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'***European Medicines Agency Parenteral Drug Association***
April 21st, 2018 - European Medicines Agency Senior Vice President Scientific and Regulatory Affairs GENERAL we would welcome a more extensive discussion of this issue"**ADAPT SMART Closing Event 21 22 March 2018 Budapest**
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April 25th, 2018 - The European Commission granted a marketing authorisation valid throughout the European Union for Nexavar Scientific Discussion European Medicines

Agency'

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April 21st, 2018 - European Medicines Agency read the Scientific Discussion medicines to treat cancer that are injected into the spinal canal'
'**Improving the Contribution of Regulatory Assessment**
April 1st, 2018 - In response to a recommendation from the Pharmaceutical Forum the European Medicines Agency and the European network for Health Technology Assessment initiated a collaboration with the aim to improve the contribution regulatory assessment reports can make to the assessment of relative effectiveness of medicinal products by health technology'

'***emainfo YouTube***
April 16th, 2018 - Video content published by the European Medicines Agency No medicine is 100 safe This video explains how side effects should be reported and what the European Medicines Agency does to monitor medicines to protect public health in Europe'

'**SOPP 8001 6 Procedures for Parallel Scientific Advice**
April 15th, 2018 - SOPP 8001 6 Procedures for Parallel Scientific Advice with European Medicines Agency EMA Version 2 Effective Date December 1 2013 I Purpose'
'***European Medicines Agency techtran co jp***
April 20th, 2018 - European Medicines Agency the CHMP decided that a thorough scientific discussion was necessary in order to establish the value and safe use of antimicrobial'

'**The ethics of biosimilars GaBI Journal**
April 17th, 2018 - the position of the European Medicines Agency or one of use of biosimilar granulocyte colony Scientific discussion 2007 cited 2012 Nov'
'***RhoVac AB participates in a discussion meeting with the***
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'**Introduction to the European Medicines Agency efanet org**
April 21st, 2018 - Introduction to the European Medicines Agency ? The main scientific principle used in the evaluation of ? Platform for discussion of public health issues"**GSK submits landmark IMPACT data to European Medicines**
February 14th, 2018 - GlaxoSmithKline plc and Innoviva Inc announce submission of landmark IMPACT data to the European Medicines Agency as part of a type II variation to support an expanded label for Trelegy Ellipta in Europe'
'**Presentation from Maria Isaac Scientific Advisor**
April 24th, 2018 - Maria Isaac Scientific Advisor European Medicines Agency presents the regulator s view'
'**11th Pharmacovigilance Conference Medicines for Europe**
April 23rd, 2018 - The 11 th Pharmacovigilance Conference will once more see leading All sessions will be followed by a panel discussion and European Medicines Agency"***SCIENTIFIC DISCUSSION European Medicines Agency***
April 26th, 2018 - Page 3 38 study has been conducted also in response to the regulatory advice given by the CPMP scientific advice October 1999 This new study is study 302 which was conducted between 2000 02'

'***Andrea Buzzi Scientific Administrator Product Team***
April 28th, 2018 - View Andrea Buzzi?s profile presso European Medicines Agency Location of independent experimental design and scientific discussion on technical"***RhoVac AB participates in a discussion meeting with the***
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'**Introduction to the European Medicines Agency efanet org**
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'**European Medicines Agency?s Postmarket Brookings**
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December 16th, 2016 - Assessment Report European Medicines Agency SCIENTIFIC DISCUSSION Information on the pharmaceutical formulation used in the study The commercial formulation"**HUMAN MEDICINES PROGRAMME TOPRA**
April 17th, 2018 - HUMAN MEDICINES PROGRAMME Anne De Bock the Heads of Medicines Agencies and European Medicines Agency NICE Scientific Advice Program'
'***RESEARCH Open Access Adaptive clinical trial designs for***
March 25th, 2018 - advice letters from the European Medicines Agency Amelie Elsäßer1 sented in Section Discussion A text search of scientific advice letters issued between'

'***GSK submits landmark IMPACT data to European Medicines***
February 14th, 2018 - GSK submits landmark IMPACT data to European Medicines Agency to The European Summary of Factors and Management s Discussion and Analysis of Financial'
'**Thorsten Vetter European Medicines Agency ResearchGate**
April 24th, 2018 - Thorsten Vetter of European Medicines Agency London EMA with expertise in Pharmacy Internal Medicine General Medicine Endocrinology Read 11 publications and contact Thorsten Vetter on ResearchGate the professional network for scientists'
'***15th EGA Regulatory and Scientific Affairs Conference***
April 9th, 2018 - 15th EGA Regulatory and Scientific European Medicines Agency EMA European Generic and Biosimilar medicines Association Panel Discussion composed of"**Public Assessment Report Scientific discussion Venastat**
April 29th, 2018 - This module reflects the scientific discussion for the approval of Venastat hard prolonged release capsule HMPC of the European Medicines Agency'

'**Re Availability of evidence of benefits on overall**
April 25th, 2018 - A detailed discussion of the examples mentioned in the but a scientific conclusion on the balance of the benefits and risks based European Medicines Agency'

'**European Medicines Agency Advagraf**
April 23rd, 2018 - European Union agency responsible for the protection of public and animal health through the scientific evaluation and supervision of medicines'

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April 15th, 2018 - SOPP 8001 6 Procedures for Parallel Scientific Advice with European Medicines Agency EMA Version 2 Effective Date December 1 2013 I Purpose'
'**3Rs in Safety Testing of Human and Veterinary Pharmaceuticals**
April 17th, 2018 - 3Rs in Safety Testing of Human and Veterinary Pharmaceuticals 9 European Medicines Agency Regulatory Perspective on 3Rs ?Under discussion''**Availability of evidence of benefits on overall survival**
April 24th, 2018 - Setting Publicly accessible regulatory and scientific reports on cancer approvals by the European Medicines Agency Discussion Summary of findings'
'**EUROPEAN MEDICINES AGENCY Journal of Medical Ethics**
April 19th, 2018 - EUROPEAN MEDICINES AGENCY Health and Medicines Agency and the Uppsala The PRACreached its scientific recommendation by consensus following plenary discussion'
'**The European Medicines Agency Review of Eribulin for the**
November 22nd, 2017 - The European Medicines Agency Review of Eribulin for the Treatment of Patients with Locally Advanced or Metastatic Breast Cancer Summary of the Scientific Assessment of the Committee for Medicinal Products for Human Use''**European Public Assessment Reports EPARs**
April 14th, 2018 - European Public Assessment Reports EPARs the European Medicines Agency Scientific discussion The EPAR may also include v EPAR'

'***Adaptive clinical trial designs for European marketing***
January 29th, 2014 - Adaptive clinical trial designs for European marketing authorization a survey of scientific advice letters from the European Medicines Agency'

'**The European Medicines Agency NPRA**
April 27th, 2018 - The European Medicines Agency Need a solid scientific base upon which to build an support outside the Committee and bring them coherently into the discussion'

'**Interview Nathalie Bere European Medicines Agency**
April 18th, 2018 - Interview Nathalie Bere European Medicines The European Medicines Agency they make an important contribution to the overall ?scientific discussion?'

'***European Medicines Agency Elaprase***
April 23rd, 2018 - European Union agency responsible for read the scientific discussion about Elaprase Every year the European Medicines Agency will review'
'**Scientific Discussion European Medicines Agency**
April 22nd, 2018 - CVMP 772 02 1 29 EMEA 2002 Scientific Discussion I SUMMARY OF THE DOSSIER This application is for an injectable solution of 0 42mg ml dexmedetomidine the dextrorotatory'
'**The European Medicines Agency Review of Ofatumumab**
April 24th, 2018 - The European Medicines Agency Review of Ofatumumab Arzerra® for the Treatment of Chronic Lymphocytic Leukemia in Patients Refractory to Fludarabine and Alemtuzumab Summary of the Scientific Assessment of the European Medicines Agency Committee for Medicinal Products for Human Use'
'**Interview Nathalie Bere European Medicines Agency**
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'**European Medicines Agency api ning com**
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April 24th, 2018 - Thorsten Vetter of European Medicines Agency European Medicines Agency EMA · Scientific Advice focusing on the key areas of discussion that included'

'**Pharmaceutical Committee European Commission**
April 3rd, 2018 - Pharmaceutical Committee ?Drug development is scientific and regulatory challenging Slide courtesy of the European Medicines Agency'
'**Clinical Trial Regulation Conference**
April 12th, 2018 - Clinical Trial Regulation Conference European Medicines Agency European Medicines Agency EMA European Union Panel Discussion with Q amp A''***European Medicines Agency Wikipedia***
April 25th, 2018 - The European Medicines Agency EMA is a European Union agency for the evaluation of medicinal products The EMA operates as a decentralised scientific agency'

'***Competences European Medicines Agency***
April 20th, 2018 - European Medicines Agency Science medicines health The network facilitate scientific exchange and discussion leading to paper preparation and publication'

'**Improving the Contribution of Regulatory Assessment**
April 1st, 2018 - In response to a recommendation from the Pharmaceutical Forum the European Medicines Agency and the European network for Health Technology Assessment initiated a collaboration with the aim to improve the contribution regulatory assessment reports can make to the assessment of relative effectiveness of medicinal products by health technology'

'**Scientific advice processes ? RWE Navigator**
April 25th, 2018 - Scientific advice processes and experts may be invited to contribute to the discussion European Medicines Agency scientific advice'

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April 22nd, 2018 - View Isabel Diaz Ugalde?s Veterinary medicines en European Medicines Agency ? Preparation of scientific advice reports for discussion at the'
'**Clinical trials for medicines apply for authorisation in**
December 17th, 2014 - responses to required CHM areas for discussion a summary of scientific advice from any member state or the European Medicines Agency'
'***11th Pharmacovigilance Conference Medicines for Europe***
April 25th, 2018 - The 11 th Pharmacovigilance Conference will once more see leading All sessions will be followed by a panel discussion and European Medicines Agency''**Stratified medicine in European Medicines Agency licensing**
January 16th, 2013 - Stratified medicine in European Medicines Agency licensing a systematic review of predictive biomarkers 20 the Scientific Discussion''**Report of EFA training for patient experts on allergy**
April 25th, 2018 - asthma and COPD on getting involved with the European Medicines Agency EMA medicines Scientific Advisory The discussion was very'

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April 9th, 2018 - Jane Moseley of European Medicines Agency European Medicines Agency EMA · Scientific We analysed the minutes of discussion meetings held at the European''**Stratified medicine in European Medicines Agency licensing**
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April 28th, 2018 - Andrea Buzzi Scientific administrator presso European Medicines Agency Location of independent experimental design and scientific discussion on technical'

'**European Medicines Agency api ning com**
April 21st, 2018 - European Medicines Agency read the Scientific Discussion also part given with certain other medicines if appropriate'

'**Critically Important Antibiotics in Veterinary Medicine**
April 14th, 2018 - As there is much discussion and the European Medicines Agency the European Medicines Agency published scientific advice on the risk to humans from''**Moxifloxacin European Medicines Agency MAFIADOC COM**
April 24th, 2018 - Public Declaration of Interests and Confidentiality Undertaking of European Medicines Agency EMA Scientific Committee Scientific Discussion European'
'**Public Assessment Report Scientific discussion Venastat**
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April 24th, 2018 - Public Declaration of Interests and Confidentiality Undertaking of European Medicines Agency EMA Scientific Committee Scientific Discussion European'

'***The European Medicines Agency Is Moving On And Theresa***
August 3rd, 2017 - Yesterday s announcement of the candidate list of cities applying to rehome the European Medicines Agency brings scientific institutes within the discussion''**European Medicines Agency International cooperation**
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April 17th, 2018 - the position of the European Medicines Agency or one of use of biosimilar granulocyte colony Scientific discussion 2007 cited 2012 Nov''***European Medicines Agency***
April 9th, 2018 - European Medicines Agency Following a change of policy the EMEA now publishes the full scientific discussion for South America and many Eastern European''**European Medicines Agency**
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'**Matthias Hofer Scientific Officer Orphan Medicines**
April 11th, 2018 - View Matthias Hofer?s profile on which are reflected in published European Medicines Agency Impact of scientific advice from the European Medicines Agency'

'***Enpr EMA European Network of Paediatric Research at the***
April 28th, 2018 - at the European Medicines Agency with the scientific support of the Enpr EMA European Network of Paediatric Research at the European Medicines'

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