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Ver.251201

# Myeloperoxidase (MPO) Activity Assay Kit

BC5710 (50T/24S)

**FOR RESEARCH USE ONLY, DO NOT USE IT IN CLINICAL DIAGNOSIS**

## Product Description

Myeloperoxidase (MPO) is a white blood cell enzyme secreted by activated neutrophils and monocytes/macrophages, mainly present in the phenylalanine blue granules of neutrophils and monocytes/macrophages. When white blood cells are activated, MPO is released into phagocytic vesicles and plasma, inducing oxidative stress and tissue damage under specific conditions, and is a biomarker of systemic inflammation and oxidative stress.

MPO catalyzes the decomposition of H<sub>2</sub>O<sub>2</sub>, while oxidizing ortho anisidine to form a colored substance. There is a characteristic absorption peak at 460nm, and the color depth is linearly related to the activity of MPO.

## Kit components

Reagent I	Volume	Storage
Reagent I	60mL	2-8°C
Reagent II	Powder x 2	2-8°C
Reagent III	4mL	2-8°C
Reagent IV	1.5mL	2-8°C
Reagent V	100µL	2-8°C
Reagent VI	1mL	2-8°C

## Reagent Preparation

**Reagent II** :15mL Reagent I before use and dissolve completely at 37°C. It could be stored at 2-8°C for twelve weeks

**Reagent III**: At low temperatures, the reagent may precipitate and solidify. It need to be heated to a clear and transparent state at 37 °C before use

**working solution**: Reagent I, Reagent IV and Reagent V are mixed by the ratio of 5.28mL : 220µL : 5.5µL (5.5mL, 6S) to make working solution according to sample number before use.

## Reagents and Equipment Required but Not Provided

Spectrophotometer, centrifuge, balance, water bath/constant temperature incubator, 1mL glass cuvette, transferpettor, mortar/homogenizer/cell ultrasonic crusher, ice and distilled water.

## Operation Procedures

**Sample Preparation (The sample size to be tested can be adjusted appropriately, and the specific proportion can be referred to in the literature)**

**1. Tissue**: According to the proportion of tissue weight (g): Reagent II volume (mL) of 1:5-10 to extract. It is suggested that 0.1 g of tissue with 1 mL of Reagent II and fully homogenized on ice bath. Centrifuge at 12000g for 10 minutes at 4°C to remove insoluble materials and take the supernatant on ice before testing.

**2. Cells**: According to the proportion of cells number (10<sup>6</sup>): Reagent II volume (mL) of 5-10:1 to extract. It is suggested that add 1 mL of Reagent II to 5 million of cells. Use ultrasonication to split cells (place on ice, ultrasonic power 200W, working time 3 seconds, interval 7seconds, repeat for 3 minutes). Centrifuge at 12000g for 10 minutes at 4°C to remove insoluble materials and take the supernatant on ice for testing.

**3. Liquid samples**: According to the proportion of liquid volume (mL): Reagent II volume (mL) of 1:1 to mix (equivalent to diluting twice). Centrifuge at 12000g for 10 minutes at 4°C to remove insoluble materials and take the supernatant on ice for testing. (If the measured value is small, the proportion of liquid volume (mL) and Reagent II volume (mL) can be adjusted, such as 2:1 or 3:1

## 2.Determination

I. Preheat spectrophotometer for 30 min, adjust the wavelength to 460 nm and set counter to zero with distilled water.

II. Add reagents in 2ml EP tube as the following:

Reagent (μL)	Test tube (T)	Control tube (C)
Sample	60	60
Reagent III	60	60
Working solution	900	-
Distilled water	-	900
Mix and react at 37 °C for 30 minutes.		
Reagent IV	15	15

Mix and react for 10min at 60°C. Taking 1mL reaction solution in 1mL glass cuvette to measure the absorbance values of each tube at 460nm and record it as A<sub>T</sub> or A<sub>C</sub>. ΔA=A<sub>T</sub>-A<sub>C</sub>. Each test tube needs to be equipped with a control tube.

Note: When the temperature is low, the reaction solution may become turbid or solidify. It needs to be heated at 37°C until the reaction solution is clear and transparent before measurement.

## Calculation

- Protein concentration:**

Definition of unit: Each mg of tissue protein per hour catalyzes the production of 1μmol oxidized ortho anisidine at 37°C is defined as an enzyme activity unit.

$$\text{MPO activity (U/mg prot)} = \Delta A \div (\epsilon \times d) \times V_{RV} \div (C_{pr} \times V_S) \div T = 3.053 \times \Delta A \div C_{pr}$$

- Sample weight:**

Definition of unit: Each g of tissue per hour catalyzes the production of 1μmol oxidized ortho anisidine at 37°C is defined as an enzyme activity unit.

$$\text{MPO activity (U/g weight)} = \Delta A \div (\epsilon \times d) \times V_{RV} \div (W \div V_T \times V_S) \div T = 3.053 \times \Delta A \div W$$

- Cells:**

Definition of unit: Each 10<sup>6</sup> Cells per hour catalyzes the production of 1μmol oxidized ortho anisidine at 37°C is defined as an enzyme activity unit.

$$\text{MPO activity (U/10}^6 \text{ cell)} = \Delta A \div (\epsilon \times d) \times V_{RV} \div (N \div V_T \times V_S) \div T = 3.053 \times \Delta A \div N$$

- Liquid:**

Definition of unit: Each mL of liquid per hour catalyzes the production of 1μmol oxidized ortho anisidine at 37°C is defined as an enzyme activity unit.

$$\text{MPO activity (U/mL volume)} = \Delta A \div (\epsilon \times d) \times V_{RV} \div V_S \times F \div T = 6.106 \times \Delta A$$

**ε** : Extinction coefficient of oxidized ortho anisidine, 11.3mL/μmol/cm;

**d** : Optical path, 1cm;

**V<sub>RV</sub>**: Total reaction volume, 1.035mL;

**V<sub>S</sub>**: Add sample volume, 0.06mL;

**V<sub>T</sub>**: The volume of reagent II added during tissue/cell sample processing, 1mL;

**F** : Dilution ratio of liquid pre-treatment, 2;

**C<sub>pr</sub>** : Sample protein concentration, mg/mL;

**W**: Sample weight, g;

**N**: Cell number, calculated as 10<sup>6</sup> ;

**T**: Enzymatic reaction time, 30min=0.5h.

**NOTE:**

1. It is recommended to check whether reagent II and reagent III are clear and transparent before each experiment. If not, they need to be heated at 37 °C until the solution is clear and transparent before use.
2. If  $\Delta A$  is less than 0.01 or the absorbance value of the test tube is close to the control tube, it is recommended to increase sample supernatant volume or extend the enzymatic reaction time at 37°C before determination. If  $\Delta A$  is more than 0.5 or the absorbance value of the test tube is more than 1.5, it is recommended to dilute the sample supernatant with Reagent II before determination. And modify the calculation formula