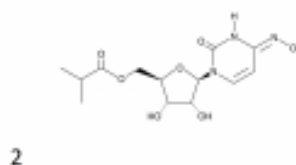
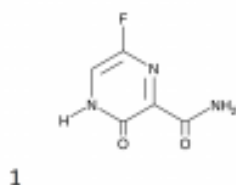
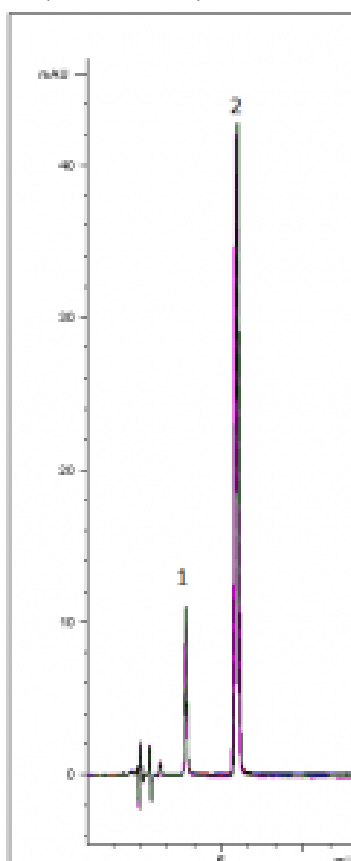


## Molnupiravir and Favipiravir Analyzed with HPLC - AppNote

### Oral Anti-Viral Medications - Easy Reversed Phase Method

Molnupiravir and Favipiravir, new anti-viral drugs were analyzed by HPLC using a simple Mobile Phase. As shown in the 10 injection overlay in the chromatogram below, the Separation, Peak Shapes and Repeatability are very good ( $\%RSD \leq 0.2$ ) .



### Peaks:

1. Favipiravir
2. Molnupiravir

### Method Conditions:

**Column:** Cogent RP C18™, 5µm, 100Å

**Catalog No.:** [68518-15P](#)

**Dimensions:** 4.6 x 150mm

**Mobile Phase:** (75:25) DI Water / Acetonitrile with 0.1% Formic Acid

**Injection Volume:** 1µL

**Flow Rate:** 1.0mL / minute

**Detection:** UV @ 254nm

**Sample Preparation:** Molnupiravir and Favipiravir are dissolved at a concentration of 0.5 mg / mL in (50:50) DI Water / Acetonitrile

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**Note :** *Molnupiravir is an oral antiviral drug that was developed for the treatment of influenza. It is a prodrug of the synthetic nucleoside derivative N4-hydroxycytidine, and exerts its antiviral action through introduction of copying errors during viral RNA replication. Favipiravir is effective against a wide range of types and subtypes of influenza viruses, including strains resistant to existing anti-influenza drugs. Of note is that favipiravir shows anti-viral activities against other RNA viruses such as arenaviruses, bunyaviruses and filoviruses, all of which are known to cause fatal hemorrhagic fever.*

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