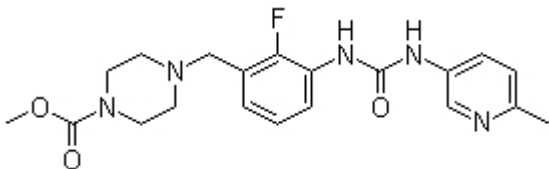


## Product Introduction

### Omecamtiv mecarbil (CK-1827452)

Omecamtiv mecarbil (CK-1827452) is a specific **cardiac myosin** activator and a clinical drug for left ventricular systolic heart failure. Phase 2.

#### Technical Data:

<b>Molecular Weight (MW):</b>	401.43	
<b>Formula:</b>	C <sub>20</sub> H <sub>24</sub> FN <sub>5</sub> O <sub>3</sub>	
<b>Solubility (25°C)</b>	DMSO 80 mg/mL	
<b>* &lt;1 mg/ml means slightly soluble or insoluble:</b>	Water 1 mg/mL	
	Ethanol 6 mg/mL	
<b>Purity:</b>	>98%	
<b>Storage:</b>	3 years -20°C Powder	
	6 months -80°C in DMSO	
<b>CAS No.:</b>	873697-71-3	

#### Biological Activity

In vitro, Omecamtiv mecarbil selectively activates cardiac myosin by increasing the myosin ATPase rate. <sup>[1]</sup>  
In isolated cardiac myocytes, Omecamtiv mecarbil results in increase of myocyte contractility and overcomes of the myosin inhibitor BDM without increasing the calcium transient or inhibiting the PDE pathway. <sup>[1]</sup>

Omecamtiv mecarbil significantly increases fractional shortening starting at 0.4 mM plasma concentrations in SD rats, sham animals and in rats with heart failure. <sup>[1]</sup> In conscious dogs with myocardial infarction

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(MI-sHF), Omecamtiv mecarbil leads to a significant increase in wall thickening (25%), stroke volume (44%), cardiac output (22%) and left ventricular (LV) systolic ejection time (26%). In addition, Omecamtiv mecarbil also results in the decreases of some hemodynamic parameters including heart rate, mean left atrial pressure, and LV end-diastolic pressure. In conscious dogs with left ventricular hypertrophy (LVH-sHF), Omecamtiv mecarbil leads to similar and not statistically different effects on hemodynamic parameters. <sup>[2]</sup>

## References

[1] Anderson RL, et al. Mol Bio Cell, 2005, 16 (Abstract #1728).

[2] Shen YT, et al. Circ Heart Fail. 2010, 3(4), 522-577.



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