

Curriculum Vitae

Informazioni personali

Nome(i) / Cognome(i)

Orazio Cantoni

Indirizzo(i)

Abitazione:
Via Cà Mignone 14, - 61029 Urbino (PU)

Lavoro:

Università degli Studi di Urbino "Carlo Bo"
Dipartimento di Scienze Biomolecolari
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Cittadinanza

Italiana

Data di nascita

25 Luglio 1954

Sesso

Maschio

Occupazione /Settore professionale

Esperienza professionale

Date

1977 - Laurea in Farmacia (cum laude) presso l'Università degli Studi di Urbino "Carlo Bo".
1981 - Laurea in Scienze Biologiche (cum laude) presso l'Università degli Studi di Urbino "Carlo Bo".
1981 - Visiting Scientist presso il Laboratory of Preclinical Pharmacology, NIMH, Washington D.C.
1982-1983 - Post-Doctoral Fellow presso il Department of Pharmacology, University of Texas, Houston.
1983 - Ricercatore presso l' Istituto di Farmacologia e Farmacognosia dell' Università degli Studi di Urbino.
1984-1985 (4 mesi) - Visiting Scientist nei Departments of Physics (1984, 4 mesi) e Experimental Radiotherapy dell'MD Anderson Hospital, Houston.
1992 - Professore Associato (BIO/14), presso la Facoltà di Farmacia dell'Università degli Studi di Urbino "Carlo Bo".
2000 - Professore Ordinario (BIO/14), presso la Facoltà di Farmacia dell'Università degli Studi di Urbino "Carlo Bo".
2009 - Preside della Facoltà di Farmacia dell'Università degli Studi di Urbino "Carlo Bo".
Attualmente il Prof Cantoni è direttore del Dipartimento di Scienze Biomolecolari dell'Università degli Studi di Urbino "Carlo Bo".
Dal 2016 ad oggi ricopre la carica di Prorettore alla Ricerca
2018: Presidente Eletto della Società Italiana di Tossicologia

Lavoro o posizione ricoperti	<p>Il Prof. Cantoni ha ricevuto diversi premi, tra cui quelli dell' "Associazione Italiana per la Lotta contro i Tumori" (1985), della "Fondazione Assicurazioni Generali" (1986), "Menzione di merito" della Commissione giudicatrice della Società Italiana di Farmacologia del "Premio Benedicenti" (1988) e, nel 1990, il "Premio Benedicenti" della Società Italiana di Farmacologia.</p> <p>Il Prof. Cantoni è stato membro del Consiglio Direttivo della Society for Free Radical Research – Europe. Attualmente è membro del direttivo della Società Italiana di Tossicologia.. E' stato reviewer per numerosi giornali scientifici, tra cui: Free Radical Research, Free Radical Biology and Medicine, Biochimica Biophysica Acta, Biochemical Pharmacology, Journal Biological Chemistry Hoppe-Seyler, Cell Death and Differentiation, Drugs, Expert Opinion on Investigational Drugs, Experimental Cell Research, Journal of Neurochemistry e The Proceedings of the National Academy of Science (USA). Attualmente membro dell Editorial Board di Pharmacological Research.</p> <ul style="list-style-type: none"> • Number of scientific articles (international journals with peer review) in extenso: 190 • H-index (Scopus – December 2017): 33 • Total citations (Scopus – December 2018): 5170 • Total impact factor of publications (5-year impact factor da ISI Web of Knowledge, Journal of Citation Reports 2016): 721,57 Average impact factor: 4,15 • Journal rankings in the subject category of scientific articles (SCImago Journal & Country Rank 2014): Q1: 140 articles; Q2: 37 articles; Q3: 13 articles.
Principali attività e responsabilità	
Nome e indirizzo del datore di lavoro	<p><i>Attività scientifica:</i> Il lavoro di ricerca condotto durante il post-doctoral training, presso il Department of Pharmacology della University of Texas, Houston, è stato svolto con il fine di individuare le lesioni del DNA mediate da specifici composti metallici cancerogeni. Dal 1984 è stato attivamente coinvolto in studi sui meccanismi di tossicità mediata da specie reattive dell'ossigeno. Il Prof. Cantoni ha contribuito in modo sostanziale allo sviluppo, ed al crescente interesse, di questo campo di ricerca con alcuni dei primi lavori sugli effetti del perossido di idrogeno. Successivamente, egli ha spostato i propri interessi sugli effetti di idroperossidi organici e, durante questo periodo ha prodotto importanti risultati che indicavano che l'accumulo mitocondriale di ioni calcio portava alla formazione di specie cito-geno-tossiche. Il Prof Cantoni ha recentemente dimostrato che l'acido arachidonico promuove fosforilazione (tirosina kinasi-dipendente) ed inattivazione della nitrossido sintasi costitutiva degli astrociti. Questo evento era critico per l'espressione di geni NFkB-dipendenti in cellule stimolate con LPS-IFNgamma. Un ulteriore contributo in questo campo è stato l'identificazione dei meccanismi attraverso cui cellule del lineaggio monocitico/macrofagico resistono al perossinitrito, il prodotto dell'interazione tra nitrossido e superossido. Questi studi sono stati stimolati dalla semplice considerazione che queste cellule, nei siti infiammatori, producono diverse specie reattive e tossiche, tra cui il perossinitrito, e debbono pertanto essere provviste di un efficiente sistema di difesa. Si è visto che, mentre "committed" alla transizione della permeabilità della membrana mitocondriale interna (MPT), queste cellule rispondono all'acido arachidonico con l'attivazione di un signalling di sopravvivenza che previene MPT e morte. Un ulteriore importante scoperta è stata che la tossicità, rilevabile solo dopo esposizione ad elevate concentrazioni di perossinitrito, era indipendente dall'entità del danno accumulato dalle cellule ed era infatti mediata dall'inibizione del signalling di sopravvivenza. Nel suo insieme, l'informazione fornita da questi studi, è consistente con la nozione che queste cellule sopravvivono al perossinitrito utilizzando molecole segnale che sono tossiche per altri tipi cellulari. Inoltre, queste cellule possono sopravvivere nonostante l'accumulo di un danno esteso ed eventualmente muoiono, quando il signalling di sopravvivenza viene inibito, attraverso un meccanismo altamente regolato di morte necrotica. Più recentemente, si è interessato del trasporto e della compartmentalizzazione sub-cellulare della vitamina C. In questo ambito, un importante contributo è rappresentato dalla scoperta del trasportatore sodio-dipendente della vitamina nei mitocondri. Questo trasportatore viene riconosciuto da anticorpi anti-SVCT2 ed è caratterizzato da una elevata affinità anche in presenza di basse concentrazioni di sodio, normalmente presenti nel distretto intracellulare (< 10 mM). Questo dato è in forte contrasto con quanto osservato con il trasportatore espresso nella membrana plasmatica, che richiede per l'attività di trasporto massimale concentrazioni di sodio superiore a 120 mM. Infine le attività dei trasportatori delle vitamina C localizzati nelle membrane plasmatiche e mitocondriali era sensibile a basse concentrazione di acido deidroascorbico, la forma ossidata della vitamina C.</p>

Tipo di attività o settore Professore Ordinario (BIO/14 - Farmacologia)

Istruzione e formazione

Date

Titolo della qualifica rilasciata

Principal tematiche/competenze professionali acquisite

Nome e tipo d'organizzazione erogatrice dell'istruzione e formazione

Livello nella classificazione nazionale o internazionale

Capacità e competenze personali

Madrelingua(e)

Italiano

Altra(e) lingua(e)

Inglese

Autovalutazione

Livello europeo (*)

Lingua

Lingua

Comprensione		Parlato		Scritto
Ascolto	Lettura	Interazione orale	Produzione orale	
C1	C1	C1	C1	C1

Capacità e competenze sociali

Capacità e competenze organizzative

Capacità e competenze tecniche

Capacità e competenze informatiche

Capacità e competenze artistiche

Altre capacità e competenze

Patente

Ulteriori informazioni

Allegati

Elenco delle pubblicazioni

Firma



Allegato

Pubblicazioni Prof. Orazio Cantoni

GUIDARELLI, A, FIORANI, M, CERIONI, L, **CANTONI, O** (2019). Calcium signals between the ryanodine receptor- and mitochondria critically regulate the effects of arsenite on mitochondrial superoxide formation and on the ensuing survival vs apoptotic signaling. *REDOX BIOLOGY*, vol. 20, p. 285-295, ISSN: 2213-2317, doi: 10.1016/j.redox.2018.10.015

GUIDARELLI, A, FIORANI, M, **CANTONI, O** (2018). Low Concentrations of Arsenite Target the Intraluminal Inositol 1, 4, 5-Trisphosphate Receptor/Ryanodine Receptor Crosstalk to Significantly Elevate Intracellular Ca²⁺. *JOURNAL OF PHARMACOLOGY AND EXPERIMENTAL THERAPEUTICS*, vol. 367 p.184-193. ISSN: 0022-3565, doi: 10.1124/jpet.118.250480

SCOTTI, M, FIORANI, M, GUIDARELLI, A, **CANTONI, O** (2018). Differentiation of Promonocytic U937 Cells to Monocytes Is Associated with Reduced Mitochondrial Transport of Ascorbic Acid. *OXIDATIVE MEDICINE AND CELLULAR LONGEVITY*, vol. 2018, p. 1-12, ISSN: 1942-0900, doi: 10.1155/2018/4194502

CANTONI, O, GUIDARELLI, A, FIORANI, M (2018). Mitochondrial Uptake and Accumulation of Vitamin C: What Can We Learn From Cell Cultures Studies?. *ANTIOXIDANTS & REDOX SIGNALING*, vol. 29 p.1502-1515. ISSN: 1523-0864, doi: 10.1089/ars.2017.7253

FIORANI, M, GUIDARELLI A, CAPELLACCI, V, CERIONI, L, CRINELLI, R, **CANTONI, O** (2018). The dual role of mitochondrial superoxide in arsenite toxicity: Signaling at the boundary between apoptotic commitment and cytoprotection. *TOXICOLOGY AND APPLIED PHARMACOLOGY*, vol. 345, p. 26-35, ISSN: 0041-008X, doi: 10.1016/j.taap.2018.03.008

GUIDARELLI, A, CERIONI, L, FIORANI, M, **CANTONI, O** (2017). Intramitochondrial Ascorbic Acid Enhances the Formation of Mitochondrial Superoxide Induced by Peroxynitrite via a Ca(2+)-Independent Mechanism. *INTERNATIONAL JOURNAL OF MOLECULAR SCIENCES*, vol. 18, p. E1686. ISSN: 1422-0067, doi: 10.3390/ijms18081686

GUIDARELLI, A, FIORANI, M, CERIONI, L, SCOTTI, M, **CANTONI, O** (2017). Arsenite induces DNA damage via mitochondrial ROS and induction of mitochondrial permeability transition. *BIOFACTORS*, vol. 43, p. 673-684, ISSN: 0951-6433, doi: 10.1002/biof.1375

PERSICHINI TIZIANA, MARIOTTO SOFIA, SUSUKI HISANORI, BUTTIRINI ELENA, MASTRANTONIO ROBERTA, **CANTONI ORAZIO**, COLASANTI MARCO (2016). Cross-Talk Between NO Synthase Isoforms in Neuro-Inflammation: Possible Implications in HIV-Associated Neurocognitive Disorders. *CURRENT MEDICINAL CHEMISTRY*, vol. 23, p. 2706-2714.

GUIDARELLI A, CARLONI S, FIORANI M, A, BALDUINI W, **CANTONI O** (2016). Mitochondrial ascorbic acid prevents mitochondrial O₂⁻ formation, an event critical for U937 cell apoptosis induced by arsenite through both autophagic-dependent and independent mechanisms. *BIOFACTORS*, vol. 42, p. 190-200.

GUIDARELLI ANDREA, FIORANI MARA, CARLONI SILVIA, CERIONI LIANA, BALDUINI WALTER, **CANTONI ORAZIO** (2016). The study of the mechanism of arsenite toxicity in respiration-deficient cells reveals that NADPH oxidase-derived superoxide promotes the same downstream events mediated by mitochondrial superoxide in respiration-proficient cells. *TOXICOLOGY AND APPLIED PHARMACOLOGY*, vol. 307, p. 35-44.

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FIORANI M, AZZOLINI C, CERIONI L, SCOTTI M, GUIDARELLI A, CIACCI C, **CANTONI O** (2015). The mitochondrial transporter of ascorbic acid functions with high affinity in the presence of low millimolar concentrations of sodium and in the absence of calcium and magnesium. *BIOCHIMICA ET BIOPHYSICA ACTA-BIOMEMBRANES*, vol. 99, p. 289-295, ISSN: 0005-2736.

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TIZIANA PERSICHINI, ROBERTA MASTRANTONIO, SILVIA DEL MATTO, LETIZIA PALOMBA, **ORAZIO CANTONI**, MARCO COLASANTI (2014). The role of arachidonic acid in the regulation of nitric oxide synthase isoforms by HIV gp120 protein in astroglial cells. *FREE RADICAL BIOLOGY & MEDICINE*, vol. 74, p. 14-20, ISSN: 0891-5849.

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GUIDARELLI A, CERIONI L, FIORANI M, AZZOLINI C, **CANTONI O** (2014). Mitochondrial acorbic acid is responsible for enhanced susceptibility of U937 cells to the toxic effects of peroxynitrite. *BIOFACTORS*, vol. 40, p. 236-246, ISSN: 0951-6433.

AZZOLINI C, FIORANI M, CERIONI L, GUIDARELLI A, **CANTONI O**. (2013). Sodium-dependent transport of ascorbic acid in U937 cell mitochondria. *IUBMB Life*, vol 65, p. 149-153, ISSN: 15216543.

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CRIMELLA C, CANTONI O, GUIDARELLI A, VANTAGGIATO C, MARTINUZZI A, FIORANI M, AZZOLINI C, ORSO G, BRESOLIN N, BASSI MT. (2011). A novel nonsense mutation in the APTX gene associated with delayed DNA single-strand break removal fails to enhance sensitivity to different genotoxic agents. HUMAN MUTATION, vol. 32, p. E2118-2133, ISSN: 1098-1004

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GUIDARELLI A, CERIONI L, FIORANI M, CANTONI O. (2009). Differentiation-associated loss of ryanodine receptors: a strategy adopted by monocytes/macrophages to prevent the DNA single-strand breakage induced by peroxynitrite. JOURNAL OF IMMUNOLOGY, vol. 183; p. 4449-4457, ISSN: 0022-1767

PALOMBA L, CERIONI L, CANTONI O. (2009). Arachidonic acid: A key molecule for astrocyte survival to peroxynitrite. GLIA, vol. 57; p. 1672-1679, ISSN: 0894-1491

CANTONI O., GUIDARELLI A (2008). Peroxynitrite damages U937 cell DNA via the intermediate formation of mitochondrial oxidants. IUBMB LIFE, vol. 60; p. 753-756, ISSN: 1521-6543

CANTONI O., GUIDARELLI A (2008). Indirect Mechanisms of DNA Strand Scission by Peroxynitrite. METHODS IN ENZYMOLOGY, vol. 440; p. 111-120, ISSN: 0076-6879

CANTONI O., PALOMBA L, PERSICHINI T, MARIOTTO S, SUZUKI H, COLASANTI M (2008). Pivotal Role of Arachidonic Acid in the Regulation of Neuronal Nitric Oxide Synthase Activity and Inducible Nitric Oxide Synthase Expression in Activated Astrocytes. METHODS IN ENZYMOLOGY, vol. 440; p. 243-252, ISSN: 0076-6879

CANTONI O., TOMMASINI I, CERIONI L (2008). The arachidonate-dependent survival signaling preventing toxicity in monocytes/macrophages exposed to peroxynitrite. METHODS IN ENZYMOLOGY, vol. 441; p. 73-82, ISSN: 0076-6879

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GUIDARELLI A, PALOMBA L, FIORANI M, CANTONI O. (2008). Susceptibility of rat astrocytes to DNA strand scission induced by activation of NADPH oxidase and collateral resistance to the effects of peroxynitrite. FREE RADICAL BIOLOGY & MEDICINE, vol. 45; p. 521-529, ISSN: 0891-5849

TOMMASINI I, CERIONI L, PALOMBA L, CANTONI O. (2008). Prostaglandin E2 signals monocyte/macrophage survival to peroxynitrite via protein kinase A converging in bad phosphorylation with the protein kinase C alpha-dependent pathway driven by 5-hydroxyeicosatetraenoic acid. JOURNAL OF IMMUNOLOGY, vol. 181; p. 5637-5645, ISSN: 0022-1767

CERIONI L, CANTONI O. (2007). ERK1/2 regulates two sequential steps promoting monocyte survival to peroxynitrite. JOURNAL OF CELLULAR PHYSIOLOGY, vol. 210; p. 177-182, ISSN: 0021-9541

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PALOMBA L, AMADORI A, CANTONI O. (2007). Early release of arachidonic acid prevents an otherwise immediate formation of toxic levels of peroxynitrite in astrocytes stimulated with lipopolysaccharide/interferon-gamma. JOURNAL OF NEUROCHEMISTRY, vol. 103; p. 904-914, ISSN: 0022-3042

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VALERIO A, CARDILE A, COZZI V, BRACALE R, TEDESCO L, PISCONTI A, PALOMBA L, CANTONI O., CLEMENTI E, MONCADA S, CARRUBA MO, NISOLI E (2006). TNF-alpha downregulates eNOS expression and mitochondrial biogenesis in fat and muscle of obese rodents. JOURNAL OF CLINICAL INVESTIGATION, vol. 116; p. 2791-2798, ISSN: 0021-9738

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PALOMBA L, PERSICHINI T, MAZZONE V, COLASANTI M, CANTONI O. (2004). Inhibition of nitric-oxide synthase-i (NO-i)-dependent nitric oxide production by lipopolysaccharide plus interferon-gamma is mediated by arachidonic acid. Effects on NFK-b activation and late inducible nos expression. THE JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 279; p. 29895-29901, ISSN: 0021-9258

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