Rapid TLC with Densitometry for Evaluation of Naproxen Stability

Wioletta Parys*, Małgorzata Dołowy, Alina Pyka-Pająk

Department of Analytical Chemistry, Faculty of Pharmaceutical Sciences in Sosnowiec, Medical University of Silesia in Katowice, Jagiellońska 4, 41-200 Sosnowiec, Poland, mdolowy@sum.edu.pl (M.D.); apyka@sum.edu.pl (A.P.-P.)

* Correspondence: wparys@sum.edu.pl

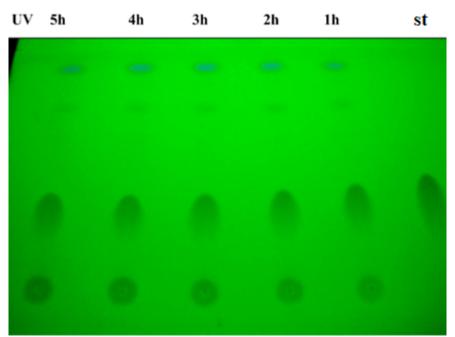


Fig. S1. Photograph of naproxen chromatogram that was irradiated with UV radiation at $\lambda = 254$ nm on silica gel for a period of 1 h, 2 h, 3 h, 4 h, 5 h before developing the chromatographic plate (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v); photo taken at $\lambda=254$ nm; where st – standard of naproxen (non-irradiated)

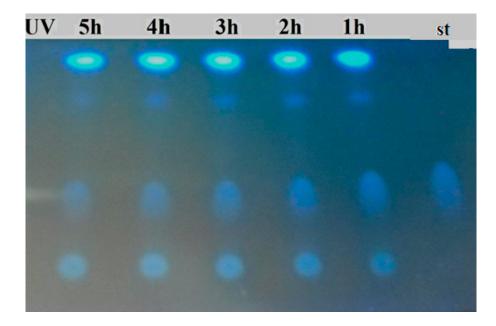


Fig. S2. Photograph of naproxen chromatogram that was irradiated with UV radiation at $\lambda = 254$ nm on silica gel for a period of 1 h, 2 h, 3 h, 4 h, 5 h before developing the chromatographic plate (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v); photo taken under UV light at 366 nm); where st – standard of naproxen (non-irradiated)

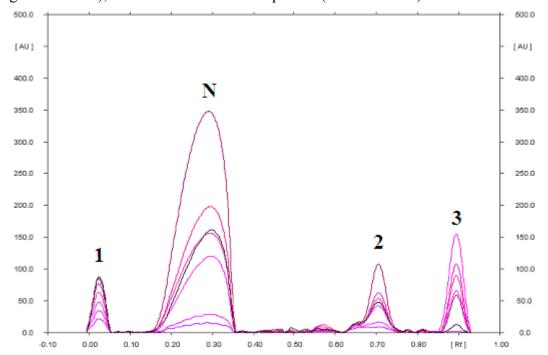


Fig. S3. Densitogram of naproxen irradiated on silica gel with UV radiation $\lambda = 254$ nm for a period of 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

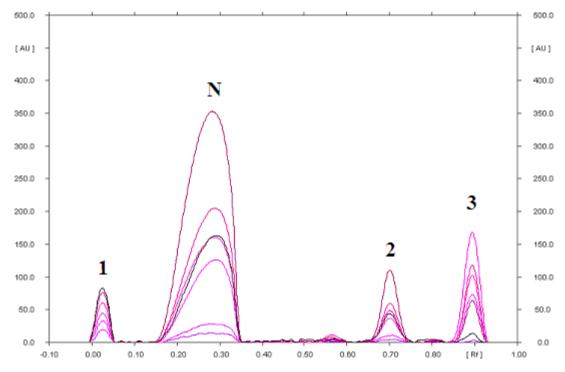


Fig. S4. Densitogram of naproxen irradiated on silica gel with UV radiation $\lambda = 254$ nm for a period of 4 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

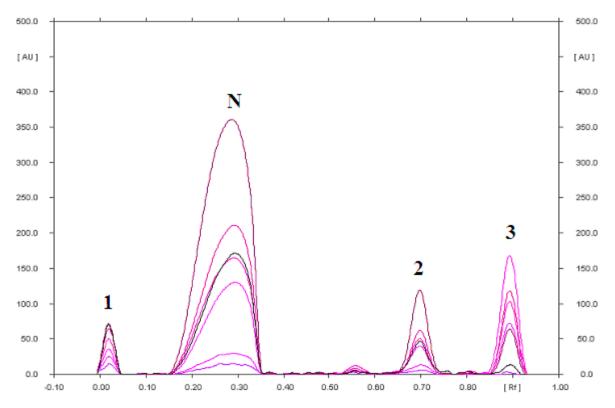


Fig. S5. Densitogram of naproxen irradiated on silica gel with UV radiation $\lambda = 254$ nm for a period of 3 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

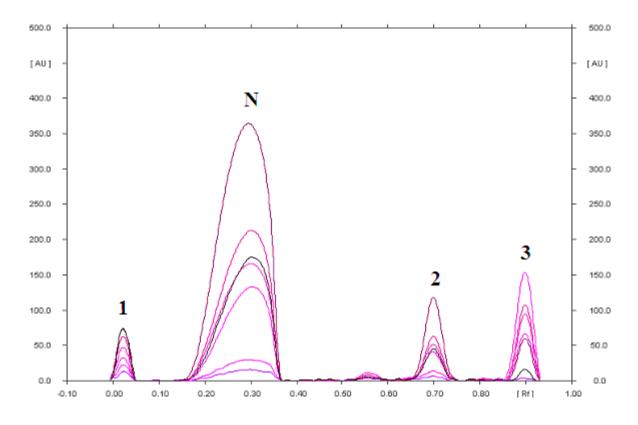


Fig. S6. Densitogram of naproxen irradiated on silica gel with UV radiation $\lambda = 254$ nm for a period of 2 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

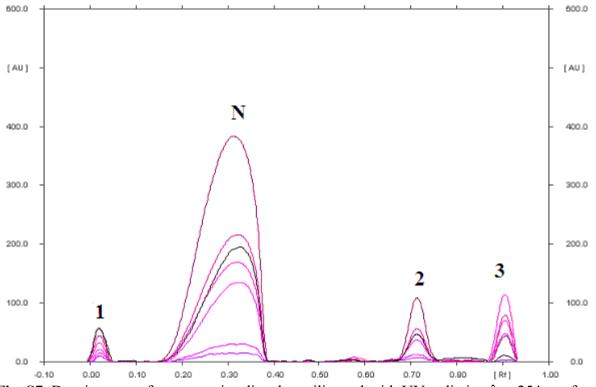


Fig. S7. Densitogram of naproxen irradiated on silica gel with UV radiation $\lambda = 254$ nm for a period of 1 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

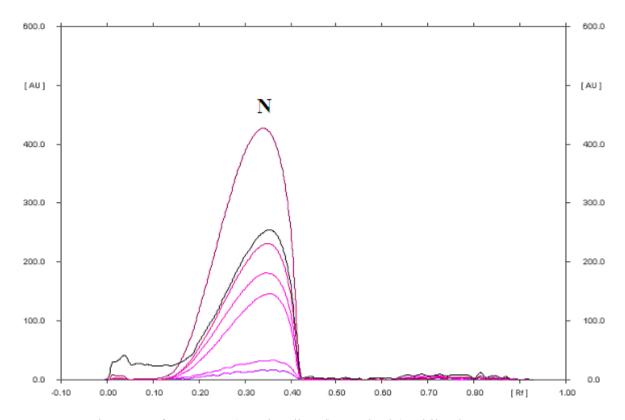


Fig. S8. Densitogram of naproxen (non-irradiated) standard (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

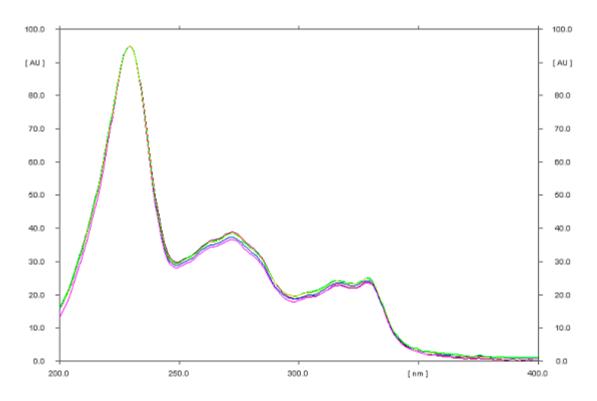


Fig. S9. Spectra of naproxen not exposed and irradiated with UV radiation at $\lambda = 254$ nm for a period of 1 h to 5 h on silica gel (mobile phase A: TOL - ACE - CHL,2:5:12 v/v/v)).

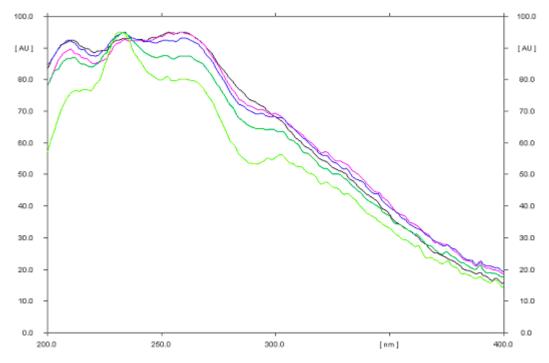


Fig. S10. Spectra of substance 1 with the value $R_F = 0.02$ formed after irradiation with UV light at $\lambda = 254$ nm naproxen on silica gel for 1 h, 2 h, 3 h, 4 h and 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

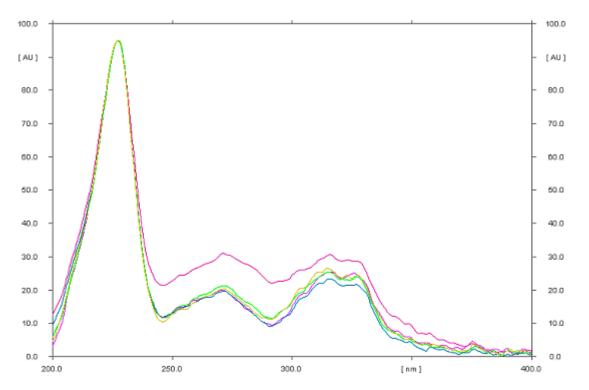


Fig. S11. Spectra of substance 2 (naproxen ethyl ester) with the value $R_F = 0.71$ formed after irradiation with UV light with $\lambda = 254$ nm naproxen on silica gel for 1 h, 2 h, 3 h, 4 h and 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

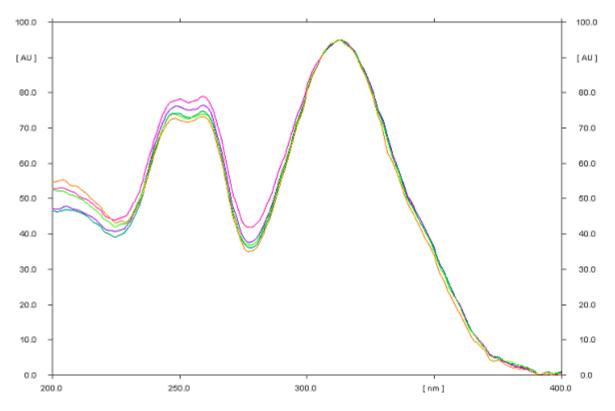


Fig. S12. Spectra of substance 3 with the value $R_F = 0.89$ formed after irradiation with UV light with $\lambda = 254$ nm naproxen on silica gel for 1 h, 2 h, 3 h, 4 h and 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

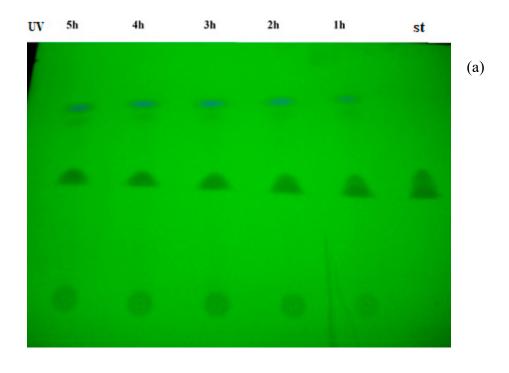
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Exposure time of naproxen on silica gel	Substance ^{a)}	R _F	Chromatographic peak area [AU] ^{b)}	% of band area
0 h	Ν	0.29	49298	100
1 h	N	0.29	36946	83.47
	S1	0.02	842	1.90
	S2	0.71	3290	7.43
	S3	0.89	3187	7.20
2 h	N	0.29	32965	77.18
	S1	0.02	1206	2.82
	S2	0.70	3814	8.93
	S3	0.89	4725	11.07
3 h	N	0.29	31668	75.92
	S1	0.02	907	2.17
	S2	0.70	3765	9.03
	S3	0.89	5373	12.88

Table	S1.	R_F	values	and	chromatographic	peak	areas	of	naproxen	and	its	chemical
		tran	sformati	on pi	oducts under the	influer	nce of	UV	light (254	nm)	on	silica gel,
		after	r separat	ion u	sing a mobile phas	e A: T	OL - A	CE	- CHL (2::	5:12,	v/v/	v).

4 h	N	0.29	30892	75.39
	S1	0.02	1278	3.12
	S2	0.71	3533	8.62
	S3	0.90	5274	12.87
5 h	N	0.29	31257	76.36
	S1	0.02	1160	2.82
	S2	0.71	3855	9.42
	S3	0.90	4666	11.39

where:

^{a)} N - naproxen S1, S2, S3 - products of chemical transformation of naproxen; and S2 was identified as naproxen ethyl ester ^{b)} Area of the chromatographic band at λ_{max}





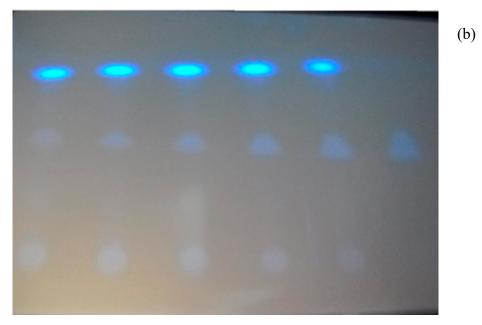


Fig S13. Photographs of chromatograms of naproxen that was irradiated with UV light (254 nm) on silica gel for 1 h, 2 h, 3 h, 4 h, 5 h before developing the chromatographic plate (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, (v/v/v); photos taken under UV light at (a) 254 nm and (b) 366 nm); where st – standard of naproxen (non-irradiated)

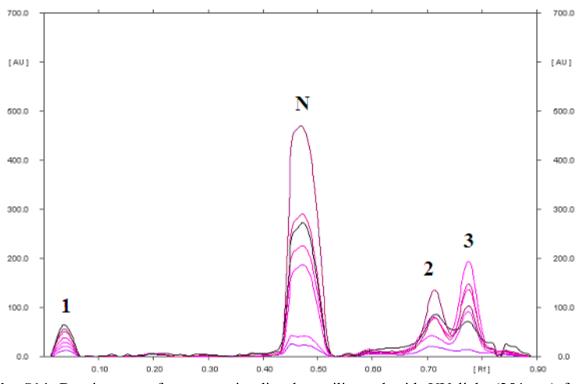


Fig. S14. Densitogram of naproxen irradiated on silica gel with UV light (254 nm) for a period of 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

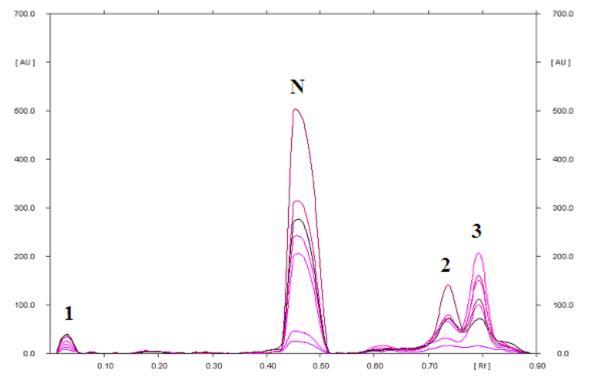


Fig. S15. Densitogram of naproxen irradiated on silica gel with UV light ($\lambda = 254$ nm) for a period of 4 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

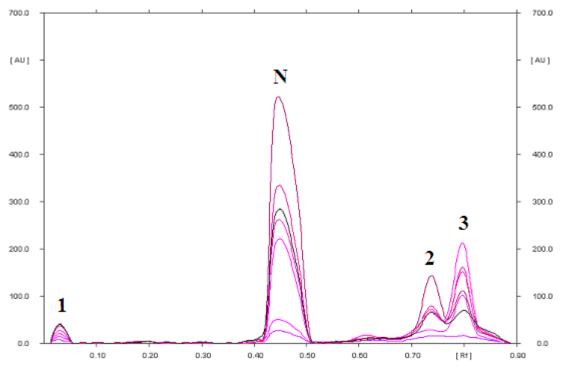


Fig. S16. Densitogram of naproxen irradiated on silica gel with UV light ($\lambda = 254$ nm) for a period of 3 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

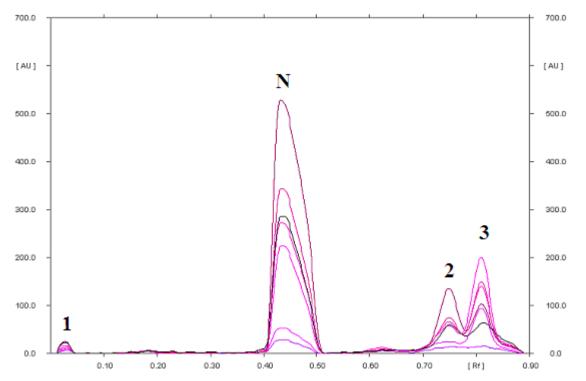


Fig. S17. Densitogram of naproxen irradiated on silica gel with UV light ($\lambda = 254$ nm) for a period of 2 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

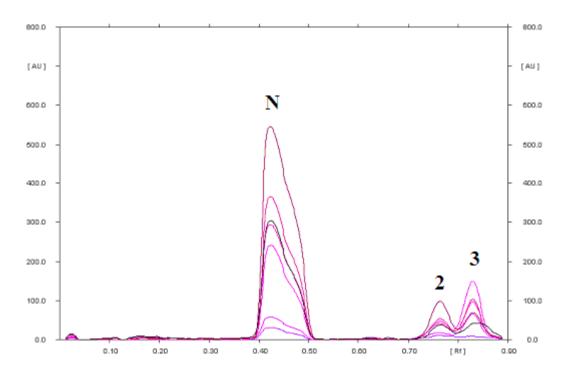


Fig. S18. Densitogram of naproxen irradiated on silica gel with UV light ($\lambda = 254$ nm) for a period of 1 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

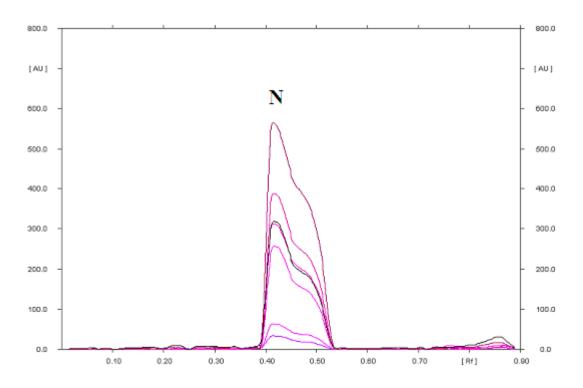


Fig. S19. Densitogram of naproxen standard (non-irradiated) (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

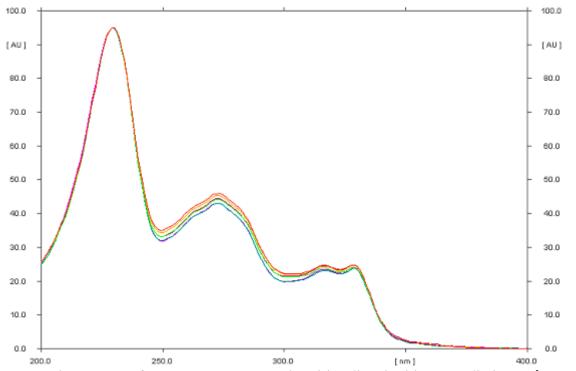


Fig. S20. The spectra of naproxen not exposed and irradiated with UV radiation at $\lambda = 254$ nm for a period of 1 h to 5 h on silica gel (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

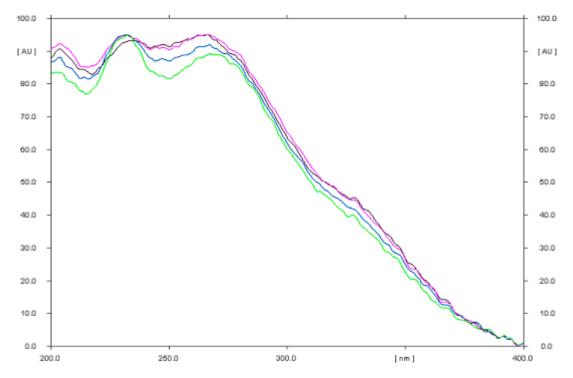


Fig. S21. The spectra of substance 1 with $R_F = 0.03$ formed after UV irradiation with $\lambda = 254$ nm naproxen on silica gel for 2 h, 3 h, 4 h and 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

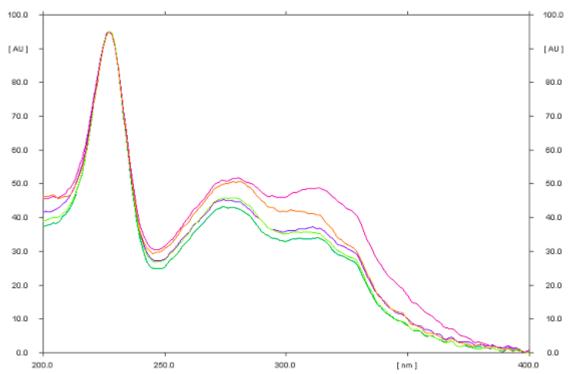


Fig. S22. Spectra of substance 2 with $R_F = 0.74$ formed after UV irradiation with $\lambda = 254$ nm naproxen on silica gel for 2 h, 3 h, 4 h and 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

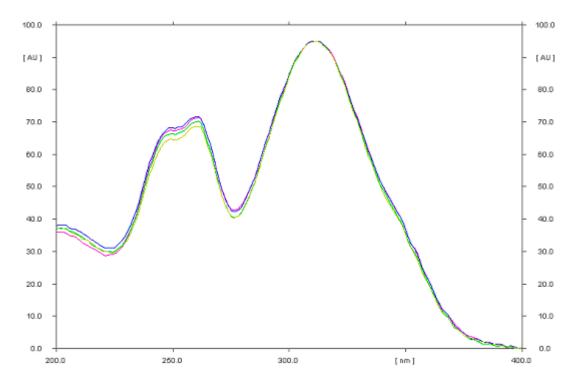


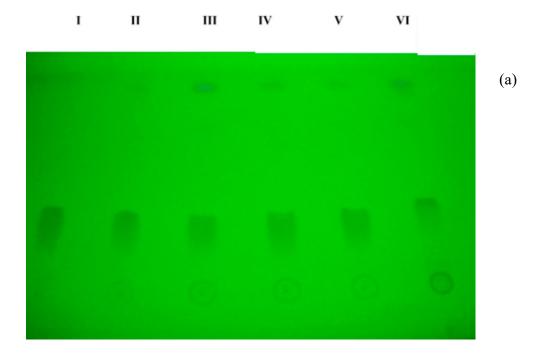
Fig. S23. Spectra of substance 3 with $R_F = 0.79$ formed after UV irradiation with $\lambda = 254$ nm naproxen on silica gel for 2 h, 3 h, 4 h and 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v)).

Exposure time of naproxen on silica gel	Substance ^{a)}	R _F	Chromatographic peak area [AU] ^{b)}	% of band area
0 h	Ν	0.44	37896	100
1 h	N	0.45	28703	90.52
	S1	-		-
	S2	0.75	2337	7.37
	S3	0.78	670	2.11
2 h	N	0.45	25327	83.60
	S1	0.03	354	1.17
	S2	0.74	2774	9.16
	S3	0.81	1839	6.07
3 h	N	0.46	22429	71.78
	S1	0.03	748	2.40
	S2	0.74	3660	11.71
	S3	0.80	4409	14.11
4 h	N	0.46	20843	69.65
	S1	0.03	615	2.05
	S2	0.74	3662	12.24
	S3	0.79	4806	16.06
5 h	N	0.47	20640	65.53
	S1	0.04	1223	3.88
	S2	0.71	4216	13.39
	S3	0.78	5416	17.20

Table S2. R_F values and of chromatographic peak areas of naproxen and its chemical transformation products under the influence of UV light with (254 nm) on silica gel, after separation using a mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v).

^{a)} N - naproxen

S1, S2, S3 - products of chemical transformation of naproxen; and S2 was identified as naproxen ethyl ester ^{b)} Area of the chromatographic band at λ_{max}



I II III IV V VI

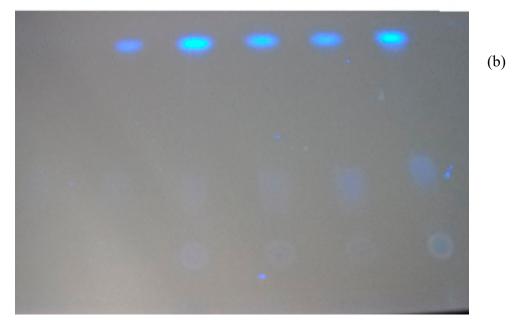


Fig. S24. Photographs of chromatograms, naproxen not exposed (I) and naproxen in solutions II, III, IV, V and VI irradiated for 5 h with UV radiation at $\lambda = 254$ nm taken in UV light at 254 nm (a) and 366 nm (b), after separation using a mobile phase A : TOL - ACE - CHL, 2:5:12 (v/v/v).

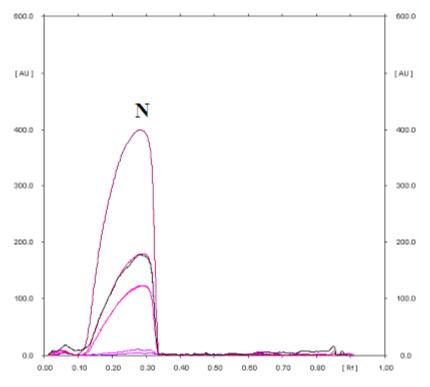


Fig. S25. Densitogram of naproxen standard (not exposed to UV) developed using mobile phase A: TOL - ACE - CHL (2:5:12, v/v/v).

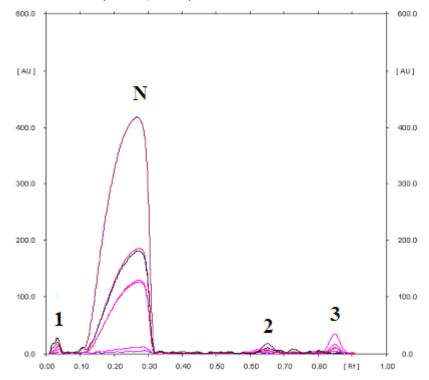


Fig. S26. Densitogram of naproxen (N) in solution of physiological salt (II), which was irradiated to UV light (254 nm) for 5h (mobile phase A: TOL-ACE-CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

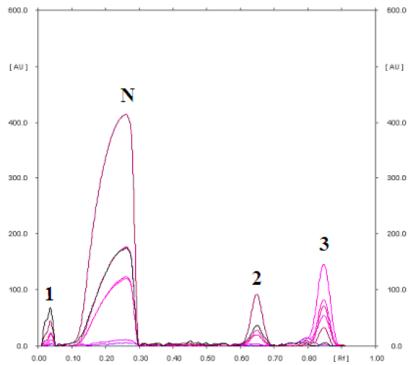


Fig. S27. Densitogram of naproxen (N) in solution of water at pH \approx 2.60 (III), which was irradiated to UV light (254 nm) for 5h (mobile phase A: TOL-ACE-CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

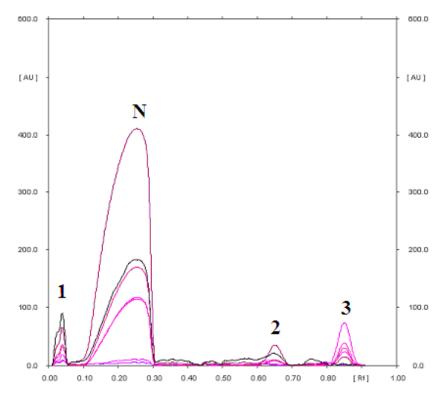


Fig. S28. Densitogram of naproxen (N) in solution of water at pH \approx 5.70 (IV), which was irradiated to UV light (254 nm) for 5h (mobile phase A: TOL-ACE-CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

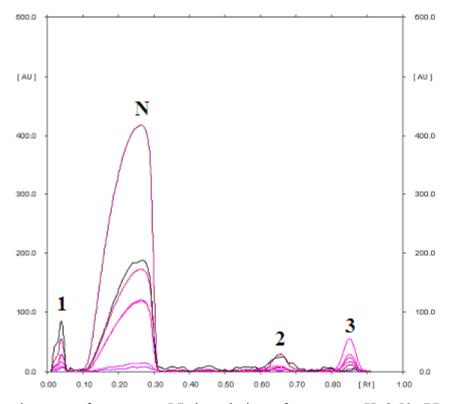


Fig. S29. Densitogram of naproxen (N) in solution of water at pH \approx 8.50 (V) which was irradiated to UV light (254 nm)for 5h (mobile phase A: TOL-ACE-CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

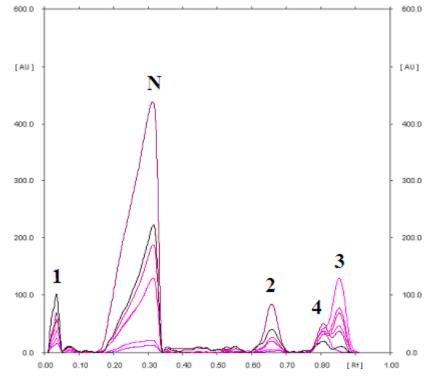


Fig. S30. Densitogram of naproxen (N) in hydrogen peroxide (VI), which was exposed to UV radiation (λ =254 nm) irradiated to UV light (254 nm) for 5h (mobile phase A: TOL-ACE-CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

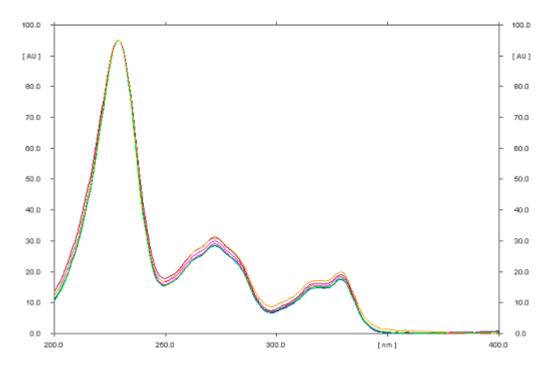


Fig. S31. Comparison of naproxen spectra ($R_F = 0.29$), which in solutions II, III, IV, V, VI was irradiated from above for 5 h and naproxen standard (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

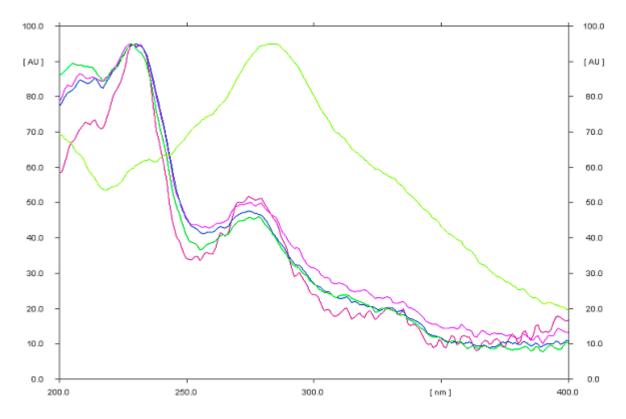


Fig. S32. Spectra of substance 1 with $R_F = 0.03$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

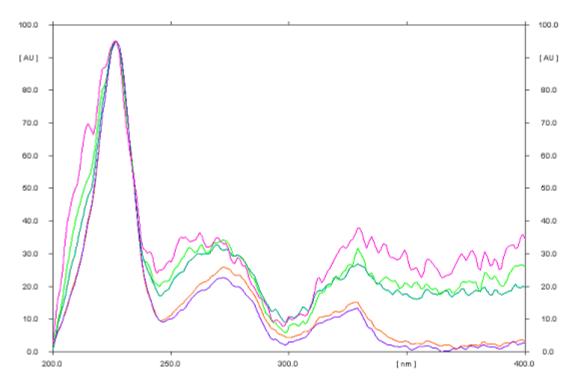


Fig. S33. Spectra of substance 2 (naproxen ethyl ester) with $R_F = 0.65$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

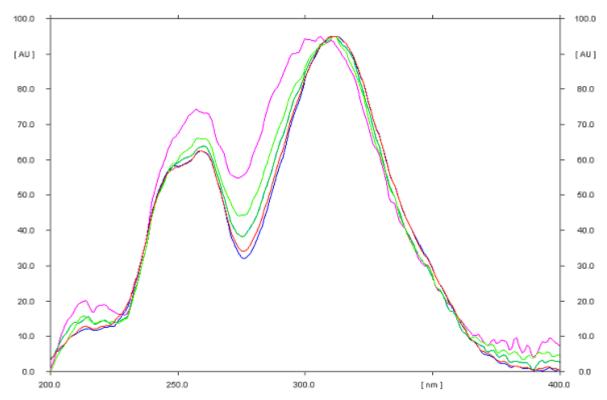


Fig. S34. Spectra of substance 3 with $R_F = 0.85$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).

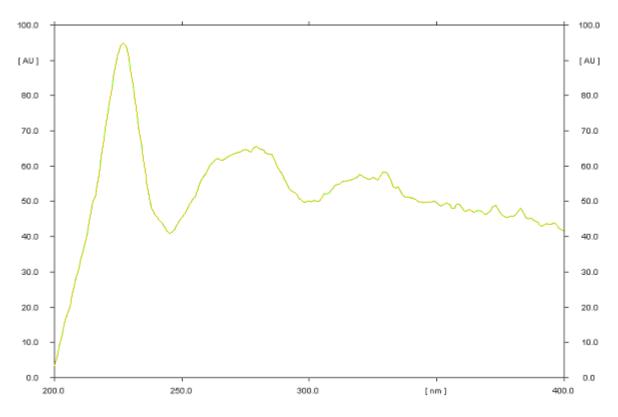
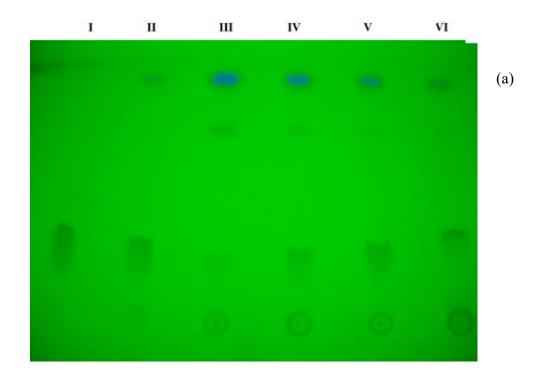


Fig. S35. Spectra of substance 4 with $R_F = 0.81$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v)).



I II III IV V VI

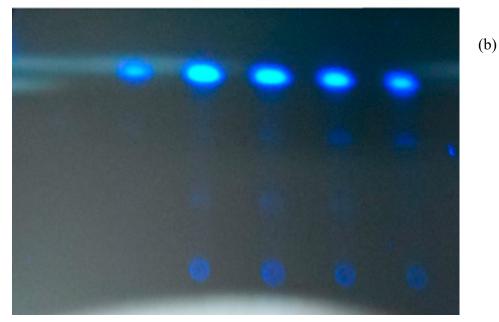


Fig. S36. Photographs of chromatograms, naproxen not exposed (I) and naproxen in solutions II, III, IV, V and VI irradiated for 10 h with UV radiation ($\lambda = 254$ nm) taken at 254 nm (a) and 366 nm (b), after separation using a mobile phase A: TOL - ACE - CHL, 2:5:12 (v/v/v).

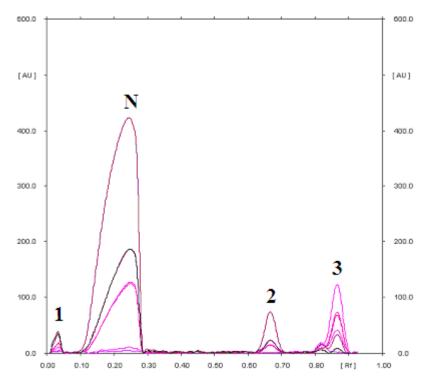


Fig. S37. Densitogram of naproxen (N) in solution of physiological salt (II), which was irradiated to UV light (254 nm)for 10 h (mobile phase A: TOL - ACE - CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

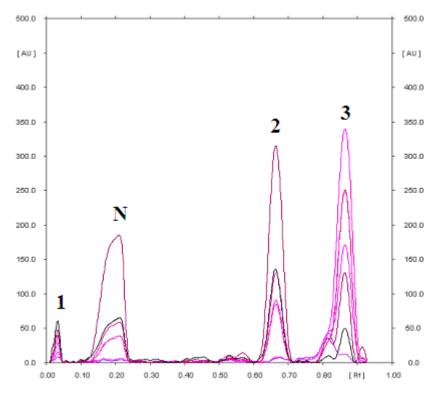


Fig. S38. Densitogram of naproxen (N) in water at pH \approx 2.60 (III), which was irradiated to UV light (254 nm) for 10 h (mobile phase A: TOL - ACE - CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

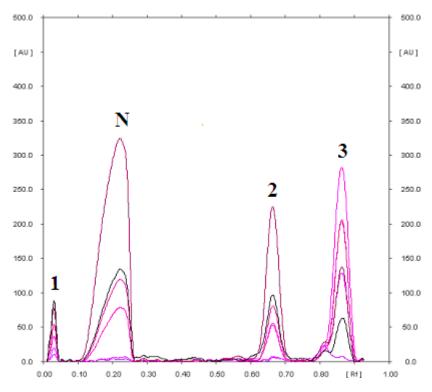


Fig. S39. Densitogram of naproxen (N) in water at pH \approx 5.70 (IV), which was irradiated to UV light (254 nm)for 10 h (mobile phase A: TOL - ACE - CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

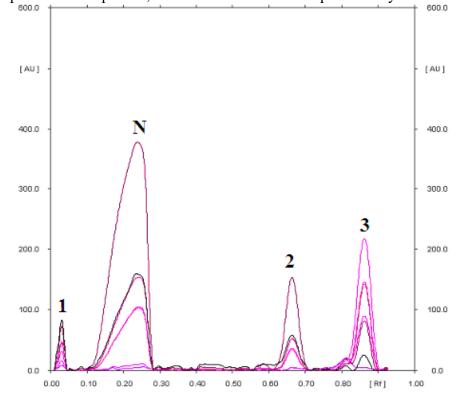


Fig. S40. Densitogram of naproxen (N) in water at pH \approx 8.50 (V), which was irradiated to UV light (254 nm) for 10 h (mobile phase A: TOL - ACE - CHL; 2:5:12, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

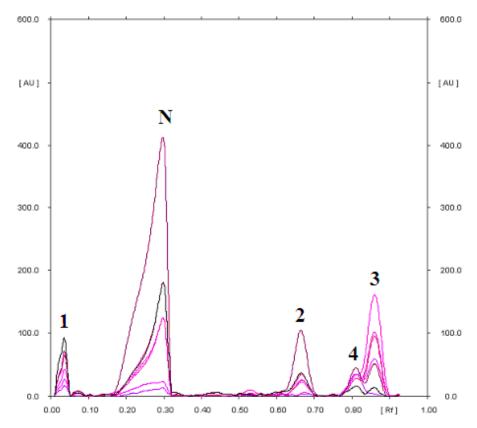


Fig. S41. Densitogram of naproxen (N) in hydrogen peroxide (VI), which was irradiated to UV light (254 nm) for 10 h (mobile phase A: TOL - ACE - CHL; 2:5:12, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

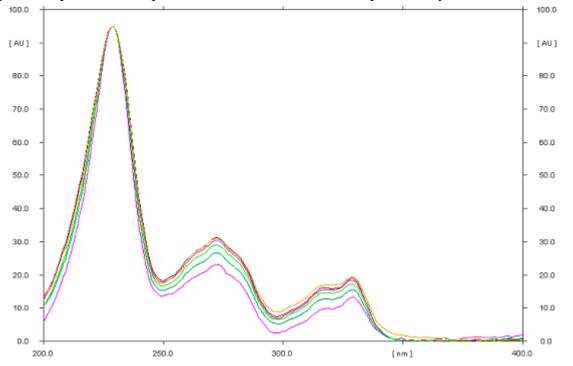


Fig. S42. Comparison of naproxen spectra ($R_F = 0.26$), which in solutions II, III, IV, V, VI was irradiated from above for 10 h and naproxen standard (mobile phase A: TOL - ACE - CHL, 2:5:12, v/v/v).

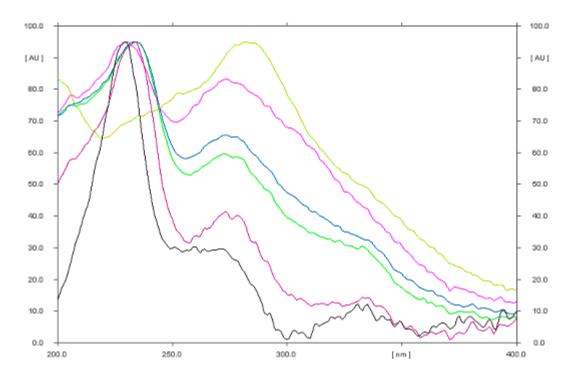


Fig. S43. Spectra of substance 1 with $R_F = 0.03$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase A: TOL - ACE - CHL, 2:5:12, v/v/v).

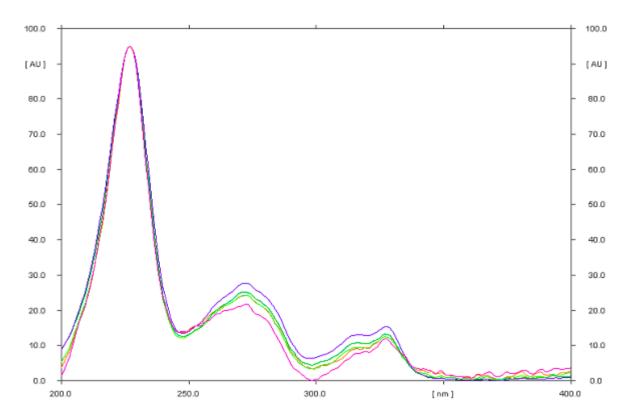


Fig. S44. Spectra of substance 2 (naproxen ethyl ester) with $R_F = 0.67$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase A: TOL - ACE - CHL, 2:5:12, v/v/v).

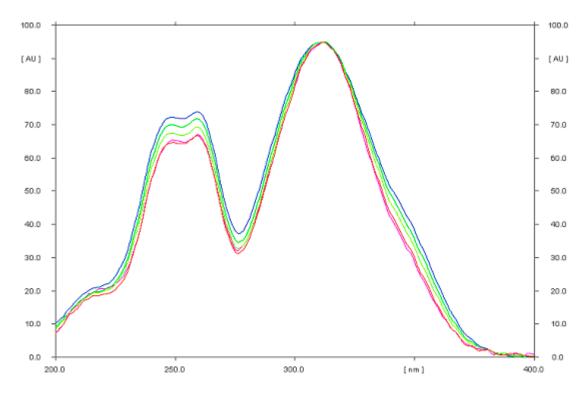


Fig. S45. Spectra of substance 3 with $R_F = 0.86$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase A: TOL - ACE - CHL, 2:5:12, v/v/v).

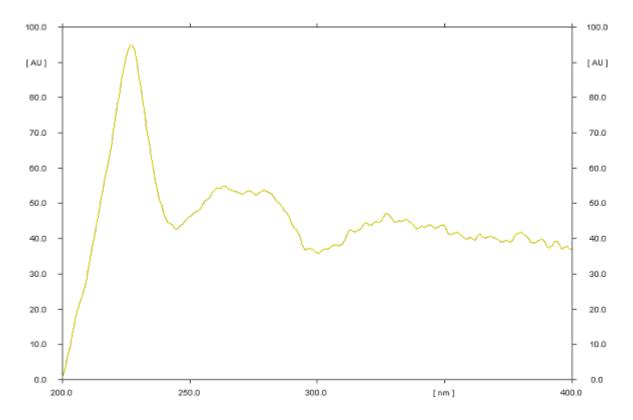


Fig. S46. Spectra of substance 4 with $R_F = 0.81$ formed after irradiation of naproxen in solution VI for 10 h (mobile phase A: TOL - ACE - CHL, 2:5:12, v/v/v).

Solution	Exposure time	Substance ^{a)}	R _F	Chromatographic peak area [AU] ^{b)}	% of band area
Ι	0 h	Ν	0.28	42457	100
		Ν	0.27	41896	95.72
	- 1	S1	0.03	271	0.62
	5 h	S2	0.65	598	1.37
		S3	0.85	1005	2.29
II		N	0.25	34277	83.05
11	101	S1	0.02	840	2.03
	10 h	S2	0.67	2150	5.21
		S3	0.87	4008	9.71
		N	0.26	37306	81.13
	5 1	S1	0.04	596	1.30
	5 h	S2	0.65	2889	6.28
		S3	0.85	5194	11.29
III		N	0.24	9513	27.64
111	101	S1	0.03	1376	4.00
	10 h	S2	0.66	10538	30.61
		S3	0.86	12993	37.75
		N	0.25	41363	89.72
	- 1	S1	0.04	1123	2.44
	5 h	S2	0.65	1204	2.61
		S3	0.85	2410	5.23
IV		N	0.24	20603	51.39
1 v	101	S1	0.03	1827	4.56
	10 h	S2	0.66	7174	17.90
		S3	0.86	10485	26.15
		N	0.26	40976	91.73
	- 1	S1	0.04	782	1.75
	5 h	S2	0.66	1079	2.41
		S3	0.85	1834	4.11
V		Ν	0.27	27115	66.26
	10 h	S1	0.03	1561	3.81
		S2	0.66	4637	11.33
		S3	0.86	7609	18.60
		N	0.31	30519	73.68
		S1	0.03	887	2.14
	5 h	S2	0.66	2832	6.84
		S3	0.85	5682	13.72
		S4	0.81	1502	3.62
		Ν	0.29	20844	62.57
VI		S1	0.03	1885	5.66
V I	10 h	S2	0.67	3297	9.90
		S3	0.86	6061	18.19
		S4	0.81	1226	3.68

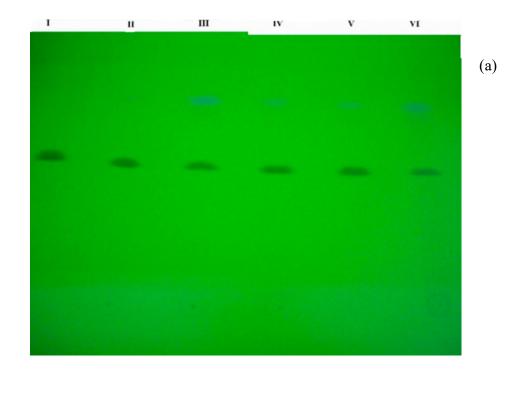
Table S3. R_F values and chromatographic peak areas of naproxen and its chemical transformation products formed in individual solutions irradiated from the top by UV light (254 nm), after separation using a mobile phase A: TOL - ACE - CHL 2:5:12 (v/v/v).

where:

 $^{a)}\,N-naproxen$

S1, S2, S3, S4 - products of chemical transformation of naproxen; and S2 was identified as naproxen ethyl ester

 $^{b)}$ Area of the chromatographic band at λ_{max}



I II III IV V VI



Fig. S47. Photographs of chromatograms, naproxen not exposed (I) and naproxen in solutions II, III, IV, V and VI irradiated for 5 h with UV radiation $\lambda = 254$ nm taken at 254 nm (a) and 366 nm (b), after separation using a mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v).

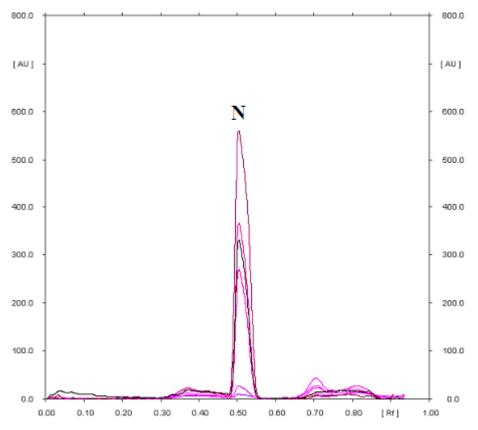


Fig. S48. Densitogram of naproxen standard (not exposed to UV) developed using mobile phase B: AcOH - Hex - ACE (0.10:10:10, v/v/v).

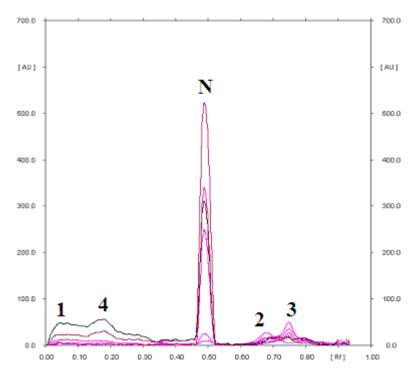


Fig. S49. Densitogram of naproxen (N) in solution of physiological salt (II), which was irradiated to UV light (254 nm) for 5h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

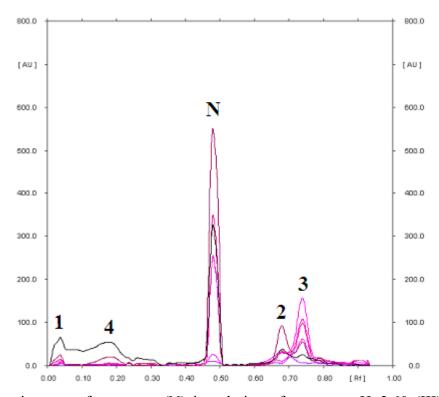


Fig. S50. Densitogram of naproxen (N) in solution of water at $pH\approx2.60$ (III), which was irradiated to UV light (254 nm) for 5h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

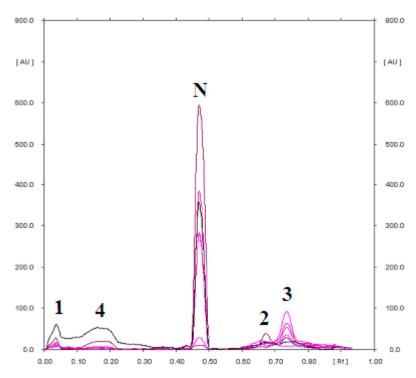


Fig. S51. Densitogram of naproxen (N) in solution of water at $pH\approx5.70$ (IV), which was irradiated to UV light (254 nm) for 5h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

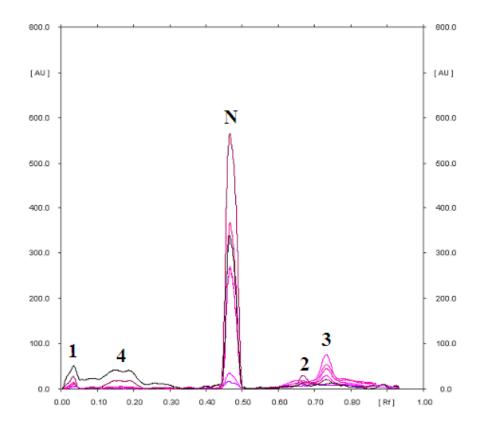


Fig. S52. Densitogram of naproxen (N) in solution of water at $pH\approx8.50$ (V), which was irradiated to UV light (254 nm) for 5h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

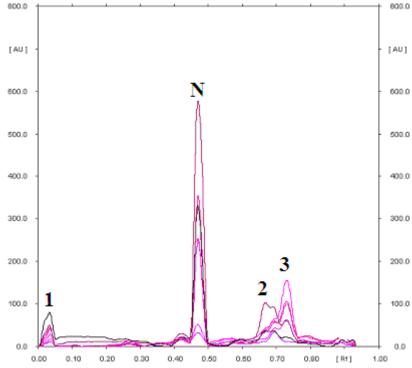


Fig. S53. Densitogram of naproxen (N) in hydrogen peroxide (VI), which was irradiated to UV light (254 nm) for 5h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

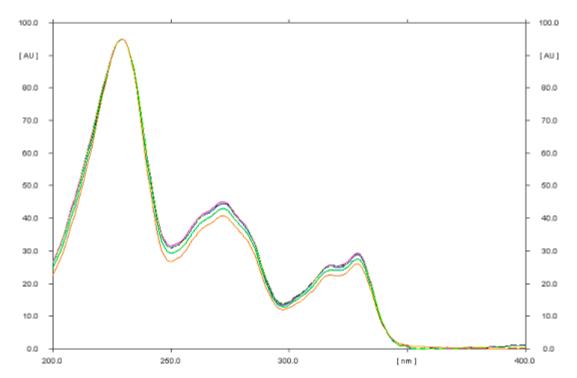


Fig. S54. Comparison of naproxen spectra ($R_F = 0.50$), which in solutions II, III, IV, V, VI was irradiated from above for 5 hours and naproxen standard (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v).

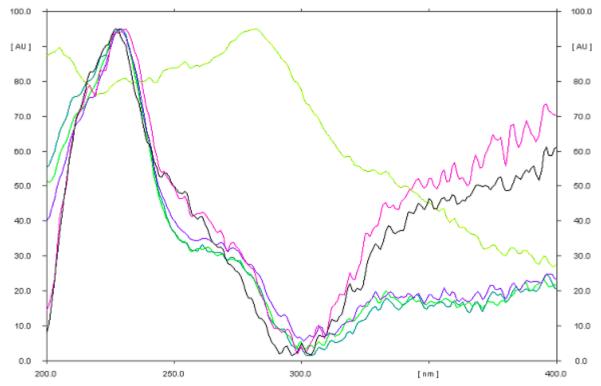


Fig. S55. Spectra of substance 1 with $R_F = 0.04$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v).

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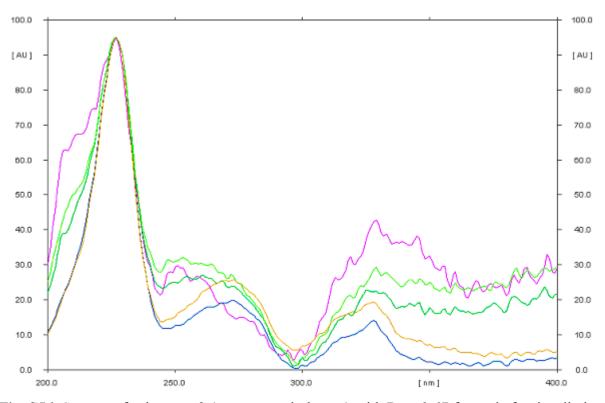


Fig. S56. Spectra of substance 2 (naproxen ethyl ester) with $R_F = 0.67$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v).

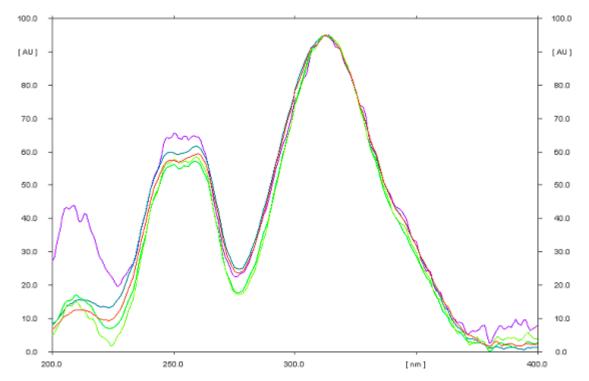


Fig. S57. Spectra of substance 3 with $R_F = 0.74$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v).

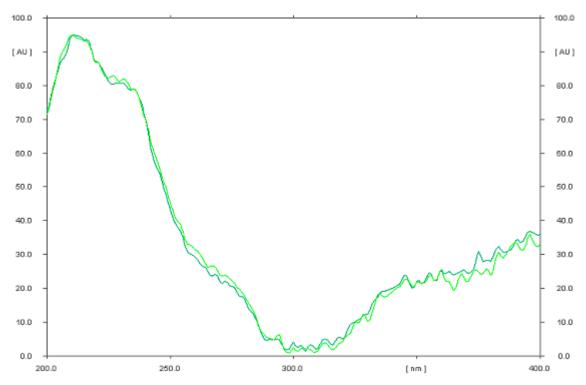
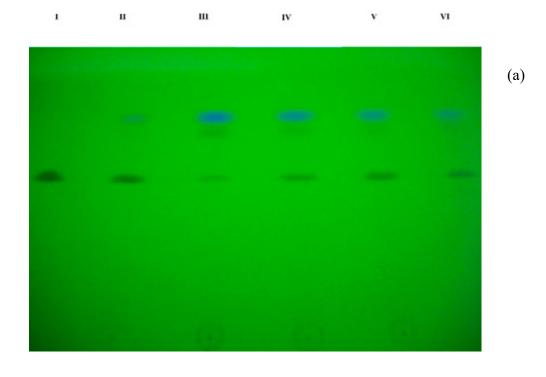


Fig. S58. Spectra of substance 4 with $R_F = 0.19$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 5 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v).



I II III IV V VI

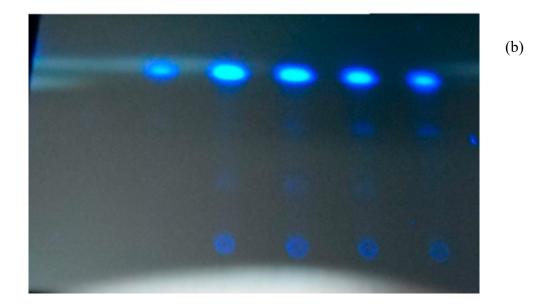


Fig. S59. Photographs of chromatograms, naproxen not exposed (I) and naproxen in solutions II, III, IV, V and VI irradiated for 10 h with UV radiation $\lambda = 254$ nm taken at 254 nm (a) and 366 nm (b), after separation using a mobile phase B: AcOH - Hex - ACE, 0.10:10:10 (v/v/v).

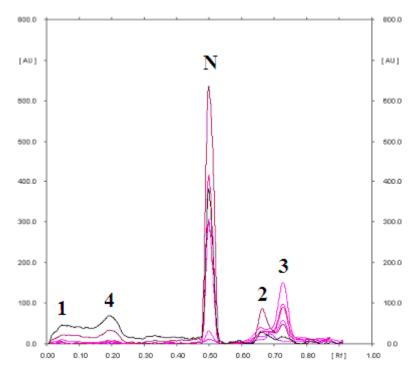


Fig. S60. Densitogram of naproxen (N) in solution of physiological salt (II), which was irradiated to UV light (254 nm) for 10h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

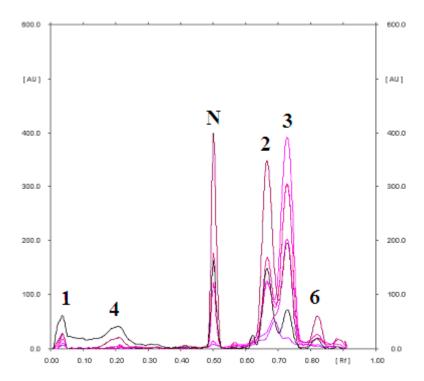


Fig. S61. Densitogram of naproxen (N) in solution of water at $pH\approx2.60$ (III), which was irradiated to UV light (254 nm) for 10h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4,6 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

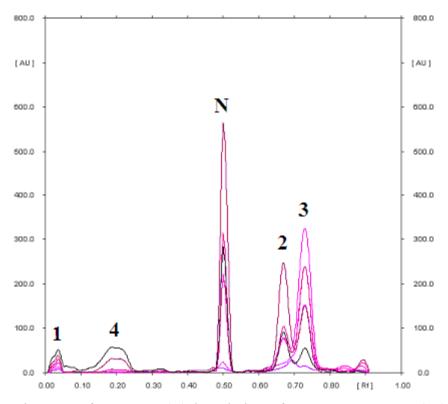


Fig. S62. Densitogram of naproxen (N) in solution of water at pH \approx 5.70 (IV), which was irradiated to UV light (254 nm) for 10h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

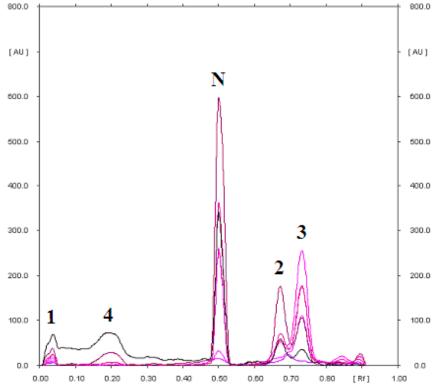


Fig. S63. Densitogram of naproxen (N) in solution of water at $pH\approx8.50$ (V), which was irradiated to UV light (254 nm) for 10h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,4 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

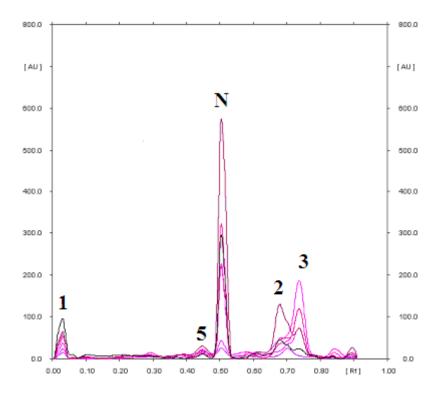


Fig. S64. Densitogram of naproxen (N) in hydrogen peroxide (VI), which was irradiated to UV light (254 nm) for 10h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v); 1, 2, 3,5 – degradation products of naproxen; and 2 was identified as naproxen ethyl ester.

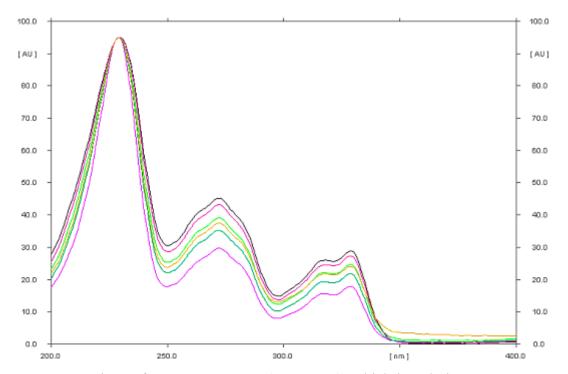


Fig. S65. Comparison of naproxen spectra ($R_F = 0.50$), which in solutions II, III, IV, V, VI was irradiated from above for 10 h and naproxen standard (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

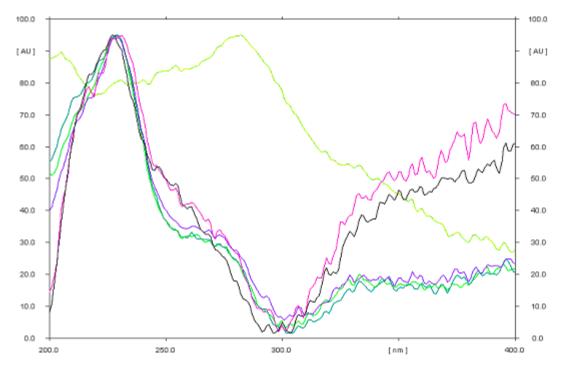


Fig. S66. Spectra of substance 1 with $R_F = 0.04$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

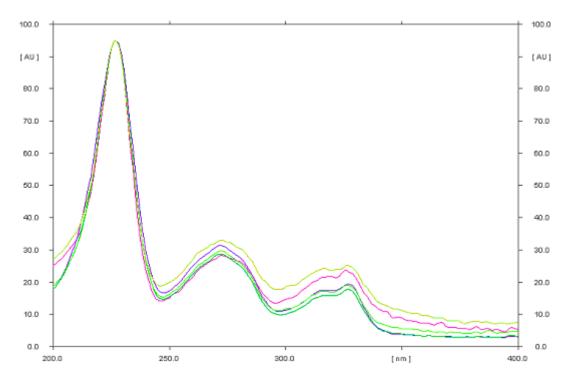


Fig. S67. Spectra of substance 2 (naproxen ethyl ester) with $R_F = 0.67$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

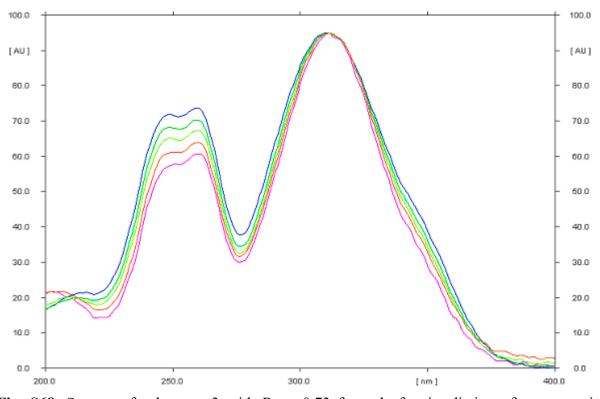


Fig. S68. Spectra of substance 3 with $R_F = 0.73$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

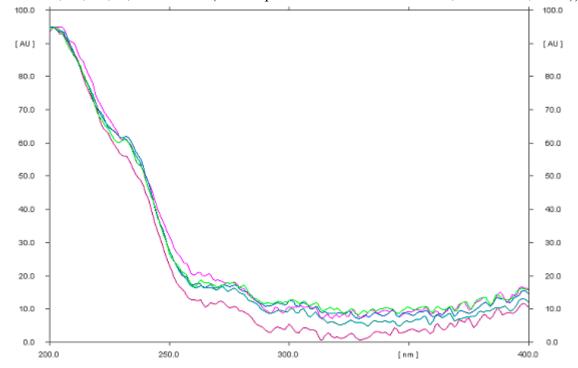


Fig. S69. Spectra of substance 4 with $R_F = 0.20$ formed after irradiation of naproxen in solutions II, III, IV, V, VI for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

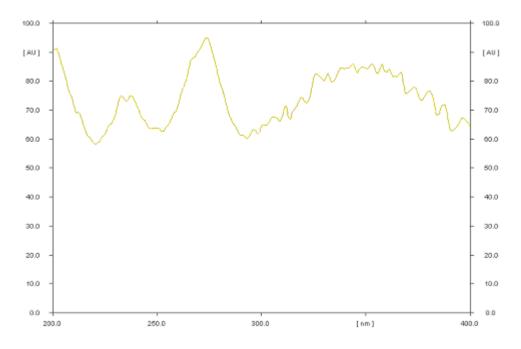


Fig. S70. Spectra of substance 5 with $R_F = 0.45$ formed after irradiation of naproxen in solution VI for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

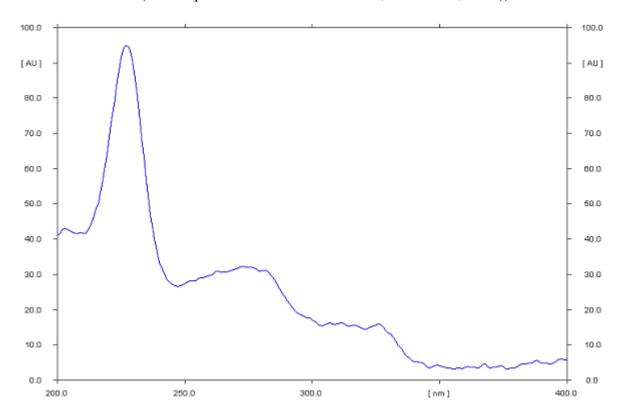


Fig. S71. Spectra of substance 6 with $R_F = 0.82$ formed after irradiation of naproxen in solution III for 10 h (mobile phase B: AcOH - Hex - ACE, 0.10:10:10, v/v/v)).

Solution	Exposure time	Substance ^{a)}	R _F	Chromatographic band area [AU] ^{b)}	% of band area
Ι	0 h	Ν	0.50	17935	100
		Ν	0.49	14622	67.59
		S1	0.04	707	3.27
	5 h	S2	0.69	975	4.51
		S3	0.75	1551	7.17
		S4	0.19	3777	17.46
II		Ν	0.50	14826	58.75
11		S1	0.04	1102	4.37
		S2	0.66		10.26
	10 h	S3	0.73		19.07
		S4	0.19		7.55
		S5	_	_	-
		S6	_	- 12790 458 2739 4943 3865 5956 936 10379 12967	_
		N	0.48	12790	51.58
		S1	0.04		1.85
	5 h	S1 S2	0.68		11.05
		S2 S3	0.74		19.93
		S4	0.19		15.59
		<u>N</u>	0.50		17.91
III		S1	0.03		2.81
		S1 S2	0.03		31.20
	10 h	S2 S3	0.07		38.98
	10 11	55 S4	0.73		
		S4 S5			4.52
		S5 S6	0.82		4.58
		<u> </u>			
			0.47		60.22
	5 h	S1	0.04		2.21
	5 11	S2	0.67		5.17
		S3	0.74	- 12790 458 2739 4943 3865 5956 936 10379	12.99
		<u>S4</u>	0.18		19.50
IV		N	0.50	$\begin{array}{r} 458\\ 2739\\ 4943\\ 3865\\ 5956\\ 936\\ 10379\\ 12967\\ 1505\\ -\\ 1519\\ 14162\\ 520\\ 1216\\ 3032\\ 4586\\ 10621\\ 880\\ 6568\\ 10132\\ 1836\\ -\\ -\\ -\\ 14496\\ 468\\ 795\\ 2506\\ \end{array}$	35.36
		S1	0.04		2.93
	10 h	S2	0.67		21.87
	10 11	S3	0.73		33.73
		S4	0.20	1836	6.11
		S5	_	—	—
		S6	_	_	_
		N	0.47		74.08
	5 1.	S1	0.03		2.39
	5 h	S2	0.67		4.06
		S3	0.73		12.81
		S4	0.18	1304	6.66
V		Ν	0.50	13412	46.06
		S1	0.04	407	1.40
	101	S2	0.67	4817	16.54
	10 h	S3	0.73	7778	26.71
		S4	0.20	2705	9.29
		S5	-	-	—
		S6	_	_	_

Table S4. R_F values and chromatographic peak area of naproxen and its chemical
transformation products formed in individual solutions irradiated from the top by
UV light (254 nm), after separation using a mobile phase B: AcOH - Hex - ACE,
0.10:10:10(v/v/v).

		Ν	0.47	12874	56.27
		S1	0.03	635	2.78
	5 h	S2	0.67	2829	12.37
		S3	0.73	6539	28.58
		S4	_	_	56.27 2.78 12.37 28.58
VI		Ν	0.51	11795	47.46
		S1	0.04	525	2.11
		S2	0.68	4720	19.00
	10 h	S3	0.74	7171	28.85
		S4	_	_	_
		S5	0.45	641	2.58
		S6	_	_	_

where:

^{a)} N - naproxen
S1, S2, S3, S4, S5, S6 - products of chemical transformation of naproxen; and S2 was identified as naproxen
^{b)} Area of the chromatographic band at λ_{max} ethyl ester