

1ª edição - Jornadas do Programa Doutoral em
Metabolismo – Clínica e Experimentação

4 de novembro de 2022

BOCK OF ABSTRACTS



COMISSÃO CIENTÍFICA

Raquel Soares	Department Of Biomedicine, Chair Of The Scientific Committee; PDMCE Scientific Committee
Fátima Martel	Department Of Biomedicine, CIM-FMUP PDMCE Scientific Committee
João Tiago Guimarães	Department Of Biomedicine, PDMCE Scientific Committee
Pedro Von Hafe	Department Of Biomedicine, PDMCE Scientific Committee
André Sousa	LaBMI-FMUP-i3S
Catarina Neves	Centro Hospitalar de Entre Douro e Vouga
Catarina Rocha	LaBMI-FMUP-i3S
Cláudia Diogo	Hospital da Luz
Filipe Araújo	FMUP
Francisca Carmo	Department of Biomedicine, CIM-FMUP
Gonçalo Soares	Centro Hospitalar Universitário do Porto
Juliana Morais	UniC-FMUP, CINTESIS@NMS FCM
Sara Sá	LaBMI-FMUP-i3S

FACULDADE DE MEDICINA | UNIVERSIDADE DO PORTO

jornadas_pdmce@med.up.pt

Índice

1. <i>Introdução ao tema das jornadas</i>	4
2. <i>Resumo dos Oradores Convidados</i>	5
2.1. <i>Into Diabetesity</i>	5
<i>Diabetesity in numbers – where are we and where to go? - Helena Trigueiro</i>	5
<i>New biomarkers for diagnosis of diabetesity - João Sérgio Neves</i>	5
<i>Mitochondria – The missing link in diabetesity - Edoardo Bertero</i>	5
2.2. <i>Diabetesity-related diseases</i>	6
<i>Type 3 Diabetes - Paula Moreira</i>	6
<i>MAFLD – Naomi Lange</i>	6
<i>Depression and anxiety – Margarida Figueiredo Braga</i>	7
2.3. <i>New therapies – issues and questions</i>	7
<i>Fecal transplant and probiotics – Diogo Pestana</i>	7
<i>Diabetic treatment by hyperbaric approach – Daniela Martins-Mendes</i>	7
<i>Industry in Diabetesity: latest trends – Jorge Caria</i>	7
3. <i>Resumos selecionados para Comunicação Oral</i>	9
3.1. <i>CO01 – Inês Vasconcelos</i>	9
3.2. <i>CO02 – Ana Luísa de Sousa Coelho</i>	9
3.3. <i>CO03 – Bárbara Mota</i>	10
3.4. <i>CO04 – Inês Castela</i>	11
3.5. <i>CO05 – Mário Fontes</i>	12
4. <i>Resumos selecionados para apresentação em Poster</i>	14
4.1. <i>P01 – Elisabete Teixeira</i>	14
4.2. <i>P02 – Carla Luís</i>	14
4.3. <i>P03 – Marco G. Alves</i>	15
4.4. <i>P04 – Catarina Isabel dos Santos Rodrigues</i>	16
4.5. <i>P05 – Sofia Martinho Dimitri Pinheiro</i>	17
4.6. <i>P06 – Manuela Meireles</i>	18
4.7. <i>P07 – Alexandra Aveiro</i>	19
4.8. <i>P08 – Sofia João dos Santos Nogueira</i>	19
4.9. <i>P09 – Patrícia Braga</i>	20
4.10. <i>P10 – Sofia Martinho Dimitri Pinheiro</i>	21
4.11. <i>P11 – Lídia Cristina Alves da Rocha</i>	22
4.12. <i>P12 – Cátia Ramos</i>	23
4.13. <i>P13 – Anna Carolina Cortez-Ribeiro</i>	23

Conclusions: Diabetes can negatively influence outcomes in Sftf patients and may impact the decision of which specific procedure technique should be employed. Further studies are necessary to define how diabetes influences response to ultrasound-guided release of the A1 pulley in Sftf, as well as the extent to which control of blood sugar levels can contribute towards the personalization and optimization of patient follow up.

Key words: collagen; diabetes; interventional radiology; trigger finger; ultrasound

4.6. P06 – Manuela Meireles

Postprandial glycemia after a high-rich carbohydrate meal: a randomized cross-over clinical trial on olive leaf tea effect

Manuela Meireles^{1,2*}; Denise Polck³; Anna Carolina Ribeiro³; Juliana Almeida-de-Souza^{1,2}; Vera Ferro-Lebres^{1,2}

Centro de Investigação da Montanha, Instituto Politécnico de Bragança, Bragança, Portugal ¹; Laboratório para Sustentabilidade e Tecnologia em Regiões de Montanha, Instituto Politécnico de Bragança, Bragança, Portugal ²; Instituto Politécnico de Bragança, Bragança, Portugal ³; *manuela.meireles@ipb.pt

Background: Infusions of olive leaves have been used in traditional herbal medicine as a way to treat and prevent many diseases, including diabetes. Olive leaves are naturally rich in oleuropein, and previous studies have shown the potential of oleuropein in mitigating diabetes and diabetes complications in vitro and in vivo. This study aimed to investigate the effect of natural olive leaves tea on postprandial glycemia in healthy volunteers, when ingested with a high-carbohydrate meal comparing with a placebo tea. The hypothesis present was that olive leaf tea would improve glycemic control and modulate postprandial glycaemia.

Methodology: Thirteen healthy adults participated in a double-blinded, randomized, placebo-controlled, and cross-over design trial. Participants ingested a test meal composed of 2 slices of wheat bread (110g) and 50g of apricot peach with olive leaf tea (OLT) or two slices of wheat bread (110g) and 50 g of apricot peach with 250 ml of placebo tea (CON) in two different moments, and after a wash-out period. Capillary blood glucose was measured at times 0, 15, 30, 60, 90 and 120 min after ingestion of each test meal.

Results: At baseline, there were no significant differences between capillary plasma glucoses measured before the CON or OLT interventions. Consumption of OLT resulted in a delay in peak time ($48,5 \pm 4,2$ min vs $35,7 \pm 4,0$ min, $p=0,03$) and a significant increase in glucose area under the curve compared to placebo ($14502,7 \pm 640,8$ vs $13633,3 \pm 869,4$, $p= 0.03$). No significant differences ($p<0.05$) between conditions at individual time points were denoted.

Conclusions: Olive leaf tea did not ameliorate a glycemic curve induced by carbohydrate rich meal ingestion, however OLT delay on glycemic peak should be further explored. Also, future studies should account for chronic consumption in order to provide a better understanding on glycemic regulation over time.

Key words: Olive leaf tea; postprandial glycemia; diabetes.

Trial registration: [ClinicalTrials.gov](https://clinicaltrials.gov) NCT05397509

Funding: This work was supported by the Foundation for Science and Technology (FCT, Portugal) for financial support through National Funds FCT/MCTES (PIDDAC) to CIMO (UIBD/00690/2020 and

UIPD/00690/2020) and SusTEC (LA/P/0007/2021); and the North Regional Operation Program funded by European Social Found – NORTE 2020 (MM, scientific- contract NORTE-06-3559-FSE-000188).

4.7. P07 – Alexandra Aveiro

The influence of adipocytes' secretome and immune response on bacterial growth

Alexandra Aveiro^{1,2*}, Carla Guedes^{1,4}, Catarina Teixeira^{1,3}, André Sousa^{1,5}, Sónia Mendo², Pilar Baylina^{1,4,6}, Ruben Fernandes^{1,4,7}, Ana Cláudia Pereira^{1,3}

Laboratory of Medical & Industrial Biotechnology (LaBMI), Porto Research, Technology & Innovation Center (PORTIC), Porto, Portugal ¹; Biology department, University of Aveiro, Aveiro Portugal ²; Institute for Research & Innovation in Health (I3S), Porto, Portugal ³; Faculty of Biology, University of Vigo, Vigo, Spain ⁴; Faculty of Medicine of Porto (FMUP), Porto, Portugal ⁵; School of Health (ESS), Polytechnic Institute of Porto (IPP), Porto, Portugal ⁶; Center of Clinical Studies (CECLIN) from Hospital Fernando Pessoa & Faculty of Health Sciences, University Fernando Pessoa, Porto, Portugal ⁷; * agouveiaaveiro@ua.pt

Background: Obesity and antimicrobial resistance are considered a threat to global public health. The state of obesity entails an imbalance in the production of proinflammatory and anti-inflammatory factors that contributes to infections' susceptibility. *Klebsiella pneumoniae* is part of the ESKAPE being one of the most virulent and antibiotic resistant bacterial pathogens, with high prevalence in hospital and community-acquired bacterial infections. Bacteria's alarming ability to gain resistance has raised concerning obstacles in antimicrobial therapies in patients with chronic inflammation, such as obese individuals.

This study aimed to understand the influence of an inflammatory obesity-mimicking environment in the growth of *Klebsiella pneumoniae* strains with different antibiotic resistance.

Methodology: The secretome of the cell lines Raw 264.7 (macrophages) and 3T3-L1 (adipocytes) were collected to serve as conditioned media for bacterial growth. Antibiotic-resistant *Klebsiella pneumoniae* strains (ATCC, carbapenem-resistance (CR), extended-spectrum beta-lactamase (ESBL) and CR/ESBL) were exposed to a variety of conditioned media (DMEM, DMEM with 10% and 50% adipocyte secretome (SA), macrophage secretome (SM), and macrophage secretome previously conditioned with 10% and 50% SA). Bacteria growth curves assessment was performed by absorbance measurement in a kinetic mode for a period of 5 days.

Results: Results showed that for ATCC strain the different mediums didn't provide distinct growth conditions between them. However, when different resistance signatures were considered, the growth profile showed distinct differences. The CR strain showed higher population in the mediums with SM, regardless the percentage of SA, while the ESBL strain presented higher population in the mediums with SA without SM. Interestingly, for 50% SA, having SM in the medium or not showed no difference. For the strain with both resistances (CR/ESBL), results were similar to the CR strain.

Conclusions: Overall, results showed that the resistance to carbapenems is a greater risk factor than resistance to extended spectrum cephalosporins in a SM environment.

Key words: Obesity; Antimicrobial Resistance; Secretome; Macrophages

4.8. P08 – Sofia João dos Santos Nogueira