

## RL0024

## TOS | Total Oxidant Status

## Summary and Explanation

Reactive oxygen and nitrogen species are produced in metabolic and physiological processes, and harmful oxidative reactions may occur in organisms that remove them via enzymatic and non-enzymatic antioxidative mechanisms. Under certain conditions, the increase in oxidants and decrease in antioxidants cannot be prevented, and the oxidative or in over 100 disorders, develops<sup>1</sup>. Serum (or plasma) concentrations of different oxidant species can be measured in laboratories separately, but the measurements are time-consuming, labor-intensive and costly and require complicated techniques<sup>2</sup>. Since the measurement of different oxidant molecules separately is not practical and their oxidant effects are additive, the total oxidant status (TOS) of a sample is measured and this is named total peroxide (TP)<sup>1,3,4,5</sup>, serum oxidation activity (SOA)<sup>6</sup>, reactive oxygen metabolites (ROM)<sup>7</sup> or some other synonyms.

## Principle of Assay

Oxidants present in the sample oxidize the ferrous ion–chelator complex to ferric ion. The oxidation reaction is prolonged by enhancer molecules, which are abundantly present in the reaction medium. The ferric ion makes a colored complex with chromogen in an acidic medium. The color intensity, which can be measured spectrophotometrically, is related to the total amount of oxidant molecules present in the sample. The assay is calibrated with hydrogen peroxide and the results are expressed in terms of micromolar hydrogen peroxide equivalent per liter ( $\mu\text{mol H}_2\text{O}_2$  Equiv./L)<sup>8</sup>.

## Reagent Composition

| Content                                  | Concentration        |
|------------------------------------------|----------------------|
| Reagent 1 Buffer Solution                |                      |
| H <sub>2</sub> SO <sub>4</sub>           | 25mM pH1.75          |
| Reagent 2 Substrate Solution             |                      |
| H <sub>2</sub> SO <sub>4</sub>           | 25mM pH1.75          |
| Ferrous ion                              | 5 mM                 |
| O-dianisidine                            | 10nM                 |
| Standard H <sub>2</sub> O <sub>2</sub>   | 10 $\mu\text{mol/L}$ |
| QC Level 1 H <sub>2</sub> O <sub>2</sub> | 5 $\mu\text{mol/L}$  |
| QC Level 2 H <sub>2</sub> O <sub>2</sub> | 20 $\mu\text{mol/L}$ |

## Storage/Stability

The kit is shipped on wet ice and storage at 2–8 °C is recommended. Stable up to expiry date when stored capped and at 2–8 °C even after start using.

## Normal Range

Human Serum: 4.00 – 6.00  $\mu\text{mol/L}$  (400 – 600  $\mu\text{mol/hL}$ )

Each laboratory is recommended to establish their own reference values.

## Performance Characteristics

## Precision

Inter-assay coefficient of variation 3.2%

Intra-assay coefficient of variation 3.9%

## Assay Range

Samples containing 0.2 – 80  $\mu\text{mol H}_2\text{O}_2$  Equiv./L can be assayed without further dilution or concentration.

## Interferences

EDTA interfere with the results.

## Sample

Blood serum, heparinised plasma, semen plasma, saliva, urine, cell lysates and tissue homogenates can be used as sample.

Serum samples are stable up to 1 week stored at 4°C, 6 months at -20°C and 1 year at -80°C.

## Safety Precautions and Warnings

1. For in vitro diagnostic use only.
2. Do not pipette by mouth.
3. Exercise the normal precautions required for handling laboratory reagents.
4. Wear disposable gloves while handling the kit reagents and wash hands thoroughly afterwards.
5. Do not use reagents beyond the expiry date.
6. The reagents must be used only for the purpose intended by suitably qualified laboratory personnel, under appropriate laboratory conditions.
7. Dispose cleaning liquid and also such used washing cloth or tissue paper with care, as they may also contain infectious agents.
8. Health and safety data sheets are available on request.

## Procedure

|                                              |        |
|----------------------------------------------|--------|
| Wavelength                                   | 530nm  |
| Pipette into cuvette as below order          |        |
| Sample OR Standard OR H <sub>2</sub> O       | 36 µl  |
| Reagent 1                                    | 240 µl |
| Mix well                                     |        |
| Read absorbance (A1)                         |        |
| Reagent 2                                    | 12 µl  |
| Mix well                                     |        |
| Read absorbance (A2) after 5 minutes at 37°C |        |
| OR                                           |        |
| Read absorbance (A2) after 10 minutes at RT  |        |

## Calculation

$A_2 - A_1 = \Delta\text{Abs of standard or sample}$

$$\text{Results} = \frac{\Delta\text{Abs Sample}}{\Delta\text{Abs Standard}} \times 10$$

## References

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