

Product Information Sheet

## WST-8 Ready Made Solution

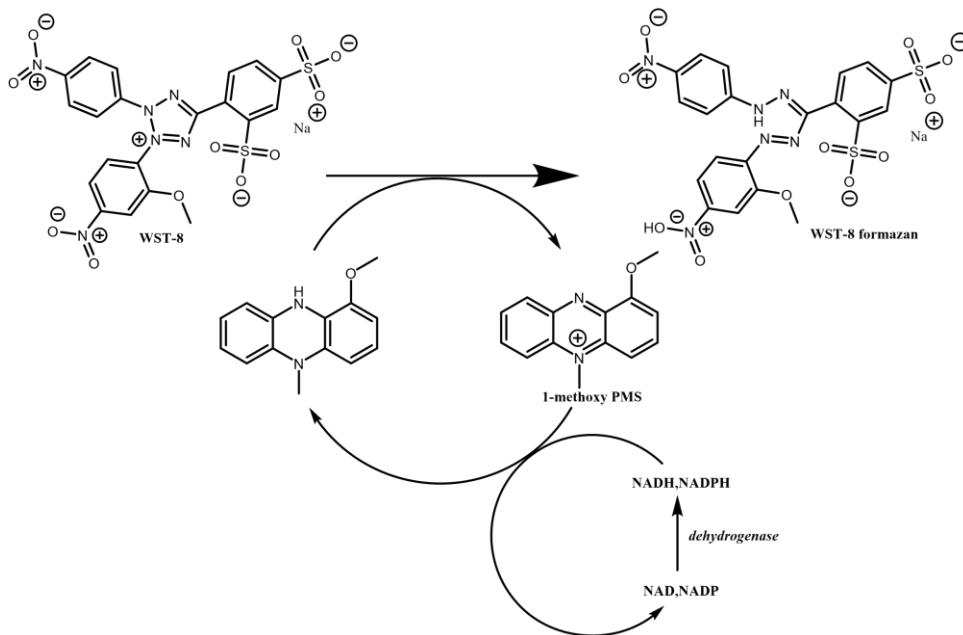
### Water-Soluble Tetrazolium 8

Catalogue Number SBR00127

### Product Description

WST-8 is a water-soluble tetrazolium salt commonly used in colorimetric assays to measure cell viability and enzyme activity.<sup>1</sup> Upon reduction, primarily by mitochondrial dehydrogenases in viable cells, it produces a yellow-orange formazan dye, which can be quantitatively measured via absorbance, reflecting the metabolic activity of the cells.<sup>2</sup>

To enhance electron transfer and boost sensitivity, 1-methoxy-5-methylphenazinium methyl sulfate (1-methoxy PMS or mPMS) is employed as an intermediate electron carrier (see Figure 1).



#### Figure 1.

Cell viability detection mechanism of WST-8 with 1-Methoxy PMS. This combination forms the foundation of the WST-8/mPMS assay, a widely used method due to its simplicity, high sensitivity, and non-radioactive nature.<sup>1-2</sup> The WST-8/mPMS system is advantageous over traditional MTT assays because the WST-8 formazan product is water-soluble, eliminating the need for organic solvents in solubilization steps.

## Applications:

- **Cell Viability and Cytotoxicity Assays:** Used extensively in drug screening, toxicology, and cancer biology to determine cell proliferation or cytotoxic effects. The pairing with mPMS ensures better electron transfer and reduces variability across assay formats.<sup>2-3</sup>
- **Dehydrogenase Enzyme Assays:** The WST-8/mPMS system efficiently monitors NAD(P)H production, making it ideal for studying enzymes such as lactate dehydrogenase and other oxidoreductases.<sup>4</sup>
- **Microbial Viability:** Rapid detection of live microorganisms through metabolic activity was demonstrated using the WST-8-mPMS assay in biosensing platforms.<sup>5</sup>
- **Comparative Studies with Other Tetrazolium Salts:** Studies have shown that WST-8, when paired with mPMS, outperforms or complements other tetrazolium-based systems (such as MTT, XTT) in terms of accuracy and biocompatibility in 2D and 3D cultures.<sup>6</sup>

## Reagents and Equipment Required but Not Provided

- 96 well flat-bottom plate
- Multiwell plate reader
- 37 °C + 5% CO<sub>2</sub> incubator
- Multichannel pipettes (10 and 100 µl)

## Precautions and Disclaimer

For R&D use only. Not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

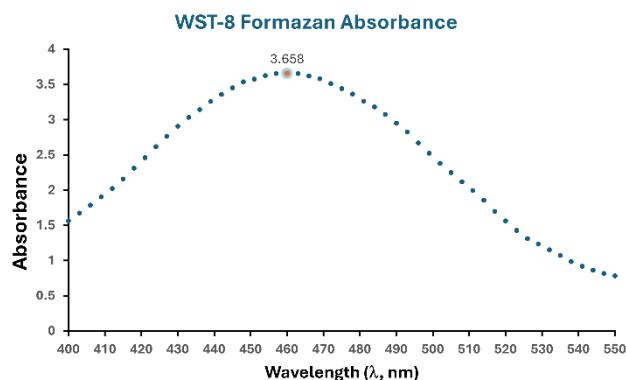
## Storage/Stability

The WST-8 RMS is stable for 4 years at 2-8 °C protected from light.

## Procedure for Cell Viability Assay

1. Grow cell suspension in a 96-well microplate (100 µl) at 37 °C in a humidified incubator with 5% CO<sub>2</sub> for the required time.
2. Add 10 µl of the WST-8 solution to the cells.
3. Incubate the plate for 0.5-4 hours at 37 °C in a humidified incubator with 5% CO<sub>2</sub>.

4. Measure the absorbance using a microplate reader at 420 to 480 nm (maximum absorption at 460 nm, see figure 2, WST-8 formazan absorbance).

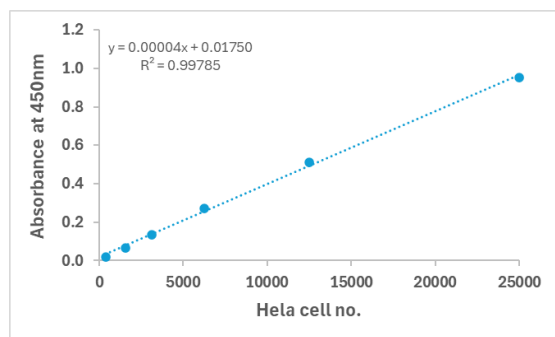


**Figure 2.**

Well plate max wavelength (λ) absorbance of WST-8 formazan analysis. The graph above illustrates the maximum absorbance peak of WST-8 solution (λ<sub>max</sub> = 460nm) after incubation (2h) with viable cells, indicating the conversion of WST-8 into soluble formazan dye.

## Results

Based on the procedure above, WST-8 solution was tested on HeLa cells ( $0.4-2.5 \times 10^4$ ) after 3 hours of incubation at 37 °C in a humidified incubator with 5% CO<sub>2</sub>.



**Figure 3.**  
WST-8 ready-made solution cell viability vs. number of HeLa cells analysis.

## References

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2. Kamiloglu, S., Sari, G., Ozdal, T., & Capanoglu, E. (2020). Guidelines for cell viability assays. *Food frontiers*, 1(3), 332-349.
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6. Lutter, A. H., Scholka, J., Richter, H., & Anderer, U. (2017). Applying XTT, WST-1, and WST-8 to human chondrocytes: A comparison of membrane-impermeable tetrazolium salts in 2D and 3D cultures. *Clinical hemorheology and microcirculation*, 67(3-4), 327-342.

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