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Supplement Table 1. Summary of RNA sequencing and mapping using the human genome as the reference.

| Sample | Raw reads | Clean reads | clean bases (G) | Error rate (%) | Q20 (%) | Q30 (%) | GC content (%) |
|--------|-----------|-------------|-----------------|----------------|---------|---------|----------------|
| C_1 | 99748474 | 96664918 | 14.50 | 0.01 | 97.43 | 93.53 | 47.49 |
| C_2 | 87971062 | 85654450 | 12.85 | 0.01 | 97.54 | 93.79 | 46.97 |
| C_3 | 96250922 | 92233510 | 13.84 | 0.01 | 97.66 | 94.02 | 47.52 |
| M_1 | 97517412 | 94017030 | 14.10 | 0.01 | 97.62 | 93.93 | 47.51 |
| M_2 | 88530574 | 84432966 | 12.66 | 0.01 | 97.87 | 94.36 | 47.32 |
| M_3 | 96093458 | 93303762 | 14.00 | 0.02 | 96.20 | 91.17 | 47.46 |

Raw reads: number of raw sequencing reads; Clean reads: filtered sequencing data; Clean bases: the number of branches of clean reads multiplied by the length and converted into the units of G; Error rate: error rate of the sequencing; Q20: the percentage of filtered bases was greater than 20% of the total bases; Q30: the percentage of filtered bases was greater than 30% of the total bases; GC content: the percentage of the total number of G and C bases out of the total number of bases (clean data)