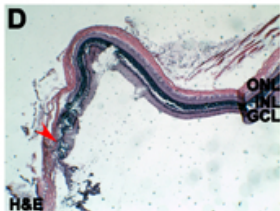
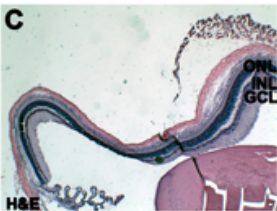
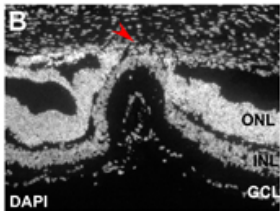
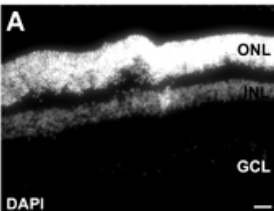


Supplemental Figure S1. FUS disrupted retinal morphology in one of six rat eyes. (A,B) DAPI (A,B) and Hematoxylin and Eosin (C,D) staining of rat retinas 30 minutes after FUS (B,D) and untreated controls (A,C). Red arrowheads in B,D point to sites of retinal dysmorphology. (Scale bars =25 μ m).

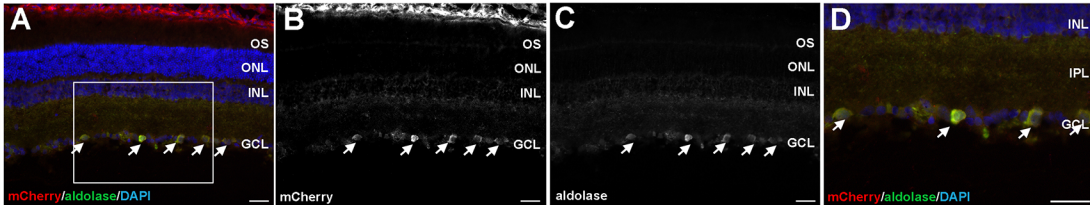
Supplemental Figure S2. FUS delivery of AAV2/8-GFAP-mCherry into retinal astrocytes. (A-D) mCherry epifluorescence in astrocyte-like cells in the nerve fiber layer (arrows, A-C). (A, C) Co-labeling of mCherry and aldolase in astrocytes in the nerve fiber layer (arrows, A-C). Blue is nuclear DAPI stain. Boxed areas in A is magnified in D. GCL, ganglion cell layer; INL, inner nuclear layer; IPL, inner plexiform layer; ONL, outer nuclear layer; OS, outer segments. (Scale bars=25 μ m).



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Tail vein injection AA2/8-GFAP-mCherry with microbubbles + FUS

GFAP promoter
in astrocytes



Touahri et al. Fig. S2

Supplemental Table 1. Pressure parameters in FUS experiments		
Animal	Enhancement	Acoustic pressures (Mpa)
#1	No enhancement	0.7- 0.538- 0.675- 0.562
#2	No enhancement	0.650- 0.562- 0.737- 0.562
#3	Medium	0.712- 0.575- 0.7- 0.675
#4	Strong	0.9- 0.825- 0.795- 0.705
#5	Weak	0.660- 0.750- 0.765- 0.840
#6	No enhancement	0.772- 0.91- 0.822- 1.295

Supplemental Table 2. List of animals used in each experiment

PROCEDURE	Number of Animals	PURPOSE
Systemic injection		
AAV9-PHP-eB	3 mice	Retinal Transduction
rAAV2/8-GFAP-mCherry 2.5×10 ⁹ GC/ml	3 mice	Peripheral Organ Transduction
PBS	3 mice	
Retinal Explants		
rAAV2/8-GFAP-mCherry 2.5×10 ⁹ GC/ml	2 mice	Retinal Transduction - in vitro
Intravitreal injection		
rAAV2/8-GFAP-mCherry	3 rat eyes	Retinal Transduction - in vivo
PBS	3 rat eyes	
MRIgFUS		
Gadolinium/Evans Blue	6 rats	Permeabilization Assay In vivo
rAAV2/8-GFAP-mCherry 2.5×10 ⁹ GC/ml	3 rats	Retinal Transduction - in vivo
rAAV2/8-GFAP-mCherry 1.25×10 ⁸ GC/ml	3 rats	