



Visible light promoted thiol-ene reactions using titanium dioxide†

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The radical addition of thiols to alkenes is reported under photo-redox conditions, using visible light and TiO₂ as a cheap and readily available photocatalyst.

Recent developments in visible light photoredox catalysis (PRC) have substantially expanded the scope and application of organic photochemistry.¹ Using a domestic lightbulb as the light source, rather than specialist UV equipment, the process uses a photo-active catalyst to mediate electron transfer between reactants, resulting in novel transformations that frequently proceed under mild conditions. The vast majority of PRC in organic synthesis to date is based on two classes of catalyst; transition metal complexes, such as the archetypal Ru(bpy)₃Cl₂ and Ir(ppy)₃ systems, or organic dyes such as methylene blue or eosin-Y. The Noble metals of the first class are expensive and a limited resource, but are highly optimisable in terms of ligand-metal tuning. Organic dyes lack this particular quality, but may be tuned *via* conventional synthesis and are usually less expensive. We were interested in exploiting a third class of photoredox catalyst; semi-conducting metal oxides such as titania (TiO₂). With 4 million tons of titania being produced per annum, the material is cheap, readily accessible, and non-toxic (commonly ingested as food additive E 171), as well as offering the process benefits of heterogeneous catalysis such as simple removal by filtration.² The UV photoredox properties of TiO₂ have been extensively investigated and applied on an industrial scale to areas such as water treatment and photovoltaics. Applications to organic synthesis, however, remain under-developed, particularly in the area of visible light photochemistry.^{3,4} We were interested in developing a titania-mediated PRC reaction using visible light, and chose to investigate the thiol-ene reaction for this purpose.

The addition of thiols to alkenes is a radical reaction that is typically initiated with UV light.⁵ Although an old reaction,⁶ its

reliability, substrate scope, and compatibility has led to renewed interest in the context of click chemistry, where the facile C–S bond formation can be used as a powerful ligation strategy in biological and materials science.⁷ Indeed, the largest field of application for thiol-ene chemistry is currently in polymer science, where thiols are cross-linked with alkenes under UV curing conditions, producing polymers with very uniform step-growth properties.⁸ Recent literature reports have established PRC for thiol-ene addition using both Ru(II) and eosin-Y photocatalysis, providing an important frame of reference for our inchoate titania process.^{9–11}

We were aware at the outset that titania visible light PRC would likely require a nanoparticulate form of the catalyst. Unmodified titania absorbs very weakly in the visible region, but there is evidence that substrate binding to the surface can shift the absorption and enable electron transfer.^{4a,c–e} Rueping and co-workers, for example, have recently demonstrated

Table 1 Titania PRC of the thiol-ene reaction^a

Entry	TiO ₂ (equiv.)	Solvent	Light source	Yield ^b (%)
1	1.0	MeCN	20 W light	98 (96 ^c)
2	1.0	DCM	20 W light	96
3	1.0	—	20 W light	97
4	1.0	MeCN	Blue LED	80
5 ^d	1.0	MeCN	No light	0
6 ^e	0	MeCN	20 W light	20
7	0	MeCN	Blue LED	0
8	0.5	MeCN	20 W light	88
9	0.1	MeCN	20 W light	82
10 ^e	0.1	MeCN	20 W light	92
11 ^f	1.0	MeCN	20 W light	50
12 ^g	1.0	MeCN	20 W light	84

^a Cyclohexene (1.0 mmol), benzyl mercaptan (4.0 mmol), titania (TiO₂ P25) and solvent (1.0 mL) at room temperature for 16 h; light source placed 10–15 cm away from reaction vials. ^b Yields determined by ¹H NMR using nitromethane as the internal standard. ^c Isolated yields. ^d Foil-wrapped flask for 16 h. ^e Reaction for 40 h. ^f Using degassed MeCN under N₂ atmosphere. ^g Using 2.5 mmol of benzyl mercaptan.

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Scheme 1 Mechanistic pathway for titania thiol-ene reaction.

tolerated at the 2- and 3-positions of the alkene (entries 3 and 4), but 1,2-disubstitution was restricted to cyclic alkenes. Vinyl acetate and acrylonitrile were both productive, interestingly undergoing no reaction at all in the absence of PRC (entries 6 and 7). More reactive alkenes such as styrenes gave very high yields of the thiol ene products, but background reaction was high for **2i** and **2j**.

Thiol scope was initially trialled (Table 3) using catalytic amounts of titania (0.1 equiv.) and vinyl acetate (**2h**), an alkene with zero background reaction for the addition of benzyl mercaptan (Table 1, entry 7). We were pleased to find that the catalytic protocol was successful, with primary alkyl thiols (entries 1–3) and thiophenols (entry 4) undergoing thiol-ene reaction under the longer reaction time of 40 h. Increasing steric hindrance around the thiol was deleterious, with cyclohexyl thiol reacting in moderate yield (entry 5) and *tert*-butyl thiol undergoing no reaction, even with stoichiometric titania (entry 6). We could extend the reaction to a double thiol-ene reaction, with ethane and propane dithiol undergoing addition to styrenes in good yield (entries 7–9). *p*-Methoxystyrene proved less reactive, and required stoichiometric quantities of titania to yield **3t** in an acceptable timeframe.

A mechanism for the reaction is outlined in Scheme 1, based upon the photo-excitation of electrons to the conduction band of the titania catalyst. The resultant holes are reductively quenched by the thiol, generating a thiyl radical cation, which can lose a proton to give a thiyl radical. Oxygen acts as a sacrificial electron acceptor, enhancing reaction efficiency by reducing hole-electron recombination in the titania. The thiyl radical can then initiate the thiol-ene cycle through addition to an alkene and generation of an alkyl radical, which propagates the reaction by abstracting a hydrogen atom from the thiol starting material.

In conclusion, we have developed a titania photoredox system for the thiol-ene reaction that uses visible light. The

photocatalyst is cheap, robust, readily available, and easily processed out of the reaction for re-use. Applications of this method to the broad remit of thiol-ene chemistry are ongoing in our laboratory.

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