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## Product Data Sheet

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[1]. Ceuleers H, Hanning N, Heirbaut J, et al. Newly developed serine protease inhibitors decrease visceral hypersensitivity in a post-inflammatory rat model for irritable bowel syndrome[J]. British Journal of Pharmacology, 2018, 175(17): 3516-3533.

[2]. Corvo I, Cancela M, Cappetta M, et al. The major cathepsin L secreted by the invasive juvenile *Fasciola hepatica* prefers proline in the S2 subsite and can cleave collagen[J]. Molecular and biochemical parasitology, 2009, 167(1): 41-47.

### Background

Tos-Gly-Pro-Arg-AMC . HCl is a fluorescent substrate for thrombin, which is widely used to detect serine protease activity. Its excitation and emission wavelengths are 380nm and 460nm, respectively.

The core structure of Tos-Gly-Pro-Arg-AMC . HCl contains a fluorescent group 7-amino-4-methylcoumarin (AMC). When the substrate is hydrolyzed by protease, AMC is released, generating a strong fluorescent signal (excitation wavelength 380nm, emission wavelength 460nm). The Gly-Pro-Arg sequence is a specific recognition site for thrombin, so this substrate is highly selective for thrombin and is also suitable for the study of other serine proteases (such as trypsin, kallikrein, etc.). Tos-Gly-Pro-Arg-AMC . HCl can also be used for high-throughput screening of thrombin inhibitors or for detecting thrombin activity in blood samples to assist in the diagnosis of coagulation abnormalities<sup>[1][2]</sup>.

### References:

[1]. Corvo I, Cancela M, Cappetta M, et al. The major cathepsin L secreted by the invasive juvenile *Fasciola hepatica* prefers proline in the S2 subsite and can cleave collagen[J]. Molecular and biochemical parasitology, 2009, 167(1): 41-47.

[2]. Wirsching F, Luge C, Schwienhorst A. Modular design of a novel chimeric protein with combined thrombin inhibitory activity and plasminogen-activating potential[J]. Molecular genetics and metabolism, 2002, 75(3): 250-259.

**Caution: Product has not been fully validated for medical applications. For research use only.**

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