

## Product datasheet

# Recombinant human TGF beta 3 protein (Animal Free) ab217402

## 2 References

### Description

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<b>Product name</b>	Recombinant human TGF beta 3 protein (Animal Free)
<b>Biological activity</b>	Determined by TGF- $\beta$ 3's ability to inhibit the mouse IL-4-dependent proliferation of mouse HT-2 cells. The expected ED <sub>50</sub> is $\leq 0.05$ ng/mL, corresponding to a specific activity of $\geq 2 \times 10^7$ units/mg.
<b>Purity</b>	> 98 % SDS-PAGE. > 98 % HPLC.
<b>Expression system</b>	Escherichia coli
<b>Accession</b>	<b><u>P10600</u></b>
<b>Protein length</b>	Full length protein
<b>Animal free</b>	Yes
<b>Nature</b>	Recombinant
<b>Species</b>	Human
<b>Sequence</b>	ALDTNYCFRNLEENCCVRPLYIDFRQDLGWKWWHEPKGY YANFCSGPCPY LRSADTTHSTVLGLYNTLNPEASASPCCVPQDLEPLTILY VGRTPKVEQ LSNMVKVSKCS
<b>Predicted molecular weight</b>	25 kDa
<b>Amino acids</b>	301 to 412
<b>Additional sequence information</b>	Recombinant Human TGF- $\beta$ 3 is composed of two identical 12.7 kDa, 112 amino acid polypeptide chains linked by a single disulfide bond.

### Specifications

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Our **Abpromise guarantee** covers the use of **ab217402** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<b>Applications</b>	Functional Studies SDS-PAGE HPLC
<b>Form</b>	Lyophilized

## Preparation and Storage

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<b>Stability and Storage</b>	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle.  This product is an active protein and may elicit a biological response in vivo, handle with caution.
<b>Reconstitution</b>	For lot specific reconstitution information please contact our Scientific Support Team.

## General Info

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<b>Function</b>	Involved in embryogenesis and cell differentiation.
<b>Involvement in disease</b>	Defects in TGFB3 are a cause of familial arrhythmogenic right ventricular dysplasia type 1 (ARVD1) [MIM:107970]; also known as arrhythmogenic right ventricular cardiomyopathy 1 (ARVC1). ARVD is an autosomal dominant disease characterized by partial degeneration of the myocardium of the right ventricle, electrical instability, and sudden death. It is clinically defined by electrocardiographic and angiographic criteria; pathologic findings, replacement of ventricular myocardium with fatty and fibrous elements, preferentially involve the right ventricular free wall.
<b>Sequence similarities</b>	Belongs to the TGF-beta family.
<b>Cellular localization</b>	Secreted.

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**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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