**Table S3. Pharmacokinetic profiles for KY19382.**

|  |  |  |
| --- | --- | --- |
| **PK Parameters** | **IV, 1 mg/kg** | **IP, 5 mg/kg** |
| **mean** | **SD** | **mean** | **SD** |
| **tmax** (hr) | N/A | - | 1.00  | 0.00  |
| **Cmax** (ng/mL) | N/A | -　 | 463.37  | 29.41  |
| **AUClast** (ng∙hr/mL) | 7832.81  | 651.28  | 6555.79  | 572.85  |
| **CL** (L/hr/kg) | 0.12  | 0.01  | 0.47  | 0.03  |
| **Vss** (L/kg) | 0.33  | 0.07  | N/A | 　 |
| **t1/2** (hr) | 3.33  | 1.34  | 16.20  | 3.86  |
| **F** (%) | N/A | -　 | 16.74 　 | - |

Pharmacokinetic parameters were based on the mean plasma concentration-time profiles of SD male rat (n=3). Pharmacokinetic parameters were obtained by non-compartmental analysis of the plasma concentration-time profiles using KineticaTM 4.4.1 (Thermo Fisher Scientific, Inc., Woburn, MA, USA). AUClast was calculated from 0 to 24 hour. IV, intravenous; IP, intraperitoneal; Tmax, Time to maximum plasm concentration; Cmax, Maximum plasma concentration after intraperitoneal injection; AUC, Area under the curve; CL, clearance; Vss, Volume of distribution at steady state; T1/2, Elimination half-life; F, bioavailability.