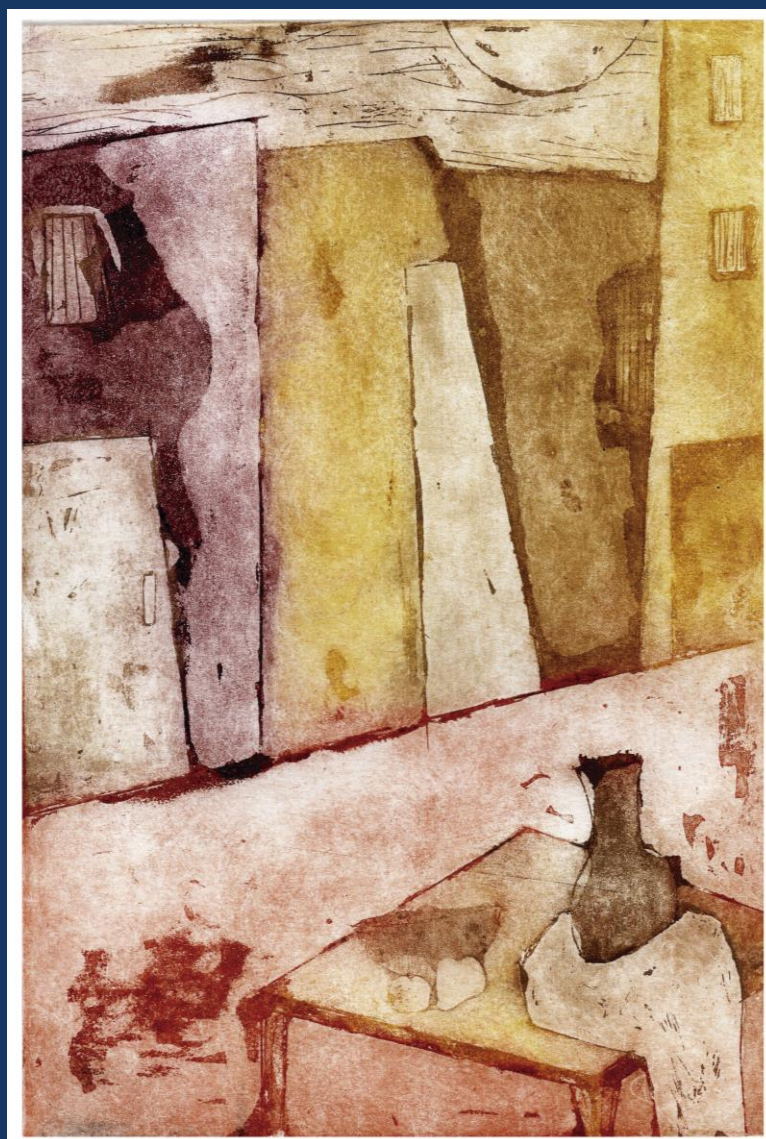


medicina

BUENOS AIRES VOL. 79 Supl. IV - 2019

80° Aniversario



medicina

BUENOS AIRES, VOL. 79 Supl. IV - 2019

COMITÉ DE REDACCIÓN

Pablo J. Azurmendi
Instituto de Investigaciones Médicas A. Lanari, UBA, Argentina
Damasia Becú Villalobos
Instituto de Biología y Medicina Experimental-CONICET, Buenos Aires, Argentina
José H. Casabé
Instituto de Cardiología y Cirugía Cardiovascular, Hospital Universitario Fundación Favaloro, Buenos Aires, Argentina
Eduardo L. De Vito
Instituto de Investigaciones Médicas A. Lanari, UBA, Argentina
Isabel Narvaiz Kantor
Organización Panamericana de la Salud (OPS/OMS) (ret.) Argentina
Basilio A. Kotsias
Instituto de Investigaciones Médicas A. Lanari, UBA, Argentina
Gustavo Kusminsky
Hospital Universitario Austral, Buenos Aires, Argentina
Isabel A. Lüthy
Instituto de Biología y Medicina Experimental (IBYME), Buenos Aires, Argentina

Daniel A. Manigot
Hospital San Juan de Dios, Buenos Aires, Argentina
Jorge A. Manni
Instituto de Investigaciones Médicas A. Lanari, UBA, Argentina
Rodolfo S. Martin
Facultad de Ciencias Biomédicas y Hospital Universitario Austral, Buenos Aires, Argentina
Guillermo D. Mazzolini
Instituto de Investigaciones en Medicina Traslacional-CONICET, Hospital Universitario Austral, Buenos Aires, Argentina
Rodolfo C. Puche
Facultad de Ciencias Médicas, Universidad Nacional de Rosario, Santa Fe, Argentina
Viviana Ritacco
Instituto Nacional de Enfermedades Infecciosas ANLIS-CONICET, Buenos Aires, Argentina
Guillermo B. Semeniuk
Instituto de Investigaciones Médicas A. Lanari, UBA, Argentina

MIEMBROS EMÉRITOS

Héctor O. Alonso
Instituto Cardiovascular Rosario, Santa Fe, Argentina
Guillermo Jaim Etcheverry
Facultad de Medicina, UBA, Argentina

María Marta de Elizalde de Bracco
IMEX-CONICET-Academia Nacional de Medicina, Argentina
Christiane Dosne Pasqualini
Academia Nacional de Medicina, Argentina

La Tapa (Ver pág. 4)
Atardecer en la tarde
Antonella Ricagni

MEDICINA (Buenos Aires) – Revista bimestral – ISSN 0025-7680 (Impresa) – ISSN 1669-9106 (En línea)

REVISTA BIMESTRAL

Registro de la Propiedad Intelectual N° 02683675

Personería Jurídica N° C-7497

Publicación de la Fundación Revista Medicina (Buenos Aires)

Propietario de la publicación: Fundación Revista Medicina

Queda hecho el depósito que establece la Ley 11723

Publicada con el apoyo del Ministerio de Ciencia, Tecnología e Innovación Productiva.

MEDICINA no tiene propósitos comerciales. El objeto de su creación ha sido propender al adelanto de la medicina argentina.

Los beneficios que pudieran obtenerse serán aplicados exclusivamente a este fin.

Aparece en MEDLINE (PubMed), ISI-THOMSON REUTERS (Journal Citation Report, Current Contents, Biological Abstracts, Biosis, Life Sciences), CABI (Global Health), ELSEVIER (Scopus, Embase, Excerpta Medica), SciELO, LATINDEX, BVS (Biblioteca Virtual en Salud), DOAJ, Google Scholar y Google Books.

Incluida en el Núcleo Básico de Revistas Científicas Argentinas del CONICET.

Directores Responsables:

Basilio A. Kotsias, Eduardo L. De Vito, Isabel Narvaiz Kantor, Guillermo B. Semeniuk

Secretaría de Redacción: Ethel Di Vita, Instituto de Investigaciones Médicas Alfredo Lanari, Combatientes de Malvinas 3150,
1427 Buenos Aires, Argentina
Tel. 5287-3827 Int. 73919 y 4523-6619
e-mail: revmedbuenosaires@gmail.com – http://www.medicinabuenosaires.com

Vol. 79, Supl. IV, Noviembre 2019

REUNIÓN ANUAL DE SOCIEDADES DE BIOCIENCIA 2019

**LXIV Reunión Anual de la
Sociedad Argentina de Investigación Clínica (SAIC)**

**LI Reunión Anual de la
Asociación Argentina de Farmacología Experimental (SAFE)**

**XXI Reunión Anual de la
Sociedad Argentina de Biología (SAB)**

**XXXI Reunión Anual de la
Sociedad Argentina de Protozoología (SAP)**

**IX Reunión Anual de la
Asociación Argentina de Nanomedicinas
(NANOMED-ar)**

**VI Reunión Científica Regional de la Asociación Argentina
de Ciencia y Tecnología de Animales de Laboratorio
(AACyTAL)**

**con la participación de
The Histochemical Society**

**13 - 16 de noviembre de 2019
Hotel 13 de Julio - Mar del Plata**

EDITORES RESPONSABLES

**Dra. Mónica Costas
Dra. Gabriela Marino
Dr. Pablo Azurmendi**

medicina

BUENOS AIRES, VOL. 79 Supl. IV - 2019

ANNUAL MEETING OF BIOSCIENCE SOCIETIES 2019

**LXIV Annual Meeting of
Sociedad Argentina de Investigación Clínica (SAIC)**

**LI Annual Meeting of
Asociación Argentina de Farmacología Experimental (SAFE)**

**XXI Annual Meeting of
Sociedad Argentina de Biología (SAB)**

**XXXI Annual Meeting of
Sociedad Argentina de Protozoología (SAP)**

**IX Annual Meeting of
Asociación Argentina de Nanomedicinas
(NANOMED-ar)**

**VI Regional Scientific Meeting of Asociación Argentina
de Ciencia y Tecnología de Animales de Laboratorio
(AACyTAL)**

**with the participation of
The Histochemical Society**

November 13th – 16th, 2019
Hotel 13 de Julio - Mar del Plata

CHIEF EDITORS

**Dra. Mónica Costas
Dra. Gabriela Marino
Dr. Pablo Azurmendi**

IFEG CONICET (1); INICSA CONICET (2); UNIVERSIDAD NACIONAL DE CÓRDOBA (3)

The *Salvia hispanica* L., whose common name is Chia, is an annual herbaceous plant belonging to the Lamiaceae family. Chia seeds have been investigated and recommended due to their high levels of essential fatty acids with the highest proportion of omega-3 fatty acid (ω 3-FA) compared to other natural sources known to date. It is well known that the consumption of ω 3-FA can slow or stop the growth of cancer cells. This work contains the preliminary results of our study about the effects of Chia oil (ChO) in a murine mammary gland adenocarcinoma by micro-X ray fluorescence (micro-XRF) analysis. The histological and multielemental analysis by micro-XRF has been combined to infer information about the mechanisms related with the anti-tumorigenic actions of ω 3-FA. It is an experimental study in which 24 mice BALB/C were used, randomly distributed between two dietetic groups: 12 with normal diet (control) and 12 with diet rich in ChO. Three months after starting the diet, they were inoculated with mammary gland cells with moderate metastatic capacity (M3). The animals were slaughtered 45-50 days after inoculation. Two adjacent cuts were made to each sample, one of 6 micrometers for conventional histological analysis and another of 30 micrometers placed in a Kapton film for measurement with micro XRF. Micro XRF measurements were performed at the Brazilian Synchrotron Light Laboratory (LNLS) in Brazil. Variation of the values of certain chemical elements of the samples treated with ChO with respect to the control was observed. Calcium (Ca) and Zinc (Zn) showed significant decreases in samples with ChO. Phosphorus (P) was located in regions of tumor cells. The results were analyzed by t-test. The distribution and concentration of Ca, Zn and P in samples can help us explain the anti-tumorigenic role of ω 3-FA as biomarkers of metabolic functions.

0953 - MICROFLUIDIC-ASSISTED SYNTHESIZED OF OXALIPLATIN NANOVEHICLES COMBINED WITH CURCUMIN POLYMERIC MICELLES APPLIED TO CHEMORESISTANT COLORECTAL CANCER TREATMENT

Rodrigo LLOYD(1) | Elena María SANMARCO(2) | Dailenys ESPINOSA MARTINEZ(1) | Mario Alberto GADAN(3) | Julia GALLINO(2) | Florencia GIANNONI(2) | Marcela MORETON(4) | Diego CHIAPPETA(4) | Juan Martín CABAILEIRO(2) | **Lucía POLICASTRO (2)**

LABORATORIO DE NANOMEDICINAS, COMISIÓN NACIONAL DE ENERGÍA ATÓMICA-ANPCYT (1); LABORATORIO DE NANOMEDICINAS, COMISIÓN NACIONAL DE ENERGÍA ATÓMICA-CONICET (2); COMISIÓN NACIONAL DE ENERGÍA ATÓMICA (CNEA) (3); DEPARTAMENTO DE TECNOLOGÍA FARMACÉUTICA, FACULTAD DE FARMACIA Y BIOQUÍMICA, UBA-CONICET (4)

Resistance and metastatic recurrence are the main barriers for the effective treatment of cancer. Administration of antineoplastic drugs in 100 nm-size nanovehicles (NV) optimizes the localization of drugs in tumor tissue. This occurs mainly due to the enhancement permeability retention effect that reduce peripheral toxicity and increase the tumor local therapeutic effectiveness. Oxaliplatin (Oxp), is a high efficiency chemotherapeutic drug but with severe adverse effects, thus this encapsulation in NV could significantly improve the effect. Moreover, the combination of this chemotherapeutic drug with sensitizing molecules, such as curcumin (Cur), could yet increase the possibilities of therapy success. This molecule is highly hydrophobic, so its encapsulation in NV also could improve its delivery. However, NV conventional synthesis technologies are inefficient with variations in the batch-to-batch procedures, which make it difficult for a rapid translation to patients. In recent years, microfluidic technology-assisted nanomedicines synthesis, where fluids are subtly controlled, significantly improve these processes and allow reproducibility between different batches. The aim of this work is to administrate Oxp in liposomes and Cur in polymeric micelles performed by

microfluidic technology in order to improve chemoresistant colorectal cancer treatment. We developed a micromixer chip for the encapsulation of Oxp in liposomes, obtaining optimal conditions in size and % of encapsulation of Oxp. Both compounds encapsulated, alone or in combination, were tested in Oxp resistant subcutaneous tumor in nude mice developed by our group and administrated by intravenous injection. The administration of the combination of encapsulated drugs produced a high effect in tumor growth with only one dose application. These results could have a high impact on the synthesis and the encapsulation of oncologic drugs in liposomes in the national and regional pharmaceutical industry.

0970 - INHIBITION OF PORCUPINE HAS A DIFFERENTIAL IMPACT ON THE GENE EXPRESSION SIGNATURE OF BREAST CANCER CELL LINES OF DIFFERENT LEVEL OF AGGRESSIVENESS.

Sofia VALLA(1) | Gianina DEMARCHI(1) | Agustina CHIMENTO(1) | **Nadia BONADEO (1)** | María Lucía ROMANO(1) | Laura Daniela ALANIZ(1) | Martin GOTTE(2) | Carolina CRISTINA(1)

CENTRO DE INVESTIGACIONES BÁSICAS Y APLICADAS (CIBA) – CITNOBA (UNNOBA – CONICET) (1); KLINIK FÜR FRAUENHEILKUNDE UND GEBURTSHILFE, UNIVERSITÄTSKLINIKUM MÜNSTER (2)

Wnt pathway is involved in cellular processes which are dysregulated in cancer as cell renewal, proliferation and EMT. In particular, in breast cancer (BC) it has a role in both tumor initiation and progression. Porcupine's palmitoylation of Wnt ligands is required for their proper signaling and release and its inhibition showed anti-tumoral effects on different BC models. In order to study if the inhibition of Porcupine (PORCN) had different effects regarding to the aggressiveness of the BC cell lines, we analyzed the impact of IWP-2, a PORCN inhibitor, in the TNBC cell line MDA-MB-231 and in the less aggressive MCF-7 cell line. Cells were treated with 5 μ M IWP-2 or DMSO as control. IWP-2 reduced the synthesis of Wnt ligands and target genes in MCF-7 but not in MDA-MB-231 cells (qPCR, 24 h, * $p < 0.05$: WNT3A*, AXIN2*, CCND1*, C-MYC*, CCNB1*). We also found a decrement in CYCLIN B1 protein levels in MCF-7 cells. However, it reduced the expression of β -CATENIN in both cell lines (WB, 48 h). Furthermore, cell number was also reduced in both cases (TB, 72 h). When analyzing stem cell markers, SOX2 mRNA was decreased and KLF-4 increased in MCF-7 and MDA-MB-231 suggesting that PORCN inhibition has a similar effect on the stem-cell phenotype in both cell lines. However, the number of colonies was reduced in MDA-MB-231 but not in MCF-7 cells (CFA, 8 d). In hanging drop assays (6 d), both cell lines formed bigger but less compact mammospheres than controls, with a stronger effect in MCF-7 cells, suggesting a loss of cell adhesion in this cell line. This was also consistent with reduced E-CADHERIN expression in MCF-7 cells. Regarding EMT, no differences were found in SNAIL-1, ZEB-1 and E-CADHERIN expression at the mRNA level with IWP-2 treatment although TGF- β synthesis was decreased in MCF-7 cells. Our results suggest that the differences in the genetic background and phenotype of breast cancer cells can modulate the effect of PORCN inhibition over tumoral properties in vitro.

1017 - MUC4 IS THE PRINCIPAL MEDIATOR OF TNF-INDUCED TRASTUZUMAB RESISTANCE AND FOSTERS AN IMMUNOSUPPRESSIVE TUMOR MICROENVIRONMENT IN HER2+ BREAST CANCER

Sofia BRUNI(1) | Mara DE MARTINO(1) | Florencia MAURO(1) | María Florencia MERCOGLIANO(2) | Lucía SANTA MARÍA DE LA PARRA(1) | Patricia ELIZALDE(1) | Roxana SCHILLACI(1)

IBYME-CONICET (1); FACULTAD DE CIENCIAS EXACTAS Y NATURALES - UBA (2)