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Supplementary Information

Detection of chymase activity using the specific peptide probe conjugated onto gold nanoparticles

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Figure S1. Illustration of the AuNPs-peptide probe catalyzed by chymase. The peptide labeled with FITC was modified onto the AuNPs in advance. Due to the quenching effect of AuNPs, there was only slight fluorescent signals detected. After the treatment of chymase, the peptide substrate was catalyzed by chymase and drifted away from the surface of AuNPs. The distance-dependent quenching effect decreased and the fluorescent signal was significantly recovered.





A. H&E stain



B. Masson's trichrome stain



Figure S3. Pathologic findings of the Sham and AAI-induced nephropathy mice. The Hematoxylin-eosin (H&E) **(A)** and Masson's trichrome **(B)** stained kidney tissue sections were from the Sham mice and AAI-treated mice. Compared with the sham group, the AAI group showed severe white blood cells infiltration, intrarenal fibrosis (blue color by trichrome stained) and severe interstitial tissue damage.

Mw of	pI of	Charge of	Attribute of
peptide	peptide	peptide	peptide
1975.06	3.98	-3	Acidic
Peptide sequence		DRVYIHPFHLDDDDDC	
Cleavage site of peptide		(F) Phe - - (H) His	
Peptide treated with chymase		DRVYIHPF - -	HLDDDDDC

Table S1. The characteristics of the peptide

Table S2. The chymase activity in different tissue homogenates of
normal C57BL/6 mice $(n = 9)$

Tissue	Chymase activity	
Heart	4.25 ± 0.25 a	
Liver	$2.02\pm0.17{}^{\mathrm{a}}$	
Spleen	3.16 ± 0.53 a	
Lung	8.31 ± 1.19 a	
Kidney	5.35 ± 1.26 a	
Testis	$4.62\pm0.94~^{\rm a}$	
Brain	12.18 ± 2.38 ^a	
Serum	10.09 ± 1.60 b	

 $^{\rm a}$ Chymase activity is $\mu g/g$ protein of tissue homogenate.

^b Chymase activity is ng/mL of blood.