**Product Information Sheet for MRA-1183**

**PARASITE**

**MR4 Number:** MRA-1183  
**Organism:** Plasmodium falciparum  
**Strain:** SenTh026.04  
**Alt. Strain designation:** T26.04  
**Geographic Origin:** Human patient isolate, Thiès, Senegal  
**Date of isolation:** 2004  
**Genotype:** CACCGCGTTTATAAAACAAGATTTG  
*(Broad 24 SNP Bar Code, Daniels et al. Malaria Journal 7:223, 2008)*  
**Date of publication:** 2012  
**Company:** BEI Resources/MR4  
**Deposit:** Dyann Wirth, Sarah Volkman, Harvard School of Public Health; Souleymane Mboup, Daouda Ndiaye  
**Media Preparation:**  
1. To make 1 liter of incomplete medium, start with either 928 ml liquid RPMI-1640 (without NaHCO₃, without L-glutamine), or, to 900 ml Tissue Culture grade water, add 10.43 g of powdered RPMI-1640 (without NaHCO₃, without L-glutamine). The RPMI-1640 media, add 25 ml of 1 M HEPES (final = 25 mM), 27 ml 7.5% sodium bicarbonate solution (final =0.2% NaHCO₃), 10 ml of 200 mM L-glutamine (final =2 mM), 10 ml 20% Glucose (final = 20 mM, optional), 27.2 mg hypoxanthine (10 ml of 20 mM hypoxanthine stock in 1M NaOH), 0.25 ml 10mg/ml Gentamicin (final =2.5 ug/ml, optional). Add TC grade water to 1.0 L. Mix thoroughly and filter with 0.22 cm sterile filter unit. Store at 4°C. Incomplete media can be used for many applications involving wash steps during preparation of parasities for culture or assay. For 500 ml complete medium, add 50 ml of appropriate heat inactivated human serum (MR4: type A is used with washed type O blood) to 450 ml incomplete medium (final =10% serum). Store at 4°C. If necessary, filter the complete media with 0.22 cm sterile filter unit and store at 4°C. Sera tend to clog the filter unit so use a pre-filter or improved filter unit able to handle human serum. (Note: complete media has a short shelf life; use possible without taking the cells.  

**MR4 Growth and Preservation Protocols**

**Recommended Blood Type for in vitro culture:** Type O blood (washed); pooled human serum Type A or Type O recommended.  
**Growth Temperature:** 37°C  
**Media Preparation:**  
1. To make 1 liter of incomplete medium, start with either 928 ml liquid RPMI-1640 (without NaHCO₃, without L-glutamine), or, to 900 ml Tissue Culture grade water, add 10.43 g of powdered RPMI-1640 (without NaHCO₃, without L-glutamine). The RPMI-1640 media, add 25 ml of 1 M HEPES (final = 25 mM), 27 ml 7.5% sodium bicarbonate solution (final =0.2% NaHCO₃), 10 ml of 200 mM L-glutamine (final =2 mM), 10 ml 20% Glucose (final = 20 mM, optional), 27.2 mg hypoxanthine (10 ml of 20 mM hypoxanthine stock in 1M NaOH), 0.25 ml 10mg/ml Gentamicin (final =2.5 ug/ml, optional). Add TC grade water to 1.0 L. Mix thoroughly and filter with 0.22 cm sterile filter unit. Store at 4°C. Incomplete media can be used for many applications involving wash steps during preparation of parasities for culture or assay. For 500 ml complete medium, add 50 ml of appropriate heat inactivated human serum (MR4: type A is used with washed type O blood) to 450 ml incomplete medium (final =10% serum). Store at 4°C. If necessary, filter the complete media with 0.22 cm sterile filter unit and store at 4°C. Sera tend to clog the filter unit so use a pre-filter or improved filter unit able to handle human serum. (Note: complete media has a short shelf life; use possible without taking the cells.  

**Cryopreservation:**  
Only immature parasite stage (rings) are viable by this method. We recommend a parasitemia of 3% or higher of ring stage parasites for cryopreservation.  
1. Centrifuge the culture at 1000 x g for 5 min.  
2. Wash the pellet once with 10 or more volumes of incomplete RPMI-1640 media. Centrifuge at 1800 for 5 min and leave sufficient supernatant to resuspend the pellet.  
3. To the volume of packed RBCs, add slowly dropwise one volume of cold (4°C) Glycerol 57. Let stand for 5 min at room temperature.  
4. Add an additional 3 volumes of cold Glycerol 57 to the pellet dropwise. Mix well and aliquot 0.5 ml into 1.5 ml sterile cryopreservation vials.  
5. Place the samples into freezing containers (e.g., Nalgene Cryo 1° C / min Freezing Container) and store at –80°C for 24-48 hr.  
6. Transfer to liquid nitrogen for long term storage.  
**Important notes:**  
This reagent was authenticated by the contributor.  
Please contact malaria@atcc.org for any comment.

**Daily Culture Maintenance and Blood Smear:**  
Changing of the culture medium every 24 hours is required for malaria-infected erythrocyte cultures.  
1. Remove flask with infected culture from 37°C incubator and place flask onto flask warmer in the biological safety hood.  
2. Carefully aspirate the medium with a sterile unplugged Pasteur pipet attached to the vacuum line. Remove as much fluid as possible without taking the cells.  
3. Add 25 ml of sterile warm (37°C) complete medium to the flask, gently mix and aerate then quickly tighten cap of the flask and place the flask in the 37°C incubator till the next change.  
4. To make a blood film: working in the biological safety hood, aspirate 0.5 –1 ml of mixed culture with a sterile pipet and put into an eppendorf tube.  
5. Centrifuge the eppendorf tube at high speed and aspirate the supernatant. Mix the pellet and put 6 ul of the suspension on a glass slide for thick film smear or 2 ul for thin film smear. Spread the drop into a thin film with the edge of a clean glass slide. Air dry for 3 min at room temperature. Fix blood smear by rinsing it with methyl alcohol. Air dry for 3 min at room temperature. Stain blood films in 5% Giemsa solution for 15 min. Rinse with distilled water, air dry.  
6. Using light microscopy at 100X magnification determine parasitemia of culture.  

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**BEI Resources/MR4**

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Biosafety Level: 2

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In Materials and Methods “P. falciparum line Dd2 (MRA-156, MR4, ATCC® Manassas Virginia),...”. In the acknowledgment portion: “We thank MR4 for providing us with malaria parasites contributed by (name of depositor).”

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