Symposium on Plasma Supply Management
29-30 January 2019

Location: EDQM premises, Strasbourg, France
Working language: English

Draft Programme (subject to change)

Tuesday, 29 January 2019

08:00-09:00 Registration

09:00-09:15 Opening Remarks
   ➢ Dr Susanne KEITEL, EDQM Council of Europe

   General Introduction
   ➢ Dr Guy RAUTMANN, EDQM Council of Europe

   Moderators: Dr Johanna CASTREN, Finnish Red Cross Blood Service
               Dr Guy RAUTMANN, EDQM Council of Europe

09:15-09:30 Review of current practices for plasmapheresis reported in the survey
          conducted by TS093 Working Party
   ➢ Dr Rut NORDA, Uppsala University Hospital, Sweden

09:30-09:45 Voice of the Patients: Patient perspective and views; Current situation and
           future needs for patients in need for plasma derived medicinal products
   ➢ Dr Frank WILLERSINN, PLUS Plasma User Group, Belgium

09:45-10:00 Is yearly collection of recovered / apheresis plasma adequate to ensure
           European self-sufficiency of essential plasma derived medicinal products?
   ➢ Dr Paul STRENGERS, International Plasma Fractionation Association
     (IPFA), the Netherlands

10:00-10:10 Panel discussion and take home messages

10:10-10:15 Break
Obstacles to strategic independence of plasma for fractionation in Europe and way forward - Is the EU legislation/regulation itself a barrier in any way?

Donor motivation - how can the obstacles be resolved?
**Moderators:** Dr Daphne THIJSSEN-TIMMER, Sanquin, Netherlands
Dr Frederic BIGEY, Etablissement Francais du Sang, France

10:15-10:30 Viewpoint of blood donor associations
➢ Ms Alice SIMONETTI, International Federation of Blood Donor Organizations (IFBDO), Italy

10:30-10:45 Learnings from the Dutch attempt to convert whole blood donors in plasmapheresis donors
➢ Ms Anna VAN KLEEF, Sanquin, the Netherlands

10:45-11:00 Learnings from the French attempt to convert whole blood donors in plasmapheresis donors
➢ Dr Frederic BIGEY, French Blood Establishment (EFS), France

11:00-11:15 Experiences on donor motivation in the private sector
➢ Mr Joshua PENROD, Plasma Protein Therapeutics Association (PPTA), USA

11:15-11:30 Landscape of deferrals and donor protective selection criteria; their impact on loss of donors and the lessons learned from the Transpose project
➢ Dr Marian VAN KRAAIJ, Sanquin, the Netherlands

11:30-11:45 Panel discussion and take home messages

11:45-12:00 Coffee / tea break

Efficiency of collection practices - how can obstacles be resolved?
**Moderators:** Dr Morten Bagge HANSEN, Copenhagen Blood Center, Denmark
Dr Giuseppe MARANO, Italian National Blood Centre

12:00-12:15 Recovered plasma for fractionation
➢ Dr Rene BUECHEL, Baxalta GmbH, Switzerland

12:15-12:30 German Red Cross in Bavaria with both whole blood and plasma-centres
➢ Dr Franz WEINAUER, Blutspendedienst des BRK, Germany

12:30-12:45 Blood establishments efforts in Denmark to expanse plasma collections
➢ Dr Jorgen GEORGSEN, South Danish Transfusion Service - Tissue Center, Odense University Hospital, Denmark

12:45-13:00 Panel discussion and take home messages

13:00-14:30 Lunch break
National regulations versus regional regulations and current developments

Moderators: Dr Joanne PINK, Red Cross Blood Service, Australia
           Dr Rut NORDA, Uppsala University Hospital, Sweden

14:30-14:45  Germany: Regulation on minimal IgG level for individualised donor management and current changes in the volumes and donation intervals
            ➢ Dr Peter HELLSTERN, Center of Hemostasis and Thrombosis Zurich, Switzerland

14:45-15:00  Hungary: New regulation for plasmapheresis donors one whole blood donation per year plus registry
            ➢ Dr Klara BAROTI-TOTH, Hungarian National Blood Transfusion Service, Hungary

15:00-15:15  Italy: Regulation of plasma self-sufficiency programme
            ➢ Dr Giancarlo LIUMBRUNO, National Blood Centre, Italy

15:15-15:30  Czech Republic: Combined system of not-for-profit and commercial plasma collection
            ➢ Dr Petr TUREK, National Blood Transfusion Committee, Czech Republic

15:30-15:45  Slovenia: The ruling in the Medisanus case
            ➢ Dr Irena RAZBORSEK, Blood Transfusion Center of Slovenia, Slovenia

15:45-16:00  Belgium: Recent programme to increase plasmapheresis for plasma-derived medicinal product
            ➢ Dr Philippe VANDEKERCKHOVE, Red Cross Flanders, Belgium

16:00-16:15  Panel discussion and take home messages

16:15-16:45  Coffee / tea break

Round Table Summary

Moderators: Dr Stefaan VAN DER SPIEGEL, European Commission DG SANTE, Belgium
           Dr Johanna CASTREN, Finnish Red Cross Blood Service
           Dr Guy RAUTMANN, EDQM Council of Europe

16:45-17:45  Solutions or ways forward to remove obstacles to strategic independence of plasma.

17:45-18:00  Close of the first day
Wednesday, 30 January 2019

**Discuss donor safety, donor selection and donor management**

**Moderators:** Dr Johanna CASTREN, Finnish Red Cross Blood Service
Dr Sheila MacLennan, NHSBT Leeds, UK

09:00-09:20 Humoral and cellular immunology in otherwise healthy persons and comparison with the donor population
  ➢ Dr Joanne PINK, Red Cross Blood Service, Australia

09:20-09:40 Management of donors using IgG levels based eligibility criteria
  ➢ Dr Stephan KIESSIG, Ruhr-Plasma-Zentrum, Germany

09:40-10:00 Strategies on protection of iron stores in plasma donors
  ➢ Dr George SCHREIBER, Plasma Protein Therapeutics Association (PPTA), USA

10:00-10:20 Data on donor adverse reactions and short introduction of utilised adverse reactions/ adverse effects classification for plasma donations: the European view
  ➢ Dr Thomas BURKHARDT, DRK Blutspendedienst Baden-Wurttemberg/Hessen and Nord-Ost, Germany

10:20-10:40 Coffee / tea break

10:40-11:00 Data on donor adverse reactions and short introduction of utilised adverse reactions / adverse effects classification for plasma donations: the USA view
  ➢ Ms Mary GUSTAFSON, Plasma Protein Therapeutics Association (PPTA), USA

11:00-11:20 Analysis of donor safety data from TS093 survey to support the pros and cons for revising the recommendation in the Guide
  ➢ Dr Joanne PINK, Red Cross Blood Service, Australia

**Data from publications: short and long-term health effects on donors of plasmapheresis**

**Moderators:** Dr Morten Bagge HANSEN, Copenhagen Blood Center, Denmark
Dr Luis Larrea, Centro de Transfusion de la Comunidad Valenciana, Spain

11:20-11:40 Sipla Study
  ➢ Dr Stephan WALSEMANN, SCINOMED GmbH, Germany

11:40-12:00 Scandat
  ➢ Dr Gustaf EDGREN, Karolinska Institutet, Sweden

12:00-12:20 Panel discussion and take home messages
Round Table Summary

Moderators: Dr Rut NORDA, Uppsala University Hospital, Sweden
Dr Harald KLUTER, DRK-Blutspendedienst Baden-Württemberg - Hessen GmbH

12:20-13:00  Does available scientific evidence support revisions of standards in the Guide that would promote plasmapheresis collection consistent with donor safety?

13:00-14:30  Lunch break

Apheresis Equipment Manufacturers

Moderators: Dr Johanna CASTREN, Finnish Red Cross Blood Service
Dr Guy RAUTMANN, EDQM Council of Europe

14:30-14:40  Discussion of differences and difficulties, design features, process characteristics, safety monitoring, EBV calculations estimations, algorithms (response to questions received).

14:40-15:10  Apheresis Equipment Supplier Presentations
Speakers to be confirmed.

15:10-15:20  Questions & Answers Session

15:20-15:30  Close of the Symposium

More information is available here

Email: prdd@edqm.eu

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