**RNase Treatment During DNeasy (Qiagen) DNA Extraction**

If you choose to RNase treat during Qiagen DNeasy DNA extraction…

1. Following Step 1 incubation add 1% volume (2uL) of RNase ONE (Promega) directly to sample
2. Mix by back-pipetting
3. Incubate at 37° C for 30 minutes
4. Immediately proceed to Step 2 (addition of Buffer AL)