Anti-BRAF (mutated V600E) antibody [RM8] ab200535

Overview

Product name
Anti-BRAF (mutated V600E) antibody [RM8]

Description
Rabbit monoclonal [RM8] to BRAF (mutated V600E)

Host species
Rabbit

Specificity
ab200535 reacts to the B Raf V600E mutant. No cross reactivity with wild type B Raf.

Immunogen
Synthetic peptide corresponding to Human BRAF (mutated V600E).

Database link: P15056

Positive control
Cell lysates prepared from cell lines expressing endogenous B Raf (mutated V600E); Human melanoma and Human colon cancer tissues; WiDr cells.

Properties

Form
Liquid

Storage instructions

Storage buffer
Preservative: 0.09% Sodium azide
Constituents: 1% BSA, 48% PBS, 50% Glycerol

Purity
Protein A purified

Purification notes
Purified from an animal origin–free culture supernatant.

Clonality
Monoclonal

Clone number
RM8

Isotype
IgG
The Abpromise guarantee

Our Abpromise guarantee covers the use of ab200535 in the following tested applications.

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

<table>
<thead>
<tr>
<th>Guaranteed</th>
<th>Predicted</th>
<th>Incompatible</th>
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<tbody>
<tr>
<td>Tested applications are guaranteed to work and covered by our Abpromise guarantee.</td>
<td>Predicted to work for this combination of applications and species but not guaranteed.</td>
<td>Does not work for this combination of applications and species.</td>
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<thead>
<tr>
<th>Application</th>
<th>Species</th>
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<tr>
<td>IHC-P</td>
<td>Human</td>
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<tr>
<th>Application</th>
<th>Abreviews</th>
<th>Notes</th>
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<tr>
<td>WB</td>
<td>⭐⭐⭐⭐⭐ (1)</td>
<td>Use a concentration of 0.5 - 2 µg/ml. Predicted molecular weight: 84 kDa.</td>
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<tr>
<td>IHC-P</td>
<td></td>
<td>Use a concentration of 2 - 10 µg/ml.</td>
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Target

Function

Involved in the transduction of mitogenic signals from the cell membrane to the nucleus. May play a role in the postsynaptic responses of hippocampal neuron.

Tissue specificity

Brain and testis.

Involvement in disease

Note=Defects in BRAF are found in a wide range of cancers.
Defects in BRAF may be a cause of colorectal cancer (CRC) [MIM:114500].
Defects in BRAF are involved in lung cancer (LNCR) [MIM:211980].
Defects in BRAF are involved in non-Hodgkin lymphoma (NHL) [MIM:605027]. NHL is a cancer that starts in cells of the lymph system, which is part of the body's immune system. NHLs can occur at any age and are often marked by enlarged lymph nodes, fever and weight loss.
Defects in BRAF are a cause of cardiofaciocutaneous syndrome (CFC syndrome) [MIM:115150]; also known as cardio-facio-cutaneous syndrome. CFC syndrome is characterized by a distinctive facial appearance, heart defects and mental retardation. Heart defects include pulmonic stenosis, atrial septal defects and hypertrophic cardiomyopathy. Some affected individuals present with ectodermal abnormalities such as sparse, friable hair, hyperkeratotic skin lesions and a generalized ichthyosis-like condition. Typical facial features are similar to Noonan syndrome. They include high forehead with bitemporal constriction, hypoplastic supraorbital ridges, downslanting palpebral fissures, a depressed nasal bridge, and posteriorly angulated ears with prominent helices. The inheritance of CFC syndrome is autosomal dominant.
Defects in BRAF are the cause of Noonan syndrome type 7 (NS7) [MIM:613706]. Noonan syndrome is a disorder characterized by facial dysmorphic features such as hypertelorism, a downward eyeslant and low-set posteriorly rotated ears. Other features can include short stature, a short neck with webbing or redundancy of skin, cardiac anomalies, deafness, motor delay and variable intellectual deficits.
Defects in BRAF are the cause of LEOPARD syndrome type 3 (LEOPARD3) [MIM:613707]. LEOPARD3 is a disorder characterized by lentigines, electrocardiographic conduction abnormalities, ocular hypertelorism, pulmonic stenosis, abnormalities of genitalia, retardation of
growth, and sensorineural deafness.

Note: A chromosomal aberration involving BRAF is found in pilocytic astrocytomas. A tandem duplication of 2 Mb at 7q34 leads to the expression of a KIAA1549-BRAF fusion protein with a constitutive kinase activity and inducing cell transformation.

**Sequence similarities**

Belongs to the protein kinase superfamily. TKL Ser/Thr protein kinase family, RAF subfamily. Contains 1 phorbol-ester/DAG-type zinc finger. Contains 1 protein kinase domain. Contains 1 RBD (Ras-binding) domain.

**Cellular localization**

Nucleus. Cytoplasm. Cell membrane. Colocalizes with RGS14 and RAF1 in both the cytoplasm and membranes.

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**Images**

**All lanes** : Anti-BRAF (mutated V600E) antibody [RM8] (ab200535) at 2 µg/ml

**Lane 1** : Cell lysates prepared from cell lines expressing endogenous B Raf (mutated V600E)

**Lane 2** : Cell lysates prepared from cell lines expressing endogenous wild-type protein

**Predicted band size**: 84 kDa

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Immunohistochemical analysis of formalin fixed and paraffin embedded WiDr cells labeling B Raf (mutated V600E) with ab200535 at 5 µg/mL.
Immunohistochemical analysis of formalin-fixed, paraffin-embedded Human melanoma tissue labeling B Raf with ab200535 at 5 µg/mL.

Why choose a recombinant antibody?
- Research with confidence
  Consistent and reproducible results
- Long-term and scalable supply
  Recombinant technology
- Success from the first experiment
  Confirmed specificity
- Ethical standards compliant
  Animal-free production

Anti-BRAF (mutated V600E) antibody [RM8] (ab200535)

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

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