

## ABSTRACT

### **BACKGROUND:**

Vaginal infections are common, frequently recur, and may increase women's risk for sexually transmitted infections (STIs). We tested the efficacy of a novel regimen to prevent recurrent vaginal infections.

### **METHODS:**

**Abstract Context:** In Kenya, most people use traditional medicine and medicinal plants to treat many diseases including malaria. To manage malaria, new knowledge and products are needed. Traditional herbal medicine has constituted a good basis for antimalarial lead discovery and drug development. **Objectives:** To determine in vivo antimalarial activity and brine shrimp toxicity of five medicinal plants traditionally used to treat malaria in Msambweni district, Kenya. **Materials and methods:** A 0.2 ml saline solution of 100 mg/kg aqueous crude extracts from five different plant parts were administered orally once a day and evaluated for their in vivo chemosuppressive effect using *Plasmodium berghei berghei*-infected Swiss mice for four consecutive days. Their safety was also determined using Brine shrimp lethality test: *Grewia trichocarpa* Hochst ex A. Rich (Tiliaceae) root, *Dicrostachys cinerea* (L) Wight et Am (Mimosaceae) root, *Tamarindus indica* L. (Caesalpiaceae) stem bark, *Azadirachta indica* (L) Burn. (Meliaceae) root bark, and *Acacia seyal* Del. (Mimosaceae) root. **Results:** Parasitaemia was as follows: *A. indica*, 3.1%; *D. cinerea*, 6.3%; *T. indica*, 25.1%; *A. seyal*, 27.8%; and *G. trichocarpa*, 35.8%. In terms of toxicity, *A. indica* root bark extract had an  $LC_{50}$  of 285.8  $\mu\text{g/ml}$  and was considered moderately toxic. *T. indica* stem bark extract and *G. trichocarpa* root extract had an  $LC_{50}$  of 516.4 and 545.8  $\mu\text{g/ml}$ , respectively, and were considered to be weakly toxic while *A. seyal* and *D. cinerea* root extracts had a  $LC_{50} >1000 \mu\text{g/ml}$  and were, therefore, considered to be non-toxic. **Discussion and conclusion:** All extracts had antimalarial activity that was not significant compared to chloroquine ( $p \geq 0.05$ ). No extract was toxic to the arthropod invertebrate, *Artemia salina* L. (Artemiidae) larvae, justifying the continued use of the plant parts to treat malaria.