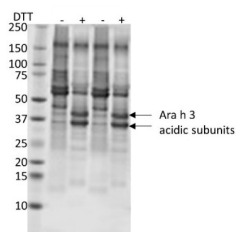


Product no **AS09 566****Anti-Peanut protein****Product information**

Anti-Peanut protein

Immunogen | *Arachis hypogaea* protein extract**Host** | Chicken**Clonality** | Polyclonal**Purity** | Immunogen affinity purified IgY in PBS pH 7.2. Contains 0.075 % sodium azide.**Format** | Liquid**Quantity** | 100 µg**Storage** | Store at -20 °C; make aliquots to avoid working with a stock. Please remember to spin the tubes briefly prior to opening them to avoid any losses that might occur from material adhering to the cap or sides of the tube.**Additional information** | Antibodies were purified on immobilized peanut proteins**Application information****Recommended dilution** | 2- 5 µg/ml (ELISA), 0,1-1 µg/ml (WB)**Confirmed reactivity** | Peanut proteins**Predicted reactivity** | Peanut proteins**Not reactive in** | No confirmed exceptions from predicted reactivity are currently known**Selected references** | [Mattison et al. \(2024\)](#). Proteomic characterization of peanut flour fermented by *Rhizopus oryzae*. Heliyon Volume 10, ISSUE 15.

Thirty (30) µg of total protein extracted freshly from defatted lightly roasted peanut flour with borate buffered saline (BBS) solution (100 mM H₃BO₄, 25 mM Na₂B₄O₇, 75 mM NaCl, and pH 8.6) for 1 hr with constant stirring at 4 °C. Samples were denatured with NuPAGE™ LDS sample buffer containing 50 mM DTT at a 1:4 (v/v) ratio and incubation at 70 °C for 5 min. Samples were separated on Novex™ 10-20% Tricine Protein Gels and blotted 7 minutes to nitrocellulose using iBlot dry transfer system. The blot was blocked with 5% milk for 1h/RT with agitation. Blot was incubated in the primary antibody at a dilution of 1: 1,000 for 1h/RT with agitation in TBS-T with agitation. The antibody solution was decanted and the blot was rinsed briefly, then washed 3 times for 5 min in TBS-T at RT with agitation. The blot was incubated in Agrisera matching secondary antibody (anti-rabbit IgG horse radish peroxidase conjugated [AS10 1489](#)) diluted to 1:25,000 in TBS-T for 1h/RT with agitation. The blot was washed as above and developed for 5 min with [AgriseraECLBright](#). Images of the blots were collected using a CCD imager and Quantity One software (Bio-Rad). Exposure time was 20 seconds.