

# NOTE on Proposals to extend pharmaceutical intellectual property incentives in reaction to US tariffs

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Medicines for Europe is aware that EFPIA is proposing an extensive prolongation of pharmaceutical regulatory protections to counter the impact of US tariffs. This will not stop pharmaceutical companies from transferring their production to the US as some have already announced. This will bankrupt public health budgets which already struggle to finance the reimbursement of expensive drugs. The note provides some important facts for consideration.

## 1. Europe has the longest pharmaceutical regulatory protection system in the world, as highlighted in a Commission Impact Assessment (page 38):

Country	Protection	Duration	
Canada	New Chemical Entity+ Market Protection	6+2 years	
EU	New Chemical Entity+ Market Protection	8+2+1 years	
Switzerland	New Chemical Entity	10 years	
USA	New Chemical Entity (small molecule)	5 years	
USA	Biosimilar Application Approval Exclusivity (biologic)	4+8 years	
Israel	Market Protection	6 or 6.5 years	
China	New Chemical Entity	6 years	
Japan	New Chemical Entity	8 years	

**Compared to the US**, Europe provides 8 years of regulatory, and 15 years of patent and supplementary protection certificate (SPC), protections compared to 5 and 14 years in the US for equivalent protections.

CURRENT SYSTEM				
REGULATORY PROTECTIONS	EUROPE	UNITED STATES	DIFFERENCE IN YEARS	
Small molecules	8DP+2MP+1MP = 11y	5DP = <b>5y</b>	+6y in EU	
Biological	8DP+2MP+1MP = 11y	4DP+8MP = <b>12y</b>	+1 in US	
Orphan medicines	10ME (equal to DP) = <b>10y</b>	7ME = <b>7y</b>	+3 in EU	
		Total n. of years	+8 in EU	

Despite these long EU protection periods that have been in force for the last 30 years, R&D investments have grown much faster in the US (with shorter protection) and China (with very little protection) than in Europe. Lengthy IP protections do not stimulate R&D investments in Europe.

Extending regulatory protection from 11 years to 13/14 years would **dramatically increase costs for healthcare systems**, which cannot afford this. The same proposals were already rejected in the European Parliament and Council because they would cost between €20 and 100 billion annually, which is equivalent to the salary of hundreds of thousands of much needed nurses or doctors.

### Simulation of the economic impact of extending regulatory data protection:

Europe is facing the risk of economic recession and needs to divert public spending to national defence. Under these circumstances, healthcare systems cannot afford €20-100 billion extra per year.

	France	Germany	Spain	EU
13.5 years of regulatory protection	5.35 BN EUR	5 BN EUR	2.25 BN EUR	19.5 BN EUR
18 years of regulatory protection	24 BN EUR	23 BN EUR	13.2 BN EUR	99.5 BN EUR

#### 2. The Bolar Exemption is important for competition and for manufacturing in Europe

The **Bolar exemption** was enacted to allow the development and approval of generics and biosimilars for immediate competition at IP expiry. Multiple studies (including Commission studies) show that the lack of harmonisation of the Bolar have led to (1) disinvestments in Active Pharmaceutical Ingredients (API) development in Europe and (2) a misuse of the IP system to systematically and unduly delay generic and biosimilar competition (well documented by national competition authorities and courts), with severe delays for patient access to medicines and massive costs for healthcare budgets, as shown in the few <u>examples</u> below:

Molecule	Treatment	Country	Originator	SPC Expiry	Generic	Delay	Cost of Delay
			approval		Entry		Lost Savings
Oxycodone/	severe pain	Germany		29/3/2017	15/11/2017	231 days	€ 51,6 MIn
Naloxone							
Ezetimibe/	high	Italy	18/11/2004	16/10/2017	9/3/2018	144 days	€ 15,4 Mln
simvastatin	cholesterol						
Ezetimibe/	high	Germany	18/11/2004	17/4/2018	15/5/2018	28 days	€ 11,3 Mln
simvastatin	cholesterol						
Lenalidomide	multiple	Hungary	14/06/2007	19/6/2022	1/6/2023	347 days	€ 1.9 Mln
	myeloma,						
	cancer						
Pirfenidone	idiopathic	Germany	27/02/2011	27/2/2021	15/11/2022	626 days	€ 32,1 Mln
	pulmonary						
	fibrosis						
Tapentadol	severe pain	Germany	19/08/2010	07/12/2020	15/1/2023	917 days	€ 184,6 Mln
Dasatinib	chronic	Poland	20/11/2006	22/5/2022	01/01/2023	224 days	€ 4,5 MIn
	myeloid						
	leukemia						
					Total:	2,517	€ 301,4 MIn

For these very well documented reasons the Bolar exemption should be clarified and harmonised in the revised Pharmaceutical Directive.

#### **MEDICINES FOR EUROPE POSITION**

Medicines for Europe is ready to work with EFPIA on a balanced regulatory incentive system while stimulating investments in API production in Europe and access to medicines by harmonising the Bolar exemption. We support a united European response to US protectionism that is fact-based, fiscally sound and focuses on patient access to medicines and European strategic autonomy.