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this book is the conference proceedings of essa 2013 the 9th conference of the european social simulation association essa conferences constitute annual events which serve as an international platform for the exchange of ideas and discussion of cutting edge research in the field of social simulations both from the theoretical as well as applied perspective this book consists of 33 articles which are divided into four themes methods for the development of simulation models applications of agent based modeling adaptive behavior social interactions and global environmental change and using qualitative data to inform behavioral rules we are convinced that this book will serve interested readers as a useful compendium which presents in a nutshell the most recent advances at the frontiers of social simulation research

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a history of u s army medical activities from the revolutionary war to 1818 the year in which congressional legislation instituted the modern medical department

the magazine that helps career moms balance their personal and professional lives

the common foreign and security policy cfsp of the european union is a highly exceptional component of the eu legal order this constitutionalised foreign policy regime with legal diplomatic and political dna woven throughout its fabric is a distinct sub system of law on the outermost sphere of european supranationalism when contrasted against other union policies it is immediately clear that eu foreign policy has a special decision making mechanism making it highly exceptional in the now depillarised framework of the eu treaties issues of institutional division arise from the legacy of the former pillar system this is due to the reality that of prime concern in eu external relations is the question of who decides by engaging a number of legal themes that cut across foreign affairs exceptionalism executive prerogatives parliamentary accountability judicial review and the constitutionalisation of european integration the book lays bare how eu foreign affairs have become highly legalised leading to ever greater coherence in how europe exerts itself on the global stage in this first monograph dedicated exclusively to the law of the eu s common foreign and security policy in modern times the author argues that the legal framework for euforeign affairs must adapt in a changing world so as to ensure the eu treaties can cater for a more assertive europe in the wider world cited in opinion of advocate general evgeni tanchev case c 730 18 p sc v eulex kosovo ecli eu c 2020 176 court of justice of the european union first chamber 5 march 2020 opinion of advocate general gerard hogan case c 134 19 p bank refah kargaran v council of the european union ecli eu c 2020 396 court of justice of the european union grand chamber 28 may 2020 opinion of advocate general evgeni tanchev case c 283 20 co me gc and 42 others v mj head of mission european commission european external action service eeas council of the european union eulex kosovo ecli eu c 2021 781 court of justice of the european union fifth chamber 30 september 2021 and opinion of advocate general tamara capeta in joined cases c 29 22 p and c 44 22 p ks kd v council of the european union european commission and european external action service eeas and european commission v ks kd council of the european union and european external action service eeas ecli eu c 2023 901 court of justice of the european union grand chamber 23 november 2023

in the european union eu its member states and the united kingdom uk post brexit as elsewhere the marketing of pharmaceuticals is subject to an ever more complex web of legislation and regulation resulting from the intense scrutiny necessary to ensure such essential products are not only efficacious but also safe this useful volume lays out this system with extraordinary clarity and logic adopting a europe wide perspective on the law governing pharmaceuticals expert authors from the law firm bird bird llp map the life cycle of a medicinal product or medical device from development to clinical trials to product launch and ongoing pharmacovigilance offering comprehensive and unambiguous guidance at every stage following a brief overview of how the exit from the eu by the uk currently affects the regulatory regime as well as an introductory overview focusing on the regulatory framework for pharmaceuticals in europe from its underlying rationales to the relevant committees and agencies each of the following twenty one incisive chapters examines a particular process or subject among the many topics and issues covered from both an eu and uk perspective are the following clinical trials stages and standards for creating a product dossier obtaining a marketing authorisation how and when an abridged marketing authorisation procedure can be used criteria for conditional marketing authorisations generic products and essential similarity paediatric use and the requisite additional trials orphan medicinal products biologicals and biosimilars homeopathic herbal and similar medicines medical devices pandemics epidemics and vaccines pharmacovigilance parallel trade advertising and relevant competition law intellectual property rights and data protection regulation in addition sample forms and urls for the most important reference materials are included pharmaceutical lawyers and regulatory advisers both in house and in private practice will welcome this unique book it offers immeasurable value for all who need to understand the process of bringing a medicinal product or medical device to market and the continuing rights and obligations

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