)

where the azimuth $\phi$ is

$$\phi = \Phi + \Phi_A - \Phi_D.$$

(16)

It is readily seen that the angles defining the orientation of $D$ are related by the following identities:

$$\sin \theta_D \cos \phi_D = \sin \theta_D \cos \phi_D - \cos \theta_D \sin \psi_D \cos \gamma_D$$

(17)

$$\cos \theta_D = \cos \theta_D \cos \phi_D + \sin \theta_D \sin \psi_D \cos \gamma_D$$

(18)

along with the equivalent identities for $A$. On substitution of Eqs. 16–18 into Eq. 15, expansion, collection of terms and averaging over the azimuthal angles $\gamma_D$ and $\gamma_A$,

$$\langle \cos \gamma \rangle = \langle \sin \gamma \rangle = \langle \cos \gamma \sin \gamma \rangle = 0,$$

(19)

$$\langle \cos^2 \gamma \rangle = \langle \sin^2 \gamma \rangle = 1/2,$$

(20)

and over the appropriate ranges of $\psi_D$ and $\psi_A$, one sees that the (dynamic) average value of the orientation factor is

$$\langle k^2 \rangle = k^{21} \langle d^1_D \rangle \langle d^1_A \rangle + \frac{1}{2} (1 - \langle d^1_D \rangle) + \frac{1}{2} (1 - \langle d^1_A \rangle)$$

$$+ \cos^2 \Theta_D \langle d^1_A \rangle (1 - \langle d^1_A \rangle) + \cos^2 \Theta_A \langle d^1_A \rangle (1 - \langle d^1_A \rangle),$$

(21)

in which

$$k^{21} = (\sin \theta_D \sin \theta_A \cos \Phi - 2 \cos \theta_D \cos \theta_A)^2$$

(22)

is the axial orientation factor defined for the axial (mean) orientations $D'$ and $A'$, while

$$\langle d^1_D \rangle = \frac{1}{2} \langle \cos^2 \psi_D \rangle - \frac{1}{2}$$

(23)

and

$$\langle d^1_A \rangle = \frac{1}{2} \langle \cos^2 \psi_A \rangle - \frac{1}{2}.$$

(24)

By using the same averaging procedure and transfer depolarization factor $d$,

$$\langle d \rangle = \frac{1}{2} \langle \cos^2 \theta \rangle - \frac{1}{2}$$

(25)

where

$$\cos^2 \Theta_T = 1$$

Eq. 25 is seen to be identical with the theorem.

**Maximum and Minimum**

It is clear from Eqs. 13 or 25 that the depolarization may in principle be measured comparing Eqs. 22 and 26, some functions of the angles $\Theta_D$, $\Theta_A$, of these variables, the third condition for maximum and minimum content.

Thus, differentiating Eq. 21 for the two axial orientations

Either $\Theta_D$ or $\Theta_A$ may then be eliminated, angle reveals that $\langle k^2 \rangle$ has the

$$\langle k^2 \rangle_{\text{max}} = \frac{1}{2} (1 - \langle d^1_D \rangle^2)$$

and

$$\langle k^2 \rangle_{\text{min}} = \frac{1}{2} (1 - \langle d^1_D \rangle^2 + \cos^2 \Phi).$$

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more detail later. Nevertheless, the majority of cases, be obtained.

of D and A or both about their EA (positive or negative) after to the isotropic value of 2/3, Dale and Eisinger, 1974, 1975).

metric D and A distributions is orientations defined, the orientation

\[
\cos \theta_A^r, \quad (15)
\]

\[
\cos \gamma_D \quad (16)
\]

D are related by the following

\[
n \psi_D \cos \gamma_D \quad (17)
\]

\[
\cos \gamma_D \quad (18)
\]

of Eqs. 16–18 into Eq. 15, the angles \( \gamma_D \) and \( \gamma_A \),

\[
> = 0. \quad (19)
\]

(20)

dynamic) average value of the

\[
<k^2> \quad (21)
\]

\[
<k^2_{\lambda} > \quad (22)
\]

\[
\cos \theta_A \quad (23)
\]

\[
\gamma_D \quad (24)
\]

\[
\cos \gamma_D \quad (25)
\]

where

\[
\cos^2 \Theta_r = (\sin \Theta_D \sin \Theta_A \cos \Phi + \cos \Theta_D \cos \Theta_A)^2 \quad (26)
\]

Eq. 25 is seen to be identical with Eq. 13, derived by the direct application of Soleillet's theorem.

**Maximum and Minimum Values of the Orientation Factor**

It is clear from Eqs. 13 or 25 that, given values of \( <d_r> \), \( <d_{\theta}^r> \), and \( <d_{\lambda}^r> \), all of which may in principle be measured experimentally, \( \Theta_r \) can be determined. As can be seen by comparing Eqs. 22 and 26, some limitation is thereby imposed on the otherwise indeterminate functions of the angles \( \Theta_D, \Theta_A, \) and \( \Phi \) appearing in Eq. 21. Differentiation with respect to two of these variables, the third being determined by the fixed value of \( \Theta_r \), will establish conditions for maximum and minimum values of \( <k^2> \).

Thus, differentiating Eq. 21 twice with respect to the azimuthal angle \( \Phi \) shows that, if the two axial depolarizations have the same sign, a minimum occurs when \( k^2 \) is identically zero. Either \( \Theta_D \) or \( \Theta_A \) may then be eliminated, and double differentiation with respect to the other angle reveals that \( <k^2> \) has the minimum value

\[
<k^2>_{\min} = \frac{1}{2} \left[ 1 - \left( <d_{\theta}^r> + <d_{\lambda}^r> \right)/2 \right.
\]

\[
+ \cos \Theta_r \left( <d_{\theta}^r> <d_{\lambda}^r> (1 - <d_{\theta}^r>) (1 - <d_{\lambda}^r>) \right)^{1/2}, \quad (27)
\]

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when

$$\cos^3 \Theta_1 = \frac{\cos \Theta_r \left[ <d_{1}^* > (1 - <d_{1}^* >) \right]^{1/2}}{3 \left[ <d_{1}^* > (1 - <d_{1}^* >) \right]}$$

(28)

where the subscripts (1, 2) refer either to (D, A) or to (A, D). As a result of the condition given in Eq. 28, the sign of the term containing $\Theta_r$ in Eq. 27 is always positive. If the axial depolarization factors have opposite signs, a maximum with respect to $\Phi$ obtains, but it leads only to a minimum with respect to $\Theta_p$ or $\Theta_s$, and then only if this angle, say $\Theta_s$, is zero, implying the unique configuration $\Theta_2 = \Theta_r = \pi/2$.

On the other hand, maxima are values of $<d_{2}^*>$ and $<d_{2}^* >$. $\Theta_p \neq 0$ so that

$$\kappa^3 = \cos \Theta_r (3)$$

in Eq. 21 for $<k^2>$. No attempt was made, in the approximation by searching through $\Theta_r \leq \pi/2$, to search both possible transfer angles $\Theta_r$ a axial depolarization factors were calculated.

---

**Figure 4** Contour plots for obtaining extreme values of $<k^2>$ consistent with the observed donor, acceptor, and transfer depolarization factors. Contours for the minimum and maximum values appear in the lower right and upper left halves of the diagrams, each for a particular value of $d_1^*$, the axial transfer depolarization factor defined in Eqs. 13 and 14. The contours are plotted as functions of $<d_{2}^* >$ and $<d_{2}^* >$, the dynamically averaged axial depolarization factors of the two luminophores between which transfer occurs. These axial depolarization factors are obtained from the observed donor and acceptor depolarization factors by using Eq. 33. If one or both of the observed depolarization factors are less than 1/2, there is a degeneracy in the sign of the axial depolarization factors; and the appropriate sign must be chosen to obtain the absolute upper and lower limit of $<k^2>$ from these contour plots, as discussed in the text.

**Figure 5** Contour plots for obt. 0, 0.1, and 0.2.
On the other hand, maxima and minima with respect to \( \Phi = 0 \) or \( \pi \) occur for all relative values of \( <d_{b}^i> \) and \( <d_{s}^j> \). \( \theta_b \) and \( \theta_s \) are related to \( \theta_f \) in both limits of \( \Phi \) by

\[
\cos \theta_f = \cos \theta_r \cos \theta_t + \sin \theta_r \sin \theta_t, \tag{29}
\]

so that

\[
x^2 = \cos \theta_f (3 \cos^2 \theta_t - 1) + 3 \sin \theta_f \cos \theta_t \sin \theta_t, \tag{30}
\]

in Eq. 21 for \( <x^2> \). No attempt was made to determine analytical maxima and minima with respect to \( \theta_b \) or, equivalently, \( \theta_s \), for this function. Instead, they were found to a good approximation by searching through the variable from 0 to \( \pi \), confining \( \theta_f \) to the range \( 0 \leq \theta_f \leq \pi/2 \), equivalent to searching the physically relevant range \( 0, \pi/2 \) of the variable for both possible transfer angles, \( \theta_f \) and \( \pi - \theta_f \). Absolute maxima for any given values of the axial depolarization factors were obtained directly in this way, as were minima when \( <d_{b}^i> \)

![Contour plots](image)

**Figure 5.** Contour plots for obtaining upper and lower limits of \( <x^2> \), as in Fig. 4, but for \( d_f = -0.1, 0, 0.1, \) and \( 0.2 \).
and \( <d_1^*> \) have opposite signs. Absolute minima when these depolarization factors have the same sign required comparison of the values derived from the search with those obtained from Eq. 27.

This searching and checking procedure was accomplished by computer, and the results were fed directly into a plotting program which, for fixed values of \( d_1^* \), generated contours of \( <k^2>_{\text{max}} \) and \( <k^2>_{\text{mn}} \) as a function of the two axial depolarization factors. Since these contour plots are invariant with respect to inversion of \( <d_5^*> \) and \( <d_4^*> \), they are symmetrical with respect to reflection on the diagonal, and the contours for \( <k^2>_{\text{max}} \) and \( <k^2>_{\text{mn}} \) may be combined in a single square diagram. They are displayed in Figs. 4–7 for steps in the axial transfer depolarization factor of 0.1 in the range \((-0.5 \leq d_1^* \leq 1)\) as functions of \( <d_1^*> \) and \( <d_5^*> \), where again the subscripts \((1, 2)\) refer to either \((D, A)\) or \((A, D)\). In Fig. 8 the same contour plots for \( d_1^* \) values of \(-0.5, 0, 0.5, \) and \(1 \) are given, and the regions in which the isotropic assumption of \( <k^2> = \frac{2}{3} \) leads to appreciable errors in the derived intramolecular separation \( R \) are also indicated.

![Figure 6: Contour plots for obtaining upper and lower limits of \( <k^2> \), as in Fig. 4, but for \( d_1^* = 0.3, 0.4, 0.5, \) and 0.6.](image)

![Figure 7: Contour plots for \( d_5^* = 0.8, 0.9, \) and 1.0.](image)

When the transfer depolarization factors are large, the contour plots in Figs. 6–8 extend the search procedure used to indicate the maxima again used to indicate the assumption leads to inaccurate values of \( <k^2> \).

When both axial depolarization factors are large, inspection of Eqs. 21 and 22 shows that values of \( <k^2> \) are

\[
<k^2>_{\text{max}} = \frac{2}{3},
\]

and

\[
<k^2>_{\text{mn}} = \frac{1}{3}.
\]

These two limiting cases correspond to perpendicular, respectively, distributions of \( D \) and \( A \) (E-
these depolarization factors have the
search with those obtained from
by computer, and the results
values of $d_1^2$, generated contours of
depolarization factors. Since these
values $<d_2^2>$ and $<d_4^2>$, they are
the contours for $<\kappa^2>_{min}$ and
they are displayed in Figs. 4-7 for
the range ($-0.5 \leq d_1^2 \leq 1$) as
for $<\kappa^2>$, as in Fig. 4, but for $d_1^2 = 0.7$, 0.8, 0.9, and 1.0.

When the transfer depolarization is not determined, it is inconvenient to use these plots. Accordingly, the contour plot of Fig. 9 displays maxima and minima of $<\kappa^2>$ obtained by extending the search procedure used above to cover the range of $\Theta$ values also. Stippling is again used to indicate the regions in the $<d_1^2>$, $<d_4^2>$ space in which the isotropic assumption leads to inaccurate values of $\kappa$.

When both axial depolarization factors are positive, particularly simple expressions apply. Inspection of Eqs. 21 and 27 shows that under these conditions the maximum and minimum values of $<\kappa^2>$ are

$$<\kappa^2>_{max} = \frac{1}{2} (1 + <d_0^2> + <d_4^2> + 3 <d_0^2> <d_4^2>)$$

and

$$<\kappa^2>_{min} = \frac{1}{2} [1 - (<d_0^2> + <d_4^2>/2)]$$

These two limiting cases correspond to $D^0$, $A^0$, and $R$ being parallel-in-line and mutually perpendicular, respectively, cases presented previously for a specific model of dynamic distributions of $D$ and $A$ (Eisinger and Dale, 1974; Dale and Eisinger, 1974, 1975).

Dale et al. Orientation Freedom and $\kappa^2$
As was mentioned before and will be demonstrated below, axial depolarization factors with values less than 0.5 may be degenerate in sign. However, because it is possible to assign a negative value in some cases, all combinations of signs have been included in all the contour plots. In general, though, each appropriate combination of signs should be checked to ensure that the absolute maximum and minimum of \(<x^2>\) consistent with the data are selected. As an example, take \(<d_f^+> = 0.064, <d_f^-> = 0.8, and <d_f^0> = \pm 0.4\). According to Eqs. 13 and 25 this corresponds to \(d_f^+ = \pm 0.2\) (Figs. 4 and 5). Minimum and maximum values of 0.35 and 1.85 are obtained when \(<d_f^+>\) and \(d_f^-\) are both positive, and with both negative the extrema are 0.20 and 1.6. The absolute limiting values for \(<x^2>\) are therefore 0.20 and 1.85, corresponding to an uncertainty \(\pm 18\%\) about the mean value. For between 0.20 and 2.1, so that \(R\) large, estimation of \(d_f^+\) becomes a emission, as is immediately evident on the diagonals of the contour plot.

**Axial and Observed Depolarization**

Because \(<d_f^+>\) and \(<d_f^->\) are determined from \(<x^2>\)\(_{ml}\), while experiments yield observed depolarization factors. Dynamic orientational averaging of depolarized, i.e., the apparent apparent time-resolved EA decay measurement.

Analogously, the same is true for the presence of energy transfer. 3

\[\text{Figure 9} \quad \text{Contour plot similar to to obtain} \quad \text{and hence} \quad d_f^+ \text{is unknown. It is obtained using}
\begin{align*}
\text{larger ranges between} \quad <x^2>_\text{max} \quad \text{and}
\text{the error in} \quad R \text{resulting from the uncertainty greater than 20%}. \quad \text{It is between 10}
\end{align*}
\]
corresponding to an uncertainty in the derived intramolecular separation of approximately \( \pm 18\% \) about the mean value. For this case, if \(<d_f>\) and therefore \(d_f^1\) is unknown, \(<k^2>\) lies between 0.20 and 2.1, so that \(R\) can be obtained with little loss of accuracy (Fig. 9). By and large, estimation of \(d_f^1\) becomes less helpful the greater the extent of depolarization of D and A emission, as is immediately evident on comparing values for \(<k^2>_{\text{max}}\) and \(<k^2>_{\text{min}}\) along the diagonals of the contour plots shown in, for example, Fig. 8.

**Axial and Observed Depolarization Factors**

Because \(<d_p^1>\) and \(<d_k^1>\) are used to parametrize the contour plots for \(<k^2>_{\text{min}}\) and \(<k^2>_{\text{max}}\), while experiments yield \(<d_p>\) and \(<d_k>\), a general relationship between axial and observed depolarization factors is needed and is derived in this section. Due to the dynamic orientational averaging postulated above, the emission of A excited directly will be depolarized, i.e., the apparent \(r_{av}\) values derived either from a Perrin plot or from nanosecond time-resolved EA decay measurements will be less than the fundamental one of 0.4. Analogously, the same is true for the emission from D alone, either in the absence or the presence of energy transfer. From the donor geometry schematized in Fig. 2, it is evident that

1A difference in the apparent \(r_{av}\) for these two conditions indicates either (a) all or part of the reorientation is on the same time scale as transfer (\(r_{av}\) is always larger, i.e., less depolarization occurs, when transfer competes with deactivation because the emission lifetime shortens) or (b) presence of the acceptor has induced some conformational change affecting the range over which dynamic reorientation can occur (\(r_{av}\) may be larger or smaller in the presence of the acceptor). The second condition obtains also for acceptor emission in the presence or absence of donor.
the directly observed depolarization factor $<d_d>$ is an average over all initial and final orientations $D$ and $D_{i}$, respectively, where it is considered that emission rather than transfer takes place at the orientation $D$. Because once again only the initial and final orientations determine the depolarization, and each orientation $D$ and $D_{i}$ is azimuthally averaged about the axial orientation $D'_{i}$, and moreover covers identically the same reorientational range, Soleillet's theorem in a degenerate form applies, so that

$$r_{0//0.4} = <d_d> = <d_d'>^2$$

(33)

obtains along with the corresponding relationships for $A$. Eq. 33 is seen to be equivalent to Eqs. 13 and 25 when $d'_{i}$ is unity ($D'$ and $A'$ parallel) and the axial depolarization factors of $D$ and $A$ are the same.

The simple relationship of Eq. 33 allows rather precise determination of the axial depolarization factors of $D$ and $A$ (bar possible uncertainty in their signs in the range $-0.5 \leq <d_{d,A}'> \leq 0.5$) independently of the reorientational distribution function as long as it possesses axial symmetry. Even in the absence of strict axial symmetry, however, eq. 33 can be expected to constitute a reasonable approximation, because, for an irregular distribution, some “center of gravity” orientation about which pseudo-axial symmetry obtains should exist in most cases.

A number of illustrative axially symmetric distributions, which may constitute reasonably realistic models for restricted dynamic reorientational relaxation of fluorophores attached or adsorbed to a macromolecular substrate, are presented in Appendix C. Among them, possibilities of negative axial depolarization factors are clearly demonstrated by those models, which start in the limit of a rigid planar oscillator (corresponding, for instance, to two degenerate orthogonal transitions), which is formally allowed to perform axially symmetric vibrations out of plane.

**SUMMARY**

In this section the conditions under which long-range resonance energy transfer experiments lend themselves to the determination of intramolecular donor-acceptor separations are summarized and experimental protocols for evaluating this distance with known uncertainty are suggested.

As has been pointed out, the dynamic averaging regime is the preferred one and lends itself to unequivocal interpretation. In the static regime, on the other hand, since there is no luminophore depolarization, no information about the orientational distribution of the donor and acceptor can be obtained. The transfer depolarization factor alone provides no useful information about the orientation factor and thereby the D-A separation.

The intermediate averaging regime, in which the reorientational motion of the probe can in principle be separated from that of the substrate as a whole, may be interpreted in terms of the dynamic averaging case to a reasonable approximation, as discussed elsewhere (Dale and Eisinger, 1974). In the intermediate case, the transfer efficiency will always be lower than that which would obtain for the isotropic example in Appendix A dynamic limit assumption will re actually increases as the distribution transitions should be avoided, for instance (Dale and Eisinger, 1975), the correlated.

Because even a modest degree of $<d'>$ appreciably, it is important respect to the substrate be measured time dependence of the overall additional measurement of $<d>$ further.

It is useful to consider three excitation energy transfer might be:

Neither $<d_d>$ nor $<d_{d}>$ under these conditions maximum separation approx.: $(<d'> = 0.5)$, but allows an arbitrary not helpful in this case.

$<d_d>$ and/or $<d_A>$

When both donor and acceptor depolarization factor is not available; important to remember that, if $<d_d>$ less than 0.25, the $<d_d'>$ and/or $<d_A>'$ unknown sign in the absence of in $<$corresponds to a planar taken into consideration in these plots. The same plot may be known, the absolute maximum defined by the known $<d'>$ the.

Note that if one of the depolarization reorientational isotropy $<d_d'>$, so that the maximum error in

$<d_d>, <d_A>$, and

The appropriate plots from among any ambiguity in the sign of $d'$, thus obtained represent estima.
that which would obtain for the same geometry in the strict dynamic limit (compare the isotropic example in Appendix A), implying that both extrema derived for \( R \) under the dynamic limit assumption will represent overestimates. The error due solely to this effect actually increases as the distributions become more isotropic.

When extrinsic labels are used, they should be chosen to be closely approximated by the limits of either a linear or planar transition moment. The chromophores' vibrational or rotational reorientation may then be estimated by the model and the experimental methods discussed here. To measure the transfer depolarization factor, excitation into higher donor transitions should be avoided, for in such cases, as discussed at length in an earlier publication (Dale and Eisinger, 1975), the orientation and transfer depolarization factors are not correlated.

Because even a modest degree of reorientational freedom can reduce the uncertainty in \( <\kappa^2> \) appreciably, it is important that the depolarization factors for both luminophores with respect to the substrate be measured as accurately as possible, preferably by determining the time dependence of the overall depolarization process directly. As discussed above, an additional measurement of \( <d_T> \) serves to narrow the range of uncertainty of \( <\kappa^2> \) further.

It is useful to consider three cases under which \( <\kappa^2> \) from polarized intramolecular excitation energy transfer might be analyzed.

**Neither \( d_D > 0 \) nor \( d_A > 0 \) Known**

Under these conditions may be arbitrarily close to its limits of 0 or 4. This sets a maximum separation approximately 35% higher than under the isotropic assumption (\( <\kappa^2> = \frac{1}{6} \)), but allows an arbitrarily small minimum separation. Knowledge of \( <d_T> \) is not helpful in this case.

**\( <d_D> \) and/or \( d_A > 0 \) Known, \( <d_T> \) Unknown**

When both donor and acceptor depolarization factors are determined, but the transfer depolarization factor is not available, the contour plot of Fig. 9 should be consulted. It is important to remember that, if one or both of the measured depolarization factors are less than 0.25, the \( <d_D^2> \) and/or \( <d_A^2> \) values obtained as their square roots (Eq. 33) are of unknown sign in the absence of independent information (e.g. as to whether one or the other of them corresponds to a planar transition). All combinations of possible signs must therefore be taken into consideration in determining the maximum and minimum values of \( <\kappa^2> \) from this plot. The same plot may be employed when only one of the depolarization factors is known, the absolute maximum and minimum being obtained by searching along the line defined by the known \( <d_D^2> \) through all values of the other depolarization factor.

Note that if one of the depolarization factors is determined to be zero, corresponding to complete reorientational isotropy of that luminophore, the extreme values of \( <\kappa^2> \) are \( \frac{1}{4} \) and \( \frac{1}{6} \), so that the maximum error in \( R \) due to the assumption of \( \frac{1}{6} \) for \( <\kappa^2> \) is only about 12%.

**\( d_D > 0 \), \( d_A > 0 \), and \( d_T > 0 \) Known**

The appropriate plots from among those presented in Figs. 4–7 are chosen with due regard to any ambiguity in the sign of \( d_T^2 \), as indicated in the body of the text. The maxima and minima thus obtained represent estimates of the true and, in the absence of additional structural
information, irreducible uncertainty associated with the relative orientations of donor and acceptor moieties.

It is worth stressing again that the protocol described here is independent of the particular orientational distribution functions that may be chosen for the luminophores and requires only that they are at least approximately axially symmetric.

It should finally be mentioned that the uncertainty caused by the experimenter's ignorance of the orientation factor can, at least in principle, be overcome in certain cases. The use of a transition metal ion with a purportedly triply-degenerate transition as acceptor already reduces the uncertainty in $<x^2>$ to about 12% in the worst case, i.e. when the donor transition is linear and fixed (Latt et al., 1970, 1972; Darnall et al., 1976; Birnbaum et al., 1977). The recent use of Tb$^{3+}$, again with a purportedly triply-degenerate transition, as the donor to Co$^{2+}$ (Berner et al., 1975; Horrocks et al., 1975) presumably completely removed all uncertainty in $<x^2>$, although the assumption of an isotropic oscillator was not reported to have been tested by a depolarization measurement or by other optical techniques. That the electronic transitions of a metal ion exhibit a single bell-shaped envelope is not sufficient ground to assume that its absorption will be isotropic. The best confirmation of isotropy is the measurement of the absorption coefficients in a single crystal of the metal-ligand complex. It is also sufficient that measurements in solution of the absorption, the circular dichroism, and the magneto-circular dichroism all can be described by a single spectral feature.

While only a few macromolecules of biological interest are likely to bind transition metal ions specifically, it should be possible to design a variety of organic chelates for these metal ions, which may then be bound to specific sites either covalently or by adsorption, as has been suggested earlier (Horrocks et al., 1975) and now accomplished (Leung and Meares, 1977). This may not always be desirable or even possible (e.g. when one or both probe moieties are intrinsic to the system). In these and all other cases, where the donor and/or acceptor lack orientational isotropy due to a degeneracy of the transition moments, the protocols described in this paper provide the narrowest possible limits to the donor-acceptor separation obtained from excitation energy transfer experiments.

APPENDIX A

Intramolecular Energy Transfer in the Limit of Static Random (Isotropic) Orientational Averaging

It can be seen from Eq. 5 that the statically averaged energy transfer efficiency between a donor and an acceptor separated by a fixed distance $R$ is

$$<T>_s = \frac{\kappa_T}{(\tau_0^{-1} + \kappa_T)} = \frac{\kappa_T^2}{(C^{-1} R^6 + \kappa_T^2)}.$$  \(34\)

As has been discussed previously (Eisinger and Dale, 1974; Dale and Eisinger, 1975, 1976), this result cannot readily be adapted, in general, to evaluate $R$ for an ensemble of D, A pairs since the $R$ and $\kappa_T$ dependence cannot be separated, each D, A pair having a different and unknown $\kappa_T^2$. The only exceptions occur when (a) $<T>_s$ is vanishingly small so that it approaches the dynamic average efficiency $<T>_d$ or (b) both D and A have isotropic orientational distributions. The latter situation will be considered in the following.

The static average transfer efficiency given by Eq. 34 may be rewritten as

$$<T>_s = 1 - \frac{1}{(1 + a x^2)},$$  \(35\)

where $a = C R^{-6}$. For the case under the coupling of the donor electric field $\omega_d$ (Steinberg, 1968):

Averaging may then be accomplished appropriate ranges:

$$<T>_s = 1 - \int_0^1 \int_0^1 [1 + a(1 + 3x^2)].$$

where $b = a(1 + 3x^2)$. $<T>_s$, an arbitrary value of $x^2$ (see Eq. 2), by Fig. 10, which also shows for completely defined by Eq. 5 with $x^2 = \frac{1}{2}$ as approx. Since in the dynamic limit all I sampled during the transfer time. The error in the intramolecular set deriving $R$ from energy transfer off 0.5 and about 50% when $<T>_s$ is negligible, as is easily shown to be distributions are isotropic or not.

APPENDIX B

Distribution of Values

It has frequently been stated or tacitly assumed that complete dynamic isotropic orienta-
relative orientations of donor and

and is independent of the particular

luminophores and requires only

by the experimenter's ignorance

time in certain cases. The use of a

cation as acceptor already

the donor

all

luminescence from a pic oscilator was not reported in

do this optical techniques. That the

shape envelope is not sufficient

estimation of isotropy is the

tion, the circular dichroism, and

gle spectral feature.

likely to bind transition metal

organic chelates for these metal

ory by adsorption, as has been

shed (Leung and Meares, 1977). 

ne or both probe moieties are

the donor and/or acceptor lack

moms, the protocols described

or-acceptor separation obtained

Figure 10 The dynamically and statically averaged transfer efficiencies, \(<T>_{d}\) and \(<T>_{s}\), as

functions of \((R_0/R)^6\), calculated under the "isotropic assumption" for the orientational freedom of both

luminophores (cf. Appendix A). The area between the curves corresponds to all averaging regimes

between the static and dynamic limits. The averaging regime is seen to make little difference in the transfer

efficiency is low.

where \(a = CR^{-4}\). For the case under consideration, it is most convenient to express \(k^2\) in terms of the

coupling of the donor electric field vector that makes an angle \(\omega\) with the acceptor transition moment

(Steinberg, 1968):

\[
k^2 = (3 \cos^2 \theta_0 + 1) \cos^2 \omega. \tag{36}\]

Averaging may then be accomplished by substituting \(x = \cos \theta_0\) and \(y = \cos \omega\) and integrating over the

appropriate ranges:

\[
<T>_{s} = 1 - \int_{0}^{1} \int_{0}^{1} [1 + a(1 + 3x^2) y^2]^{-1} \, dy \, dx, \tag{37}
\]

where \(b = a(1 + 3x^2)\). \(<T>_{s}\) may be obtained as a function of \((R_0/R)\), where \(R_0\) contains an

arbitrary value of \(k^2\) (see Eq. 2), by numerical integration of Eq. 37. The result for \(k^2 = \frac{5}{3}\) is displayed in

Fig. 10, which also shows for comparison the corresponding dynamically averaged efficiency \(<T>_{d}\)

defined by Eq. 5 with \(k^2 = \frac{5}{3}\) as appropriate for the isotropic case considered.

Since in the dynamic limit all D, A orientations, including those favorable for rapid transfer, are

sampled during the transfer time, \(<T>_{d}\) always exceeds \(<T>_{s}\) for a given intramolecular separation.

The error in the intramolecular separation \(R\), introduced by assuming the wrong averaging regime in

deriving \(R\) from energy transfer efficiencies, in this case is readily seen to be about 15% when \(<T>_{d}\) is

0.5 and about 50% when \(<T>_{d}\) is 0.9 (cf. Fig. 10). At low transfer efficiencies the error becomes

negligible, as is easily shown to be true, e.g. from Eq. 5, independently of whether the orientational

distributions are isotropic or not.

APPENDIX B

Distribution of Values of the Orientation Factor

It has frequently been stated or tacitly assumed that the average value of \(\frac{5}{3}\) for \(k^2\), which corresponds to

complete dynamic isotropic orientational averaging of both D and A, is a reasonable approximation for
D-A pairs in which the D and A orientations are either fixed or incompletely averaged. It is of course
doubtful whether the use of a value equal to that for a statistical average of all possible relative D and A
orientations in the dynamic limit is justifiable in a particular situation. Leaving this question aside, the
claims for the approximate validity of the “two-thirds” assumption can be tested by investigating the
probability density and probability distribution of \( \kappa^2 \) under the assumption that all spatial orientations of
both D and A are equally likely and that the orientation of the donor is independent of that of the
acceptor.

The probability density of \( \kappa^2, p(\kappa^2) \), will be the probability density of \( (3 \cos^2 \theta_0 + 1) \cos^2 \omega \) taken over
all values of \( \theta_0 \) and \( \omega \) in Eq. 36. This is isomorphous with the probability density, \( p(z) \), of \( z = (3x^2 + 1) y^2 \) taken over all values of \( x \) and \( y \) between 0 and 1. For any fixed value of \( x \), \( p(z) \), will be
proportional to \( 1/\sqrt{z} \) for \( 0 < y < (3x^2 + 1) \) as \( p(z) = 1/2 \sqrt{az} \) over the interval \( 0 < y < a \) when
\( z = y^2 \). In the interval \( 1 < z < 4, p(z) \) may be obtained from \( p(z) \), by integration. Substituting \( v = 3x^2 + 1 \), one obtains

\[
p(z) \propto (1/\sqrt{z}) \int_z^4 (p(v)/\sqrt{v}) dv.
\]

(38)

As \( p(v) = \sqrt{3/(v-1)} \), the integral is \( \ln (\sqrt{v-1} + \sqrt{v}) \). Evaluating the constant of integration and
normalizing to unit total probability gives

\[
p(\kappa^2) = \left[ 2/\sqrt{3} \kappa^2 \right] \left[ \ln (2 + \sqrt{3}) - g(\kappa^2) \right].
\]

(39)

where \( g(\kappa^2) = 0 \) for \( 0 < \kappa^2 < 1 \) and \( g(\kappa^2) = \ln (\sqrt{\kappa^2} - 1 + \sqrt{\kappa^2}) \) for \( 1 < \kappa^2 < 4 \). This probability
density function is plotted in Fig. 11 and is equivalent, though simpler in form, to an expression derived
elsewhere (Jones, 1970). It can be seen at a glance that the most probable value of \( \kappa^2 \) is zero.

The probability distribution of this density function is obtained by integrating from zero to any
arbitrary value of \( \kappa^2 \):

\[
P(\kappa^2) = \int_0^{\kappa^2} p(z) dz.
\]

(40)

Substituting Eq. 39 into Eq. 40 and integrating gives

\[
P(\kappa^2) = \frac{\sqrt{3} \kappa^2 \ln (2 + \sqrt{3})}{\sqrt{3} \kappa^2 - \ln \left( \sqrt{\kappa^2 + \sqrt{\kappa^2 - 1}} - \sqrt{\kappa^2 - 1} \right)} \frac{1}{(1/\sqrt{3})[\sqrt{\kappa^2}]^2} \frac{\ln (2 + \sqrt{3})}{\sqrt{\kappa^2}}
\]

(41)

This probability distribution function is also plotted in Fig. 11. Inspection of the latter curve shows that,
for instance, there is approximately a 60\% probability that \( \kappa^2 \) has a value outside the limits \( 1/2 \) to \( 1/2 \), and an
almost 45\% probability that it is less than \( 1/2 \). In addition, there is approximately a 20\% probability that \( \kappa^2 \) has a value less than 0.1 of its average value of \( 1/2 \) and almost 12\% probability that it is less than 0.02.

The corresponding values derived for the D-A separation in the latter two cases are 0.11\% and 0.03\%, i.e.
about 0.68 and 0.56, of that obtained under the assumption of an average \( \kappa^2 \). It has in the past been argued that the “isotropic” value of \( \kappa^2 \) for \( \langle \kappa^2 \rangle \) has statistical validity for D-A
pairs with unknown relative orientations, since extreme values for \( \kappa^2 \) (0 or 4) are unlikely to be obtained for a
pair of independent transition dipoles. The foregoing discussion dispels the credibility of this argument.
Indeed, it shows that the opposite is true, since values of \( \kappa^2 \) near zero are seen to predominate in a
random distribution. There exists, moreover, a fundamental objection to obtaining an estimate for the
orientation factor on statistical grounds. A fluorescent label, be it intrinsic or extrinsic, is found at a
specific site of a macromolecule because of the unique structural or functional properties of its
interaction with the site. It is therefore likely that the resultant distribution of dipole orientations, while
somewhat heterogeneous, will be characterized by a narrow range of geometries. It is quite unlikely that
any particular geometry will obtain when it is calculated on the basis of probability distributions, as has
been proposed recently (Hillel and Wu, 1976), even when the calculations include, as there, estimates of

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Eq. 39

CHANGE [2/\sqrt{3}k^2]

TO READ (1/2\sqrt{3}k^2)

Eq. 41

CHANGE \sqrt{3k^2} \ln(2 + \sqrt{3}) (0 < k^2 < 1)

\[ P(k^2) = \sqrt{3k^2} \ln(2 + \sqrt{3}) - \]

TO READ

\[ P(k^2) = \left[ k^2/3 \right] \ln(2 + \sqrt{3}) \left[ (1/\sqrt{3})[k^2]^2 \right] \ln(2 + \sqrt{3}) \]

where \( \psi \) is the angle between the pl.
the orientational distribution. The au.

Average axial depolarization factors
and 44, weighted by the appropriate

\[ < d^* > = \left( \int_0^1 \right) \]

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CORRECTIONS

Page 180

Eq. 39

CHANGE [2/√3k]

TO READ (1/2√3k)

Eq. 41

CHANGE √3k^2 ln(2 + √3) (0<k^2<1)

P(κ^2) = √3k^2 [ln(2 + √3) - √k^2 ln(√k^2 + √k^2 - 1) - √k^2 - 1] (1<k^2<4).

TO READ

P(κ^2) = √3k^2 [ln(2 + √3) - √k^2 ln(√k^2 + √k^2 - 1) - √k^2 - 1] (1<k^2<4).

where ψ is the angle between the planar transition dipole and the plane normal to the symmetry axis of the orientational distribution. The axial depolarization factors for a given ψ or ψ' are

\[ d'(ψ) = \frac{1}{2} \cos^2 ψ - \frac{1}{2}, \]

or, correspondingly,

\[ d'(ψ') = \frac{1}{2} \sin^2 ψ' - \frac{1}{2}. \]

Average axial depolarization factors for the various distributions may be obtained by integrating Eqs. 43 and 44, weighted by the appropriate radial density function and geometrical element:

\[ <d'> = \left( \int_0^\pi d'(ψ) g(ψ) \sin ψ \, dψ \right) \left/ \left( \int_0^\pi g(ψ) \sin ψ \, dψ \right) \right.. \]
incorporates the case of all possible relative $D$ and $A$ vectors. Leav- ing this question aside, the con- can be tested by investigating the prin- nected that all spatial orientations of donor is independent of that of the

of $(3 \cos^2 \theta + 1) \cos^2 \omega$ taken over nity density, $p(z)$, of $z = (3x^2 + x_0$ fixed value of $\theta, (\nu)$ will be.

![Figure 11](image.png)

**Figure 11** The curve $P(x^2)$ gives the probability density that any particular value of $x^2$ obtains between two dipoles, each of which may adopt any orientation independently of the other. $P(x^2)$ is the probability distribution of $x^2$ having a value between 0 and $x^2$. The probability of $x^2$ being less than 0.1, for example, is seen to be 0.25.

restricted dynamic orientational averaging. Only a range of possible $x^2$ values can be defined as outlined in this work. Some value within this range will be the appropriate one, but it cannot be estimated more closely.

**APPENDIX C**

**Depolarization Factors for Some Axially Symmetric Orientational Distributions**

In this Appendix a number of models for a luminophore attached to a macromolecular substrate with some orientational freedom will be used to calculate the corresponding dynamically averaged axial and observed depolarization factors. Only axially symmetric distribution functions are considered, and the transition moments are assumed to be either linear, as would obtain for the lowest nondegenerate transition of a luminophore, or planar, corresponding, for instance, to two degenerate orthogonal transitions. These two limiting cases, here examined in parallel, correspond respectively to centrally and peripherally weighted angular distributions of a linear transition. In the former case the angular distribution is best defined by a radial density function $g(\psi)$, where $\psi$ is the polar angle between the transition moment and the axis of symmetry of the distribution. It is more convenient to give the results for planar oscillators in terms of

$$\psi' = \pi/2 - \psi,$$

where $\psi'$ is the angle between the planar transition dipole and the plane normal to the symmetry axis of the orientational distribution. The axial depolarization factors for a given $\psi$ or $\psi'$ are

$$d'(\psi) = \frac{1}{2} \cos^2 \psi - \frac{1}{2}.$$  \hspace{1cm} (43)

or, correspondingly,

$$d'(\psi') = \frac{1}{2} \sin^2 \psi' - \frac{1}{2}.$$  \hspace{1cm} (44)

Average axial depolarization factors for the various distributions may be obtained by integrating Eqs. 43 and 44, weighted by the appropriate radial density function and geometrical element:

$$<d'> = \left( \int^\pi_0 d'(\psi) g(\psi) \sin \psi \, d\psi \right) / \left( \int^\pi_0 g(\psi) \sin \psi \, d\psi \right).$$  \hspace{1cm} (45)

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or, correspondingly,

\[ <d^* > = \left( \int_0^{r/2} d^*(\psi') g(\psi') \cos \psi' \, d\psi' \right) / \left( \int_0^{r/2} g(\psi') \cos \psi' \, d\psi' \right) \]  \hspace{1cm} (46)

The dynamically averaged observed depolarization factors \(<d^*>\) are then, according to Eq. 33, simply the squares of these axial depolarization factors.

**Free Rotation about One or More Bonds**

This model is useful for examining the cumulative effects of rotation about various bonds between the luminophore and the macromolecular substrate. The radial density function for rotation about a single bond is

\[ g(\psi) = \begin{cases} 1 & (\psi = \Psi) \\ 0 & (\psi \neq \Psi) \end{cases} \]  \hspace{1cm} (47)

where \(\Psi\) is the angle between the transition dipole and the bond about which free rotation is assumed to occur. In other words, the transition moment is constrained to lie on the surface of a cone with half-angle \(\Psi\). Then

\[ <d^* > = d^*(\Psi) = \frac{1}{2} \cos^2 \Psi - \frac{1}{2}, \]  \hspace{1cm} (48)

so that

\[ <d^*> = <d^* >^2 = \left( \frac{1}{2} \cos^2 \Psi - \frac{1}{2} \right)^2. \]  \hspace{1cm} (49)

These equations have been given elsewhere (Dale and Eisinger, 1974, 1975) and are also obtainable as the limit of the expression for the time-dependence of depolarization derived much earlier for this model (Gottlieb and Wahl, 1963). Figs. 12a and b (with \(n = 1\)) display these depolarization factors as functions of \(\Psi\) and illustrate the fact that they vanish, corresponding to complete depolarization, at the "magic angle"

\[ \Psi_m = \cos^{-1} \left( 1/\sqrt{3} \right) \approx 54.7^\circ. \]  \hspace{1cm} (50)

If free rotation about several bonds is possible, Soleilh's theorem (cf. Eq. 8) permits the evaluation of the cumulative depolarization

\[ <d^*_n > = d(\Psi) \prod_{i=1}^{n} d(\Psi_i) = d(\Psi) <d^*_1>, \]  \hspace{1cm} (51)

where \(\Psi\) is, as before, the angle between the transition moment and the proximal bond about which rotation occurs, and \(\Psi_i\) are the angles between each of the \(n\) successive rotatable bonds. As always for the axially symmetric distributions considered here, the observed depolarization factor \(<d^*_n >\) is the square of this expression. Figs. 12a and b depict \(<d^*_n >\) and \(<d_n^* >\) for \(n = 1-6\) and \(\Psi = \Psi\) between 0 and \(\pi/2\), since these functions are symmetrical about the latter angle. The time dependence of depolarization caused by multiple bond rotations has also been considered previously (Wallach, 1967).

It is of interest to generalize the foregoing model to cases in which a degree of radial vibration of amplitude \(\epsilon\) about a mean angle \(\Psi\) occurs. The corresponding radial density function is then

\[ g_{\epsilon}(\psi) = \begin{cases} 1 & (\max (0, \Psi - \epsilon) \leq \psi \leq \min (\pi, \Psi + \epsilon)) \\ 0 & (\text{elsewhere}) \end{cases} \]  \hspace{1cm} (52)

where \(\Psi\) and \(\epsilon \leq \pi/2\). Upon substituting

\[ <d^* > = \begin{cases} \frac{1}{2} \epsilon \hspace{1cm} \epsilon < 0 \\ \frac{1}{2} \epsilon \hspace{1cm} \epsilon \geq 0 \end{cases} \]  \hspace{1cm} (53)

These functions and the observed \(<d^*>\) and \(<d^*_n >\) for a set of \(\epsilon\) values. Figs.
\[
\left( \int_{0}^{2\pi} g(\psi') \cos \psi' \, d\psi' \right). \tag{46}
\]

are then, according to Eq. 33, simply

rotation about various bonds between the

\[
\Psi = \frac{1}{2} \tag{48}
\]

\[
\Psi = \frac{1}{2} \cos^2 \tag{49}
\]

1974, 1975) and are also obtainable as

ion derived much earlier for this model

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ding to complete depolarization, at the

\[
\psi \tag{50}
\]

m (cf. Eq. 8) permits the evaluation of

\[
\Psi <d^*_n>, \tag{51}
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and the proximal bond about which

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epolarization factor <d^*_n> is the

> for n = 1-6 and \Psi, = \Psi between 0

er angle. The time dependence of

ered previously (Wallach, 1967),

lich a degree of radial vibration of

density function is then

\[
\sin (\pi, \Psi + \epsilon) \tag{52}
\]

\textbf{Figure 12} The curves labeled n = 1 in a and b give the axial and observed depolarization factors, respectively, for a luminophore constrained to the surface of a cone of half-angle \Psi. This orientational constraint corresponds to the existence of a single, freely rotatable bond between the luminophore and the substrate. The remaining curves, labeled n = 2, 3, 4, 5, 6, correspond to n equivalent rotatable bonds between luminophore and substrate and are seen to give rise to ever-decreasing depolarization factors.

\[
<d^* > = \begin{cases} 
\frac{1}{2} \cos^2 \Psi \cos^2 \epsilon - \cos^2 \Psi - \cos^2 \epsilon & (\epsilon \leq \Psi) \\
\frac{1}{2} \cos (\Psi + \epsilon) [1 + \cos (\Psi + \epsilon)] & (\epsilon \geq \Psi).
\end{cases} \tag{53}
\]

These functions and the observed depolarization factors derived from them are presented in Figs. 13 a and b for a set of \epsilon values. Frechland (1976) has recently considered the time-dependence of the

\textbf{Dale et al. Orientational Freedom and } \epsilon^2 \]
Space-Filling Models

Only a few space-filling models with axial symmetry will be considered here. While these were chosen for illustration, some correspond to constraints imposed by simple potential wells. The radial density functions $g(\psi)$ and $g(\psi')$ for these models are displayed in Fig. 14. Analytical expressions can be derived for the depolarization factors for all except the Gaussian model. Some of these expressions are rather complex and lengthy, and in practice the Gaussian model, by numerical integration, which forms an analogue of the infinite space-filling model considered previous (1974, 1975; Eisinger, 1976). It corresponded in connection with the time dependence of lipid bilayer membranes (Kawato et al. for which the axial depolarization factor

$$<d>$$

The depolarization factors computed incorporating peripheral weighting, i.e., the limit of a linear transition rotating distribution function

$$g_\psi(\psi') =$$

and is characterized by the very simple

This function is displayed, along with Fig. 15 b.

An error in a previous calculation of $<d>$ (Dale and Eisinger, 1975) is apparent and...
complex and lengthy, and in practice the corresponding depolarization factors were calculated, as for the Gaussian model, by numerical integration.

**Model I. Random Walk Within the Confinements of a Cone of Half-Angle \( \Psi \)** This model, which forms an analogue of the infinitely deep square well potential in two dimensions, is the only space-filling model considered previously in this connection (Eisinger and Dale, 1974; Dale and Eisinger, 1974, 1975; Eisinger, 1976). It corresponds to a random walk over a conical segment and has also been invoked in connection with the time dependence of restricted rotational motion of fluorescent probes in lipid bilayer membranes (Kawato et al., 1977; Kinosita et al., 1977). The radial density function is

\[
g_r(\Psi) = \begin{cases} 
1 & (0 \leq \Psi \leq \min(\Psi, \pi)) \\
0 & (\Psi > \Psi) 
\end{cases}
\]  

(54)

for which the axial depolarization factor reduces to

\[
<d^a> = \frac{1}{2} \cos \Psi (1 + \cos \Psi) .
\]  

(55)

The depolarization factors computed on this basis are presented in Fig. 15a. The corresponding model incorporating peripheral weighting, i.e., starting in the limit of a planar transition or, equivalently, in the limit of a linear transition rotating rapidly in a plane (cf. Kinosita et al., 1977), has the radial distribution function

\[
g_l(\Psi') = \begin{cases} 
1 & [0 \leq \Psi' \leq \min(\Psi', \pi/2)] \\
0 & (\Psi' > \Psi') 
\end{cases}
\]  

(56)

and is characterized by the very simple axial depolarization factor

\[
<d^a> = -\frac{1}{2} \cos^2 \Psi' .
\]  

(57)

This function is displayed, along with its square, which represents the observed depolarization factor, in Fig. 15b.

---

\(^{3}\) An error in a previous calculation of \(<d^a>\) (designated \(<d'^a>\); there) for this model in the range \((\pi/2 < \Psi < \pi)\) (Dale and Eisinger, 1975) is apparent and is corrected here.
and the equivalent expression in $\psi'$

$$<d'> = \{$$

and

$$<d'> = \left\{ \begin{array}{l}
-\frac{1}{24} \\
-\frac{1}{3} \\
\end{array} \right.$$ 

along with the observed depolarization factors,

\( g_e \)

and the equivalent for $\psi'$ in $(0, \pi/2)$ the axial depolarization factors,

and

$$<d'> =$$

for centrally and peripherally weighted presented in Figs. 18 a and b. Since the peripherally ($\psi'$) than in the exponent "characteristic" angles $\psi$ and $\psi'$ are $17 a$ and $b$.

Of course, an infinity of distribution limited number correspond to physical energy transfer experiments, it is not only to know that they have at least a

It should finally be noted that the beyond its use in connection with the resonance order parameter for rapid the distribution models discussed above.
and the equivalent expression in $\psi'$, $\psi$, and min ($\psi$, $\pi/2$). The resulting axial depolarization factors

$$
<d^*> = \begin{cases} 
\frac{1}{24} \left( \frac{3 \sin \psi - \sin (3\psi)}{\psi - \sin \psi} \right) & (0 \leq \psi \leq \pi) \\
0 & (\psi \geq \pi)
\end{cases}
$$

(59)

and

$$
<d^*'> = \begin{cases} 
-\frac{1}{24} \left( \frac{4 - 3 \cos \psi' - \cos (3\psi')}{{1 - \cos \psi'}} \right) & (0 \leq \psi' \leq \pi/2) \\
-\frac{1}{3} \left( \frac{2\psi' - \pi + 2}{{2\psi' - \pi + 2}} \right) & (\psi' \geq \pi/2)
\end{cases}
$$

(60)

along with the observed depolarization factors $<d'>$ derived from them are given in Figs. 16a and b.

**Model III. Exponential Distribution** This model corresponds to an analogue of the two-dimensional triangular well potential. The radial density function is

$$
g_e(\psi) = e^{-(\psi/\Psi)} \quad (0 \leq \psi \leq \pi),
$$

(61)

and the equivalent for $\psi'$ in $(0, \pi/2)$. In this case, again, integration gives rise to quite simple forms for the axial depolarization factors,

$$
<d^*> = 1/(1 + 9\Psi^2)
$$

(62)

and

$$
<d^*> = -\frac{1 + \psi'(3\psi' - 2e^{-\psi'/\Psi})}{2(1 + 9\Psi^2)(1 + \psi' e^{-\psi'/\Psi})},
$$

(63)

for centrally and peripherally weighted models, respectively. These and the corresponding $<d'>$'s are presented in Figs. 17a and b, in which the ranges of $\psi$ and $\psi'$ are arbitrarily restricted to $\pi$ radians for comparison with the previous distribution models.

**Model IV. Gaussian Distribution** A Gaussian radial density function may constitute the most realistic of the models considered here and corresponds to an analogue of the two-dimensional parabolic well potential. The radial density function is

$$
g_G(\psi) = e^{-(\psi/\Psi)^2} \quad (0 \leq \psi \leq \pi),
$$

(64)

along with the equivalent expression for $\psi'$ in $(0, \pi/2)$. The corresponding depolarization factors are visualized in Figs. 18a and b. Since this distribution function is weighted more heavily centrally ($\psi$) or peripherally ($\psi'$) than in the exponential model, the absolute depolarization factors as functions of the "characteristic" angles $\psi$ and $\psi'$ are much smaller than those for the exponential model shown in Figs. 17a and b.

Of course, an infinity of distributions could be invoked in this context, although presumably only a limited number correspond to physical reality. It is important to emphasize that, for the analysis of energy transfer experiments, it is not necessary to know the precise form of the actual distributions but only to know that they have at least approximate axial symmetry.

It should finally be noted that the axial depolarization factor $<d^*>$ defined here has significance beyond its use in connection with fluorescence depolarization. Thus, it appears as the electron spin resonance order parameter for rapid anisotropic tumbling of spin labels, described in terms of some of the distribution models discussed above, e.g. the random walk model (Van et al., 1974; Israelachvili et
al., 1975) and the Gaussian distribution (Gaffney and McConnell, 1974). A model not considered here, that corresponding to a Gaussian spread about a mean bond rotation angle $\Psi$,

$$
g(\Psi) = e^{-[(\Psi - \Psi_0)/\sigma]^2} \quad (0 \leq \Psi \leq \pi),
$$

(compare Eqs. 52 and 64), has also been utilized by the last-named authors.

**Figure 16**

- **a.** Dynamically averaged axial and observed depolarization factors, $\langle d' \rangle$ and $\langle d \rangle$, for a luminophore whose linear transition moment has a triangular radial density function $g_\gamma (d')$ in Fig. 14 of characteristic angle $\Psi$. $b.$ Same as $a$ except that the transition moment is considered to be planar, and $\Psi'$ is the angle between it and the plane normal to the symmetry axis of the orientational distribution, i.e. $\Psi' = (\pi/2) - \Psi$.

**Figure 17**

- **a.** Dynamically averaged luminophore whose linear transition of characteristic angle $\Psi$. $b.$ Same as $a$ is the angle between it and the plane $\Psi' = (\pi/2) - \Psi$.

**ADDENDUM**

Since completion of the above intramolecular separations using data appeared in the literature (Shepherd, 1978; Pober et al., 1978; Tu et al., 1 another of the reorientational model complete energy transfer study of including an extensive set of polariz appeared (Haas et al., 1978).
Figure 17  a. Dynamically averaged axial and observed depolarization factors, $<d^a>$ and $<d>$, for a luminophore whose linear transition moment has an exponential radial density function ($g_r(\psi)$ in Fig. 14) of characteristic angle $\psi$. b. Same as a except that the transition moment is considered to be planar, and $\psi'$ is the angle between it and the plane normal to the symmetry axis of the orientational distribution, i.e. $\psi' = (\pi/2) - \psi$.

ADDENDUM

Since completion of the above in final manuscript form, a number of contributions reporting intramolecular separations using depolarization measurements to delimit upper and lower bounds have appeared in the literature (Shepherd and Hammes, 1977; Holowka and Hammes, 1977; Dockter et al., 1978; Pober et al., 1978; Tu et al., 1978; Hahn and Hammes, 1978). These have all made use of one or another of the reorientational models previously reported (Dale and Eisinger, 1974, 1975). A more complete energy transfer study of the distribution of end-to-end distances in a series of oligopeptides, including an extensive set of polarization measurements over a wide range of solvent viscosities, has also appeared (Haas et al., 1978a). Most recently, Haas et al. (1978b) have detailed a treatment of the
REFERENCE


FIGURE 18 a. Dynamically averaged axial and observed depolarization factors, \( <d'> \) and \( <d> \), for a luminophore whose linear transition moment has a Gaussian radial density function \( <g_r (\psi) \) in Fig. 4) of characteristic angle \( \psi \). b. Same as a except that the transition moment is considered to be planar, and \( \psi \) is the angle between it and the plane normal to the symmetry axis of the orientational distribution, i.e. \( \psi = (\pi/2) - \psi. \)

orientation factor problem formally equivalent to that presented here, but using directly the concept of three-dimensional transition dipole moments and formulating their results in vector form. Finally, in both that article and a recent energy transfer review (Stryer, 1978) as well as in work previously quoted (Hillel and Wu, 1976), a "statistical" argument is invoked in attempts to predict a "probable" narrower range of orientation factor values than given by the absolute upper and lower bounds obtainable directly. The applicability of this kind of approach to real situations is questionable and has been discussed here in Appendix B.
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