DNase procedure for quantitative RT-PCR RNA
-->use Rnase-free DNase

1.) Use 1 μL DNase < 20 μg RNA.

Use 2 μL DNase > 20 μg, but < 45 μg RNA.

Use 3 μL DNase > 45 μg RNA.

2.) Use 10X buffer supplied by manufacturer.

3.) Use appropriate volume, ~50 μL.

4.) 37°C 30m.

5.) Clean-up with Qiagen Rneasy.