



Monoclonal Antibody against α B-Crystallin's Function, A Heat Shock Protein implicated in Cancer & Fibrosis

Category

Technology: Monoclonal Antibody
Developed: 2017

Problem

α B-Crystallin is a potential therapeutic target in many diseases

Technology Overview

A monoclonal antibody that selectively inhibits the function of α B-Crystallin through ABCP binding

IP Status

- ▶ Available for Exclusive or Non-Exclusive Licensing

Value Proposition

- ▶ Monoclonal antibody
- ▶ Inhibits function

Market Attractions

- ▶ Detection of protein
- ▶ Protein modification
- ▶ Therapeutics

Contact

Heather Callahan

Heather.Callahan@ucdenver.edu

Ref# CU3971H

CU Innovations
13001 E. 17th Place
Suite W5130, Aurora, CO
80045

t. 303-724-0221

f. 303-724-0816

innovations.ucdenver.edu

Problem: α B-crystallin, a heat shock protein, is implicated in the pathogenesis of several diseases or disease processes. In triple negative basal-type breast cancers, higher expression levels of α B-crystallin are directly related to poor prognosis. α B-crystallin is highly expressed and obligatory in fibrosis in lungs, retina and lens and is involved in pathological angiogenesis in the retina. As a result, α B-crystallin is thought to be a potential therapeutic target for many medical conditions including cancer and fibrosis.

Technical Solution and Key Value Propositions:

α B-crystallin functions as a molecular chaperone and has anti-apoptotic effects mediated through its core domain peptide, ABCP. Dr. Nagaraj's team at the University of Colorado has developed and characterized a monoclonal antibody against ABCP (ABCP Ab). Using western blotting, ABCP Ab was shown to selectively recognize the core domain peptide of α B-crystallin, but not the core domain of α A-crystallin and Hsp27 (Fig. 1). It was also shown to detect α B-crystallin in both human and wild type mouse lenses, but not in α B-crystallin knock-out mouse lenses (Fig. 2). Subsequent *in vitro* functional assays showed that ABCP Ab efficiently inhibits the chaperone function of α B-crystallin for several proteins *in vitro*.

Through their ability to selectively recognize a target, monoclonal antibodies are widely used to study mechanisms of action and in high-throughput screens for new therapeutics.

Highlights:

- ▶ Target protein highly implicated in diseases
- ▶ Recognizes both human and mouse proteins in western blots, ELISA, immunohistochemistry, and immunocytochemistry.
- ▶ Inhibits protein function

Key Documents and Sources:

"A Monoclonal Antibody Targeted to the Functional Peptide Inhibits the Chaperone Activity of Human α B-Crystallin" Rooban B. Nahomi, Sandip K. Nandi, and Ram H. Nagaraj is available under CDA.

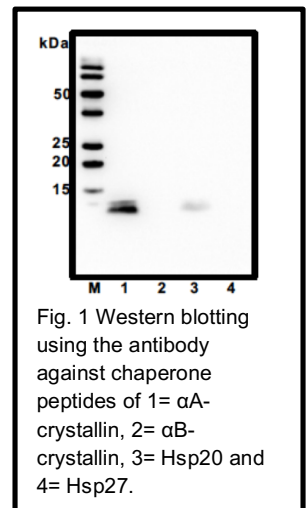


Fig. 1 Western blotting using the antibody against chaperone peptides of 1= α A-crystallin, 2= α B-crystallin, 3= Hsp20 and 4= Hsp27.

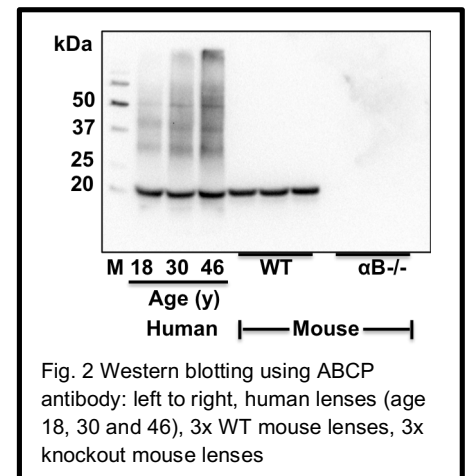


Fig. 2 Western blotting using ABCP antibody: left to right, human lenses (age 18, 30 and 46), 3x WT mouse lenses, 3x knockout mouse lenses