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# **Product Datasheet**

### CD25 antibody (orb758826)

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| Descriptionnts.        | Rat monoclonal antibody to CD25  |
|------------------------|--|
| Species/Host           | Rat  |
| Reactivity             | Mouse  |
| Conjugation            | Unconjugated   |
| Tested<br>Applications | ELISA, FA, FC, IF, WB  |
| Immunogen              | This antibody was raised by immunising rats with<br>the B6.1 mouse cytotoxic T cell line, followed by<br>fusing spleen cells with P3X63Ag8.653 myeloma<br>cells.   |
| Target                 | CD25   |
| Preservatives          | PBS with 0.02% Proclin 300.  |
| Concentration          | 1 mg/ml  |
| Storage                | Store at 4°C for up to 3 months. For longer storage, aliquot and store at -20°C.   |
| Note                   | For research use only  |
|                        | This antibody has been used in numerous FACS<br>analyses, for instance, to demonstrate that<br>glycolysis and glutaminolysis cooperatively<br>control T cell function by limiting metabolite<br>supply to N-glycosylation (Araujo et al, 2017), to<br>suggest that IL-2C could be a potential<br>therapeutic method to alleviate excessive<br>inflammation and maintain blood vessel stability<br>after traumatic brain injury (Gao et al, 2017),<br>and to evaluate how alternative splicing of<br>MALT1 controls signalling and activation of<br>CD4(+) T cells (Meininger e al, 2016). This<br>antibody has also been used in<br>immunofluorescence as part of the DNA A $\beta$ 42<br>immunization studies in mice (Lambracht-<br>Washington et al, 2015), in ELISA assays to<br>investigate the dynamics of T cell receptor<br>(TCR)-dependent signaling networks<br>(Brockmeyer et al, 2011), and in Western Blot to<br>determine the effects of increased p300<br>expression on glucocorticoid receptor (GR)-T-<br>cell-receptor (TCR) crosstalk between<br>thymocytes (Yu et at, 2002). In addition, this<br>antibody has been used in various in vivo<br>functional studies, for instance, to suggest that a<br>combination of local anti-CTLA-4 antibody<br>production with systemic Treg depletion could<br>enhance antitumor immune responses (Tuve et |

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glomerulonephritis during the preactive phase in