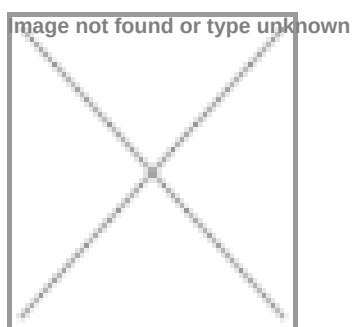


KRIT1 loss-of-function and enhanced cell predisposition to oxidative damage



A recent research work published in *Free Radical Biology & Medicine* ^[1] and supported by the **Telethon Foundation** ^[2] sheds new light on the mechanistic relationship between KRIT1 loss-of-function and enhanced cell predisposition to oxidative damage, thus providing valuable new insights into CCM pathogenesis and novel options for the development of preventive and therapeutic strategies.

Free Radical Biology & Medicine
Epub 2017 Nov 21.

^[3]*Free Radic Biol Med.* 2018 Feb 1;115:202-218.

KRIT1 loss-of-function induces a chronic Nrf2-mediated adaptive homeostasis that sensitizes cells to oxidative stress: implication for Cerebral Cavernous Malformation disease.

Cinzia Antognelli, Eliana Trapani, Simona Delle Monache, Andrea Perrelli, Martina Daga, Stefania Pizzimenti, Giuseppina Barrera, Paola Cassoni, Adriano Angelucci, Lorenza Trabalzini, Vincenzo Talesa, Luca Goitre, and Saverio Francesco Retta.

Abstract ^[4]

Full article ^[5] (PDF ^[6])

Highlights

KRIT1 loss causes a chronic adaptive redox response based on the JNK-Nrf2-Glo1 axis

Phospho-JNK, Nrf2 and Glo1 are upregulated in endothelial cells lining human CCMs

Defective autophagy contributes to the sustained upregulation of the Nrf2-Glo1 axis

Nrf2-Glo1 upregulation causes a drop of AP-modified Hsp70 and Hsp27 proteins

Sustained Nrf2-Glo1 activation sensitizes cells to oxidative stress and apoptosis

Graphical abstract

Graphical abstract

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Related article:

Data in Brief

Data Brief. 2017 Dec 13;16:929-938. eCollection 2018 Feb.

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Data in support of sustained upregulation of adaptive redox homeostasis mechanisms caused by KRIT1 loss-of-function.

Cinzia Antognelli, Eliana Trapani, Simona Delle Monache, Andrea Perrelli, Claudia Fornelli, Francesca Retta, Paola Cassoni, Vincenzo Nicola Talesa, and Saverio Francesco Retta.

Abstract ^[7]

Full article ^[8] (PDF ^[9])

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Source URL: <http://www.ccmitalia.unito.it/it/content/krit1-loss-function-and-enhanced-cell-predisposition-oxidative-damage>

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