

Product Information Sheet for MRA-904

Plasmodium berghei, Strain pv25DR (15cy1Pb25-/PD28-)

Catalog No. MRA-904

For research use only. Not for human use.

Contributor:

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Manufacturer:

BEI Resources

Product Description:

Protozoa Classification: *Plasmodiidae*, *Plasmodium*

Species: *Plasmodium berghei*

Strain: pv25DR

Phenotype: 15cy1Pb25-/PD28-¹

Original Source: *Plasmodium berghei* (*P. berghei*), strain pv25DR is a transgenic clone (29E), of the wild-type *P. berghei* clone 15cy1(hp), created by double knockout of endogenous *pb25* and *pb28* genes and replacement with the *pv25* gene from *P. vivax*.^{1,2} The selectable markers used were *Toxoplasma gondii* dihydrofolate reductase (TgDHFR) and pyrimethamine in water, pH 5.0.

Comments: *P. berghei*, strain pv25DR expresses the *P. vivax* antigen P25 and was developed to simplify transmission-blocking activity assays for *P. vivax* vaccine studies.^{1,2}

P. berghei is a protozoan parasite that infects mammals other than humans, especially rodents, and is commonly used in rodent model studies of malaria.³ *P. berghei* preferentially invades reticulocytes, typically producing infections in mice that induce severe pathology.⁴

Material Provided:

Each vial contains approximately 0.5 mL of *P. berghei*-infected mouse blood in Glycerolyte 57 solution (1:2). **This item is host restricted and must be amplified in rodents.** Please refer to Appendix I for cryopreservation instructions.

Packaging/Storage:

MRA-904 was packaged aseptically in cryovials. The product is provided frozen and should be stored at -80°C or colder immediately upon arrival. For long-term storage, the vapor phase of a liquid nitrogen freezer is recommended (-130°C or colder). Freeze-thaw cycles should be avoided.

Growth Conditions:⁵

In vivo, Swiss Webster mouse

Note: Some strains of mice may require dietary or drug pretreatment protocols for successful infection as *P. berghei* strains have a strong predilection for invasion of reticulocytes.

Inoculation:

1. Thaw a frozen cryovial of MRA-904 in a 35°C to 37°C water bath for approximately 2 to 3 minutes. Do not allow the vial to immerse near the cap line seal while thawing.
2. Once thawed, wipe the outside of the vial with 70% ethanol before opening. Using a 1 mL syringe equipped with a 27-gauge 1/2-inch needle, remove approximately 200 µL to 300 µL from the vial.
3. Wipe the injection site of the mouse with 70% ethanol and inject the sample intraperitoneally at 50 µL to 100 µL per mouse (approximately 3 mice for most applications).

Monitoring parasitemia:

1. Starting 3 days post-inoculation, monitor the growth of parasites by tail vein bleed sampling and Giemsa-stained thin blood smear microscopy at 1- to 2-day intervals.
2. Passage the strain when the infection is at or near the first peak of parasitemia (> 5%). This will normally occur within one week of inoculation.

$$\% \text{ parasitemia} = (\text{Infected RBC} / \text{Total RBC}) \times 100$$

Passaging:

1. Anesthetize infected mice by CO₂/O₂ inhalation. Collect the blood by orbital bleeding or from the tail vein into 25 mL of cold 1× PBS-heparin anticoagulant solution (please refer to Appendix I for preparation instructions).
2. Inject the sample into each of the uninfected mice (approximately 10 mice) as described in Inoculation step #3 above.
3. Monitor parasitemia as described above and passage as needed.

Please refer to Appendix I for cryopreservation instructions.

Note: Do not directly inject freshly thawed parasites from cryopreserved stocks by the intravenous (IV) route, as these samples contain cryoprotectant, anticoagulant and may contain traces of lysed or coagulated red blood cells. Direct IV inoculation from cryopreserved stock may result in pulmonary embolism or shock in mice.

Citation:

Acknowledgment for publications should read "The following reagent was obtained through BEI Resources, NIAID, NIH: *Plasmodium berghei*, Strain pv25DR (15cy1Pb25-/PD28-), MRA-904, contributed by Robert E. Sinden."

Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the following publication: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, and National Institutes of Health. Biosafety in Microbiological and Biomedical Laboratories. 5th ed. Washington, DC: U.S. Government Printing Office, 2009; see www.cdc.gov/biosafety/publications/bmb15/index.htm.

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All blood cultures should be handled with appropriate safety precautions necessary for the handling of bloodborne pathogens. Personnel must be trained in accordance with their institutional policy regarding bloodborne pathogens.

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References:

1. Sinden, R. E., Personal Communication.
2. Ramjane, S., et al. "The Use of Transgenic *Plasmodium berghei* Expressing the *Plasmodium vivax* Antigen P25 to Determine the Transmission-Blocking Activity of Sera from Malaria Vaccine Trials." *Vaccine* 25 (2007): 886-894. PubMed: 17049690.
3. Hall, N., et al. "A Comprehensive Survey of the *Plasmodium* Life Cycle by Genomic, Transcriptomic, and Proteomic Analyses." *Science* 307 (2005): 82-86. PubMed: 15637271.
4. Otto, T. D., et al. "A Comprehensive Evaluation of Rodent Malaria Parasite Genomes and Gene Expression." *BMC Biol.* 12 (2014): 86. PubMed: 25359557.
5. Peters, W. and B. L. Robinson (1999), "Chapter 92 -- Malaria." In *Handbook of Animal Models of Infection*. Eds. O. Zak and M. Sande, Academic Press: London, pp. 757-773.
6. McCaffery, J. N., et al. "A Multi-Stage *Plasmodium vivax* Malaria Vaccine Candidate Able to Induce Long-Lived Antibody Responses Against Blood Stage Parasites and Robust Transmission-Blocking Activity." *Front. Cell. Infect. Microbiol.* 9 (2019): 135. PubMed: 31119106.

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APPENDIX I: CRYOPRESERVATION

Note: Only immature blood-stage parasites (rings) are viable by this method. Parasitemia of 1% or higher of ring-stage parasites is recommended for cryopreservation. All steps should be carried out in a biosafety cabinet under proper air flow.

1. Prepare a 1× PBS-heparin anticoagulant solution using sterile 1× PBS (pH ~ 7.2) without calcium or magnesium (ATCC® 30-2200™) adjusted to contain 30 Units/mL sterile heparin.
2. Harvest parasitized mouse blood into 25 × volume ice cold sterile 1× PBS-heparin anticoagulant solution and place on ice.
3. Centrifuge the diluted blood culture at 1000 × g for 5 minutes at 4°C.
4. Aspirate the supernatant carefully. Measure the volume of packed red blood cells using centrifuge tube graduations or standard volume controls.
5. To the volume of packed red blood cells, add dropwise one volume of cold (4°C) Glycerolyte 57 solution (Fenwal, Cat. No. 4A7831, or equivalent). Let stand for 5 minutes at room temperature.
6. Add dropwise an additional volume of cold Glycerolyte 57 solution to the pellet. Mix well and aliquot 0.5 mL into 1.5 mL sterile cryopreservation vials.
7. Place the vials in a controlled-rate freezing unit. From room temperature, cool the vials at -1°C/min to -40°C. If the freezing unit can compensate for the heat of fusion, maintain rate at -1°C/min through this phase. At -40°C, plunge vials into liquid nitrogen. Alternatively, place the vials in a Nalgene 1°C freezing container. Place the container at -80°C for 24 to 48 hours and then plunge vials into liquid nitrogen.
8. Store in either the vapor or liquid phase of a nitrogen refrigerator (-130°C or colder).