



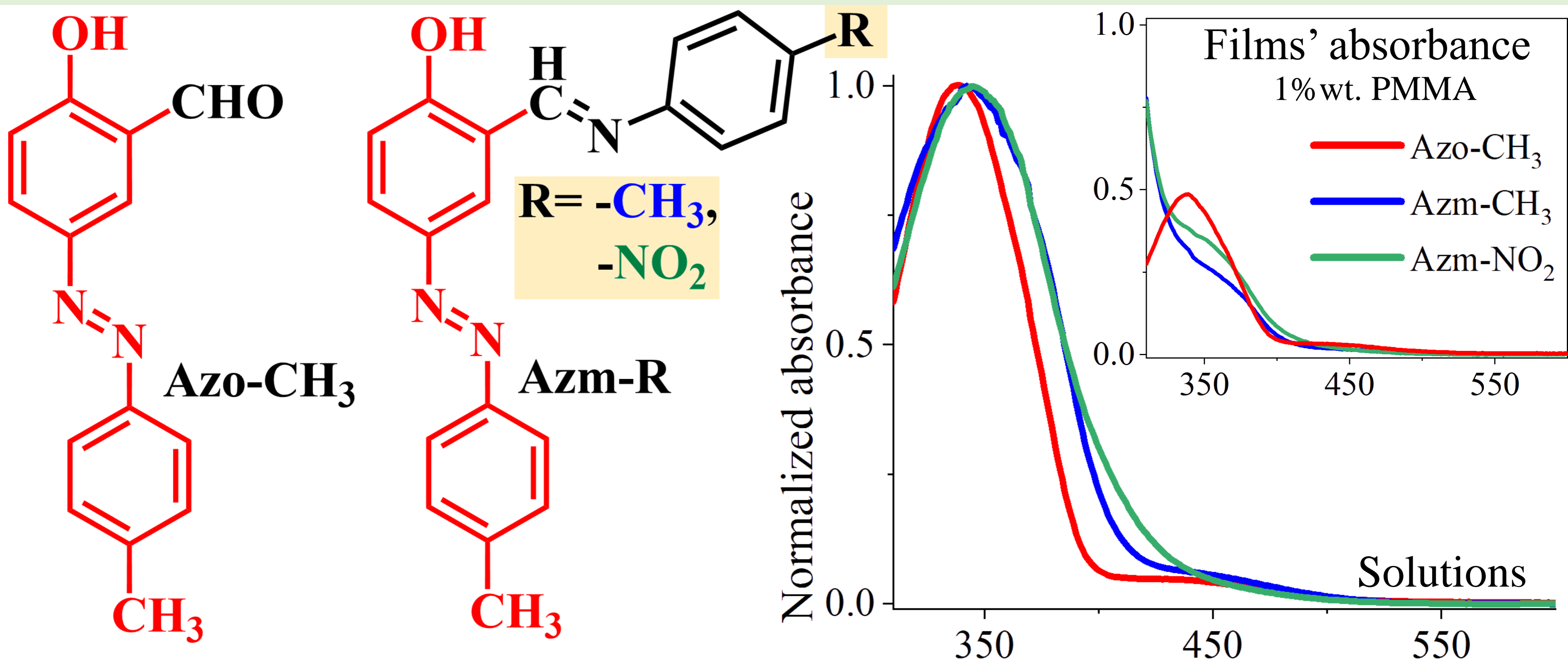
Effect of $-\text{CH}_3/-\text{NO}_2$ substitution on the self-action in azo- and azo-azomethines derivatives polymeric thin films under picosecond laser excitation at 1064 nm

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Advanced light-responsive polymeric materials containing $-\text{A}=\text{B}-$ chromophore groups undergo reversible *trans-cis-trans* photo-isomerization cycle, like $-\text{N}=\text{N}-$ or $-\text{CH}=\text{N}-$, are used in a number of modern electronic photoswitchable systems like sensors, liquid crystal displays [1], elements of optical information storage devices, etc.



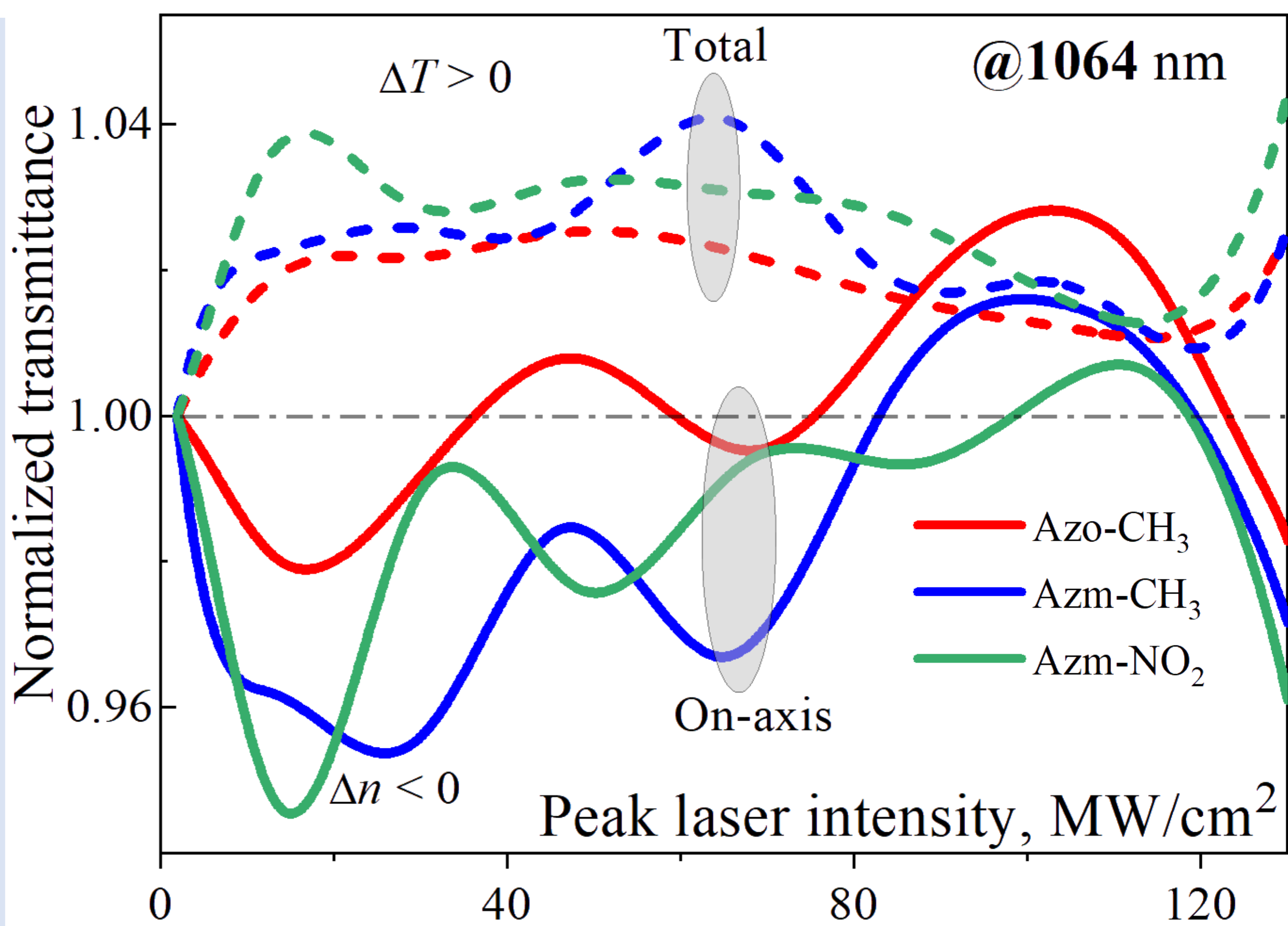
In present work composite “guest-host” films with “guest” azo- (*Azo-CH₃*) or azo-azomethine (*Azm-CH₃* / *Azm-NO₂*) and “host” PMMA were prepared. Photoinduced self-action effects in thin films were investigated using pulsed ps laser radiation at wavelength 1064 nm [2].

Conclusions:

1. Initial self-defocusing effect in *Azo-CH₃* rose twice after azomethine group incorporation without major donor/acceptor substituents impact. *Azo-CH₃* and *Azm-*

NO₂ saturate at lower intensity ($I \sim 15 \text{ MW/cm}^2$) than *Azm-CH₃* ($I \sim 30 \text{ MW/cm}^2$).

2. It was shown: efficiency of the refractive NLO response $|\text{Re}(\chi^{(3)})| \sim 10^{-8} \text{ esu}$ increases in the following order: *Azm-NO₂* > *Azm-CH₃* > *Azo-CH₃*; all the dyes demonstrated slight photobleaching ($\sim 2\%$ for *Azo-CH₃* and *Azm-CH₃* and $\sim 4\%$ for *Azm-NO₂*) that saturated at about $I \sim 25 \text{ MW/cm}^2$.



1. Uklein, A. et al. // Opt. Commun. **296** (2014) P. 79. <https://dx.doi.org/10.15407/fm22.01.020>

2. Uklein, A. et al. // Appl. Phys. B. **122** (2016) P. 287. <https://doi.org/10.1007/s00340-016-6561-2>