

*accepted abstracts*

Chairs:

Pietro Lampertico and Heiner Wedemeyer

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## DIAGNOSIS, STAGING, HCC RISK AND ANTIVIRAL THERAPY

### P1 Nearly One-Third of Veterans with Hepatitis Delta Virus Infection in the United States Have Already Developed Cirrhosis or Hepatocellular Carcinoma at Time of Diagnosis

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### P2 Quantification of plasma HDV RNA in untreated and bulevirtide-treated patients with CHD: a comparison between robogene 2.0, Eurobioplex and altostar

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### P3 Value and kinetics of virological markers in the natural course of chronic hepatitis D virus infection

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### P4 Hepatitis delta virus (HDV) replication through HBV integrants in HCC recurrence after liver transplantation

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### P5 Development and implementation of an hepatitis D detection and linkage to care program in Catalonia. Preliminary results.

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### P6 THE WHO HDV RNA International Standard does not reflect variability of real-world samples

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### P7 Bulevirtide: a success case after pegINF $\alpha$ failure

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### P8 The prevalence and characteristics of hepatitis B/D in 48,522 HBsAg tested individuals in Mongolia

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**P9 Validation of Streamlined serodiagnosis of hepatitis delta virus**

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**P10 Limited use of established risk scores for the prediction of hepatocellular carcinoma in patients with chronic hepatitis D virus infection**

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**P11 Is there a Hepatitis Delta Virus variant in West Africa, not detected by current serological diagnostic tests?**

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**P12 Novel anti-HDV therapies require reliable quantification of plasma HDV RNA: A European multicenter study**

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**P13 The asymmetry of the liver and spleen stiffness measures between patients with chronic hepatitis B and D reflects important clinic-pathologic differences**

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**P14 Diagnostic performances of different HDV RNA quantification assays used in clinical practice in Italy: results from a national quality control multicenter study**

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**P15 Hepatitis Delta Virus RNA quantification: a story about fruitful collaboration between private company and academic laboratory**

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**P16 Clinical evaluation of the Altostar HDV RT-PCR Kit 1.5**

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**P17 Hepatitis Delta infection in individuals living with HIV – multicentric portuguese study**

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**P18 HDV RNA assay sensitivity is critical for determining a correct outcome during Bulevirtide antiviral therapy**

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**P19 Treatment with pegIFNα inhibits cell division-mediated spread of HDV in a humanized mouse model supporting cell proliferation**

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**P20 Chronic delta hepatitis without efficient treatment poses a greater risk for hepatocellular cancer development than chronic hepatitis B with efficient treatment**

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**P21 Complexities of Hepatitis Delta Virus Testing in a High HBV Prevalence Setting in London**

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**P22 High frequency of liver cirrhosis in European patients with hepatitis D: Data from a large multicentre study (D-SOLVE and HDV-1000 consortia)**

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## CURRENT AND NEW ANTIVIRAL TREATMENTS

### P23 Bulevirtide for patients with chronic hepatitis d (CHD) in Italy: a multicenter prospective nationwide real-life study (D-SHIELD)

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### P24 Virological and clinical outcomes of patients with HDV-related cirrhosis treated with bulevirtide monotherapy for up to 96 weeks: a multicenter european study (SAVE-D)

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## P25 Bulevirtide monotherapy prevents liver decompensation in patients with hdv-related cirrhosis: a case control study with propensity score weighted analysis

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## P26 Results From an Integrated Analysis at Week 96: Continued Treatment of Early Virologic Non-responders or Partial Responders With Bulevirtide Monotherapy for Chronic Hepatitis Delta Leads to Improvement in Virologic and Biochemical Responses

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**P27 Efficacy and Safety of 144 Weeks of Bulevirtide 2 mg or 10 mg Monotherapy From the Ongoing Phase 3 Study MYR301**

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**P28 Forty-Eight-Week Off-Therapy Efficacy and Safety of Bulevirtide in Combination With Pegylated Interferon Alfa-2a in Patients With Chronic Hepatitis Delta: Final Results From the Phase 2b, Open-Label, Randomised, Multicentre Study MYR204**

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**P29 Treatment of chronic hepatitis D with bulevirtide: 1st year report**

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**P30 Bulevirtide monotherapy in patients with compensated cirrhosis and CSPH: a 96-week interim kinetic analysis of real-life setting**

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**P31 Efficacy and safety of tobeviribart (VIR-3434) alone or in combination with elebsiran (VIR-2218) in participants with chronic hepatitis delta virus infection: preliminary results from the Phase 2 SOLSTICE trial in non-cirrhotic and compensated cirrhotic participants**

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**P32 Improvement in Liver Histology Is Observed in Most Patients With Chronic Hepatitis Delta After 48 Weeks of Bulevirtide Monotherapy**

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**P33 Rapid reductions of HDV RNA and ALT with the monoclonal antibody, BJT-778: results from a phase 2 study**

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**P34 Treatment of chronic hepatitis delta with bulevirtide in Portugal: data from a real-life cohort**

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**P35 Detection and characterization of anti-preS1 antibodies in HDV-infected patients under Bulevirtide treatment**

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**P36 Treatment of HDV infection in Solid Organ Transplant with Bulevirtide: a case report**

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**P37 Patient-Reported Outcomes Among Patients With Chronic Hepatitis Delta Treated With Bulevirtide 2 mg: A Long-Term Analysis of the Phase 3 MYR301 Trial at 96 Weeks**

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**P38 Real life experience of HBV/HDV-related compensated cirrhosis treatment in an Italian prison. A case report.**

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**P39 Undetectable HDV RNA at 24 Weeks of Treatment With Bulevirtide and Pegylated Interferon Alfa-2a Combination Therapy Is an Important Predictor of Maintained Response Off-Therapy**

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**P40 Undetectable HDV RNA Defined as Target Not Detected at the End of Treatment With Bulevirtide and/or Pegylated Interferon Alpha-2a Is an Important Predictor of 48 Weeks Sustained Virologic Response in Chronic Hepatitis Delta**

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**P41 High Rates of Adherence to Bulevirtide Monotherapy for Chronic Hepatitis Delta Through 96 Weeks: Results From an Interim Analysis of the Phase 3 Study MYR301**

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**P42 Improvements in Fibrosis and Necroinflammation With Bulevirtide Combined With Pegylated Interferon for Chronic Hepatitis Delta**

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**P43 Efficacy and safety of BLV monotherapy for chronic hepatitis delta: post treatment results through 24 weeks after the end of treatment from an interim analysis of a randomized Phase 3 study MYR301**

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## VIROLOGY AND PATHOGENESIS

### P44 HIV-HBV-HDV co-infection. Case presentation.

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### P45 Differential patterns of HBV rna and HBcrAG levels in a large european cross-sectional study of untreated patients with chronic hepatitis delta

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### P46 HELZ2 is an interferon stimulated gene with antiviral properties against Hepatitis Delta Virus

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### P47 Hospitalised Adults With Hepatitis Delta Virus Infection Have Higher Risk of Disease Progression Than Those With Hepatitis B Virus Mono-infection in Italy

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### P48 Baseline characteristics and risk of liver-related events in hepatitis B and C coinfection with and without hepatitis D infection

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### P49 TIGIT-expression on natural killer cell subsets correlate with liver inflammation and bulevirtide treatment response in chronic hepatitis D

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### P50 Adenine Base editing of Hepatitis B surface antigen potently inhibits HDV

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### P51 Sequence Analysis of the Hepatitis D Virus (HDV) Across Genotypes shows Highly Conserved Regions and Evidence of Recombination

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**P52 Deciphering the cellular response to IFN treatment in HDV-infected cells**

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**P53 Epidemiology and natural history of chronic hepatitis B and D infections in France from 2013 to 2022**

T. Asselah<sup>1</sup>, V. Loustaud-Ratti<sup>2</sup>, J.P. Bronowicki<sup>3</sup>, E. Maugain<sup>4</sup>, A. Le Blevec<sup>4</sup>, C. Idelovici-Marchal<sup>4</sup>, C. Lacueille<sup>5</sup>, E. Lambourg<sup>5</sup>, S. Larrieu<sup>5</sup>, Charlotte Colombi<sup>4</sup>

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**P54 Healthcare Resource Utilisation and Costs Among Terminal, Hospitalised Adults With Hepatitis Delta Virus or Hepatitis B Virus Monoinfection in Italy**

P. Lampertico<sup>1,2</sup>, V. Perrone<sup>3</sup>, L.D. Esposti<sup>3</sup>, M. Leogrande<sup>3</sup>, C. Kim<sup>4</sup>, M. Rock<sup>4</sup>

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**P55 Rational immunotherapy design for clinical development against HDV using a synthetic DNA-prime and protein-booster strategy**

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**P56 Development of a rapid test for HDV-specific T cell characterization in whole blood**

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**P57 HDV persistence can be independent from the extent of HBV reservoir and can be sustained by HBsAg production mainly derived from HBV-DNA integration**

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**P58 Low expression of circulating liver-enriched miRNAs in anti-HDAg patients in absence of active viral replication.**

M.F. Corfese<sup>1,2</sup>, J. Pérez Garreta<sup>1</sup>, B. Pacín Ruiz<sup>1</sup>, A. Palom<sup>2,4</sup>, D. Tabernero<sup>1,2</sup>, A. Rando Segura<sup>1,2,4</sup>, E. Vargas Accarino<sup>2,3</sup>, J.C. Ruiz Cobo<sup>2,3</sup>, M. Riveiro Barciela<sup>2,3,5</sup>, M. Buti<sup>2,3,5</sup>

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**P59 Real world patient profile for individuals with hepatitis delta virus infection treated with bulevirtide 2mg in Europe**

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**P60 High diversity of the TCR repertoire in hepatitis delta virus patients with undetectable viral RNA**

M.F. Cortese<sup>1,2</sup>, A. Palom<sup>2,3</sup>, B. Pacín Ruiz<sup>1,2</sup>, F. Rudilla<sup>4,4</sup>, E. Enrich Randé<sup>4,5</sup>, M. Antón Iborra<sup>5</sup>, J.C. Ruiz-Cobo<sup>2,3</sup>, D. Tabernero<sup>1,2,6</sup>, M. Riveiro Barciela<sup>3,7</sup>, A. Rando-Segura<sup>1,2,8,9</sup>, M.J. Herrero<sup>4,5</sup>, M. Buti<sup>2,3,7</sup>

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**P61** Dichotomy between HBcrAg and pre-genomic HBV RNA in relation to HDV RNA response in patients with chronic hepatitis delta during bulevirtide treatment

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**P62** Plasma pre-S1 HBsAg levels during antiviral therapy with Bulevirtide in chronic hepatitis delta patients – any help in predicting response to therapy?

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**P63** HDV co-infection and HCV eradication in persons with HIV (PWH): data from the ICONA cohort

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**P64** Role of intrahepatic HDV reservoir in potentially modulating response to bulevirtide treatment at 24 weeks.

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**P65** Integrative transcriptomics and epigenomics reveals a viral footprint of chronic HDV infection in HBV co-infected chimeric livers

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**P66** Adaptive immunity may drive treatment outcome of pegylated IFN alpha in patients with chronic hepatitis D

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