

## Curriculum vitae CLAUDIA BOCCA

### **Personal details**

Born in Biella, 31/01/1968

Nationality: Italian

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### **Educations**

1992. Degree in Biological Sciences, University of Torino

1997. PhD in Experimental and Molecular Pathology, University of Torino

1998. Qualification as a Professional Biologist

2001. Post-graduated degree in Clinical Pathology

2001-2003. Post-Doctoral training in Medical Sciences

### **Professional experiences and current position**

10/2022 – Present. Associate Professor of General Pathology, Dept Clinical and Biological Sciences, University of Torino

01/2005–09/2022. Assistant Professor of General Pathology, Dept Clinical and Biological Sciences, University of Torino

2008. Visiting Fellow at Dept Neuroscience, University of Roma Tor Vergata

2003-2004. CoCoCo at Pathological Anatomy of ASO Mauriziano Umberto I of Torino with a role in a national project founded by Compagnia di San Paolo regarding the expression of prognostic factors in thyroid neoplasia.

### **Participation to Directive Boards of Scientific Societies and/or Institutions:**

From 2022. Member of Editorial Board team of the journal "Antioxidants"

From 2022. Guest Editor of the journal Antioxidants for the special issue "The Role of Oxidative Stress in Liver Cancer"

### **Honors**

2001-2002. Post-doc fellowships in Medical Sciences, Post-doc fellowships, University of Torino

1997-2001. Full scholarship from University of Torino to attend the Post Graduate Specialization School in Clinical Pathology, University of Torino.

1992-1996. Full scholarship from the Italian Ministry of Public Education for the PhD course, University of Torino.

### **Teaching activity:**

- General Pathology with Elements of Pathophysiology (9 credits) - Pharmacy, University of Torino
- General Pathology and Medical Terminology (5 credits) - Pharmaceutical Chemistry and Technology, University of Torino
- General Pathology and Immunology (1 credits) - School of Specialization in Hospital Pharmacy, University of Torino
- Molecular and genetic basis of diseases (1 credits) - Pharmaceutical Chemistry and Technology, University of Torino

### Research main topics

**Pathology/Chronic liver diseases – basic and translational studies:** main research interests are related to the analysis of molecular, cellular and tissue events involved in the progression of chronic liver disease (CLD) towards hepatocellular carcinoma (HCC), with a focus on chronic inflammation and phenotypic and functional responses of macrophages during CLD progression. Other recent point of interest is on the identification of new drug-delivery system to improve/modulate the uptake of new liposome formulations by liver cancer cells and human monocytic cells.

### Main projects as PI:

2012. Collaboration with Istituto Zooprofilattico Sperimentale del Piemonte, Liguria e Valle d'Aosta: "Le terre rare: una nuova generazione di promotori di crescita?" (IZS PLV 22/12 RC - BOCC01AE13)

2009. Ricerca Sanitaria Finalizzata, regione Piemonte funds

2008. Ricerca Sanitaria Finalizzata, regione Piemonte funds

2005-2008, 2012, 2016-2021: Local Research funds, University of Torino

### Bibliometry (1993-present)

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### Publications

1. Gadoni E, Olivero A, Miglietta A, **Bocca C**, Gabriel L, Cytoskeletal modifications induced by 4-hydroxynonenal, *Cytotechnology* 1993;1: S62-64.
2. Gadoni E, Gabriel L, Olivero A, **Bocca C**, Miglietta A. Antimicrotubular effect of calvatic acid and of some related compounds, *Cell Biochemistry and Function*, 1995;13:231-238.
3. Miglietta A, **Bocca C**, Gadoni E, Gabriel L, Rampa A, Bisi A, Valenti P, Da Re P. Interaction of geiparvarin and related compounds with purified antimicrotubular protein, *Anti-Cancer Drug Design* 1996; 11:35-48.
4. Miglietta A, Gadoni E, **Bocca C**, Gabriel L. Prooxidant agents induce modifications in the cytoskeletal organisation of fibroblasts, *ATLA* 1996;24:557-565.
5. Miglietta A, **Bocca C**, Rampa A, Bisi A, Gabriel L. Geiparvarin and derivatives in combination with taxol: effect on microtubular organization in 3T3 fibroblasts, *Anti-Cancer Drug Design* 1997; 12:607-620.
6. **Bocca C**, Caputo O, Cavalli R, Gabriel L, Miglietta A, Gasco MR. Phagocytic uptake of fluorescent stealth and non-stealth solid lipid nanoparticles, *International Journal of Pharmaceutics* 1998;175:185-193.
7. **Bocca C**. Taxol: a short history of a promising anticancer drug, *Minerva Biotechnologica* 1998;10:81-83.
8. Cavalli R, **Bocca C**, Miglietta A, Caputo O, Morel S, Gasco MR. Albumin adsorption on stealth and non-stealth solid lipid nanoparticles, *S.T.P. Pharma Sciences* 1999;9:183-189.
9. Miglietta A, Cavalli R, **Bocca C**, Gabriel L, Gasco MR. Cellular uptake and cytotoxicity of solid lipid nanospheres (SLN) incorporating doxorubicin or paclitaxel, *International Journal of Pharmaceutics* 2000;210:61-67.
10. Miglietta A, **Bocca C**, Gabriel L, Rampa A, Bisi A, Valenti P. Antimicrotubular and cytotoxic activity of geiparvarin analogs, alone and in combination with paclitaxel, *Cell Biochemistry and Function* 2001;19:181-189.
11. **Bocca C**, Gabriel L, Miglietta A. Cytoskeletal-interacting activity of geiparvarin, diethylstilbestrol and conjugates, *Chemico-Biological Interactions* 2001;137:285-305.
12. Miglietta A, **Bocca C**, Gabriel L. Comparative studies on biological activity of certain microtubule-interacting taxanes, *Chemico-Biological Interactions* 2002;139:283-299.

13. **Bocca C**, Gabriel L, Bozzo F, Miglietta A. Microtubule-interacting activity and cytotoxicity of the prenylated coumarin ferulenol, *Planta Medica* 2002;8:1135-1137.
14. Miglietta A, Gabriel L, Appendino G, **Bocca C**. Biological Properties of jatrophone polyesters, new microtubule-interacting agents, *Cancer Chemotherapy and Pharmacology* 2003;51:67-74.
15. **Bocca C**, Gabriel L, Bozzo F, Miglietta A. A sesquiterpene lactone, costunolide, interacts with microtubule protein and inhibits the growth of MCF-7 cells, *Chemico-Biological Interactions* 2004;147:79-86.
16. Miglietta A, Panno ML, Bozzo F, Gabriel L, **Bocca C**. Insulin can modulate MCF-7 cell response to paclitaxel, *Cancer Letters* 2004;209:139-145.
17. Miglietta A, Bozzo F, Gabriel L, **Bocca C**. Microtubule-interfering activity of parthenolide, *Chemico-Biological Interactions* 2004;149:165-173.
18. Marino G, Motta E, Mosso L, **Bocca C**, Ravarino N, Torchio B. Primary signet ring cell adenocarcinoma of the bladder, *Minerva Urol Nefrol* 2005;57:125-7.
19. Miglietta A, Bozzo F, **Bocca C**, Gabriel L, Trombetta A, Belotti S, Canuto RA. Conjugated linoleic acid induces apoptosis in MDA-MB-231 breast cancer cells through ERK/MAPK signalling and mitochondrial pathway, *Cancer Letters* 2006;234:149-57.
20. Miglietta A, Bozzo F, Gabriel L, **Bocca C**, Canuto RA. ERK1/2 and protein phosphatase 2A are involved in the antiproliferative activity of conjugated linoleic acid in MCF-7 cells, *British Journal of Nutrition* 2006;96:22-27.
21. **Bocca C**, Bozzo F, Gabriel L, Miglietta A. Conjugated linoleic acid inhibits Caco-2 cell growth via ERK-MAPK signaling pathway, *Journal of Nutritional Biochemistry* 2006;18: 332-340.
22. **Bocca C**, Bozzo F, Francica S, Colombatto S, Miglietta A. Involvement of PPAR $\gamma$  and E-cadherin/ $\beta$ -catenin pathway in the antiproliferative effect of conjugated linoleic acid in MCF-7 cells, *International Journal of Cancer* 2007;121:248-56.
23. Bozzo F, **Bocca C**, Colombatto S, Miglietta A. Antiproliferative effect of conjugated linoleic acid in Caco-2 cells: involvement of PPAR $\gamma$  and APC/ $\beta$ -catenin pathways, *Chemico-Biological Interactions* 2007;169:110-121.
24. **Bocca C**, Bozzo F, Martinasso G, Canuto RA, Miglietta A. Involvement of PPAR $\alpha$  in the growth inhibitory effect of arachidonic acid on breast cancer cells, *British Journal of Nutrition* 2008;100:739-750.
25. Bozzo F, Bassignana A, Lazzarato L, Boschi D, Gasco A, **Bocca C**, Miglietta A. Novel nitrooxy derivatives of celecoxib for the regulation of colon cancer cell growth, *Chemico-Biological Interactions* 2009;182:183-190.
26. **Bocca C**, Bozzo F, Cannito S, Colombatto S, Miglietta A. CLA reduces breast cancer cell growth and invasion through ER $\alpha$  and PI3K/Akt pathways, *Chemico-Biological Interactions* 2010;183:187-193.
27. **Bocca C**. Anti-cancer effect of CLA: a matter of multiple interactions between signalling pathways involving Src, IGF and PPAR $\gamma$  pathway? *Chemico-Biological Interactions* 2010;186:252-253.
28. Miglietta A, Toselli M, Ravarino N, Vencia W, Chiecchio A, Bozzo F, Motta M, Torchio B, **Bocca C**. COX-2 expression in human breast carcinomas: correlation with clinicopathological features and prognostic molecular markers, *Expert Opinion on Therapeutic Targets* 2010;14:655-64.
29. **Bocca C**, Bozzo F, Bassignana A, Miglietta A. Antiproliferative Effect of a Novel Nitro-oxy Derivative of Celecoxib in Human Colon Cancer Cells: Role of COX-2 AND Nitric Oxide, *Anticancer Research* 2010;30:2659-66.
30. **Bocca C**, Bozzo F, Bassignana A, Miglietta A. Antiproliferative effects of COX-2 inhibitor celecoxib on human breast cancer cell lines, *Molecular and Cellular Biochemistry* 2011;350:59-70.
31. **Bocca C**, Bozzo F, Cannito S, Parola M, Miglietta A. Celecoxib inactivates epithelial-mesenchymal transition stimulated by hypoxia and/or epidermal growth factor in colon cancer cells, *Molecular Carcinogenesis* 2012;51:783-795.

32. **Bocca C**, Bozzo F, Ievolella M, Miglietta A. A novel nitro-oxy substituted analogue of rofecoxib reduces human colon cancer cell growth, *Molecular and Cellular Biochemistry* 2012; 361:105-110.
33. Cannito S, Paternostro C, Busletta C, **Bocca C**, Colombatto S, Miglietta A, Novo E, Parola M. Hypoxia, hypoxia-inducible factors and fibrogenesis in chronic liver diseases, *Histology and Histopathology*, 2014; 29:33-44.
34. **Bocca C**, Ievolella M, Autelli R, Motta M, Mosso L, Torchio B, Bozzo F, Cannito S, Paternostro C, Colombatto S, Parola M, Miglietta A. Expression of COX-2 in human breast cancer cells as a critical determinant of epithelial-mesenchymal transition and invasiveness, *Expert Opinion on Therapeutic Targets*, 2014;18:121-35.
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36. **Bocca C**, Bozzo F, Miglietta A. COX2 Inhibitor NS398 Reduces HT-29 Cell Invasiveness Modulating Signalling Pathways Mediated by EGFR and HIF1- $\alpha$ , *Anticancer Research*, 2014;34: 1793-1800.
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39. Cannito S, Morello E, **Bocca C**, Foglia B, Benetti E, Novo E, Chiazza F, Rogazzo M, Fantozzi R, Povero D, Sutti S, Bugianesi E, Feldstein AE, Albano E, Collino M, Parola M. Microvesicles released from fat-laden cells promote activation of hepatocellular NLRP3 inflammasome: A pro-inflammatory link between lipotoxicity and non-alcoholic steatohepatitis. *PLoS One* 2017;1:1-22.
40. Novo E, Villano G, Turato C, Cannito S, Paternostro C, Busletta C, Biasiolo A, Quarta S, Morello E, **Bocca C**, Miglietta A, David E, Sutti S, Plebani M, Albano E, Parola M, Pontisso P. SerpinB3 Promotes Pro-fibrogenic Responses in Activated Hepatic Stellate Cells. *Sci Rep.* 2017; 7:3420-3430.
41. Morello E, Sutti S, Foglia B, Novo E, Cannito S, **Bocca C**, Rajsky M, Bruzzi S, Abate ML, Rosso C, Bozzola C, David E, Bugianesi E, Albano E, Parola M. Hypoxia-inducible factor 2 $\alpha$  drives nonalcoholic fatty liver progression by triggering hepatocyte release of histidine rich glycoprotein. *Hepatology* 2018;67:2196-2214.
42. Benedetto A, **Bocca C**, Brizio P, Cannito S, Abete MC, Squadrone S. Effects of the rare elements lanthanum and cerium on the growth of colorectal and hepatic cancer cell lines. *Toxicol In Vitro* 2018;46:9-18.
43. Foglia B, Cannito S, **Bocca C**, Parola M, Novo E. ERK Pathway in Activated, Myofibroblast-Like, Hepatic Stellate Cells: A Critical Signaling Crossroad Sustaining Liver Fibrosis. *Int J Mol Sci.* 2019;20:2700.
44. Pandolfi L, Frangipane V, **Bocca C**, Marengo A, Tarro Genta E, Bozzini S, Morosini M, D'Amato M, Vitulo S, Monti M, Comolli G, Scupoli, Fattal E, Arpicco S, Meloni F. Hyaluronic Acid-Decorated Liposomes as Innovative Targeted Delivery System for Lung Fibrotic Cells. *Molecules* 2019;24:3291.
45. Foglia B, Sutti S, Pedicini D, Cannito S, **Bocca C**, Maggiora M, Bevacqua MR, Rosso C, Bugianesi E, Albano E, Novo E, Parola M. Oncostatin M, A Profibrogenic Mediator Overexpressed in Non-Alcoholic Fatty Liver Disease, Stimulates Migration of Hepatic Myofibroblasts. *Cells* 2019;9: 28.
46. Novo E, **Bocca C**, Foglia B, Protopapa F, Maggiora M, Parola M, Cannito S. Liver fibrogenesis: an update on established and emerging basic concepts. *Arch Biochem Biophys.* 2020;689:108445.
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48. Foglia B, Novo E, Protopapa F, Maggiora M, **Bocca C**, Cannito S, Parola M. Hypoxia, Hypoxia-Inducible Factors and Liver Fibrosis. *Cells.* 2021;10:1764.
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Prevents NASH-Related Liver Carcinogenesis by Decreasing Cancer Cell Proliferation. *Cell Mol Gastroenterol Hepatol.* 2022;13:459-482.

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51. Novo E, Cappon A, Villano G, Quarta S, Cannito S, **Bocca C**, Turato C, Guido M, Maggiora M, Protopapa F, Sutti S, Provera A, Ruvoletto M, Biasiolo A, Foglia B, Albano E, Pontisso P, Parola M. SerpinB3 as a Pro-Inflammatory Mediator in the Progression of Experimental Non-Alcoholic Fatty Liver Disease. *Front Immunol.* 2022 Jul 8;13:910526. doi: 10.3389/fimmu.2022.910526.

52. Bordano V, Kinsella GK, Cannito S, Dianzani C, Gigliotti CL, Stephens JC, Monge C, **Bocca C**, Rosa AC, Miglio G, Dianzani U, Findlay JBC, Benetti E. G protein-coupled receptor 21 in macrophages: An in vitro study. *Eur J Pharmacol.* 2022;926:175018. doi: 10.1016/j.ejphar.2022.175018.

53. **Bocca C**, Protopapa F, Foglia B, Maggiora M, Cannito S, Parola M, Novo E. Hepatic Myofibroblasts: A Heterogeneous and Redox-Modulated Cell Population in Liver Fibrogenesis. *Antioxidants* 2022,28;11::1278. doi: 10.3390/antiox11071278.

### **Books**

1. Miglietta A, **Bocca C**, Gadoni E, Gabriel L *Relationship between cytoskeletal alterations and cell proliferation in oxidative cell conditions*, in: *Advances in Free Radicals in Disease IV* (casa editrice *La Goliardica Pavese, Pavia, Italia*, editori Pizzala R, Tomasi A, Vannini V), 1999, 19-23.

2. Parola M, Autelli R, **Bocca C**, Costelli P, Muzio G, Novo E, Penna F, Tamagno E. *Patologia Generale ed Elementi di Fisiopatologia*, EdISES Università, 2020