

## Product datasheet

# Recombinant Human FOXC2 protein ab114305

### 画像数 1

#### 製品の概要

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|--------|---------------------------------|
| 製品名    | Recombinant Human FOXC2 protein |
| タンパク質長 | Protein fragment                |

#### 製品の詳細

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|           |  |
|-----------|--|
| 由来        | Recombinant  |
| 由来        | Wheat germ   |
| アミノ酸配列    |  |
| アクセッション番号 | <a href="#">Q99958</a>   |
| 生物種       | Human  |
| 配列        | AASWYLNHSGDLNHLPGHTFAAQQTFPNVREMFNSHRLGIENSTLGESQ<br>VSGNASCQLPYRSTPPLYRHAAPYSYDCTKY |
| 分子量       | 35 kDa including tags  |
| 領域        | 421 to 501   |

#### 特性

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Our [Abpromise guarantee](#) covers the use of **ab114305** in the following tested applications.

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

|          |   |
|----------|---|
| アプリケーション | Western blot<br>ELISA<br>SDS-PAGE   |
| 製品の状態    | Liquid  |
| 備考       | Protein concentration is above or equal to 0.05 mg/ml.<br>This protein is best used within three months from the date of receipt. |

#### 前処理および保存

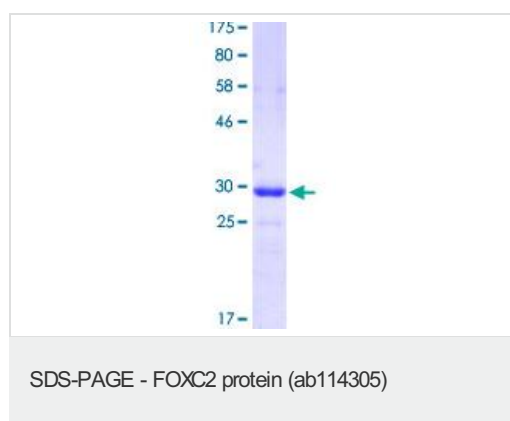
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|            |   |
|------------|---|
| 保存方法および安定性 | Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.<br>pH: 8.00 |
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## 関連情報

|       |  |
|-------|--|
| 機能    | Transcriptional activator. Might be involved in the formation of special mesenchymal tissues.  |
| 関連疾患  | <p>Defects in FOXC2 are the cause of lymphedema hereditary type 2 (LMPH2) [MIM:153200]; also known as Meige lymphedema. Hereditary lymphedema is a chronic disabling condition which results in swelling of the extremities due to altered lymphatic flow. Patients with lymphedema suffer from recurrent local infections, and physical impairment.</p> <p>Defects in FOXC2 are a cause of lymphedema-yellow nails (LYYN) [MIM:153300]. LYYN is characterized by yellow, dystrophic, thick and slowly growing nails, associated with lymphedema and respiratory involvement. Lymphedema occurs more often in the lower limbs. It can appear at birth or later in life. Onset generally follows the onset of ungual abnormalities.</p> <p>Defects in FOXC2 are a cause of lymphedema-distichiasis (LYD) [MIM:153400]. LYD is characterized by primary limb lymphedema usually starting at puberty (but in some cases later or at birth) and associated with distichiasis (double rows of eyelashes, with extra eyelashes growing from the Meibomian gland orifices).</p> |
| 配列類似性 | Contains 1 fork-head DNA-binding domain.   |
| 細胞内局在 | Nucleus.   |

## 画像



12.5% SDS-PAGE image showing ab114305  
Stained with Coomassie Blue.

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

## Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
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- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

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