Supporting Information

Expansile Nanoparticles Encapsulate Factor Quinolinone Inhibitor 1 and Accumulate in Murine Liver upon Intravenous Administration

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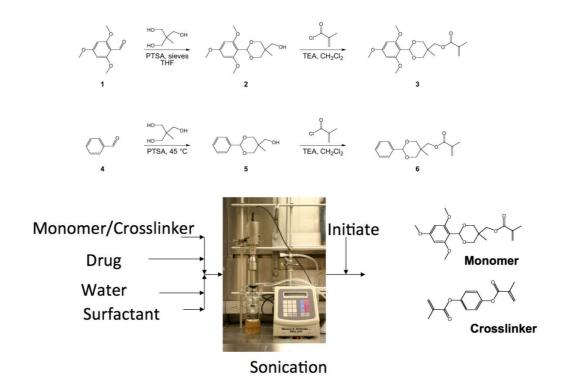
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Scheme S1. Scheme for the preparation of the polymer/nanoparticles.



			white blood cells				neutrophils				lymphocytes			monocytes			eosinophils				basophils			
	Mouse #	Treatment	result (in K/µl)	norma			result n K/µl)		natological ormalities	result (in K/µl)		nematological abnormalities	result (in K/µl)		ematological bnormalities	res (in K			atological ormalities	result (in K/μl)	normal range	hemato		
Leukocytes	1	PBS	high	high high high 1.8 -	leukocyto	sis 1	76.14	neu	utrophilia		ly	mphocytosis	9.47	mo	onocytosis	120	.45	eosi	nophilia	53.21		baso	ohilia	
	2	PBS	high		leukocyto	sis 1	35.31	neu	trophilia		lym	mphocytosis	8.73	m	onocytosis	152	.58	eosi	nophilia	85.37 71.05		basophilia		
	3	PBS	high		leukocyto	sis 1	77.98	neu	utrophilia	29.62	ly	mphocytosis	9.42	0.0 - mor 0.4 r	onocytosis	138	138.64	eosi	nophilia			baso	ohilia	
	4	eNP	high		leukocyto	sis 1	176.71 1.96 2.61	2.4 no	trophilia	0.8	oo ly	mphocytosis	0.33 0.48		onocytosis	0.12 0.2	eosinophilia normal			0.0 -	baso	ohilia		
	5	eNP	6.28						normal		9.3	normal			normal					0.0 -	nor			
	6	eNP	8.66		normal				utrophilia	4.77		normal			onocytosis		-	eosinophilia		0.24		basophilia		
	7	eNP		9.6		2.48			6.73						0.0		511-4603065		0.02		The second second			
		-	1000		normal			ıtrophilia	4.33		normal	0.32		normal	(3000		normal				normal			
	8	eNP	10.28		normal		3.72	neu	neutrophilia			normal	0.8	m	onocytosis	s 1.05		eosinophilia		0.38		basophilia		
				red blood cells				lobin		hema	tocrit	mean	comosc	ular volume	mean con	ooscular h	emoglobin	mean corposcular hem		noglobin concentration	on	red cell distribution		
				normal	hematological	7		hematologic	al result	normal	hematolog		normal	hematologic	_	normal	hematological	result normal		<u> </u>	result			
	Mouse #	Treatment	(in M/µl)	range	abnormalities					range	abnormali		range	abnormalitie			abnormalities	(in g/dl)		abnormalities	(in %)	range	abnormalities	
Erythrocytes	1	PBS	6.84		normal	high		high	38.8		norma	56.7	4.2	normal	high	1	high	high		high	18		normal	
	2	PBS	9.39		normal	high		high	50.9	35.1 - 45.4	polycythe	mia 54.2		normal	high		high	high		high	19.5 19.4 18.9 14.7		normal	
	3	PBS	8.56		normal	high		high	high 46 high 47.5 normal 50 normal 38.8		polycythe	mia 53.7		normal	high		high	high		high			normal	
	4	eNP	8.25	6.36 -	normal	high	11.0	high			. polycythe	mia 57.6	45.4 -	normal	high	14.1 -		high	high 30.2 - 30.2 34.2 28.6 30 29.8	high		12.4 -	normal	
	5	eNP	8.34	9.42	normal	15.1	15.1	normal			polycythe	mia 59.9		normal	18.1	19.3		30.2		normal		27.0	normal	
	6	eNP	6.91		normal	11.1		normal			norma	56.2		normal	16.1			28.6		low	14.6		normal	
	7	eNP	8.47		normal	14.5		normal	48.4		polycythe	mia 57.1		normal	17.1		normal	30		low	14.7		normal	
	8	eNP	9.61		polycythemia	16.1		high	54		polycythe	mia 56.2		normal	16.8		normal	29.8		low	17.1		normal	
					platelet cour	ıt		mean platelet v		rolume	olume													
	Mouse #	Treatment	result (in K/µl)		normal range			result normal (in fl) range		hematological abnormalities														
Thrombocytes	1	PBS	17	23		norn	nal	- 25 - 25	5.7		nal													
	2	PBS	17	55		normal		6.1	3.1	normal														
	3	PBS	20	77		norn	nal	5.7 6.1		norr	mal													
	4	eNP	13		592 -	norn				norr														
	5	eNP	22	22		rombocytopenia		5	.0 - 20.0	norr	mal													
	6	eNP	52	21	thi	romboci	ytopenia	6		поп	mal													
	7	eNP	25				ytopenia			non														
	8	eNP	38	34			ytopenia	222		norr	mal													

Table S1. Hematological analysis of eNP- *versus* PBS-treated CD-1 mice 24 h after injection. In eNP-injected mice numbers 5-8, little toxicity was observed with only a slight increase in neutrophil, monocyte, eosinophil, and basophil counts and minimal polycythemia; noticeably, these mice experienced thrombocytopenia. In mice numbers 1-4 (three control mice and one eNP-treated mouse), the initial tail-vein injection was inefficient, therefore resulting in a second attempt of tail vein injection; this is likely the cause for abnormalities observed in leukocyte counts and other profiles.

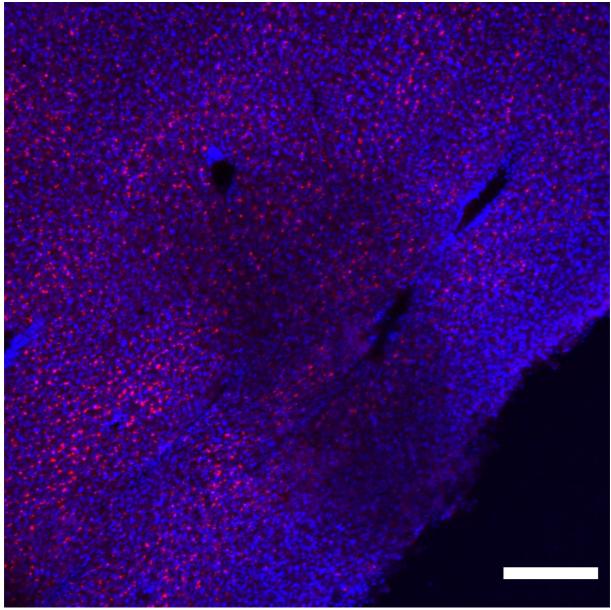


Figure S1. Accumulation of rho-eNPs in mouse liver after 48 hours. Image of liver section of CD-1 mouse treated with rho-eNP intravenously at a dosage of 62.5mg/kg. Animal was sacrificed 48 hours after injection and liver was fixed, cryosectioned and stained with Hoechst. Red and blue denote rho-eNPs and Hoechst, respectively. Scale bar is 200 μ m

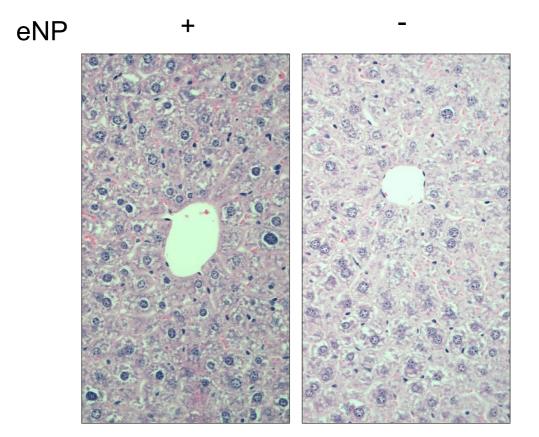


Figure S2. Higher magnification of H&E stained liver sections of eNP- versus PBS-treated CD-1 mice 24 h after injection. Images of H&E stained sections of liver of eNP- and vehicle-treated male CD-1 mice 24 hours after injection. Both treatment groups exhibited minor toxicological abnormalities, including Kupffer cell hyperplasia, enlarged or lost nuclei, and vacuolation. Because these observations occurred in both treatment conditions, it suggests that these abnormalities resulted from damage incurred by the tail-vein injection, the animal euthanasia, or the organ processing.

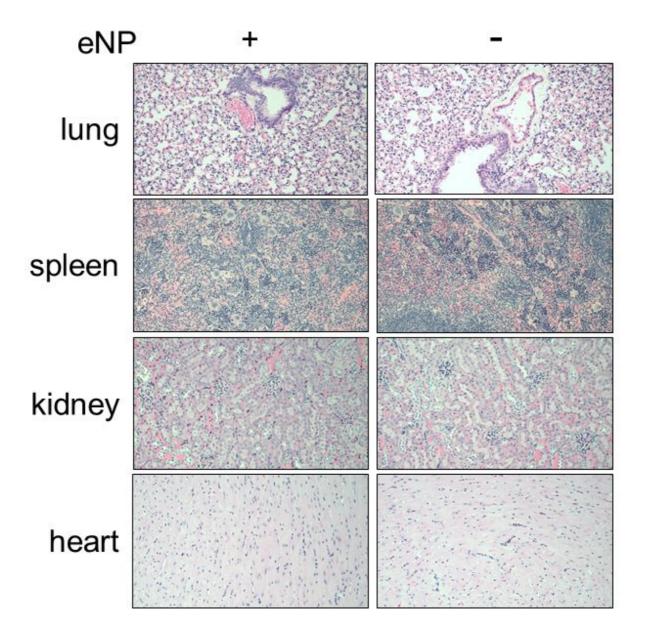


Figure S3. Pathological examination of H&E stained organs sections of eNP-versus PBS-treated CD-1 mice 24 h after injection. Images of H&E stained sections of heart, lung, kidney, and spleen of eNP- and vehicle-treated male CD-1 mice 24 hours after injection. No toxicity was detectable in treated versus control organs.