

4TH ANNUAL PHARMACBIOTECH PATENT I ITIGATION EUROPE

21-22 January, 2025 (UPC Litigation Forum on 20 January) Amsterdam, Netherlands, Beurs van Berlage

Join Europe's Premier Healthcare IP Forum to Define, Shape and Create **Your Patent Litigation Strategy**



Gain Invaluable Industry Insights From **Our 55+ Expert In-House Speakers**



Benchmark Alongside 450+ Attendees From 35+ Different Countries



Build Strong Long-Lasting Connections During Dedicated Networking Times (15+ Hours)

Connect With The Right Company Through Our 50% In-House Counsel Audience

Kisaco has grown the conference over the last few years and it is one of the key conferences to go to in life sciences patent space. The organisation has been great and variety of topics covered also. Great event to attend"

SHOHTA UENO, REGENERON

LEADING IN HOUSE INSIGHTS FROM:



MANUEL NEETZ Head of IP - Diagnostic Imaging Siemens Healthineers

KRISTIN COOKLIN Global Head IP Counsel Recordati

NATALIA

WRIGHT

RAQUEL

Novo Nordisk

FRISARDI

Associate General

Counsel

Counsel

Assistant General Patent

KORA KNUCHEL Vice President IP & Legal Brucker

MATTHEW **O'NEIL** Director of European IP Glenmark Pharmaceuticals



Global Head of Patents Pierre Fabre Group



SHOHTA UENO Senior Director -Dispute Resolution

OUR INSTITUTIONAL AND JUDICIAL SPEAKING



KUPECZ Honourable Judge **Central Division Munich**

HEIKE Section





SERGIO NAPOLITANO General Counsel edicines for Europe

European Commi MARGOT KOKKE Honourable Judge

Case Handler

DG Competition

GIANLUCA

VASSALLO

Local Division Hague

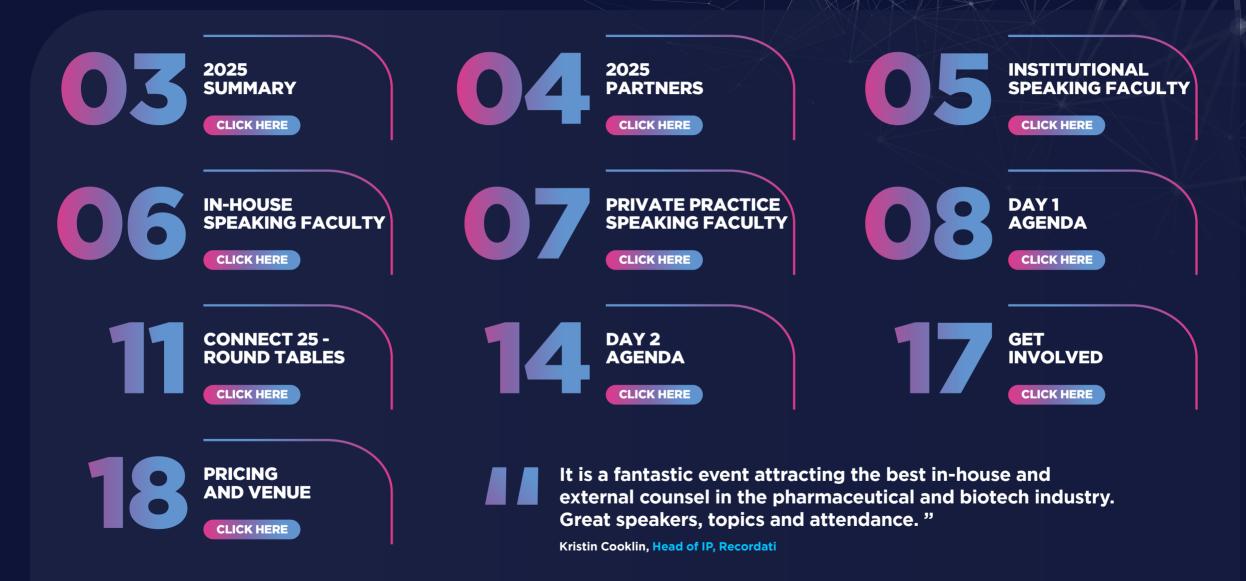


THOMAS Honourable Judge Local Division Dusseldorf - UPC

STEFAN LUGINBUEHL

Head Of Department -European Legal Affairs

ATTEND OUR PRE-EVENT UPC LITIGA-TION FORUM ON MONDAY 20 JANUARY!



WILL YOU BE ATTENDING THIS YEAR?

Stay ahead of the curve with insights on the first UPC decisions on the merits, second medical uses, injunctions, SPCs, competition law, patentability issues, and cross-border case law. This premier gathering of European life science patent litigation professionals returns for its 2025 edition. Join this elite community to stay updated on the latest legal developments in Europe and beyond, while forging essential connections with key industry players.

WHAT IS NEW FOR



Attend

Attend the industry's first-ever Diversity & Inclusion at the UPC workshop during our UPC Litigation Forum, offering targeted discussions and networking opportunities.



Benefit

Benefit from specialized and intimate litigation sessions tailored to the unique challenges of the MedTech, large molecule, and small molecule sectors, providing focused insights and strategies.



Discover

Discover our crossborder life science litigation review for 2025 and understand the trends taking place across the globe.

CONTENT FOR 2025

ANTITRUST IN LIFE SCIENCES

Explore the intersection of patent quality and divisional patenting with antitrust considerations, crucial for navigating competitive landscapes.

PATENT LITIGATION STRATEGY

Delve into cross-border case law, settlements, and UPC life science reviews to enhance your litigation approach.

REGULATORY UPDATES

Stay informed with key sessions on the latest European Commission reforms, Bolar Exemption case law, and a unitary SPC review, providing essential regulatory insights.

SECTOR -SPECIFIC FOCUS

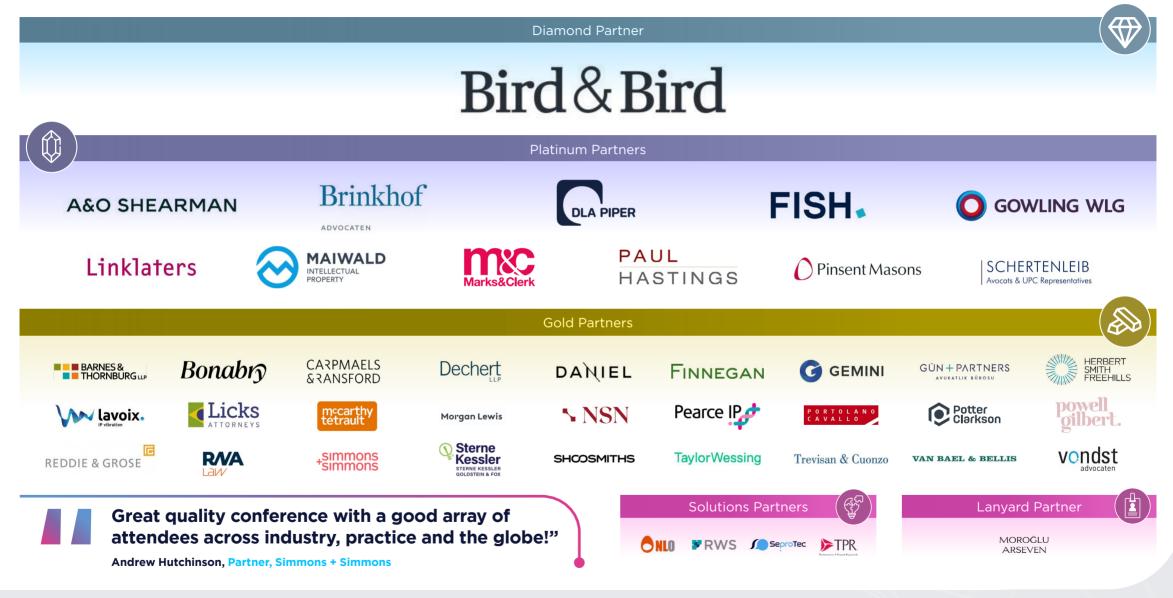
Access tailored sessions on patent litigation for the MedTech, chemical, and biopharmaceutical industries, ensuring targeted insights relevant to your field.

UPC LITIGATION FORUM 2025

Don't miss the cross-industry UPC Litigation Forum on Monday, January 20th, for comprehensive discussions on the latest UPC developments.



OUR 2025 PARTNERS



INSTITUTIONAL SPEAKING



ANDRÁS KUPECZ

Honourable Judge **Central Division Munich**

ELISABETTA EDGER BRINKMAN PAPA Honourable Judge, Technically Qualified

Local Division Hague Judge



FLORENCE BUTIN Honourable Judge-

Central Division Paris

GERBEN

HARTMAN

Lawyer



GIANLUCA VASSALLO Case Handler

DG Competition **European Commission**

ACCORNERO Legal Officer- IP Disputes Section

CHIARA

KAI HÄRMAND Nordic-Baltic Regional Division

PETER

THOMSEN

President

个

MARGOT KOKKE

Honourable Judge. Local Division Hague

MARTIN SCHMIDT Technically Qualified

Judge

MICHAEL

SWITA

Head of IP



RAINER FRIEDRICH

Technically Qualified Judge & Director Global IP UPC & CSL Behring



Division Dusseldordf

SERGIO THOMAS NAPOLITANO Honourable Judge, Local

General Counsel **Medicines For Europe**



SVEN BOSTYN Associate Professor of

Innovation Law University of Copenhagen





Head of Department &

European Legal Affairs

IN-HOUSE SPEAKING FACULTY

CLICK HERE TO BE INVOLVED!



PRIVATE PRACTICE AND CONSULTANCY SPEAKERS



INTERACTIVE ROUNDTABLE SPEAKERS



VIEW OUR UPC LITIGATION FORUM SPEAKERS HERE

CLICK HERE TO BE INVOLVED!

个

21 January 2025

8.00

Registration and Light Networking Breakfast

09:00

Chairs' Opening Remarks



09:10

Cross-Border Patent Litigation Considerations in the Life Sciences Sector 2025

Cross-border litigation poses unique challenges and opportunities for the pharmaceutical and biotech industries. This panel will explore the complexities of managing patent litigation across multiple jurisdictions, strategies for harmonizing legal approaches. and the impact of international regulatory environments.

Part 1 - Case Law

Review the major cross-border case law for these cases and discuss the resulting IP effects of these suggested decisions:

- Tecfidera Apixaban
 Rivaroxaban
- Covid 19 Cases Glucose monitoring devices •

Part 2

- Discuss common challenges faced in cross-border patent litigation, including jurisdictional issues, differing legal standards, and enforcement of judgments.
- Understand strategies for harmonising litigation approaches and minimising conflicting decisions.



Andrew Hutchinson Partner Simmons & Simmons



Kristin Cooklin

Recordati

Global Head IP Counsel

Julia Schönbohm

Partner

Linklaters



Chiara Accornero Legal Officer- IP **Disputes Section WIPO**



Rob Rodrigues

Partner **RNA** Law

09.55 - 10.50

Global Review of Preliminary Injunctions: the UPC. National Courts. the USA and Further Afield

The biggest hammer that a patent holder has is an injunction. therefore meaning that case law, updates and strategies concerning PIs for the healthcare sector are crucial in determining your life science strategy for 2025. This cross-jurisdictional panel will discuss the power of a preliminary injunction and the different nuances occurring across the globe.

- Receive an overview of recent developments in preliminary injunctions across the UPC. European jurisdictions, and the USA.
- Discover significant recent cases and their implications for pharma and biotech patent litigation.
- Understand strategies for obtaining and defending against preliminary injunctions in different jurisdictions.
- Audience Q&A to address specific questions and practical concerns related to preliminary injunctions.



Carsten Richter Head of Division IP EUROIMMUN



Martina Hufnal

Matthew O'Neill Director of European IP **Glenmark Pharmaceuticals**

Tim Powell Partner

Networking Break

11:20

The Effect of the UPC Upon the Life Sciences Industry

The UPC is becoming an established venue for Pharma and Biotech Patent Litigation and with this, the life science industry has more established and nuanced strategies towards litigation strategy at the UPC. This industry-led panel session will discuss the key cases within the life sciences sector and help you understand the ramifications for the wider industry.

Understand the functioning of the UPC for the life sciences sector with key takeaways from cases including:

- Amgen vs Sanofi/Regeneron
- 10x Genomics vs Nanostring
- Sanofi vs Generics

Discuss perspectives towards opt-in and opt-out, the UP and transparency at the court in relation to the life sciences sector.

Discover the considerable number of medical device cases at • the UPC and the takeaways from these cases.









Fish & Richardson





Morgan Lewis

Principal IP Counsel



Senior Director- IP **Polpharma Biologics**



21 January 2025

12.15

Competition Law and IP: Updates from the EC and the FTC

An important cross-sectional area of competition law and life science patenting law are divisional practices. In this realm, 2024 will see several high-level decisions form the European commission and the FTC regarding this topic. This panel session will review these decisions and provide generic and innovator perspectives towards competition law and IP.

Part 1

- Discuss the recent decision in the Teva Copaxone case adopted by the EC.
- Discover the innovator and generic perspective towards evergreening patents.

Part 2

Determine the US status regarding divisional patent applications:

How will policy affect the life science industry within the US?



Beatriz Díaz de Escauriaza Head of IP Legal

Insud Pharma



Enzo Marasa Partner Portolano Cavallo



Michael Clancy

Partner Van Bael & Bellis



Amalia Athanasiadou Lead Counsel **CSL Vifor**

Networking Lunch

Advisory Board Private Lunch

14.00

Litigate, Oppose, Licence or Settle? Patent Litigation **Considerations for the Healthcare Industry**

Choice of dispute resolution is an important topic and never more so than with the introduction of the UPC. This multi-iurisdictional mix of in-house counsel members and litigators will discuss the best forum for patent revocation in Europe and further afield.

Discover in-house counsel perspectives between choosing to licence innovation or litigate:

- Does this differ between industry?
 - Understand the advantages and disadvantages between using national courts, the UPC or the EPO for life science disputes.
- Is there a trend towards settlements in patent litigation proceedings in the life science industry?
 - How is this best achieved?



Katherine Helm

Frank Landolt Chief Counsel- IP & Legal

Nick McDonald Partner Potter Clarkson

Raquel Frisardi Associate General Counsel Novo Nordisk

Toni Santamaria Senior Vice President IP & Legal Adalvo

14:45

Global Judges Session: Multi-Jurisdictional Perspectives for Life Science Patents

With the UPC now being becoming an established venue for litigation in the life science arena, the healthcare industry has a variety of different forums for litigation. This expert-led panel will discuss the jurisdictional differences and key cases in 2025 from some of Europe's top courts.



András Kupecz Honourable Judge **Central Division Munich**

Denis Schertenleib Partner Schertenleib Avocats



Kai Härmand Honourable Judge - Nordic-Baltic Regional Division



Rainer Friedrich Technically Qualified Judge & Director of Global IP **UPC & CSL Behring**







Confo Therapeutics



21 January 2025

15:30

Navigating Patent Strategies: Second Medical Use and Skinny Labels

The intersection of second medical use claims and skinny labeling remains a crucial battleground for both generic and innovator drug manufacturers. This expert panel will delve into the latest case law, providing actionable insights for refining your patent litigation strategy in 2025.

Part 1 - Second Medical Use

- Explore how recent rulings, particularly those concerning the Novartis product Everolimus, have influenced the interpretation and enforcement of second medical use claims across Europe.
- Gain insights into the strategic use of second medical use claims in protecting pharmaceutical, biologic, and biosimilar innovations.
- Assess how second medical use claims are being addressed within the framework of the UPC, and what this means for your litigation strategies.

Part 2 - Skinny Labels

- Understand the ongoing impact of the GSK vs. Teva decisions on skinny labeling practices, particularly the implications of the 2017 Supreme Court ruling.
- Discuss how this precedent affects generic manufacturers in their competition with innovator drug companies, and what this means for your approach to skinny label litigation.



Alexander de Leeuw Partner Brinkhof

.....

Paul Ainsworth

Director Sterne Kessler



Philippe Bessiere Global Head of Patents PIERRE FABRE GROUP

Sam Woodley Lead Patent Counsel

Zentiva

Viviane Kunisawa Partner Daniel Law

Networking Break

16:45 Connect 25

Roundtable discussions are back! Led by a private practice expert within a jurisdiction, you can discover, question and gain solutions to your most pressing challenges in patent litigation strategy.

Each roundtable session is 45 minutes long with x1 rounds of changes required.

Click here for details

17:25 Roundtable Swap Break

17:30

Interactive Roundtables: Connect 24 Continued

8:10

Chair Closes on Day 1

Drinks Reception





21 January 2025



Roundtable discussions are back!

Led by a private practice expert within a jurisdiction, you can discover, question and gain solutions to your most pressing challenges in patent litigation strategy.

> Each roundtable session is 45 minutes long with x1 rounds of changes required.

United States 1 Patent Litigation Considerations for Biosimilar Litigation Robert Cerwinski, Partner, GEMINI

Mike Cottler, Partner, GEMINI Join this roundtable to explore strategic insights and practical solutions for navigating the complexities of global patent litigation.

United States 2 Discovery in the US & Coordinating that with European, UK & Asian Counter-Part Cases

John Cox, Partner, Barnes and Thornburg Lauren Baker, Associate, Barnes and Thornburg

Examine strategies for managing discovery in U.S. patent litigation while coordinating with parallel cases in Europe, the UK, and Asia. This session will address cross-border discovery challenges, data protection, and aligning patent litigation strategies across multiple jurisdictions.







United Kingdom 1 AI and IP – patentability of AIrelated and AI-derived inventions

Rebecca Lawrence, Partner, DLA Piper Dr Philipp Cepl, Partner, DLA Piper

Is the patent system fit for purpose in the age of AI? Considering protection of AI-related and AI-derived inventions in light of the Emotional Perception case in the UK, EPO decisions and beyond. Understand the evolving IP framework and its impact on AI innovations and patentability.



United Kingdom 2

Arrow Declaration Update in the UK

Chris Freeth, Managing Associate, GOWLING

Get the latest updates on Arrow Declarations and their strategic use in the UK. Discover how these declarations can serve as a defensive tool in patent litigation.

Day One 21 January 2025



Netherlands The Status and Effect of G2/21 in the Netherlands and Further Afield

Carlos Van Staveren, Counsel, Bird & Bird Andreas Obermeier, Counsel, Bird & Bird

Explore the implications of the G2/21 decision on the doctrine of equivalence in Dutch patent law. Discuss how this impacts your patent litigation strategies in the Netherlands.





Germany 1 Comparison of German patent litigation and UPC proceedings

Christian Meyer, Partner, Maiwald Heike Röder-Hitschke, Counsel, Maiwald Anja Fux, Associate, Maiwald

Compares traditional German patent litigation with UPC litigation in the life sciences field. Discusses procedural nuances, material law and strategical considerations for the different forums. recent case law regarding saisie-contrefacon, a crucial evidence-gathering procedure in intellectual property disputes. Experts will discuss how the UPC is addressing this tool in the context of life sciences, offering insights for effectively navigating

France The Latest Case Law on

"Saisie-Contrefaçon" (in the field

of life science) Before the UPC

This session will provide an in-depth analysis of

infringement litigation.
Camille Pecnard, Partner, LAVOIX

Aude Veinante. Partner, LAVOIX Pierre-Emmanuel Meynard, Partner, LAVOIX

Germany 2

Patent Litigation

Considerations in the Life

Science Sector

Daniel Hoppe, Partner, Bonabry







21 January 2025



Global Roundtable **Regulatory Data Exclusivity: Global Litigation Strategies**

Kamleh Nicola, Partner, Marks & Clerk Kevin Yurkerwich. Senior IP Counsel. Novartis

Regulatory data exclusivity and market protection regimes around the world are similar but different. These regimes operate separately from the patent system but are no less important in the promotion of innovation and the efficient entry of subsequent entry products. Join this Roundtable to discuss the latest strategies, anticipated changes, and the impact of recent court decisions.



Europe 2 **Patent Search in Litigation. The** Crowd: the unbeatable prior art search

Andre Andrade. VP Partnerships. RWS

Explore how crowdsourced prior art searches are transforming litigation strategies in the pharmaceutical and biotech sectors. This roundtable delves into the effectiveness of leveraging collective expertise to uncover critical prior art, strengthen patent challenges, and bolster defense arguments. Join industry leaders to discuss the advantages, limitations, and future potential of crowdpowered solutions in high-stakes patent disputes.





Europe 3 The Latest on PI **Cases at the UPC**

Charlotte Garnitsch, Partner, Taylor Wessing Christian Dekoninck, Partner, Taylor Wessing Andrew Payne, Partner, Taylor Wessing

Stay updated on the latest developments in preliminary injunction (PI) cases at the UPC. Learn about recent trends and effective strategies for securing PIs in life sciences patent disputes



Global **Managing Patent Litigation Global Challenges Outside Europe**

Naomi Pearce, Founder, Pearce IP

This roundtable session will be led by elite life science litigators from outside of Europe to help you successfully navigate a global litigation strategy



The Interaction Between the **SPC Manufacturing Waiver and** the Bolar Exemption

Aoife Murphy, Partner, DLA Piper Kokularajah Paheenthararajah, Partner, DLA Piper

A review of the Bolar Exemption and any recent case

law (including the recent decision of the Italian Supreme court), the interplay of the Bolar Exemption with the SPC Manufacturing Waiver including the recent cases in Ireland, Germany and the Netherlands relating to the SPC Manufacturing Waiver. The Research and Bolar Exemption under the UPCA and the latest position relating to the proposed EU Pharma Reforms of Bolar and what the envisaged modification of the Bolar exemption could mean for IP rights holders etc.



Europe 1 **Managing Patent Litigation Proceedings** in the UK. UPC and National Courts

Laëtitia Bénard, Partner and Global Co-Head of IP Litigation, A&O Shearman Rafi Allos, Partner, A&O Shearman Stephan Neuhaus, Partner, A&O Shearman

Gain insights into managing patent litigation across the UK. UPC, and national courts. Discuss coordination strategies to navigate multi-jurisdictional disputes effectively.



Europe 5 **Defensive and Offensive strategies in** originator v. generics disputes at the UPC

Frédéric Chevallier, Partner, Herbert Smith

Insights into the use of revocation and infringement actions, and the involvement of multiple defendants and forums in actions at the UPC

22 January 2025

07:30 Registration

8:00 **Breakfast Briefing:**

Will the UPC Get the Doctrine of Equivalents Right? Comparing National Case Law with UPC Case Law in the Life Science Sector

This session explores how the UPC's approach to equivalence compares to national practices in key jurisdictions such as the UK. Turkey, and Brazil. Panellists will critically assess the four key guestions guiding equivalence in UPC cases and their application in the life sciences sector. Join us to evaluate whether the UPC is striking the right balance between patentee protection and third-party certainty.

- Compare the UPC's equivalence framework to national approaches in the UK. Turkey, and Brazil.
- Are the four UPC questions appropriate for life sciences disputes, or are adjustments needed?

1) Technical equivalence: does the variation solve (essentially) the same problem that the patented invention solves and perform (essentially) the same function in this context?

2) Fair protection for patentee: Is extending the protection of the claim to the equivalent proportionate to a fair protection for the patentee?

3) Reasonable legal certainty for third parties: does the skilled person understand from the patent that the scope of the invention is broader than what is claimed literally?

4) Is the allegedly infringing product novel and inventive over the prior art?"

- Balance fair protection for patentees with legal certainty for third parties
- Understand the implications of UPC decisions for global life sciences patent strategies.



UPC

Eduardo Hallak Founding Partner Licks Attorneys









Senior Partner NSN Law

09:00 Chair Recap of Day 1 and Intro to Day 2

Kristin Cooklin Group Head of IP Recordati



Pauline Debré Partner Linklater

Track 1: Pharma and Biotech Patent Litigation

09:10

Global Biosimilar Molecule Patent Litigation and Launch Review

There is a suggestion that a second wave of biosimilars have appeared. This global panel session will review the state of biosimilar litigation in Europe the US and further afield. By taking looking at individual molecules, this industry-led panel session will see assess the state of the biosimilar IP industry.

- Understand the state of biosimilar litigation related to molecules including: - Evlea, Inflectra, Soliris, Stelara
- Discover strategies concerning BPCIA. PIs and second medical use/ skinny labels for biosimilar products.



10:00

Bolar Exemption and Safe Harbor Update Across Europe and the USA

The Bolar exemption and safe harbor regulations set out the legal provisions allowing generic manufacturers to perform experiments and tests necessary for regulatory approval. This industry-led session will highlight some of the core case law in Europe and the USA and determine your optimal IP strategy.

Part 1 - The Bolar Exemption

- Discover the divergences between the Bolar exemptions across Europe and the UPC.
- Understand the interaction between the SPC manufacturing waiver and the Bolar exemption.
- Determine the latest update from the European Commission on the potential for continent-wide harmonisation.
- Case Study Highlight: Italian Supreme Court Rules on Bolar Exemption in Boehringer Ingelheim Dispute (2024).

Track 2: MedTech Patent Litigation

09.10

AI Considerations for Life Science Litigation

With AI becoming a more and more common feature of the healthcare industry. IP attorneys are having to face up to different question relating to Al patents. This MedTech focused panel will discover the regulatory concerns of the industry and the potential for litigation.

- Understand the consequences for AI driven patents and the industry after the Emotional Perception AI decision in the UK (2024).
- Discuss the consequences for clinical trials, patentability and litigation with increased AI use in healthcare.



Georgia Roussou Senior Legal Counsel

Jason Raeburn Partner **Paul Hastings**

Philip Vuley Senior Patent Attorney TomTom

10:00 Vaccine Patent Litigation Review in Europe and Further Afield

The Covid 19 patent battles are ongoing over multiple jurisdictions. These battles centre on MRNA advances, with there being parallel proceedings taking place over the world. This expert panel session will discuss the state of vaccine litigation and suggest the ramifications for the wider IP industry.

- Discuss the status of the Pfizer/ BioNTech vs Moderna cross border litigation cases.
- Understand the impact of other MRNA litigation cases: Arbutus/Genevant vs Moderna - Alnyam vs Pfizer - Curevac vs BioNTech







Paul Inman Partner Gowling











22 January 2025

Track 1:

Part 2 - Safe Harbor

- Understand the scope and history of protection afforded by safe harbor in the USA.
- Compare the interpretations and applications of safe harbor regulations between the USA and Europe.



General Counsel Atalanta Therapeutics



Head of IP Zentiva

Cecile Teles



Christoph Rehfuess Head of IP Sotio

Doug Mccann



Principal Fish & Richardson

Marco Stief Partner Maiwald



Özge Atılgan Karakulak Partner Gun + Partners

Networking Break

11:15

SPC and PTE Litigation Review: Re: OCD, the manufacturing waiver and Article 3A and 3C

Updates and challenges surrounding SPCs (and PTEs) continue to be of significant interest for the life science sector. This industry-led panel session will discuss the latest regulatory

changes and the key cases which will allow you to update your patent extension strategy.

Part 1

• Discuss the impact of the Article 3a and 3c Advocate General opinion at the CJEU.

 How have the latest cases on the manufacturing waiver impacted its status: - Stelara case in Germany (2023).

- Stelara case in the Netherlands (2024)

Part 2

Discover the global PTE updates. Understand the impact of re Cellect (2024) concerning the status of PTEs and PTAs.



James Horgan Chief IP Counsel MSD



Senior Patent Manager-Patent Litigation **STADA**



End of track



Track 2:

Networking Break

11:15

MedTech at the UPC: Opt-in Opt-out, **Case-Law and Strategy**

The MedTech industry have been one of the front runners at the UPC, with there being multiple major cases operating in Europe. This industry focused panel sessions will discuss the key takeaways from these cases and look at the status of the MedTech industry in Europe. - Gain insights and key takeaways from cases includina:

- Abbott Diabetes Care vs SiBio (2024) Edwards Life Sciences Vs Meril (2024)
- Understand the impact of Germany in the UPC, potential inclusion of Ireland, Spain, and Poland, and UK's post-Brexit role.
- Discuss MedTech views on patent enforcement. EPO decisions. and UPC. balancing innovation with enforcement.



Christopher Sharp **Pinsent Masons**



Manuel Neetz Head of IP- Diagnostic Imaging **Siemens Healthineers**

End of track





12:00

Appeal Procedures, Patentability Reviews and Commentary for the Life Sciences Sector

Appeals to the BOA have wide ranging effects on guestions of patentability and by consequence, litigation proceedings across Europe. This industry-led panel session will look at some of these decisions and assess how they are affecting healthcare litigation strategy across Europe.

Part 1 - G2/21

- Discover the decisions from T116/18 in Sumitomo vs Syngenta (2023).
- Compare this with T681/21 in Unilever vs Procter & Gamble (2023) and understand whether the case law is diverging from the G2/21 decision from the EPO BOA.

Part 2 - G1/23

Discuss this case in relation to the state of the art in view of the commercial availability of a product prior • to the filing date/ priority date of a patent.

Part 3 - G1/24

٠ Understand the potential impact of this referral on claim interpretation.

Part 4 - Appeal Procedures at the EPO

Understand how the EPO's Administrative Council can overrule the highest judiciary in the system (EBoA). which is not possible in other judicial systems

Natalia Wright

UCB

Robin Ellis

Partner **Reddie & Grose**

Assistant General Patent Counsel



Corinna Sundermann Senior VP IP **Fresenius Kabi**



Daan de Lange Partner **Brinkhof**



Rob Aerts Patent Counsel Europe ADM

Networking Lunch



13.55

New Pharmaceutical Regulations in Europe: Perspectives from the Innovator and **Generic Industries**

The new pharmaceutical regulations are a point of political and IP interest for the life science industry. with there being the potential for a wide-ranging impact on the innovator, generic, biotech and biosimilar industries. This industry focused panel session will discuss the latest update and the potential effect of these reforms.

- Determine the status of the new pharmaceutical regulations in Europe and the impact of the new elections.
- Understand the effect of the new reforms and the potential effect for the IP departments of the life sciences industry.





Michael Swita

EFPIA

Director- IP Policy



Serena ter Kuile Genmab

Senior Legal Counsel, Corporate Strategy

14:40

The Unitary SPC: Update and Strategy for Your Life Science Product

The UPC's launch in summer 2023 reshaped European patent litigation, though it initially excluded a unitary SPC. The European Commission has proposed a unitary SPC, bringing new considerations for centralized examinations, national litigation, and legislative impacts. Join our panel discussion with industry leaders.

- What are the potential benefits and challenges of a centralised approach?
- How will the unitary SPC interact with the strategy towards the UPC for life science companies.



Jiri Salvik Director of IP Adalvo



Filip de Corte Head of IP- Crop Protection Syngenta

Chair Closing Remarks

Conference End

INTERESTED IN HOW YOU CAN GET INVOLVED WITH PHARMA & BIOTECH PATENT LITIGATION 2025?

Partnering with us presents a unique opportunity to establish your company as an industry leader in the life science IP space. We provide unparalleled access to key decision-makers through tailored introductions, industry panel sessions, jurisdictional roundtables and bespoke branding opportunities, ensuring you connect with the most influential stakeholders in the field.

By collaborating with us, your brand will be positioned in front of hundreds of in-house executives, creating valuable conversations and networking with the elite of the IP industry. Additionally, our multi-channel approach offers a powerful platform to amplify your announcements, share expertise, and effectively engage your target market both during and leading up to the event.

Bigger and better than ever this year. The only place to be if you want the most up to date discussions on what is happening in life science IP and a chance to meet some great people from across the industry." Robin Ellis, Partner, Reddie & Grose



Contact

Duncan Henderson

Commercial Partnership Director Kisaco Research duncan.henderson@kisacoresearch.com LISTEN TO WHAT OUR PREVIOUS PARTNERS HAVE TO SAY ABOUT PARTNERING WITH US:





CLICK HERE TO BE INVOLVED!







PATENT LITIGATION EUROPE

January 21-22, 2025 Amsterdam, Netherlands, Beurs van Berlage



REGISTER NOW



VENUE INFORMATION Beurs van Berlage

Damrak 243, 1012 ZJ Amsterdam, Netherlands

Click the link below to book local hotels, we recommend booking the Anantara Grand Hotel Krasnapolsky Amsterdam.

BOOK NOW