Supplementary Materials for

Polymer Filters for Ultraviolet-Excited Integrated Fluorescence Sensing

Marc Dandin, Pamela Abshire, and Elisabeth Smela

Photopolymerization Model for Curing BTA-Doped Photosensitive Polymers

We begin by calculating the exposure dose, i.e. the energy, required for photopolymerization. To do so, we first consider the penetration depth D_p . This parameter is defined as the location at which the ultraviolet energy incident on a photosensitive film decays to a value of 1/e of its maximum value. The thickness of the film that is cured is related to the penetration depth as follows:

(1)
$$C_p = D_p \ln\left(\frac{E_{\max}}{E_c}\right),$$

where E_{max} is the energy at the surface of the film and E_c is the minimum energy required to induce photopolymerization¹.

Furthermore, the curing energy as a function of film thickness follows the Beer-Lambert law:

(2)
$$E_u(z) = E_{\max} \cdot e^{-\frac{z}{D_p}}$$
.

The subscript u indicates the case of an undoped film. Substituting Equation (1) into Equation (2) yields

(3)
$$E_u(z) = E_{\max} \cdot e^{-z \frac{1}{C_p} \ln\left(\frac{E_{\max}}{E_c}\right)}$$

The term $\frac{1}{C_p} \ln\left(\frac{E_{\text{max}}}{E_c}\right)$ has inverse length units. Thus, this term can be considered as an effective absorption coefficient and Equation (3) can be requirited.

effective absorption coefficient and Equation (3) can be re-written

(4)
$$E_u(z) = E_{\max} \cdot e^{-\alpha_{eff} z}$$

where

(5)
$$\alpha_{eff} = \frac{1}{C_p} \ln\left(\frac{E_{\max}}{E_c}\right).$$

Equation (4) evaluated at $z = n \times \delta$, where δ is the physical thickness of the film and *n* its refractive index, is the *minimum* energy required to fully cure the film. The total exposure dose (E_u) as a function of film thickness is typically supplied by the manufacturer.

Let us now consider the case of a doped photosensitive pre-polymer film. The energy absorbed by the UV-absorbing compound reduces the energy available for curing. This effect is additive in terms of the absorption coefficient. The amount of energy available for curing a doped film becomes

(6)
$$E_d(z) = E_{\max} e^{-\alpha_{eff} z} e^{-\alpha_d z} = E_u(z) e^{-\alpha_d z},$$

where the parameter α_d is the additional attenuation resulting from introducing the dopant.

As Equation (6) shows, for large α_d as in the case for BTA, $E_d \ll E_u$. This implies that the energy received by the SU-8, E_d , will not be sufficient to induce polymerization. The loss of energy resulting from BTA absorption must be compensated by increasing the exposure time.

Thus, the energy required to cure the doped film, E_d^* , where the subscript "*" denotes the adjustment in exposure time, is:

(7)
$$E_d^*(z) = \mathbf{I}_{\max}(t_E + \Delta t) \cdot e^{-(\alpha_{eff} + \alpha_d)z}$$

 I_{max} is the irradiance of the UV lamp, t_E is the time it would take to cure an undoped film of optical thickness z, and Δt is the requisite additional time.

Simplifying Equation (7) we obtain

(8)
$$E_d^*(z) = \mathbf{I}_{\max} t_E e^{-\alpha_{eff} z} e^{-\alpha_d z} + \mathbf{I}_{\max} \Delta t e^{-\alpha_{eff} z} e^{-\alpha_d z}.$$

Since $E_{\text{max}} = I_{\text{max}} t_E$, we can write

(9)
$$E_d^*(z) = E_{\max} e^{-\alpha_{eff} z} e^{-\alpha_d z} + \frac{E_{\max}}{t_E} \Delta t e^{-\alpha_{eff} z} e^{-\alpha_d z}$$

(10)
$$E_d^*\left(z\right) = E_{\max} e^{-\alpha_{eff} z} e^{-\alpha_d z} \left(1 + \frac{\Delta t}{t_E}\right) = E_u\left(z\right) e^{-\alpha_d z} \left(1 + \frac{\Delta t}{t_E}\right).$$

Recall that for polymerization to occur, the undoped film must be exposed with a minimum dosage of E_u . In order to have $E_d^* \ge E_u$, the condition required for curing the BTA-doped film, the following condition must hold:

(11)
$$e^{-\alpha_d z} \left(1 + \frac{\Delta t}{t_E} \right) \ge 1.$$

Thus, the additional time required for successful curing must satisfy

(12)
$$\Delta t \ge t_E \left(e^{\alpha_d z} - 1 \right).$$

The total exposure time is thus:

(13)
$$t_E^* = t_E + \Delta t \ge t_E e^{\alpha_d z}$$

The parameter α_d is a concentration and wavelength dependent absorption coefficient and it is given by the following formula:

(14)
$$\alpha_d = \frac{\varepsilon \cdot c}{M},$$

where ε is the extinction coefficient¹, M is the molecular weight of the doping chromophore, and c is its concentration in the pre-polymer. For a model particular to the photolithography of SU-8-BTA composites at the i-line wavelength, we re-write Equation (13) as

(15)
$$t_E^* \ge t_E e^{\left(\frac{\mathcal{E}_i \cdot \mathcal{C}_{BTA}}{M}\right) n_{SU8} \delta}.$$

The i-line extinction coefficient ε_i is obtained experimentally using the molecular formulation of the Beer-Lambert law, namely, $A = \varepsilon_i cl$, where A is the absorbance of the chromophore in solution at concentration c across an optical thickness l. The chromophore concentration c_{BTA} is obtained by dividing the number of moles of BTA in the doping aliquot by the pre-polymer volume. (The solvent volume can be neglected because it evaporates during curing).

Scaling Equation (15) by I_{max} yields the minimum exposure dose needed for polymerizing the composite:

(16)
$$E_d^*(\delta) \ge E_u(\delta) e^{\left(\frac{\varepsilon_i \cdot c_{BTA}}{M}\right) n_{SUS}\delta}.$$

References

1. J. H. Lee, R. K. Prud'Homme and I. A. Aksay, "Cure depth in photopolymerization: experiments and theory", *J. Mater. Res.*, 2001, **16**, 3536-3544.