



Current mechanistic insights into the CCCP-induced cell survival response

Submitted by Guy Lenaers on Sat, 12/22/2018 - 12:12

Titre	Current mechanistic insights into the CCCP-induced cell survival response
Type de publication	Article de revue
Auteur	Kane, Mariame-Selma [1], Paris, Aurelien [2], Codron, Philippe [3], Cassereau, Julien [4], Procaccio, Vincent [5], Lenaers, Guy [6], Reynier, Pascal [7], Chevrollier, Arnaud [8]
Editeur	Elsevier
Type	Article scientifique dans une revue à comité de lecture
Année	2018
Langue	Anglais
Date	Février 2018
Pagination	100-110
Volume	148
Titre de la revue	Biochemical pharmacology
ISSN	1873-2968
Mots-clés	Animals [9], Carbonyl Cyanide m-Chlorophenyl Hydrazone [10], Carbonyl Cyanide p-Trifluoromethoxyphenylhydrazine [11], Cell Survival [12], mitochondria [13]
Résumé en anglais	<p>The ring-substituted derivatives of carbonyl cyanide phenylhydrazine, CCCP and FCCP, are routinely used for the analysis of the mitochondrial function in living cells, tissues, and isolated mitochondrial preparations. CCCP and FCCP are now being increasingly used for investigating the mechanisms of autophagy by inducing mitochondrial degradation through the disruption of the mitochondrial membrane potential ($\Delta\Psi_m$). Sustained perturbation of $\Delta\Psi_m$, which is normally tightly controlled to ensure cell proliferation and survival, triggers various stress pathways as part of the cellular adaptive response, the main components of which are mitophagy and autophagy. We here review current mechanistic insights into the induction of mitophagy and autophagy by CCCP and FCCP. In particular, we analyze the cellular modifications produced by the activation of two major pathways involving the signaling of the nuclear factor erythroid 2-related factor 2 (Nrf2) and the transcription factor EB (TFEB), and discuss the contribution of these pathways to the integrated cellular stress response.</p>
URL de la notice	http://okina.univ-angers.fr/publications/ua18488 [14]
DOI	10.1016/j.bcp.2017.12.018 [15]
Lien vers le document	https://www.sciencedirect.com/science/article/abs/pii/S000629521730730X?... [16]
Titre abrégé	Biochem. Pharmacol.

Identifiant
(ID) 29277693 [17]
PubMed

Liens

- [1] <http://okina.univ-angers.fr/publications?f%5Bauthor%5D=32231>
- [2] <http://okina.univ-angers.fr/publications?f%5Bauthor%5D=32225>
- [3] <http://okina.univ-angers.fr/pcodron/publications>
- [4] <http://okina.univ-angers.fr/julien.cassereau/publications>
- [5] <http://okina.univ-angers.fr/v.procaccio/publications>
- [6] <http://okina.univ-angers.fr/guy.lenaers/publications>
- [7] <http://okina.univ-angers.fr/pascal.reynier/publications>
- [8] <http://okina.univ-angers.fr/arnaud.chevrollier/publications>
- [9] <http://okina.univ-angers.fr/publications?f%5Bkeyword%5D=964>
- [10] <http://okina.univ-angers.fr/publications?f%5Bkeyword%5D=26647>
- [11] <http://okina.univ-angers.fr/publications?f%5Bkeyword%5D=26648>
- [12] <http://okina.univ-angers.fr/publications?f%5Bkeyword%5D=6698>
- [13] <http://okina.univ-angers.fr/publications?f%5Bkeyword%5D=984>
- [14] <http://okina.univ-angers.fr/publications/ua18488>
- [15] <http://dx.doi.org/10.1016/j.bcp.2017.12.018>
- [16] <https://www.sciencedirect.com/science/article/abs/pii/S000629521730730X?via%3Dihub>
- [17] <http://www.ncbi.nlm.nih.gov/pubmed/29277693?dopt=Abstract>

Publié sur *Okina* (<http://okina.univ-angers.fr>)