The Role of Polarizabilities in the Fluorescence Behaviour of 4-cyano-N,N-dimethylaniline

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The effect of an external electric field on the absorption and the double fluorescence of 4-cyano-N,N-dimethylaniline can be understood, taking into account reaction field induced polarizability effects. If a TICT state conformation emits the a-fluorescence in dioxane, the permanent dipole moment in this state is only slightly larger than in the equilibrium ground state.

I. Introduction

Lippert et al. [1] have first reported that with 4-cyano-N,N-dimethylaniline (dimethylaminobenzo-nitrile≡DMABN) in polar solvents there are two fluorescence bands, called a- and b-fluorescence, but only the b-fluorescence in nonpolar solvents. He ascribed this effect to a solvent induced reversal of the two lowest excited singlet states having different dipole moments.

Later on other models have been proposed and discussed extensively in the literature [2—7]. Besides the level crossing model it seems now that there is only one other model remaining: The TICT model introduced by Krystyna Rotkiewicz, K. H. Grellmann, Z. R. Grabowski [7] and discussed in a series of papers [7—12], where this idea is shown to be applicable to a lot of molecules consisting of a donor and an acceptor subassembly. Also, results of theoretical investigations on such molecules have been shown to be consistent with the TICT model [11, 13—17]. The TICT model ascribes the two fluorescence bands to two rotational isomers with coplanar or twisted donor and acceptor subunits of these molecules. In some recent papers Rettig, Wermuth and Lippert [18—20] have shown that a combination of the level crossing model and the TICT model might account for all observed effects in this class of molecules.

Starting from the idea that the TICT state is expected to have a very much larger dipole moment than the primarily excited Franck-Condon (FC) state, the electrooptical investigations described in this paper have been performed. The comparison of

the dipole moment of the excited FC state and the equilibrium excited state (equilibrated with respect to intramolecular as well as to intermolecular relaxations after excitation) then should give valuable insight into the nature of the excited state emitting the longwavelength a-fluorescence in polar solvents. On the other hand, from similar investigations on 4-(9-anthryl)-N,N-dimethylaniline and related molecules came the idea [21] that the polarizability of these molecules could stand for at least part of the observed effects. Also, Wermuth, Rettig and Lippert in their recent paper [19] found that the solvatochromism of p-N,N-diethylaminobenzoic acid ethylester could be better described taking into account polarizabilities. So reaction field induced polarizability effects on the experimentally determined dipole moments of DMABN in solution will be considered in this paper, too.

II. Theoretical and Experimental

Experimental methods used to determine dipole moments in solutions in Franck-Condon and in equilibrated excited states are electrooptical absorption measurements (EOAM) and electrooptical emission measurements (EOEM). EOAM has been developed in a lot of papers by Labhart [22—24] and Liptay [25—30], whose most elaborated theoretical model has been taken for EOEM [31], too. In this way results from both methods are comparable on the basis of the same model. This model, founded on the work of Onsager [32] and Scholte [33] approximates the solute molecule as a polarizable point dipole situated in the center of a cavity, the shape and size of which may be approximated by the shape and size of the solute. In this paper the most simple spherical approximation is used,

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where the cavity is approximated by a sphere of radius a . The theory developed in Ref. [4] is only valid if the life time of the molecules investigated by EOEM is much longer than all reorientation relaxation times in solution. This condition is fulfilled with DMABN, as is shown below.

With EOEM [31] the relative change of the fluorescence photon current density is measured, when an external electric field E_a is applied. It is measured as a function of the fluorescence wavenumber $\tilde{\nu}_e$ and the angle between the external electric field and the polarization direction, given by an analyzing polarizer. Then the following quantities can be derived [31] from EOEM, if smaller contributions are neglected:

$$E^e = \beta^2 [3(\mathbf{m}_e \vec{\mu}_a)^2 - \mu_a^2], \quad (1)$$

$$F^e = \beta \vec{\mu}_a \Delta^e \vec{\mu}, \quad (2)$$

$$G^e = \beta (\mathbf{m}_e \vec{\mu}_a) (\mathbf{m}_e \Delta^e \vec{\mu}), \quad (3)$$

$$H^e = (\Delta^e \mu)^2, \quad (4)$$

$$I^e = (\mathbf{m}_e \Delta^e \vec{\mu})^2, \quad (5)$$

with $\beta = (kT)^{-1}$.

Quite similar with EOAM [27, 30] the relative change of the absorbance is measured, when an external electric field is applied. It is measured as a function of the absorption wavenumber $\tilde{\nu}_a$ and the angle between the external electric field and the electric field vector of the linearly polarized light, defined by a polarizer. From these measurements the following quantities can be derived if smaller contributions are neglected:

$$E^a = \beta^2 [3(\mathbf{m}_a \vec{\mu}_g)^2 - \mu_g^2], \quad (6)$$

$$F^a = \beta \vec{\mu}_g \Delta^a \vec{\mu}, \quad (7)$$

$$G^a = \beta (\mathbf{m}_a \vec{\mu}_g) (\mathbf{m}_a \Delta^a \vec{\mu}), \quad (8)$$

$$H^a = (\Delta^a \mu)^2, \quad (9)$$

$$I^a = (\mathbf{m}_a \Delta^a \vec{\mu})^2. \quad (10)$$

$\mathbf{m}_e$ and $\mathbf{m}_a$ are unit vectors in the direction of the transition moment involved with the emission and absorption process, respectively.

$$\vec{\mu}_g = f_e (1 - f \alpha_g^0)^{-1} \vec{\mu}_g^0, \quad (11)$$

$$\vec{\mu}_a = f_e (1 - f \alpha_a^0)^{-1} \vec{\mu}_a^0, \quad (12)$$

* Anm. d. Red.: Aus satztechnischen Gründen mußte der Vektor $\vec{\mu}$ mit Pfeil statt fett, kursiv gesetzt werden.

$$\Delta^a \vec{\mu} = (1 - f' \alpha_a^{0FC})^{-1} (1 - f \alpha_a^{0FC}) (1 - f \alpha_g^0)^{-1} \cdot (1 - f' \alpha_g^0) (\vec{\mu}_a^{FC} - \vec{\mu}_g), \quad (13)$$

$$\Delta^e \vec{\mu} = (1 - f' \alpha_g^{0FC})^{-1} (1 - f \alpha_g^{0FC}) (1 - f \alpha_a^0)^{-1} \cdot (1 - f' \alpha_a^0) (\vec{\mu}_a - \vec{\mu}_g^{FC}), \quad (14)$$

where $\vec{\mu}_g^{FC}$ and $\vec{\mu}_a^{FC}$ are defined analogous to (11) and (12):

$$\vec{\mu}_g^{FC} = f_e (1 - f \alpha_g^{0FC})^{-1} \vec{\mu}_g^{0FC}, \quad (15)$$

$$\vec{\mu}_a^{FC} = f_e (1 - f \alpha_a^{0FC})^{-1} \vec{\mu}_a^{0FC}. \quad (16)$$

In this paper the spherical approximation for the Onsager cavity is used, so

$$f_e = \frac{3\varepsilon}{2\varepsilon + 1} \cdot 1 = f \cdot 1, \quad (17)$$

$$f = \frac{1}{4\pi\varepsilon_0} \frac{2}{a^3} \frac{\varepsilon - 1}{2\varepsilon + 1} \cdot 1 = f \cdot 1, \quad (18)$$

$$f' = \frac{1}{4\pi\varepsilon_0} \frac{2}{a^3} \frac{n^2 - 1}{2n^2 + 1} \cdot 1 = f' \cdot 1 \quad (19)$$

with a being the radius of the sphere, ε_0 the permittivity of the vacuum, ε the dielectric constant and n the refractive number.

In nonpolar solvents f equals f' and therefore (13) and (14) become very simple:

$$\Delta^a \vec{\mu} = \vec{\mu}_a^{FC} - \vec{\mu}_g, \quad (20)$$

$$\Delta^e \vec{\mu} = \vec{\mu}_a - \vec{\mu}_g^{FC}. \quad (21)$$

$\vec{\mu}^0$ and α^0 denote dipole moments and polarizability tensors of the free molecule.

$\vec{\mu}_g^0$, α_g^0 and $\vec{\mu}_a^0$, α_a^0 are dipole moments and polarizabilities of the equilibrium ground state and of the equilibrated excited state of the molecule under consideration, $\vec{\mu}_a^{0FC}$ and α_a^{0FC} are dipole moment and polarizability of the Franck-Condon (FC-) excited state, reached by the absorption process and $\vec{\mu}_g^{0FC}$ and α_g^{0FC} those of the FC ground state reached by the emission process.

The experimental setup used in both methods is described in [34] and [35].

III. Experimental Results

III.1. Results of Electrooptical Emission Measurements on the α -fluorescence of DMABN

EOEM have been performed on DMABN in dioxane at 298 K. The solutions could not be deaerated in the measuring cell used. Dioxane has been

Table 1. Results of EOEM on solutions of DMABN in dioxane. $\Delta\bar{\nu}_e$ shows the spectral range used for the determination of E^e to G^e , $\delta\bar{\nu}_e$ is the spectral bandwidth of the fluorometer used for EOEM. Excitation bandpass has been $(34-37) \cdot 10^5 \text{ m}^{-1}$.

a-fluorescence	
$E^e/10^{-20} \text{ m}^2 \text{ V}^{-2}$	51500 ± 300
$F^e/10^{-40} \text{ CV}^{-1} \text{ m}^2$	10100 ± 400
$G^e/10^{-40} \text{ CV}^{-1} \text{ m}^2$	10100 ± 400
$\Delta\bar{\nu}_e/10^5 \text{ m}^{-1}$	$20.4 - 24.4$
$\delta\bar{\nu}_e/10^5 \text{ m}^{-1}$	0.6

used as a pseudopolar solvent that withstands very well breakdown at high electric fields.

In order to assure that all orientational relaxations have come to an end before emission takes place, the fluorescence decay time τ_f has been measured for DMABN in those nondeairated dioxane solutions used for EOEM. Excitation at $(34-37) \cdot 10^5 \text{ m}^{-1}$ and monitoring the a-fluorescence at wavenumbers smaller than $24.0 \cdot 10^5 \text{ m}^{-1}$, resulted in $\tau_f = (3.2 \pm 0.1) \text{ ns}$.

This decay time is much longer than the solvent and solute reorientational relaxation times in dioxane. So (1) to (5) apply for the evaluation of the molecular properties from EOEM.

The results of EOEM on DMABN are shown in Table 1. Errors shown are single standard deviations as derived from a regression analysis.

III.2. Determination of Dipole Moments from the a-Fluorescence of DMABN in Dioxane

With the values from Table 1 using (1)–(3) the dipole terms in Table 2 are calculated.

Table 2 shows that from the a-fluorescence one finds

$$\vec{\mu}_a \Delta^e \vec{\mu} = (\mathbf{m}_e \vec{\mu}_a) (\mathbf{m}_e \Delta^e \vec{\mu}). \quad (22)$$

Table 2. Dipole moments of DMABN as determined by EOEM from the a-fluorescence. The value of μ_a^{FC} is put into brackets, because in pseudopolar dioxane Eq. (21) is not valid exactly.

	a-fluorescence
$[3(\mathbf{m}_e \vec{\mu}_a)^2 - \mu_a^2]/10^{-60} \text{ C}^2 \text{ m}^2$	8718 ± 50
$\vec{\mu}_a \Delta^e \vec{\mu}/10^{-60} \text{ C}^2 \text{ m}^2$	4160 ± 160
$(\mathbf{m}_e \vec{\mu}_a) (\mathbf{m}_e \Delta^e \vec{\mu})/10^{-60} \text{ C}^2 \text{ m}^2$	4160 ± 160
$\mu_a/10^{-30} \text{ Cm}$	66 ± 0.2
$\Delta^e \mu/10^{-30} \text{ Cm}$	63 ± 2.5
$\mu_a^{\text{FC}}/10^{-30} \text{ Cm}$	(3 ± 2.7)

This is a very good hint that there is $\mathbf{m}_e \parallel \vec{\mu}_a \parallel \Delta^e \vec{\mu}$. Then the dipole moments given in Table 2, too, can be evaluated.

III.3. Results of Electrooptical Absorption Measurements

To all knowledge the first absorption band of DMABN starting around $31 \cdot 10^5 \text{ m}^{-1}$ is a superposition of a strong transition to a singlet excited electronic state $^1A^*$ and a transition to a $^1B^*$ state that is very close to the $^1A^*$ state, if C_2 symmetry is assumed. EOAM clearly shows this superposition of transitions at the longwavelength tail of the absorption in n-heptane and in dioxane. Thus EOEM in this spectral range cannot be utilized. On the other hand, from measurements over the main part of the first broad absorption band, which corresponds to the nearly unperturbed $A^* \leftarrow A$ transition, the terms E^a to I^a are determined without complications. The results are listed up in Table 3. All errors are single standard deviations from a regression analysis.

F^a is found to be a little bit larger than G^a . This might be due to experimental errors, as only single standard deviations are given, or it might be due to the neglect of minor polarizability terms, as discussed in Chapter VI.

III.4. Determination of Dipole Moments from EOAM

Taking the appropriate values of Table 3, as is indicated in the last row of Table 4, and using (6) to (10), the values of Table 4 are found for some dipole moments of DMABN.

Table 4 shows that the moment μ_g as determined by EOAM in dioxane is found to be in good agreement with the values reported in the literature [36–39]. The value used here for μ_g was taken

Table 3. Results of EOAM on the $A^* \leftarrow A$ transition of DMABN in dioxane and heptane solutions at 298 K. $\delta\bar{\nu}_a$ is the spectral bandwidth used with EOAM.

Solvent	Dioxane	Heptane
$E^a/10^{-20} \text{ V}^{-2} \text{ m}^2$	10300 ± 100	8420 ± 50
$F^a/10^{-40} \text{ CV}^{-1} \text{ m}^2$	2100 ± 100	1240 ± 30
$G^a/10^{-40} \text{ CV}^{-1} \text{ m}^2$	1700 ± 50	1140 ± 30
$H^a/10^{-60} \text{ C}^2 \text{ m}^2$		350 ± 120
$I^a/10^{-60} \text{ C}^2 \text{ m}^2$		450 ± 120
$\delta\bar{\nu}_a/10^5 \text{ m}^{-1}$	0.05	0.04

Table 4. Some dipole moments of DMABN.

Solvent	Dioxane	Heptane	taken from:
$\mu_g/10^{-30}$ Cm	29.5 ± 0.5	26.7 ± 0.1	E^a
$\Delta^a\mu/10^{-30}$ Cm	29.3 ± 1.9	19.1 ± 0.6	E^a, F^a
$\Delta^a\mu/10^{-30}$ Cm		20 ± 3	H^a, I^a
$\mu_a^{FC}/10^{-30}$ Cm	(58.8 ± 1.4)	45.8 ± 0.5	E^a, F^a
$\mu_g/10^{-30}$ Cm	30 ± 4		diel. *

* μ_g has been determined by dielectric investigations.

from [37] where the same model has been used for the description of the effective field in a solution.

Note that μ_a^{FC} of the A^* state is found to be about 25% larger in dioxane than in heptane. μ_a from a-fluorescence in dioxane is still larger than μ_a^{FC} in this state.

IV. Discussion of the Results from Electrooptical Studies on DMABN

IV.1. Discussion assuming $f\alpha \ll 1$

With the assumption that all terms $f\alpha$ are negligible compared to 1, (20) and (21) are valid. Then with $f_e = 1.2$ the dipole moments of the free molecule given in Table 5 are calculated from the dipole moments in Tables 2 and 4.

Under these assumptions the difference found for μ_a^{FC} in heptane and dioxane for the A^* state can only be understood if different ground state conformations exist in dioxane and heptane. If the TICT model is taken into account [7, 12], a conformer more twisted in the ground state in dioxane than in heptane would be in agreement with the experimental results.

The even more polar equilibrium state with $\mu_a = 55 \cdot 10^{-30}$ Cm also fits well to this model, as after excitation a further twist of the dimethylamino

Table 5. Dipole moments of free DMABN determined with the assumption that reaction field induced moments are negligible.

	From a-fluorescence	from $A^* \leftarrow A$ absorption	
Solvent	Dioxane	Dioxane	Heptane
$\mu_a^0/10^{-30}$ Cm	55 ± 0.2		
$\mu_g^0/10^{-30}$ Cm	2.5 ± 2.3		
$\mu_a^{FC}/10^{-30}$ Cm		49.0 ± 1.2	39.2 ± 0.2
$\mu_g^0/10^{-30}$ Cm		24.6 ± 0.4	22.3 ± 0.1

group according to the TICT model might occur. But the very small FC-ground state dipole moment cannot be understood even in the TICT theory. This is because in that model μ_g^{FC} should have a value similar to that of benzonitrile ($\mu_g^0 \approx 12$ to $15 \cdot 10^{-30}$ Cm). So the whole set of dipole moments given in Table 5 cannot be understood consistently if $f\alpha$ is assumed to be negligible compared to 1. On the other hand all the results in Table 5 can be understood, if reaction field induced moments are taken into account, which is shown in the next section.

IV.2. Discussion Assuming that the Permanent Dipole Moment of DMABN in its A^* FC-state and in its Ground State is not Solvent Dependent

Here and in the following the left superscripts D and H label dioxane and heptane as solvents, respectively.

It is assumed that the permanent dipole moment and the polarizability of the ground state and the excited FC state are independent from the solvent, that means

$$\begin{aligned} {}^H\mu_g^0 &= {}^D\mu_g^0, & {}^H\alpha_g^0 &= {}^D\alpha_g^0, \\ {}^H\mu_a^{FC} &= {}^D\mu_a^{FC}, & {}^H\alpha_a^{FC} &= {}^D\alpha_a^{FC}. \end{aligned}$$

Because of the symmetry axis (z -axis) all $(1-f\alpha)$ like terms must be considered as $(1-f\alpha_{zz}^0)$ terms, furtheron.

From (13) a term S_1 may be defined:

$$S_1 = \frac{{}^D\Delta^a\mu}{{}^D\mu_a^{FC} - {}^D\mu_g} = \frac{1 - {}^Df\alpha_{azz}^{FC}}{1 - {}^Df'\alpha_{azz}^{FC}} \frac{1 - {}^Df'\alpha_{gzz}^0}{1 - {}^Df\alpha_{gzz}^0}. \quad (23)$$

Using ${}^Df' = {}^Hf$, (23) can be rewritten to give

$$S_1 = \frac{{}^H\mu_a^{FC}}{{}^H\mu_g} - \frac{{}^D\Delta^a\mu}{{}^D\mu_g}. \quad (24)$$

Taking the values of these dipole moments from Table 4, S_1 can be calculated:

$$S_1 = 0.725 \pm 0.012.$$

From ${}^D\mu_g$ and ${}^H\mu_g$ a term S_2 can be defined:

$$S_2 = \frac{{}^D\mu_g}{{}^H\mu_g} = \frac{1 - {}^Hf\alpha_{gzz}^0}{1 - {}^Df\alpha_{gzz}^0}. \quad (25)$$

With the values from Table 4, S_2 is found to be

$$S_2 = 1.105 \pm 0.025.$$

Combining (23) and (25), a term S_3 may be defined:

$$S_3 = \frac{1 - {}^H f \alpha_{azz}^{0FC}}{1 - {}^D f \alpha_{azz}^{0FC}} \quad (26)$$

With the values for S_1 and S_2 determined above, S_3 is found to be $S_3 = 1.52 \pm 0.06$.

Note that S_1 , S_2 and S_3 have been calculated with dipole moments determined only by one method, namely EOAM. So calibration errors etc. do not influence these results.

With these values for S_2 and S_3 at first $\alpha_{gzz}^0/2\pi\epsilon_0 a^3$ and $\alpha_{azz}^{0FC}/2\pi\epsilon_0 a^3$ and then terms $(1 - f\alpha_{zz}^0)$ and the dipole moments of the free molecule can be calculated, using ${}^D\epsilon = 6$ and ${}^H\epsilon = 2$ for the description of the reaction field. ${}^D\epsilon = 6$ as an effective dielectric constant of dioxane is used in the following, because in all solvatochromism measurements dioxane behaves as a solvent with ${}^D\epsilon \approx 6$. The results are shown in Table 6 and 7 together with the ratio $\alpha_{azz}^{0FC} : \alpha_{gzz}^0$.

With the values of Tables 4 and 6 and with $f_e = 1.2$, the dipole moments in Table 7 result, using (11), (13) and (16).

The results in Table 7 show:

μ_g^0 determined from EOAM in dioxane and heptane and from dielectric investigations agree very well. The excited A^* FC-state has a slightly larger dipole moment than the ground state. Clearly from the assumptions, μ_a^{0FC} in dioxane and heptane and

$\alpha_{gzz}^0/2\pi\epsilon_0 a^3$	0.465 ± 0.1	Table 6. $\alpha_{gzz}^0/2\pi\epsilon_0 a^3$, $\alpha_{azz}^{0FC}/2\pi\epsilon_0 a^3$, $\alpha_{azz}^{0FC} : \alpha_{gzz}^0$ and $(1 - f\alpha_{zz}^0)$ type terms, using ${}^D\epsilon = 6$.
$\alpha_{azz}^{0FC}/2\pi\epsilon_0 a^3$	1.35 ± 0.08	
$\alpha_{azz}^{0FC} : \alpha_{gzz}^0$	2.9 ± 0.7	
$1 - {}^H f \alpha_{azz}^{0FC}$	0.730 ± 0.015	
$1 - {}^D f \alpha_{azz}^{0FC}$	0.480 ± 0.03	
$1 - {}^H f \alpha_{gzz}^0$	0.907 ± 0.02	
$1 - {}^D f \alpha_{gzz}^0$	0.821 ± 0.034	

Table 7. Dipole moments of the free molecule DMABN, as determined from EOAM.

		Taken from:
$\mu_g^0/10^{-30}$ Cm	20.5 ± 0.9	dielectric investigations
$\mu_g^0/10^{-30}$ Cm	20.2 ± 0.6	μ_g , dioxane
$\mu_g^0/10^{-30}$ Cm	20.2 ± 1.3	μ_g , heptane
$\mu_a^{0FC}/10^{-30}$ Cm	27.9 ± 6	μ_g , $\Delta^a\mu$, heptane
$\mu_a^{0FC}/10^{-30}$ Cm	27.9 ± 1	μ_g , $\Delta^a\mu$, dioxane

μ_g^0 in these solvents must be found identical within their error margins.

IV.3. Discussion on the Permanent Dipole Moment of the Excited A^* -State, Emitting a-Fluorescence

Table 5 shows that with the assumptions $f\alpha \ll 1$ an FC ground state dipole moment μ_g^{0FC} is determined from the experimental values for μ_a and $\Delta^e\mu$ that seems to be unacceptably small, as discussed in Chapter IV.1.

On the other hand in Chapter IV.2 it has been shown that the solvent dependence of μ_g and $\Delta^a\mu$ can be fully understood, if $(1 - f\alpha)$ type terms are taken into account. So one could ascribe the discrepancy between μ_a and $\Delta^e\mu$ at least to a great deal to the neglect of these terms, too.

Two limiting cases may be considered:

case a) μ_g^{0FC} and α_g^{0FC} of the species emitting a-fluorescence in dioxane are the same as μ_g^0 and α_g^0 of the molecule in its equilibrium ground state. Then it follows

$${}^D\mu_g^{FC} = {}^D\mu_g = (29.5 \pm 0.5) \cdot 10^{-30} \text{ Cm};$$

case b) μ_g^{0FC} and α_g^{0FC} of the emitting species in dioxane are the same as those of the TICT ground state, the dipole moment of which might be very roughly taken as that of benzonitrile, which gives

$${}^D\mu_g^{FC} = {}^D\mu_{g,TICT} = (20 \pm 3) \cdot 10^{-30} \text{ Cm} [40].$$

These two limiting cases are considered parallelly, furtheron.

From (14) a term S_1' can be defined:

$$S_1' = \frac{{}^D\Delta^e\mu}{{}^D\mu_a - {}^D\mu_g^{FC}} = \frac{1 - {}^D f' \alpha_{azz}^0}{1 - {}^D f \alpha_{azz}^0} \cdot \frac{1 - {}^D f \alpha_{gzz}^{0FC}}{1 - {}^D f' \alpha_{gzz}^{0FC}} \quad (27)$$

With the values of Table 2 and those discussed above, S_1' can be calculated:

$$S_1' = 1.726 \pm 0.1 \quad \text{case a)}, \\ 1.370 \pm 0.16 \quad \text{case b)}.$$

The term $(1 - {}^D f' \alpha_{gzz}^{0FC})/(1 - {}^D f \alpha_{gzz}^{0FC})$ can be taken equal to S_2 of (25) in both cases in good approximation, because even if there are different values for $f\alpha_{gzz}^{0FC}$ in postulated FC ground states with TICT conformation, or with planar FC ground state conformation, or with equilibrium ground state conformation, the influence of such differences on S_2

Table 8. The dipole moment μ_a^0 of the A* state emitting a-fluorescence.

	Case a)	Case b)
$\alpha_{azz}^0/2\pi\epsilon_0 a^3$	1.70 ± 0.1	1.34 ± 0.2
$1 - Df\alpha_{azz}^0$	0.347 ± 0.036	0.483 ± 0.08
$\mu_a^0/10^{-30}$ Cm	19.1 ± 1.9	26.6 ± 4.4

is small, as S_2 itself is nearby 1. So the error introduced by this uncertainty in $f\alpha_{gzz}^{0FC}$ clearly is small.

With S_2 from Chapter IV.2 and S_1' taken from above, S_3' may be calculated:

$$S_3' = \frac{1 - Df'\alpha_{azz}^0}{1 - Df\alpha_{azz}^0} = \begin{matrix} 1.907 \pm 0.016 & \text{case a),} \\ 1.514 \pm 0.02 & \text{case b).} \end{matrix}$$

This value for S_3' may be compared to the value found for S_3 Section IV.2. Although the errors of S_3' are pretty high, S_3' resembles S_3 very well, especially in case b).

With S_3' the values of $\alpha_{azz}^0/2\pi\epsilon_0 a^3$ and then $(1 - Df\alpha_{azz}^0)$ and μ_a^0 can be calculated. Using $f_e = 1.2$ one gets the results shown in Table 8.

The dipole moment μ_a found for either case a) or b) is in between μ_g and μ_a^{0FC} , as can be seen by comparison with the results in Table 7, and α_{azz}^0 is nearly equal to α_a^{0FC} within the errors, as may be seen by comparison with the results in Table 6.

So from these results it may be deduced that if there is a TICT formation after excitation in dioxane to give a-fluorescence, this TICT state has a dipole moment not much larger than the ground state and very similar to that of the A* FC excited state, and a polarizability also very similar to that of the A* FC excited state.

V. The Onsager Cavity Radius a

V.1. The Solvent Dependent Spectral Shift of the a-Fluorescence and the Determination of the Onsager Cavity Radius a

Lippert [1] has shown the solvatochromism of the a-fluorescence of DMABN. These results seem to oppose the results reported here. But if the terms $(1 - f\alpha)$ differ markedly from 1, Lippert's [41] or Mataga's [42] formulae describing solvent shifts are no more valid. Instead, the more elaborated formulae of Liptay [43] must be used. In good approxi-

mation this formula is written as

$$hc\tilde{\nu}_e = \text{const} - \frac{1}{2\pi\epsilon_0} \cdot \frac{1}{a^3} F(g) \cdot \{ \bar{\mu}_a^0(\bar{\mu}_a^0 - \bar{\mu}_g^{0FC}) + (1 - f'\alpha_g^{0FC})^{-1} \cdot (1 - f\alpha_a^0)^{-1} \bar{\mu}_a^0 \cdot [\frac{1}{2}f(\alpha_a^0 - \alpha_g^{0FC}) \cdot (1 - f'\alpha_a^0)\bar{\mu}_a^0 - f'(\alpha_a^0 - \alpha_g^{0FC}) \cdot (1 - f\alpha_a^0)(\bar{\mu}_a^0 - \bar{\mu}_g^{0FC})] \}, \quad (28)$$

where

$$F(g) = \left(g^{-1} - \frac{1}{2\pi\epsilon_0} \frac{\alpha_{azz}^0}{a^3} \right)^{-1} \quad (29)$$

with

$$g = (\epsilon - 1)/(2\epsilon + 1). \quad (30)$$

In order to get a simple but sufficiently accurate plot of $\tilde{\nu}_e$ against $F(g)$, the total term in braces is calculated, using the results from Tables 5 and 6 and estimating the solvent dependent terms

$$(1 - f\alpha_{azz}^0) \quad \text{and} \quad (1 - f\alpha_{gzz}^{0FC})$$

in the minor part of the sum in braces to be 0.55 ± 0.2 and 0.85 ± 0.06 , respectively.

Then using Lippert's data [1], from a plot in Fig. 1 according to (28), a^3 is found to be $(49.5 \pm 20) \cdot 10^{-30} \text{ m}^3$, and from this $a = (3.7 \pm 0.5) \cdot 10^{-10} \text{ m}$.

This is a relatively small value, but it seems to be plausible as a mean value over the three molecular axis.

So the results from solvent shift measurements supply consistently the results from electrooptical investigations.

Fig. 1. Plot of $\tilde{\nu}_e$ against $F(g)$.

V.2. Determination of the Onsager Radius by use of α_g^0

From refraction measurements [37] $\text{tr } \alpha_g^0$ is known

$$\text{tr } \alpha_g^0 = (64.5 \pm 3) \cdot 10^{-40} \text{ CV}^{-1} \text{ m}^2.$$

From group polarizabilities together with this value, α_{gzz}^0 may be estimated to be

$$\alpha_{gzz}^0 = (27 \pm 4) \cdot 10^{-40} \text{ CV}^{-1} \text{ m}^2.$$

With this value a^3 can be estimated using $\alpha_{gzz}^0 / 2\pi\epsilon_0 a^3$ from Table 6:

$$a^3 = (104 \pm 40) \cdot 10^{-30} \text{ m}^3$$

or

$$a = (4.7 \pm 0.7) \cdot 10^{-10} \text{ m}.$$

This value is much larger than that determined from solvent shift measurements, but overlaps within the errors.

Although the value for a^3 determined from solvent shift measurements fits better to the molecular dimensions that might be estimated as

$$\begin{aligned} a^3 &= (5 \times 3.5 \times 2.5) \cdot 10^{-30} \text{ m}^3 \\ &= 44 \cdot 10^{-30} \text{ m}^3, \end{aligned}$$

this is not an argument to favour that value, if a^3 is considered to be a parameter which comprises a lot of effects.

Thus these results also must be seen as consistent with those from electrooptical investigations.

VI. Critical Check on the Neglectability of Explicit Polarizabilities

In Chapter II explicit polarizability terms [31] have been neglected as minor terms. Now, with the data calculated on the basis of this simplification, the explicit polarizability terms $\frac{1}{2} \text{tr } \Delta^a \alpha$ and $\frac{1}{2} \text{tr } \Delta^e \alpha$ may be calculated and can be compared to the experimental values of F^a and F^e according to Eqs. (100) and (110) from Ref. [31]. With the definition of $\Delta^a \alpha$ and $\Delta^e \alpha$ from Eqs. (37) and (40) of Ref. [31] and with the values from Tables 6 and 8 and from Section IV.4 $\frac{1}{2} \text{tr } \Delta^a \alpha$ and $\frac{1}{2} \text{tr } \Delta^e \alpha$ are found to be

$$\begin{aligned} \frac{1}{2} \text{tr } \Delta^a \alpha &= (133 \pm 55) \cdot 10^{-40} \text{ CV}^{-1} \text{ m}^2 \\ &\quad \text{in heptane} \\ &= (161 \pm 100) \cdot 10^{-40} \text{ CV}^{-1} \text{ m}^2 \\ &\quad \text{in dioxane,} \\ \frac{1}{2} \text{tr } \Delta^e \alpha &= (468 \pm 300) \cdot 10^{-40} \text{ CV}^{-1} \text{ m}^2 \\ &\quad \text{in dioxane, case b).} \end{aligned}$$

Comparing $\frac{1}{2} \text{tr } \Delta^e \alpha$ with F^e in Table 1 immediately shows that the results from EOEM reported are by no way changed if $\frac{1}{2} \text{tr } \Delta^e \alpha$ is taken into account. But a comparison of $\frac{1}{2} \text{tr } \Delta^a \alpha$ with F^a in Table 3 shows that $\frac{1}{2} \text{tr } \Delta^a \alpha$ amounts up to 10 to 15% of F^a . A brief recalculation of the results of EOAM reported here shows that the results are not changed to a substantial amount, as S_1 to S_3 are not changed at all by taking into account $\frac{1}{2} \text{tr } \Delta^a \alpha$. As the influence of polarizabilities on G^a and G^e (cf. Ref. [31]) is similar, it may be seen that the results reported here on polarizabilities are correct. The dipole moment $\mu_a^{0\text{FC}}$ as reported in Table 7 is changed from $27.9 \cdot 10^{-30} \text{ Cm}$ to $26.7 \cdot 10^{-30} \text{ Cm}$ taking into account $\frac{1}{2} \text{tr } \Delta^a \alpha$, and this clearly is within the errors. So a recurrence cannot improve the results, or even change the interpretation of the results from EOAM and EOEM on DMABN.

VII. Concluding Remarks

It has been shown that the results from EOEM could only be understood taking into account terms like $(1 - f\alpha)^{-1}$. Then EOEM give a permanent dipole moment μ_a^0 for the excited state emitting the a-fluorescence that is in between 19 and $27 \cdot 10^{-30} \text{ Cm}$. $\mu_a^{0\text{FC}}$ of the FC excited A* state is determined from EOAM to be around $28 \cdot 10^{-30} \text{ Cm}$. This means that the permanent dipole moment increases with excitation but does not increase further with the following solvent or intramolecular relaxation.

So the results presented here do neither interfere with the TICT model nor with the level crossing model in their essential points. But in the case of the TICT model, the postulated twisted rotamer does not show a nearly full charge transfer, and in the case of the level crossing model, the level crossing is to an essential amount due to polarizabilities.

In order to elucidate the part of polarizabilities in the total effective dipole moments in solvents further, electrooptical investigations on sterically hindered molecules related to DMABN are under way. Such investigations also suggest themselves because electrooptical investigations on ADMA [21] and on a class of merocyanine dyes show strong evidence that polarizabilities cannot be neglected.

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