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## Product Datasheet

### Human Heme Oxygenase 1 (HO1) ELISA Kit BYT-ORB1807471

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|--------------------------|---|
| Artikelname              | Human Heme Oxygenase 1 (HO1) ELISA Kit    |
| Artikelnummer            | BYT-ORB1807471                            |
| Hersteller Artikelnummer | orb1807471                                |
| Alternativnummer         | BYT-ORB1807471-48,BYT-ORB1807471-96       |
| Hersteller               | Biorbyt                                   |
| Kategorie                | Kits/Assays                               |
| Spezies Reaktivität      | Human                                     |
| Produktbeschreibung      | Human Heme Oxygenase 1 (HO1) ELISA Kit... |
| Detektionsbereich        | 0.31-20ng/mL                              |
| Sensitivitaet            | 0.16 ng/mL                                |
| UniProt                  | <a href="#">P09601</a>                    |
| Proben                   | serum                                     |
| Target-Kategorie         | HO1                                       |

Anwendungsbeschreibung

Application Notes: This ELISA kit uses the Sandwich-ELISA principle. The micro ELISA plate provided in this kit has been pre-coated with an antibody specific to Human HO1. Standards or samples are added to the micro ELISA plate wells and combined with the specific antibody. Then a biotinylated detection antibody specific for Human HO1 and Avidin-Horseradish Peroxidase (HRP) conjugate are added successively to each micro plate well and incubated. Free components are washed away. The substrate solution is added to each well. Only those wells that contain Human HO1, biotinylated detection antibody and Avidin-HRP conjugate will appear blue in color. The enzyme-substrate reaction is terminated by the addition of stop solution and the color turns yellow. The optical density (OD) is measured spectrophotometrically at a wavelength of 450 nm. The OD value is proportional to the concentration of Human HO1. You can calculate the concentration of Human HO1 in the samples by comparing the OD of the samples to the standard curve