

# Validation Report #029805

Validation Date: 08/26/14

## Summary

Antigen	Frizzled Family Receptor 7 (FZD7)
Catalog number	<a href="#">ABIN710051</a>
Supplier	Bioss
Supplier catalog number	<a href="#">bs-5125R</a>
Lot number	120426
Method validated	<a href="#">Western Blot</a>
Laboratory	<a href="#">Alamo Laboratories Inc</a>
Validation number	<a href="#">29805</a>
Positive Control	<a href="#">MDA-MB-231 cells</a>
Negative Control	<a href="#">C6/36 cells (non-reactive species)</a>
Notes	A strong band was observed in the positive control sample at the expected molecular weight of ~64 kDa. The band is also faintly observed in the negative control sample, which may indicate species cross-reactivity.



# Full Methods

## **Primary Antibody**

- Antigen: Frizzled Family Receptor 7 (FZD7)
- Catalog number: ABIN710051
- Supplier: Bioss
- Supplier catalog number: bs-5125R
- Lot number: 120426
- Antibody Dilution: 1:100

## **Loading Control Antibody**

- Antigen: Mouse Anti-Actin
- Supplier: BD Transduction Laboratories
- Catalog number: 612657
- Antibody Dilution: 1:6,000

## **Secondary Antibody**

- Antigen: Goat Anti-Rabbit IgG (H + L)-HRP Conjugate
- Supplier: Bio-Rad
- Catalog number: #170-6515
- Lot number: L170-6515
- Antibody Dilution: 1:10,000

## **Controls**

- Positive control: MDA-MB-231 cells
- Negative control: C6/36 cells

## **Protocol**

1. The cell extracts were heated at 95°C for 5 minutes in 1X SDS Sample Buffer containing 1% SDS and 1.25%  $\beta$ -mercaptoethanol.
2. 15  $\mu$ l of heated culture-media were loaded and resolved on 8-16% SDS-polyacrylamide gel.
3. The Thermo Scientific - Spectra Multicolor Broad Range (Cat # 26634) were used as molecular mass markers.
4. Proteins were then transferred onto PVDF membrane by wet transfer and protein transfer was confirmed with Ponceau-S staining.
5. The PVDF membrane was incubated with 25 ml of blocking buffer [Tris Buffered Saline, pH 7.4 plus 0.1% TW20 (TBST)] containing 5% (W/V) BSA at room temperature for 1 hour.
6. The membrane was rinsed with TBST once.
7. The membrane was immersed with the protein side up in the primary antibody solution in TBST containing 5% (W/V) BSA and incubated for 24 hours at 4°C.
8. The membrane was rinsed in TBST thrice for 5 minutes each.
9. The membrane was incubated in the HRP-conjugated secondary antibody solution in TBST containing 5% (W/V) BSA and incubated for 1 hour at room temperature (~26°C) with gentle agitation.
10. The membrane was rinsed thrice TBST thrice for 5 minutes each.
11. The membrane was rinsed in TBS twice for 30 seconds each.
12. Signals were detected with ECL-2 Substrate. The blot was scanned for 45 minutes.
13. The membrane was rinsed three times TBST.
14. Incubated in Acidic Glycine Stripping Buffer at room temperature with gentle agitation for 3 times, 10 minutes each.
15. The membrane was washed in TBST 2 times for 10 minutes each.
16. Repeated Steps 5-12 with the loading control antibody (for Anti-actin) and its matching secondary antibody.

## **Experimental Notes**

- No experimental challenges noted.

## Figures

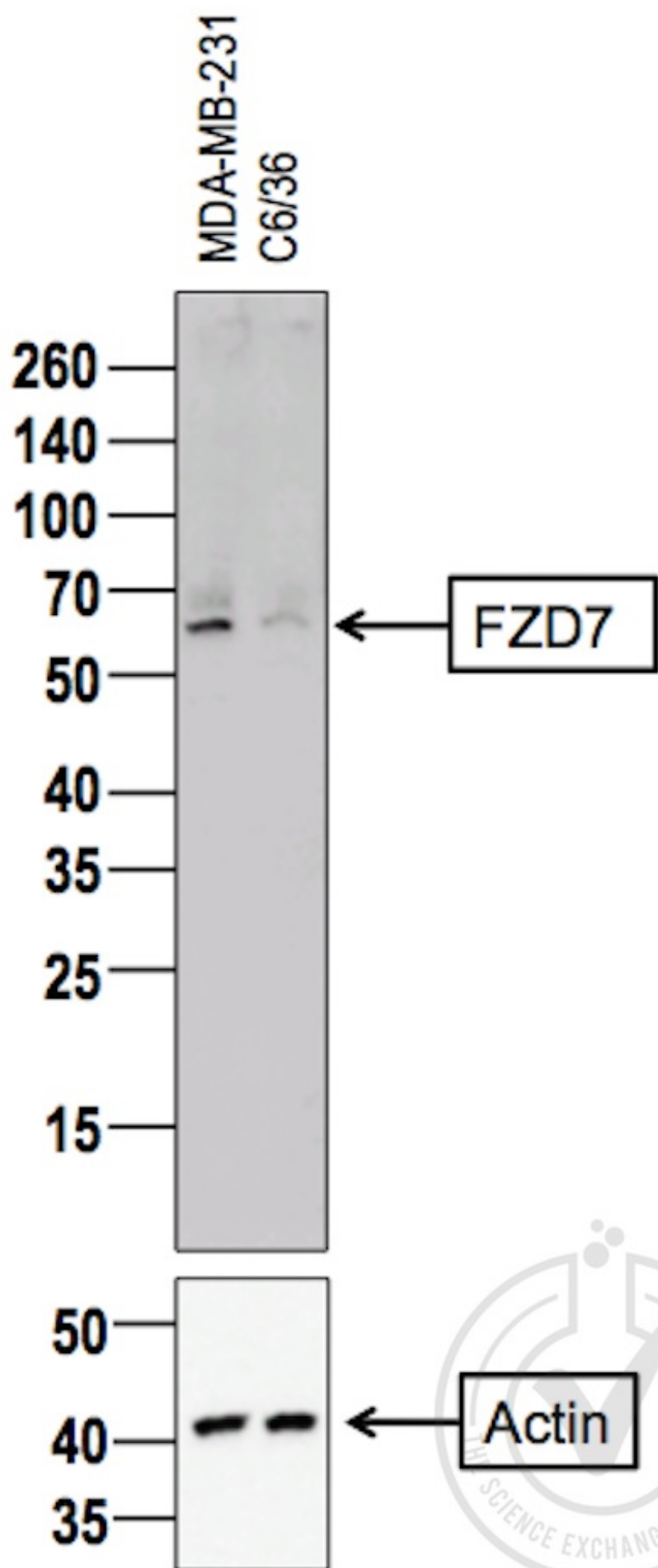


Figure 1: Western Blot for FZD7. Arrowhead indicates the expected molecular weight of ~64 kDa.