

## Product datasheet for TA361171

### DMP1 Rabbit Polyclonal Antibody

#### Product data:

|                         |   |
|-------------------------|---|
| Product Type:           | Primary Antibodies  |
| Applications:           | WB  |
| Reactivity:             | Human   |
| Host:                   | Rabbit  |
| Clonality:              | Polyclonal  |
| Immunogen:              | The immunogen is a synthetic peptide directed towards the middle region of human DMP1   |
| Specificity:            | <b>Expected reactivity:</b> Human   |
| Formulation:            | Liquid. Purified antibody supplied in 1x PBS buffer with 0.09% (w/v) sodium azide and 2% sucrose.                                       |
| Concentration:          | lot specific  |
| Purification:           | Affinity purified   |
| Conjugation:            | Unconjugated  |
| Storage:                | For short term use, store at 2-8°C up to 1 week. For long term storage, store at -20°C in small aliquots to prevent freeze-thaw cycles. |
| Stability:              | Shelf life: one year from despatch.   |
| Predicted Protein Size: | 54 kDa  |
| Gene Name:              | dentin matrix acidic phosphoprotein 1   |
| Database Link:          | <a href="#">NP_001073380.1</a><br><a href="#">Entrez Gene 1758 Human</a><br><a href="#">Q13316-2</a>                                    |

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**Background:**

Dentin matrix acidic phosphoprotein is an extracellular matrix protein and a member of the small integrin binding ligand N-linked glycoprotein family. This protein, which is critical for proper mineralization of bone and dentin, is present in diverse cells of bone and tooth tissues. The protein contains a large number of acidic domains, multiple phosphorylation sites, a functional arg-gly-asp cell attachment sequence, and a DNA binding domain. In undifferentiated osteoblasts it is primarily a nuclear protein that regulates the expression of osteoblast-specific genes. During osteoblast maturation the protein becomes phosphorylated and is exported to the extracellular matrix, where it orchestrates mineralized matrix formation. Mutations in the gene are known to cause autosomal recessive hypophosphatemia, a disease that manifests as rickets and osteomalacia. The gene structure is conserved in mammals. Two transcript variants encoding different isoforms have been described for this gene.

**Synonyms:**

ARHP; ARHR; DMP-1

**Protein Families:**

Secreted Protein

**Product images:**


Host: Rabbit  
Target Name: DMP1  
Sample Type: Leiomyosarcoma Tumor Lysate  
Antibody Dilution: 1.0µg/ml

Host: Rabbit  
Target Name: DMP1  
Sample Tissue: Human Leiomyosarcoma Tumor lysates  
Antibody Dilution: 1ug/ml