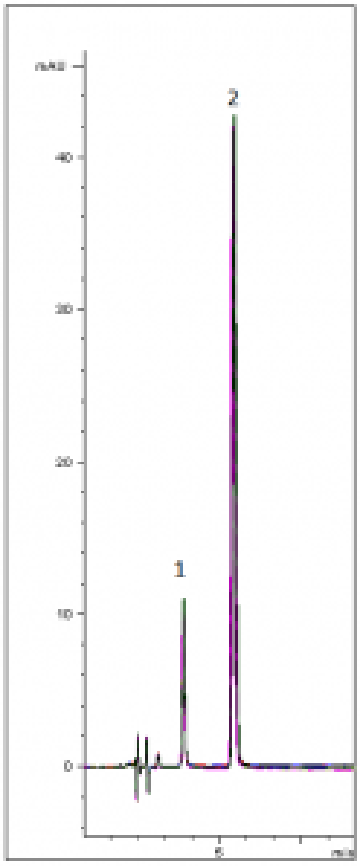
## Oral Anti-Viral Medications – Easy Reversed Phase Method

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 ≤ 0.2)*.




1 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



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**Flow Rate**: 1.0mL / minute

**Detection**: UV @ 254nm

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

***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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