

DeNovix dsDNA Broad Range 2 Point Assay Instructions

Technical Note 142

Introduction

The DeNovix dsDNA Broad Range Assay enables accurate detection of double-stranded DNA (dsDNA) samples with a standard detection range from 2 to 2000 ng total mass in 200 μ L volumes. This equates to sample concentrations of 0.1 ng/ μ L to 2000 ng/ μ L when using 1 to 20 μ L sample volumes in a 200 μ L total assay volume.

The upper detection limit can be extended to 4000 ng/ μ L by adding 1 μ L of a 4000 ng/ μ L sample to 199 μ L of working reagent. There is some loss of linearity with this assay when adding more than 2000 ng total mass per assay tube.

Kit Contents

Kits are available in 1000, 250, and 50 (evaluation) assays and include these components:

Component	1000	250	EVAL
DeNovix dsDNA Broad Range Dye (100x)	2 x 1 mL	0.5 mL	100 μ L
DeNovix dsDNA Broad Range Buffer	250 mL	50 mL	10 mL
DeNovix dsDNA Broad Range Enhancer (100x)	2 x 1 mL	0.5 mL	100 μ L
200 ng/ μ L dsDNA Standard (calf thymus)	2 mL	1 mL	0.5 mL
0 ng/ μ L dsDNA Standard	2 mL	1 mL	0.5 mL

Best Practices


- Use calibrated pipettes and DNase-free pipette tips.
- Prepare the working solution fresh for each assay.
- Ensure all samples and standards are treated identically in terms of incubation times and temperature.
- Avoid introducing air bubbles when mixing.
- Generate a new standard curve for each assay.
- Assay total mass must be considered when deciding how much sample to use. This assay is appropriate for 2-2000 ng total mass per tube.
- Label the top, not the sides of the assay tubes.

See *Technical Note 143* for additional information regarding solvent compatibility and multi-point curves.

Sample Prep

1. Equilibrate all solutions to room temperature before use. Vortex, then centrifuge vials briefly to minimize reagent loss on the cap.
2. Prepare working solution by mixing 10 mL of the assay buffer with 100 μ L of the dye and 100 μ L of the enhancer. Scale volumes as needed to make enough volume to aliquot 190 μ L of the mixture for each standard and unknown.
3. For each standard or unknown sample, add 190 μ L of the working solution to a labeled tube. Adjust volume when adding more or less than 10 μ L of the unknown sample.
4. Use thin-walled, clear UV-transparent 0.5 mL PCR tubes for assay measurements (DeNovix cat# TUBE-PCR-0.5-500 or equivalent).
5. Add 10 μ L of the 0 ng/ μ L and 200 ng/ μ L standards and 1-20 μ L of unknown DNA samples to the respective tubes and mix well.
6. Incubate assay tubes at room temperature for 5 minutes.

Sample Measurement

1. Launch the Fluoro dsDNA app using a DeNovix fluorometer.
2. Use the drop-down menu to select the **DeNovix dsDNA Broad Range Assay**.
3. Select **Preconfigured 2 Standards** and then choose **Generate New Standard Curve**.
4. Insert the 0 ng/ μ L dsDNA standard, lower the lid and tap **Measure**.
5. Insert the 200 ng/ μ L dsDNA standard, lower the lid and tap **Measure**.
6. After both standards are measured, tap the **Samples**  button, insert a sample tube and tap **Measure**.

Reagent Storage

Component	Protect from Light	Temperature
DeNovix dsDNA Broad Range Dye (100x)	Yes	4°C - Room Temperature
DeNovix dsDNA Broad Range Buffer	Optional	4°C - Room Temperature
DeNovix dsDNA Broad Range Enhancer (100x)	Optional	4°C - Room Temperature
dsDNA Standards	Yes	4°C

