

Catalog Number: CM01580

## 产品信息

**Catalog Number:**  
CM01580

**CAS号:**  
2262452-06-0

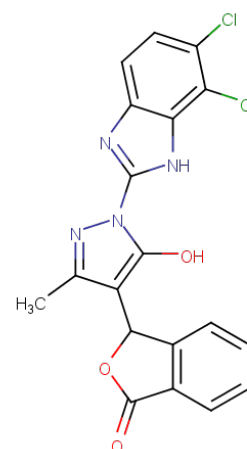
**分子式:**  
 $C_{19}H_{12}Cl_2N_4O_3$

**主要靶点:**  
cGAS|DNA

**主要通路:**  
DNA 损伤和修复|免疫与炎症

**分子量:**  
415.23

**溶解度:**  
DMSO:125 mg/mL (301.04 mM)



## 靶点活性

dsDNA:700 nM

## 体内活性

将初代BMDMs处理RU.521或其类似物，可减少小鼠IFNB1表达，表明它们在抑制缺乏细胞质DNA外切酶的细胞中固有的、构成性激活的I型干扰素表达方面的有效性。

## 动物实验

the chronically elevated levels of cytokines observed in Trex1 null mice are a consequence of constitutively activated cGAS, due to the inability to eliminate aberrantly localized self-DNA. We harvested BMDMs from 6–8-week old Trex1  $\Delta/\Delta$  mice, treated them with each compound, and measured expression levels of IFNB1 by quantitative reverse transcription PCR (qRT-PCR). Treatment of primary BMDMs with RU.521 or its analogs reduced IFNB1 expression, indicating their effectiveness in suppressing intrinsic DNA-dependent, constitutively-activated type I interferon expression in cells deficient of a cytoplasmic DNA exonuclease.

## 细胞实验

Small-molecule compounds were serially diluted to concentrations spanning the range tested in the response curves were added to  $6.7 \times 10^5$  RAW-Blue macrophages plated 16h prior in 96-well dishes, then harvested 72h after compound addition. ATP was measured using CellTiter Glo Viability Assay using  $50 \mu M$  Tamoxifen as a positive control for cytotoxicity. Viability values were generated using vehicle (DMSO) or the first dose as 100% and Tamoxifen as 0%. Outliers were removed.

## 储存

Powder: -20°C for 3 years | In solvent: -80°C for 1 year | Shipping with blue ice.