

Colorimetric protease activity assay method using engineered procaspase-3 enzymes

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Table S1. Primers used in this study

Oligonucleotide	Sequence (5' -> 3')
Primer 1	AAGGTTCATATGTCTGGAATATCCCTGGACAACAG
Primer 2	GTGGTGCTCGAGGTGATAAAAATAGAG
Primer 3	GTGGCCCCTGGGTGTGCGTGGTGTGATGATGACATGGCGTGTC
Primer 4	CACCACGCACACCCAGCGGGCCACAGTCCAGTTCTGTACCAC
Primer 5	ACAGGCCCTCTGGGTGTGCGTAGTGGTGTGATGATGACATGGCGTG
Primer 6	ACTACGCACACCCAGAGGGCCTGTCTCAATGCCACAGTCCAGTTC
Primer 7	ACAGGTAGCGGCCCTCTGGGTGTGCGTGGCAGTGGTGTGATGATGAC
Primer 8	ACTGCCACGCACACCCAGAGGGCCGCTACCTGTCTCAATGCCACAGTC
Primer 9	ACAGGCGGTAGCGGCCCTCTGGGTGTGCGTGGCGGTAGTGGTGTGATG
Primer 10	ACTACCGCCACGCACACCCAGAGGGCCGCTACCGCCTGTCTCAATGCCACA GTC
Primer 11	ACAGGTGGCGGTAGCGGCCCTCTGGGTGTGCGTGGTGGCGGTAGTGGTG TTGATG
Primer 12	ACTACCGCCACCACGCACACCCAGAGGGCCGCTACCGCCACCTGTCTCAAT GCCACAGTC

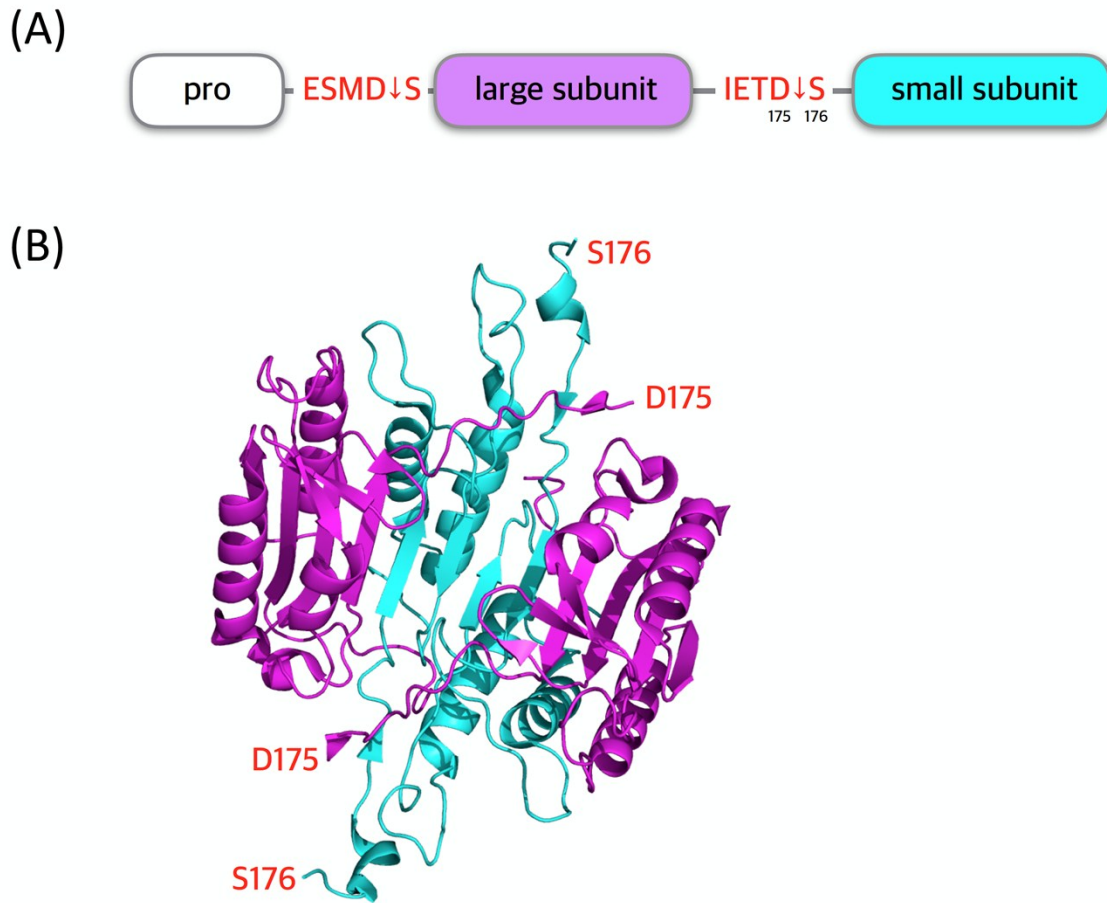


Fig. S1. (A) Structure of procaspase-3. The processing sequences in red (P4-P1' with a down arrow to denote the scissile bond) are shown in the context of caspase-3 domain structure. (B) Structure of dimeric caspase-3 (PDB: 1I3O). The processed site between the large and small subunits is shown in red (D175 and S176).