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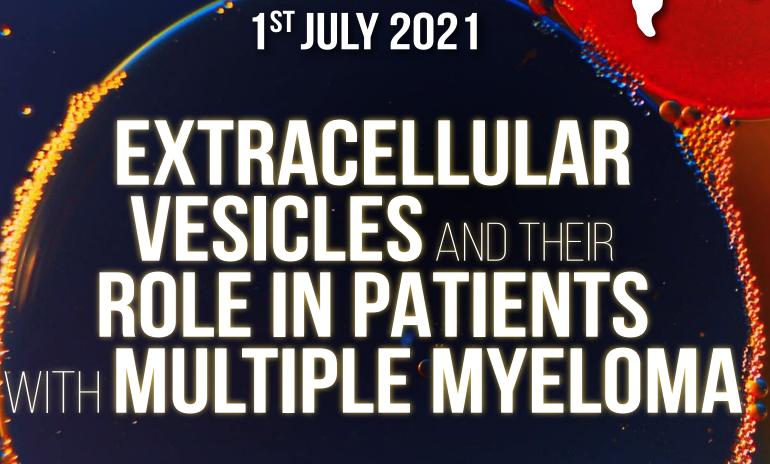
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10,5 CME/ECM FOR: PHYSICIANS, BIOLOGISTS, BIOMEDICAL LABORATORY TECHNICIANS, NURSES

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IN ASSOCIATION WITH





IN THE PAST DECADE, THERE HAVE BEEN MAJOR ADVANCES IN THE TREATMENT OF THE BLOOD CANCER MULTIPLE MYELOMA (MM). THE INTRODUCTION OF NOVEL AGENTS SUCH AS IMMUNE-MODIFYING AGENTS (IMIDS), PROTEASOME INHIBITORS, MONOCLONAL ANTIBODIES, WITH OR WITHOUT STEM CELL TRANSPLANTATION, HAS RESULTED IN SIGNIFICANTLY IMPROVED PATIENT SURVIVAL. MEANWHILE, THE INCREASED UNDERSTANDING OF MM TUMOR BIOLOGY HAS PROVIDED A RATIONALE FOR NEW COMBINATIONS OF DRUGS AND RISK-ADAPTED AND INDIVIDUALIZED TREATMENTS TO FURTHER IMPRO-VE PATIENT MANAGEMENT.

EXTRACELLULAR VESICLES (EVS) ARE CELL-DERIVED MEMBRANOUS PARTICLES THAT MEDIATE CELL-TO-CELL COMMUNICATION BY TRANSFERRING PROTEINS, LIPIDS AND NUCLEIC ACIDS LOCALLY AND THROUGH SYSTEMIC CIRCULATION. EVS ARE ACTIVE REGULATORS IN THE CROSS-TALK BETWEEN MM TUMOUR CELLS AND BONE MARROW MICROENVIRONMENT, WITH THE CAPACITY TO ALTER ANGIOGENESIS, OSTEOCLAST DIFFERENTIATION AND IM-MUNOSUPPRESSION, PROMOTING TUMOUR PROGRESSION AND DRUG RESISTANCE. CIRCULATING EVS CONTAINING TUMOUR-SPECIFIC MOLECULAR SIGNA-TURES (ONCOPROTEINS, RNAS, DNA FRAGMENTS) HAVE POTENTIAL CLINICAL UTILITY AS NEXT-GENERATION LIQUID BIOPSY BIOMARKERS IN CANCER DIAGNOSIS ANDMANAGEMENT, WITH THE POTENTIAL TO CHARACTERISE BOTH SPATIAL HETEROGENEITY AND CLONAL EVOLUTION THUS INFORMING NEW MODALITIES FOR DIAGNOSIS, RISK STRATIFICATION, MONITORING AND THERAPEUTIC INTERVENTION IN MM. HOWEVER, THE NANO-SCALE NATURE OF EVS AND THE COMPLEXITY OF BIOFLUIDS PRESENT CHALLENGES THAT NEED TO BE ADDRESSED BEFORE THE POTENTIAL OF EVS AS BIOMARKERS AND THERAPEUTIC TARGETS CAN BE ACHIEVED.

THE ITALIAN SOCIETY OF HEMATOLOGIC ONCOLOGY (SOHO ITALY) WAS ESTABLISHED AS A NON-PROFIT ORGANIZATION IN 2019 TO PROMOTE WORLDWI-DE RESEARCH (EDUCATION, PREVENTION, PRECLINICAL AND CLINICAL STUDIES AND PATIENT CARE) OF HEMATOLOGIC MALIGNANCIES AND RELATED DISORDERS. IN THIS SCENARIO, SOHO ITALY TOGETHER WITH AUSTRALIAN COLLEAGUES AIM TO BRING TOGETHER INTERNATIONAL EXPERTS TO DISCUSS THE LATEST ADVANCES IN THE PATHOPHYSIOLOGY AND THERAPY OF MM AND TO BETTER UNDERSTAND THE ROLE OF EVS IN PATIENTS WITH MM.



07.50 OPENING REMARKS

C. CERCHIONE D. W. GREENING G. MARTINELLI A. REALE A. SPENCER A. VACCA

SESSION 1 - THE MULTIPLE MYELOMAS

H. EINSELE A. SPENCER G. MARTINELLI **08.00** WHAT IS SOHO ITALY C. CERCHIONE G. MARTINELLI **08.20** THE MULTIPLE MYELOMAS - BIOLOGY, DIAGNOSIS, RISK STRATIFICATION J.L. HAROUSSEAU **08.40** ROLE OF MICROENVIRONMENT IN MM A. VACCA 09.00 IMMUNE SYSTEM IN MM P. NERI **09.20 ABSTRACT SUBMISSION 09.30** LECTURE LIQUID BIOPSY IN MM **A. SPENCER**

BREAK

SESSION 2 - UNDERSTANDING EXTRACELLULAR VESICLES

M. BEBAWY A. VACCA D. W. GREENING **10.20** EXTRACELLULAR VESICLES - OVERVIEW, UPDATE K. WITWER 10.40 EXTRACELLULAR VESICLES IN CANCER—IMPLICATIONS FOR FUTURE IMPROVEMENTS IN CANCER CARE A. RAI **11.00** EVS AS CANCER DIAGNOSTICS **A. MÖLLER 11.20** EV BYSTANDER SIGNALING AND CANCER RESISTANCE P. SAMUEL **11.40** TOOLS FOR TRACKING BIODISTRIBUTION OF CANCER EVS B. SUNG **12.00 ABSTRACT SUBMISSION**

12.10 STUDENT/ECR NETWORK ON EVS (SNEV), OVERVIEW A. NASIRI KENARI

SESSION 3 - HOW I MANAGE MULTIPLE MYELOMA

K.C. ANDERSON C. CERCHIONE M.V. MATEOS 12.55 HOW I MANAGE FRONTLINE MM M. V. MATEOS **13.15** HOW I MANAGE RELAPSED/REFRACTORY MM C. CERCHIONE **13.35** BIOLOGICALLY BASED THERAPIES FOR MM K.C. ANDERSON **13.55** NEW TREATMENT AVENUES IN MM H.C. LEE 14,15 MANAGING INFECTIONS IN MM R. RIA **14.35 ABSTRACT SUBMISSION**

BREAK

SESSION 4 - ROLE OF EXTRACELLULAR VESICLES IN MYELOMA A. REALE G. SIMONETTI B. SUNG 14.55 EVS IN MM BONE DISEASE K. VANDERKERKEN 15.15 EVS IN MM PROGRESSION A. ROCCARO 15.35 MM FIBROBLASTS ENHANCE BONE MARROW ANGIOGENESIS VIA SMALL EVS RELEASE I. SALTARELLA 15.55 MM-SMALL EVS, OMICS, PLASMA A. REALE **16.15 ABSTRACT SUBMISSION 16.25** | FCTURE PROTFOMIC INSIGHTS IN FVS: KEY PLAYERS IN CANCER AND POTENTIAL THERA-PFUTIC STRATEGY D. W. GREENING 16.55 CONCLUDING REMARKS C. CERCHIONE G. MARTINELLI A. VACCA

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