

## Supplementary data

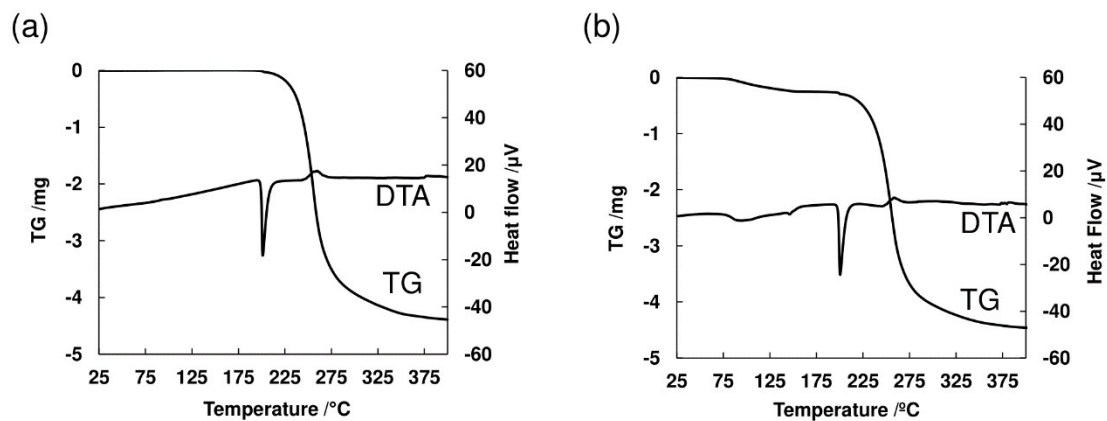


Fig.S1 The thermogravimetric (TG) analysis and differential thermal analysis (DTA) of PX-AH (a) and yellow solid precipitate (b). In (b), about 5.01% of mass loss was observed around 100 °C.

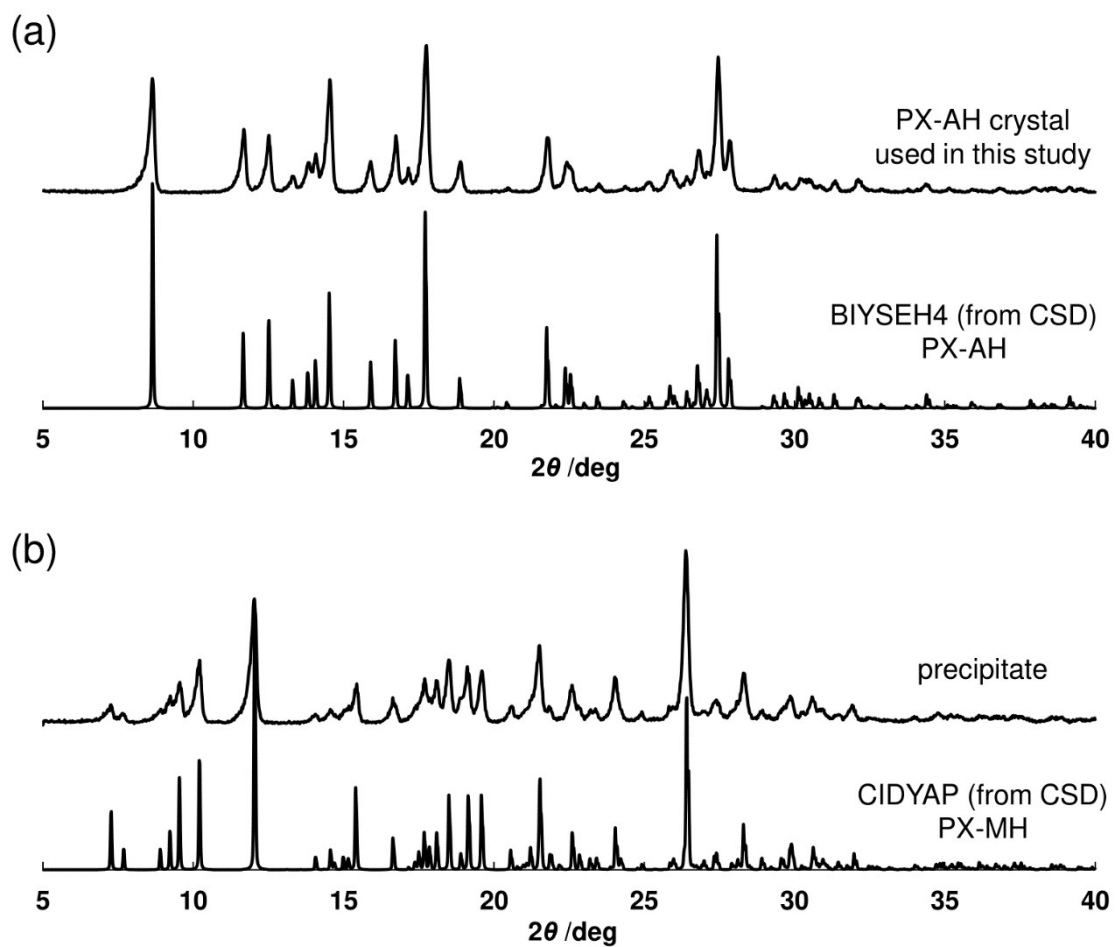


Fig. S2 The powder X-ray diffraction patterns of PX-AH (a) and yellow solid precipitate obtained from the dissolution study (b). The patterns of the PX-AH used in this study and the precipitate were compared to that of the PX-AH and PX-MH structure with a reference code of BIYSEH4 and CIDYAP indexed in the Cambridge Structural Database (CSD).

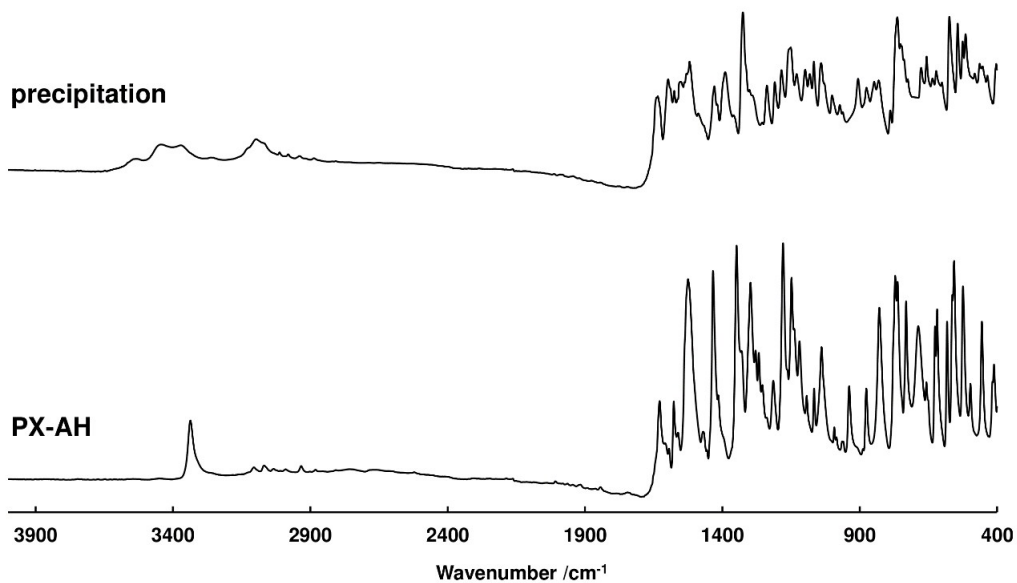


Fig.S3 The ATR-FTIR spectra of PX-AH (a) and yellow solid precipitate (b) obtained from the dissolution study of PX-AH in the absence of LAs.

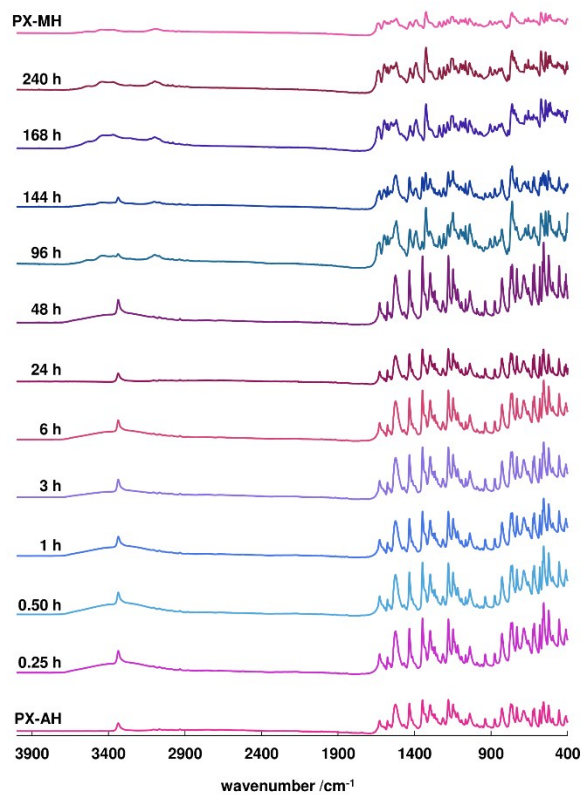


Fig.S4 The ATR-FTIR spectra of the solid phase of the samples when the dissolution study of PX-AH in 5 mM OXY solution was conducted in section 3.1.2.

Table S1. The values  $[PX]_{24h}$ ,  $\Delta[PX]_{24h}$ ,  $\gamma$ , and  $\Delta\gamma$  of for each LAs.

	$[PX]_{24h}$	$\Delta[PX]_{24h}$	$\gamma$	$\Delta\gamma$
control	1.32	0.00	71.24	0.00
5 mM OXY	0.73	-0.59	46.86	-24.38
5 mM TET	0.71	-0.60	46.05	-25.19
5 mM LID	1.33	0.02	62.88	-8.36
5 mM DIB	0.06	-1.25	32.04	-39.20

a:  $[PX]_{24 h}$  is the concentration of PX at 24h in buffer or 5 mM basic drug solutions

b:  $\Delta[PX]_{24 h} = [PX]_{5 \text{ mM basic drugs}}_{24 h} - [PX]_{\text{control}}_{24 h}$

c: Surface tension of 5 mM basic drug solutions. Control is the buffer solution without basic drugs.

d:  $\Delta\gamma = \gamma_{5 \text{ mM basic drug}} - \gamma_{\text{control}}$

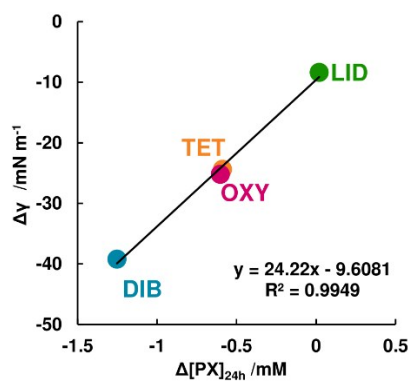


Fig. S5  $\Delta[\text{PX}]_{24\text{h}}$  and  $\Delta\gamma$  of LAs. For  $\Delta[\text{PX}]_{24\text{h}}$  and  $\Delta\gamma$ , the values in Table S1 of each LAs are used.

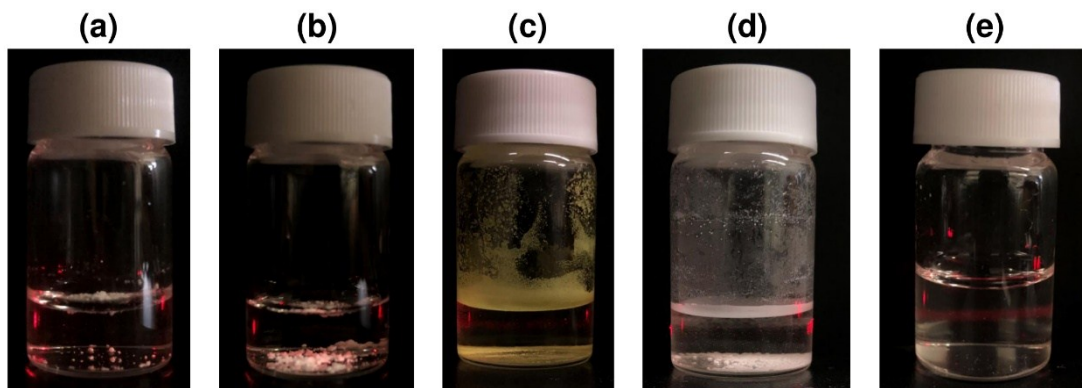


Fig. S6 The photos of the samples shined with light to observe the Tyndall effect. (a) PX-AH in buffer (before stirring), (b) PX-AH in 5 mM OXY solution (before stirring), (c) PX-AH in buffer (after stirring 6 h), (d) PX-AH in 5 mM OXY (after stirring 6 h) and (e) hydrogel of agarose (before stirring). The Tyndall effect only showed in (e).

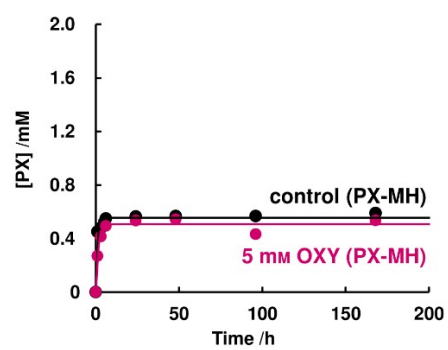


Fig. S7 The dissolution behavior of PX-MH in the absence and presence of OXY.



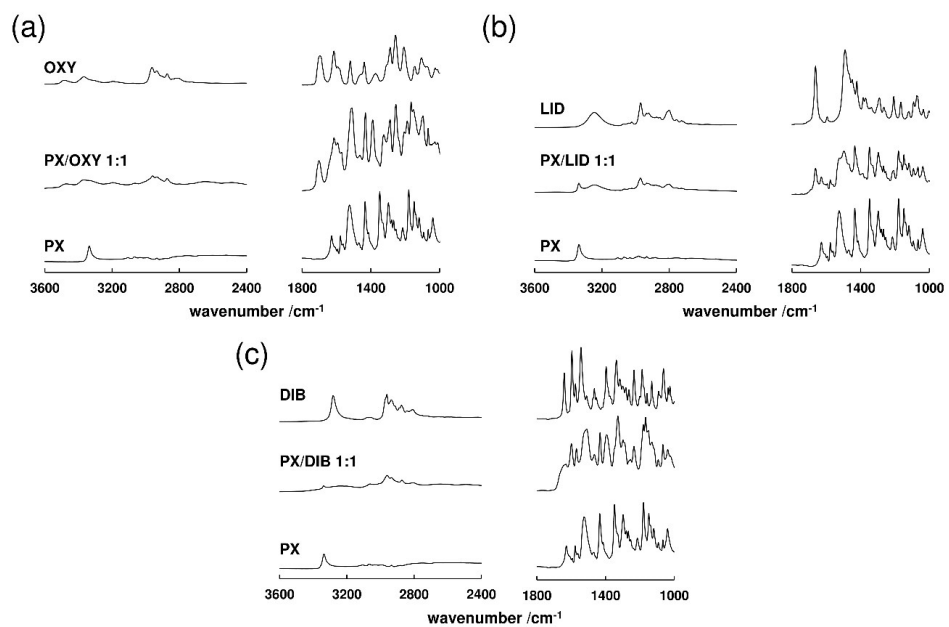


Fig. S8 The ATR-FTIR spectra of PX/OXY (a), PX/LID (b), and PX/DIB (c) mixtures in the molar ration of 1:1.

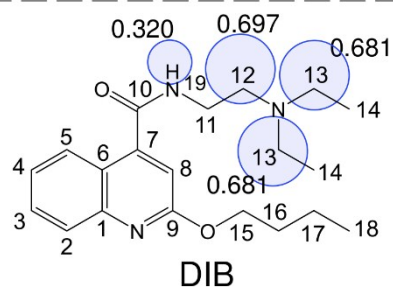
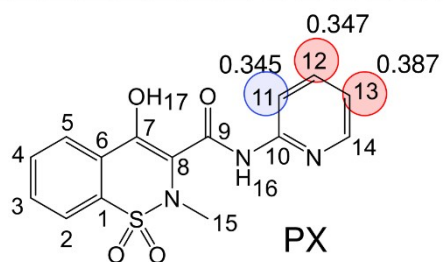
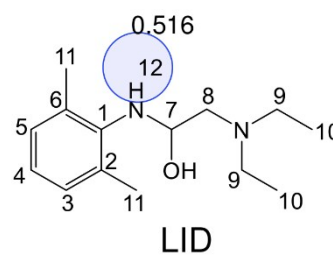
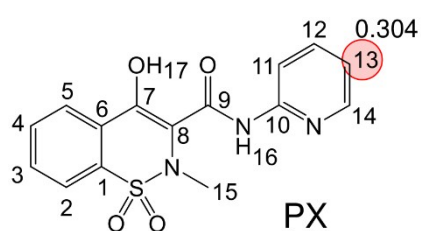
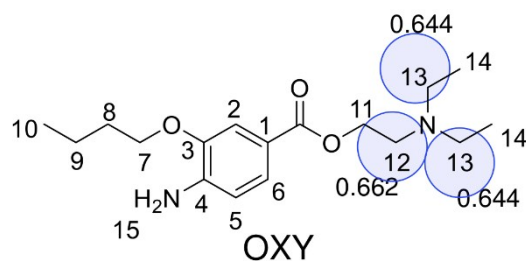
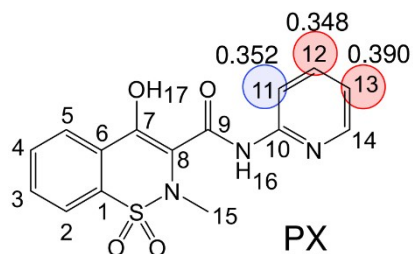
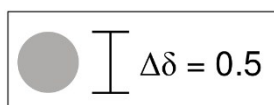


Fig. S9 The results of  $^1\text{H-NMR}$  analysis of PX/OXY, PX/LID, and PX/DIB mixtures in  $\text{DMSO-}d_6$ . The size of the circle shows how much the chemical shift shifted ( $\Delta\delta$ ) to lower (red) or higher (blue) magnetic field due to mixing PX and each LAs. The value written near each circle is the value of  $\Delta\delta$ .