# antibodies

# Datasheet for ABIN6698839 Fluorescent TrueBlot®: Anti-Rabbit IgG DyLight<sup>™</sup> 680

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2 Im	ages
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Publications



#### Overview

Quantity:	100 µL
Target:	lgG
Reactivity:	Rabbit
Host:	Mouse
Conjugate:	DyLight 680
Application:	Flow Cytometry (FACS), Immunohistochemistry (IHC), Western Blotting (WB), FLISA,
	Fluorescence Microscopy (FM)

## Product Details

Brand:	TrueBlot®
Characteristics:	Synonyms: DL680, TrueBlot, DL680 TrueBlot ULTRA, DyLight™ 680 TrueBlot, TrueBlot for IP/WB
	TrueBlot for immunoprecipitation, TrueBlot for western blotting, Fluorescent TrueBlot, Rb
	TrueBlot
	Background: Rabbit IgG TrueBlot® is a unique DyLight™ 680 conjugated Anti-rabbit IgG
	immunoblotting (second step) reagent. Rabbit IgG TrueBlot® enables detection of
	immunoblotted target protein bands, without hindrance by interfering immunoprecipitating
	immunoglobulin heavy and light chains. It is easy to generate publication-quality IP/Fluorescent
	Western Blot data with Rabbit IgG TrueBlot $^{ m (B)}$ , simply substitute the conventional DL680 Anti-
	rabbit IgG blotting reagent with Fluorescent Rabbit TrueBlot® Antibody DyLight™ 680 and
	follow the prescribed protocol for sample preparation and immunoblotting. Rabbit IgG
	TrueBlot® is ideal for use in protocols involving immunoblotting of immunoprecipitated
	proteins. TrueBlot preferentially detects the non-reduced form of rabbit IgG over the reduced,
	SDS-denatured form of IgG. When the immunoprecipitate is fully reduced immediately prior to

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#### Product Details

	SDS-gel electrophoresis, reactivity of Rabbit IgG TrueBlot $^{\ensuremath{\mathbb{B}}}$ with the 55 kDa heavy chains and
	the 23 kDa light chains of the immunoprecipitating antibody is minimized thereby eliminating
	interference by the heavy and light chains of the immunoprecipitating antibody in IP/Western
	blot applications. Applications include studies examining post-translational modification (e.g.,
	phosphorylation or acetylation) or protein-protein interactions.
Purification:	Fluorescent Rabbit TrueBlot® Antibody DyLight™ 680 Conjugate was prepared from tissue
	culture supernatant by Protein G affinity chromatography. Assay by Immunoelectrophoresis
	resulted in a single precipitin arc against Anti-Rabbit Serum. Reactivity is observed against
	native Rabbit IgG by both Western blot and ELISA.
Labeling Ratio:	2.3

### Target Details

Target:	lgG
Abstract:	IgG Products
Target Type:	Antibody

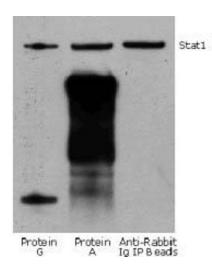
## Application Details

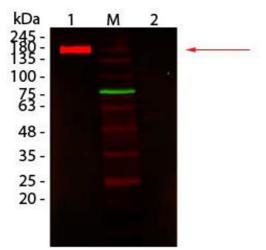
Application Notes:	Immunohistochemistry Dilution: User Optimized
	Application Note: Fluorescent Rabbit TrueBlot® Antibody DyLight™ 680 may also be used for
	detection in immunoassays that do not employ immunoprecipitation. Fluorescent Rabbit
	TrueBlot® Antibody DyLight™ 680 is provided as a lyophilized powder. To conserve reagent, we
	recommend incubating the blots from minigels in sealed bags (removing as much air as
	possible) with minimal volume (2-3 mLs). If used conservatively at 2.5 mLs/blot will yield
	enough reagent for 40 blots. Note that there are three key procedural considerations: 1. Protein
	A or G should not be used for the immunoprecipitation. Use of protein A or G beads with the
	rabbit TrueBlot will result in contaminating bands. For immunoprecipitation, anti-rat IgG beads,
	or anti-rabbit IgG beads should be used for rat or rabbit immunoprecipitating antibodies,
	respectively. 2. Immunoprecipitate should be completely reduced. 3. MB-070 Blocking Buffer
	for Fluorescent Western Blotting should be used as the blocking protein for the immunoblot. All
	recommended dilutions for listed applications are intended as an initial recommendation,
	specific conditions for each protein and antibody combination should be specifically optimized
	by the end user.
	FLISA Dilution: User Optimized

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Application Details	
	Flow Cytometry Dilution: 1:2,000 - 1:10,000
	Western Blot Dilution: 1:1000
	IF Microscopy Dilution: 1:500 - 1:2,500
Restrictions:	For Research Use only
Handling	
Format:	Lyophilized
Reconstitution:	Reconstitution Volume: 100 µL
	Reconstitution Buffer: Restore with deionized water (or equivalent)
Buffer:	Buffer: 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2
	Stabilizer: 10 mg/mL Polyethylene Glycol (PEG-8000)
	0.01 % (w/v) Sodium Azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
	should be handled by trained staff only.
Storage:	RT,4 °C,-20 °C
Storage Comment:	Store vial at 4 °C prior to restoration. For extended storage aliquot contents and freeze at -20 °C
	or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after
	standing at room temperature. This product is stable for several weeks at 4 $^\circ C$ as an undiluted
	liquid. Dilute only prior to immediate use.
Expiry Date:	12 months
Publications	
Product cited in:	Tian, Li, Gao, Li, Yang, Wang: "Identification and validation of the role of matrix
	metalloproteinase-1 in cervical cancer." in: International journal of oncology, Vol. 52, Issue 4,
	pp. 1198-1208, (2018) (PubMed).
	Zu, Wang, Ping, Sun: "Tan IIA inhibits H1299 cell viability through the MDM4-IAP3 signaling
	pathway." in: Molecular medicine reports, Vol. 17, Issue 2, pp. 2384-2392, (2018) (PubMed).

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#### Western Blotting

**Image 1.** Rabbit IP / Western Blot: Jurkat cell lysate (0.5 ml of 1x10e7 cells/ml) was incubated with rabbit anti-human Stat1 and immunoprecipitated using Protein G, Protein A and Anti-Rabbit Ig IP Beads. Precipitate from 5x10e5 cells was subjected to electrophoresis, transferred to a PVDF membrane, and Western blotted with anti-Stat1 using Rabbit: Anti-Rabbit IgG HRP

#### Western Blotting

**Image 2.** Fluorescent: Anti-Rabbit IgG DyLight 680 Conjugated - Western Blot. Western Blot of Fluorescent: Anti-Rabbit IgG DyLight 680 Conjugated. Lane 1: Rabbit IgG, Non-denatured. Lane 2: Rabbit IgG, Denatured. Load: 50 ng per lane. Primary antibody: none. Secondary antibody: Fluorescent: Anti-Rabbit IgG DyLight 680 Conjugated antibody at 1:1,000 for 60 min at RT. Block: ABIN925618 for 30 min at RT. Predicted: 160 kDa for non-denatured; observed: 170-180 kDa for non-denatured. Band migrates at slightly higher molecular weight.

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