

Supporting Information

Chiral microenvironment promotes retinal progenitor cells proliferation by activating the Akt and ERK pathways

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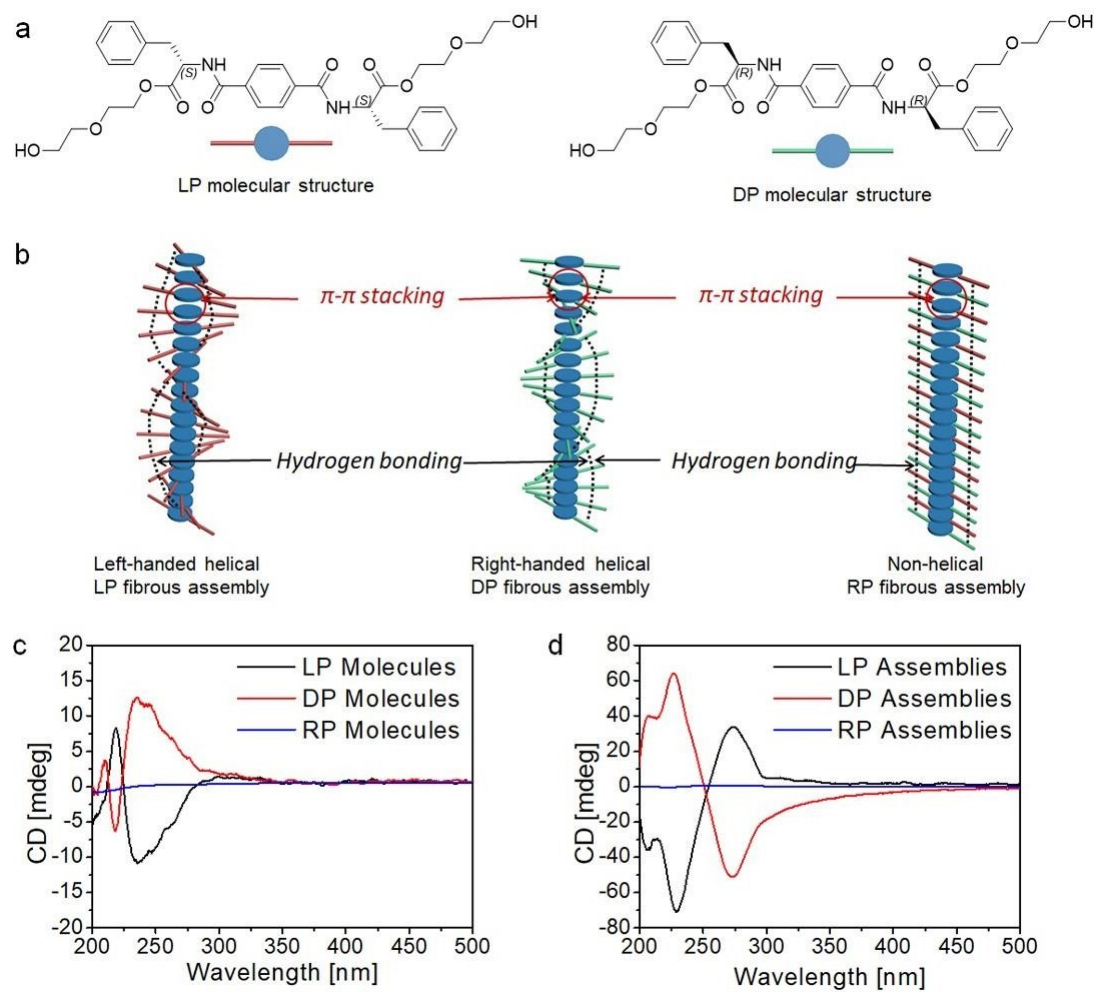


Figure S1. (a) The molecular structures of LP and DP. (b) The assembly was driven by π - π stacking and hydrogen bonding interactions. (c) CD spectra of LP, DP, and RP molecules in MeOH. (d) CD spectra of LP, DP, and RP fibrous assemblies.

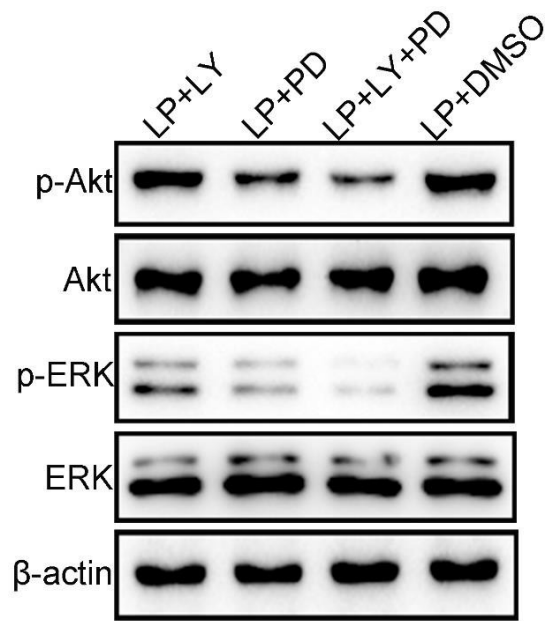


Figure S2. L-chiral nanofibrous films regulated RPC proliferation via Akt and ERK signaling pathways. RPCs were cultured with LY294002, PD98059, or LY294002 plus PD98059 in LP-treated group. Western blot assays revealed the temporal changes of phosphorylated (p-Akt, p-ERK) and total proteins.

Table 1. Primer pairs for qPCR analysis

Genes	Accession number	Forward (5'–3')	Reverse (5'–3')	Annealing Temperature (°C)
IL-6	NM_031168	aggagtggctaaggac	ataacgcactag	60
	.1	caaga	gtttgccga	
MCP-1	NM_011333	acctgctgctactcattc	attccttctgggggt	60
	.3	acc	cagca	
Annexin A5	NM_009673	tgctcaggagttaaga	taatctcggtaaat	60
	.2	ctctgttt	actttctcgtc	
β -actin	NM_007393	agccatgtacgtagcca	ctctcagctgtggt	60
		tcc	ggtgaa	
Ki-67	NM_001081	cagtactcggaatgcag	cagtctcagggg	60
	117.2	caa	ctctgtc	
Vimentin	NM_011701	tggttgacaccactca	gctttgggggtgctc	60
n	4	aaa	agttgt	

Table 2. Primary antibodies used for western blot assays

Antibodies	Type	Source	Dilution
β -actin	Mouse monoclonal	Millipore	1:5000
Vimentin	Mouse monoclonal	Millipore	1:1000
Akt	Rabbit monoclonal	CST	1:1000
Erk1/2	Rabbit monoclonal	CST	1:1000
p-Akt	Rabbit monoclonal	CST	1:1000
p-Erk1/2	Rabbit monoclonal	CST	1:1000