

DESCRIPTION

Species Reactivity	Mouse/Hamster
Specificity	Detects mouse GDF-9 in direct ELISAs and both mouse and hamster GDF-9 in Western blots. In direct ELISAs, less than 1% cross-reactivity with recombinant mouse (rm) GDF-1, rmGDF-5, rmGDF-6, rmGDF-8, and rmGDF-15 is observed.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
Immunogen	<i>E. coli</i> -derived recombinant mouse GDF-9
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS.

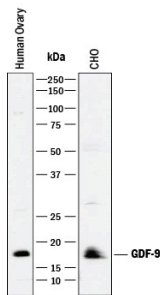
APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. *General Protocols* are available in the *Technical Information* section on our website.

	Recommended Concentration	Sample
Western Blot	1 µg/mL	See Below
Immunohistochemistry	5-15 µg/mL	See Below

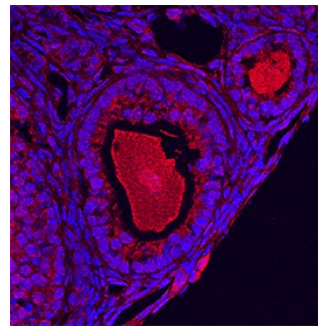
DATA

Western Blot



Detection of Mouse and Hamster GDF-9 by Western Blot. Western blot shows lysates of human ovary tissue and CHO Chinese hamster ovary cell line. PVDF membrane was probed with 1 µg/mL of Goat Anti-Mouse/Hamster GDF-9 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF739) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF017). A specific band was detected for GDF-9 at approximately 18 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 1.

Immunohistochemistry



GDF-9 in Mouse Ovary. GDF-9 was detected in perfusion fixed frozen sections of mouse ovary using Goat Anti-Mouse/Hamster GDF-9 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF739) at 5 µg/mL overnight at 4 °C. Tissue was stained using the NorthernLights™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). Specific staining was localized to developing oocytes. View our protocol for [Fluorescent IHC Staining of Frozen Tissue Sections](#).

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> ● 12 months from date of receipt, -20 to -70 °C as supplied. ● 1 month, 2 to 8 °C under sterile conditions after reconstitution. ● 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Growth differentiation factor-9 (GDF-9) is a member of the transforming growth factor- β (TGF- β) superfamily, and is an oocyte secreted paracrine factor essential for mammalian ovarian folliculogenesis (1-2). Mouse GDF-9 is synthesized as a 441 amino acid (aa) prepropeptide that contains a 29 aa signal sequence, a 277 aa propeptide, and a 135 aa mature chain. Residues 340-441 constitute a TGF- β like domain. In addition, there is one potential site of N-linked glycosylation in the mature chain. Unlike other members of the TGF- β superfamily, GDF-9 lacks the conserved cysteine residue that is believed to form the sole disulfide linkage between subunits in other family members (3). Mature mouse GDF-9 shares 90% aa sequence identity with mature human GDF 9. The protein is expressed throughout the development of the maturing follicle (2). GDF-9 functions as a paracrine factor in the regulation of granulosa cell proliferation and differentiation, and is essential for fertility (2, 4). Studies on GDF-9 null mice have demonstrated arrested follicular development at the primary follicle stage (5). Mouse GDF-9 induces Smad2 phosphorylation and inhibin production in rat diethylstilbestrol treated granulosa cells (6) and in human granulosa-luteal cells (7). The downstream signaling actions of GDF 9 are mediated by the type I receptor, activin receptor-like kinase 5 (ALK5), initiating the subsequent activation of Smad2 and Smad3 (2, 8). GDF 9 uses the BMP type II receptor (BMPRII) as its other signaling receptor (2, 9).

References:

1. McGrath, S.A. *et al.* (1995) *Mol. Endocrinol.* **9**:131.
2. Mottershead, D.G. *et al.* (2008) *Mol. Cell. Endocrinol.* **283**:58.
3. McPherron, A.C. and S.-J. Lee (1992) *J. Biol. Chem.* **268**:3444.
4. Gilchrist, R.B. *et al.* (2006) *J. Cell. Sci.* **119**:3811.
5. Dong, J. *et al.* (1996) *Nature* **383**:531.
6. Roh, J.S. *et al.* (2003) *Endocrinology* **144**:172.
7. Kaivo-Oja, N. *et al.* (2003) *J. Clin. Endocrinol. Metab.* **88**:755.
8. Mazerbourg, S. *et al.* (2004) *Mol. Endocrinol.* **18**:653.
9. Vitt, U.A. *et al.* (2002) *Biol. Reprod.* **67**:473.