

The impact of pharmaceutical innovation on mortality, disability, and hospitalization: evidence from 4 studies

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Three types of health outcomes

Pharmaceutical innovation

(development and use of new drugs)







Longevity

Disability

(work- and school-loss days, quality of life

Hospitalization

and nursing home use



4 studies of impact of prescription drugs on health

study	health outcome	drug measure	region(s)	data
				19 cancer sites,
1	mortality	number of drugs	36 countries	36 countries
				31 diseases, 11
2	disability	number of drugs	11 European countries	countries
				X diseases, 15
3	hospitalization	vintage of drugs	USA	years
				500,000
4	mortality	vintage of drugs	Puerto Rico	patients



3 alternative research designs

- Studies 1 and 2: is *relative* mortality and disability from a disease in a country lower when the *relative* number of drugs available to treat that disease in that country is higher?
- Study 3: has hospitalization in the USA declined more for diseases that have experienced more pharmaceutical innovation?
- Study 4: do Puerto Rican Medicaid patients using new drugs live longer than patients using older drugs?



Two alternative measures of (access to) pharmaceutical innovation

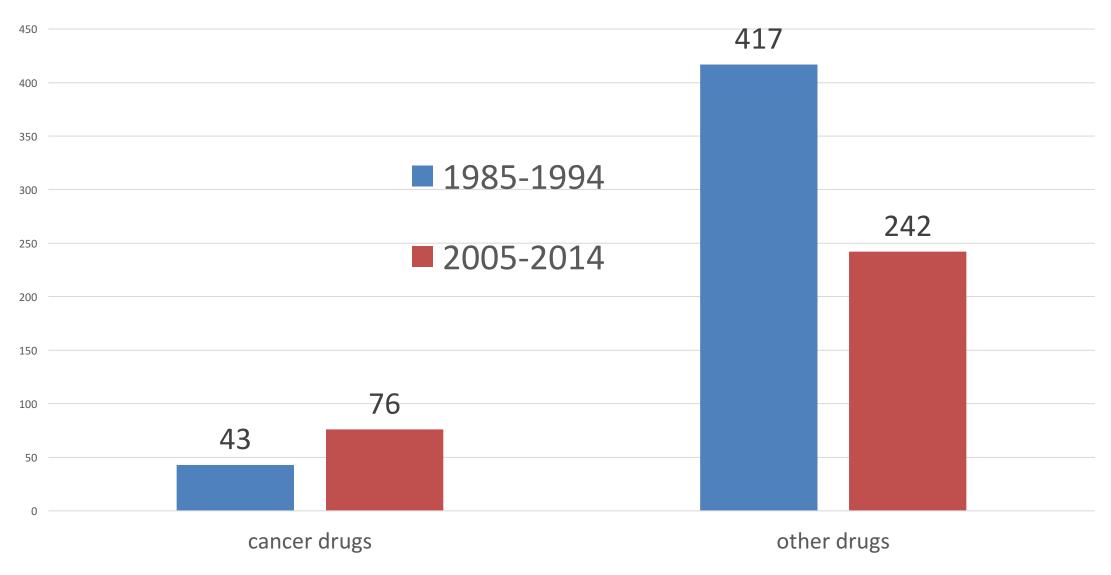
- Number of drugs for a disease that have been launched
- Mean vintage (e.g. FDA approval year) of drugs used to treat a patient or a medical condition



4 studies of impact of prescription drugs on health

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Number of new cancer drugs and other new drugs launched worldwide, 1985-1994 and 2005-2014

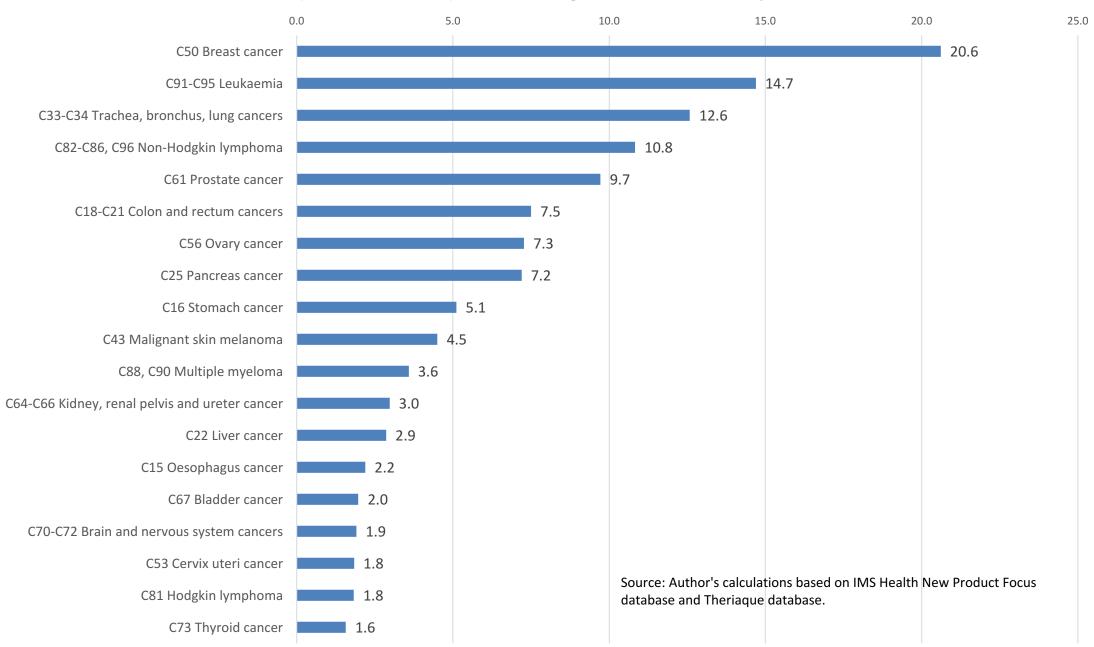


Source: author's calculations based on IMS Health New Product Focus database
"Cancer NMEs" are NMEs in EphMRA/PBIRG Anatomical Classification L (ANTINEOPLASTIC AND IMMUNOMODULATING AGENTS)



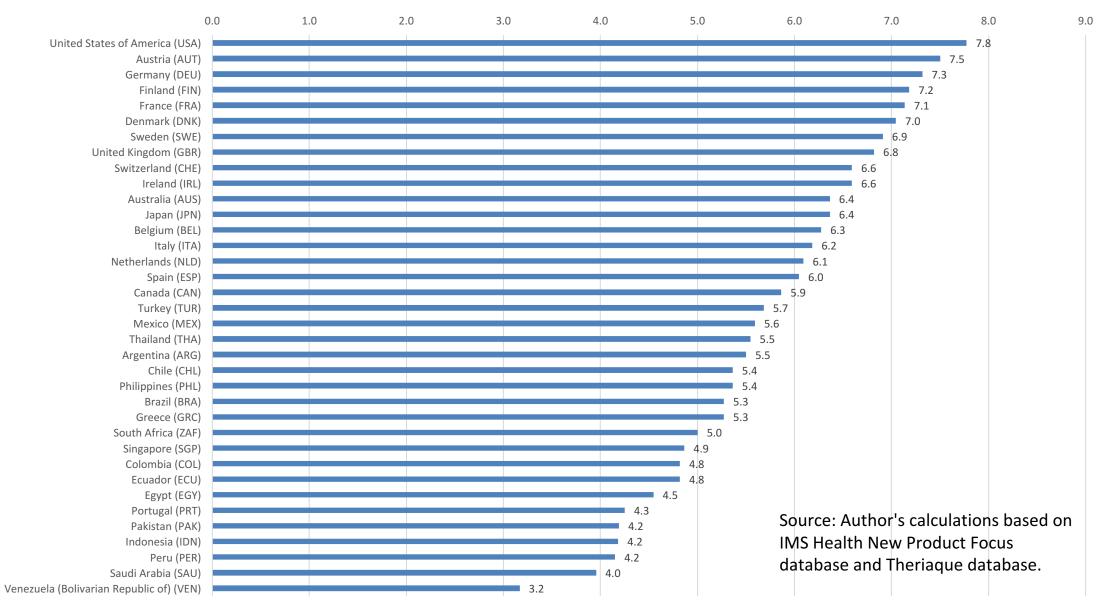
Launch of a drug in a country is a necessary condition for access, not a sufficient condition

Mean (across 36 countries) number of drug launches, 1982-2015, by cancer site



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Figure 3
Mean (across 19 cancer sites) number of drug launches, 1982-2015, by country





Objective

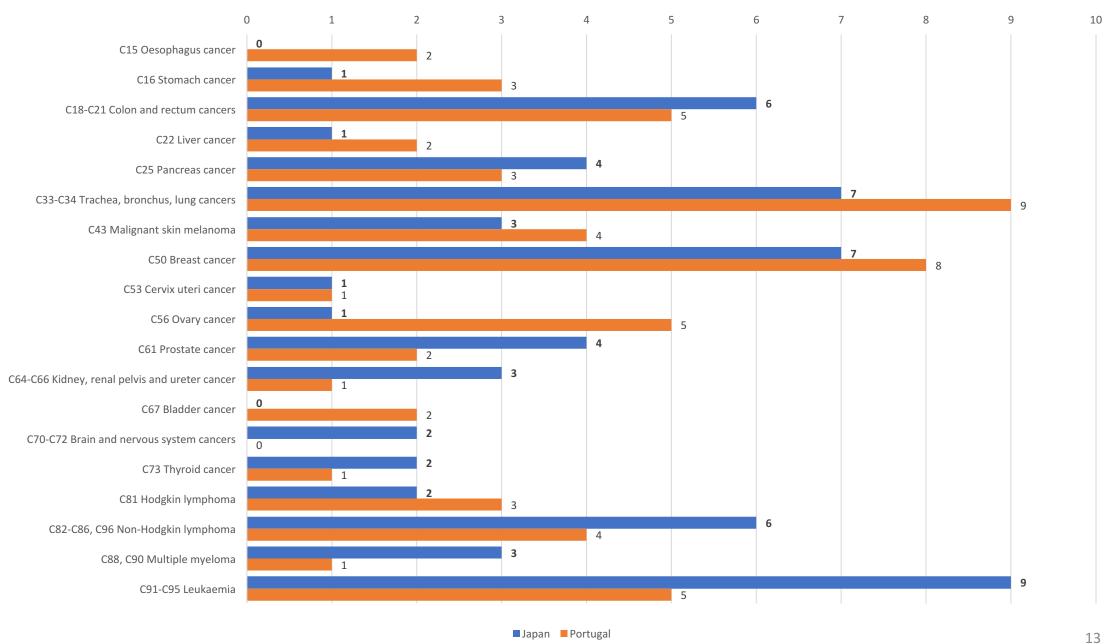
- This study seeks to determine the extent to which the number of years of life lost (YLL) due to cancer in 36 countries in 2015 was reduced by previous launches of new cancer drugs, and to measure the average cost-effectiveness of (cost per life-year gained from) those drugs.
- If 100 people die from lung cancer at age 60, they have collectively lost 500 (= 100 * (65 60)) years of life before age 65, and 1500 (= 100 * (75 60)) years of life before age 75. Hence YLL depends on the number of deaths, age at death, and the age cutoff that is used.



Research design

- I will analyze the correlation across countries between *relative* mortality from each type of cancer in 2015 and the *relative* number of drugs previously launched in that country to treat that type of cancer, controlling for relative incidence.
- The mortality models I will estimate will include both **country fixed effects**, which control for potential determinants of mortality (e.g. national per capita income, education, and average health expenditure) that are invariant across cancer sites within a country, and **cancer-site fixed effects**, which control for potential determinants of mortality that are invariant across countries within a cancer site.
- This approach is feasible because the *relative* number of drugs launched for different types of cancer has varied considerably across countries.

Number of drugs launched during 2006-2015 in Japan and Portugal for 19 types of cancer





Econometric model of life-years lost from cancer

$$\begin{split} & \ln(\text{DALYS}_2015_{sc}) = \beta_{0\text{-}4} \ \text{LAUNCHES}_2011_2015_{sc} + \beta_{5\text{-}9} \ \text{LAUNCHES}_2006_2010_{sc} \\ & + \beta_{10\text{-}14} \ \text{LAUNCHES}_2001_2005_{sc} + \beta_{15\text{-}33} \ \text{LAUNCHES}_1982_2000_{sc} \\ & + \gamma \ln(\text{CASES}_2012_{sc}) + \alpha_s + \pi_c + \epsilon_{sc} \end{split} \tag{1}$$

DALYS 2015sc	= the number of disability-adjusted life years (DALYs) lost due to cancer at site s in country c in 2015
	site s in country c in 2015
LAUNCHES 2011 2015	= the number of post-1981new chemical entities used to treat cancer at site s launched in country c during 2011-2015
	launched in country c during 2011-2015
CASES_2012 _{sc}	= the number of people diagnosed with cancer at site s in country c in 2012
$\alpha_{ m s}$	= a fixed effect for cancer at site s
$\pi_{ m c}$	= a fixed effect for country c

Eq. (1) will be estimated by weighted least squares, weighting by Y_{sc} . The disturbances of eq. (1) will be clustered within countries or within cancer sites. When eq. (1) is estimated without weighting, the residuals clearly exhibit heteroskedasticity: the variance of the residuals is strongly inversely related to $Y_{sc,2015}$.



Other indicators of disease burden

DALYS_2015 _{sc}	= the number of disability-adjusted life years (DALYs) lost due to cancer at site s in country c in 2015
YLL_2015 _{sc}	= the number of years of life lost (as measured in the WHO Global Burden of Disease Estimates) due to cancer at site s in country c in 2015
YLD_2015 _{sc}	= the number of years lost to disability due to cancer at site s in country c in 2015
YLL75_2015 _{sc}	= the number of years of life lost before age 75 due to cancer at site s in country c in 2015
YLL65_2015 _{sc}	= the number of years of life lost before age 65 due to cancer at site s in country c in 2015

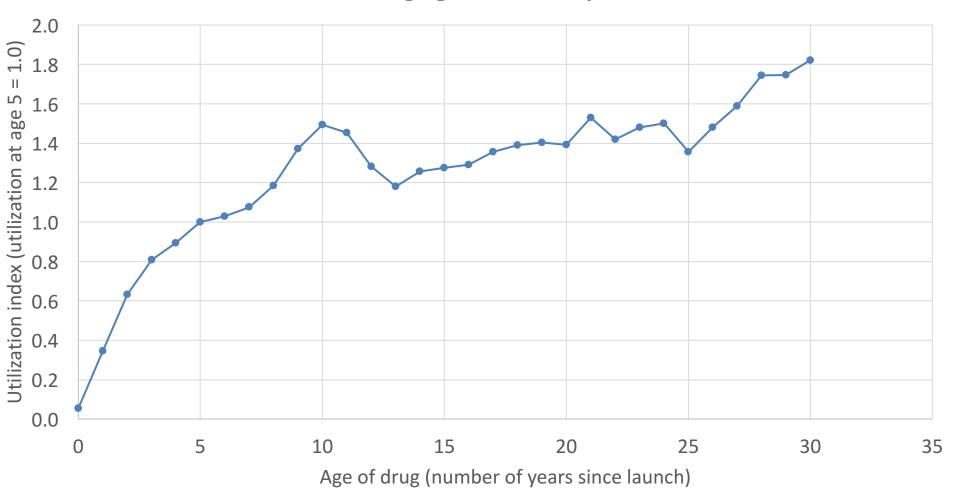


Impact on drug launch on YLL depends on both drug quality and drug quantity

- The effect of a drug's launch on mortality is hypothesized to depend on both the *quantity* and the *quality* (or effectiveness) of the drug.
- Indeed, it is likely to depend on the *interaction* between quantity and quality: a quality improvement will have a greater impact on mortality if drug utilization (quantity) is high.
- Drugs launched in the 4 different periods are likely to vary (in opposite ways) with respect to both quantity (in 2015) and quality:
 - Newer drugs are likely to be of higher quality than older drugs.
 - On the other hand, utilization of new drugs tends to be much lower than utilization of old drugs.



Figure 5
Cancer drug age-utilization profile





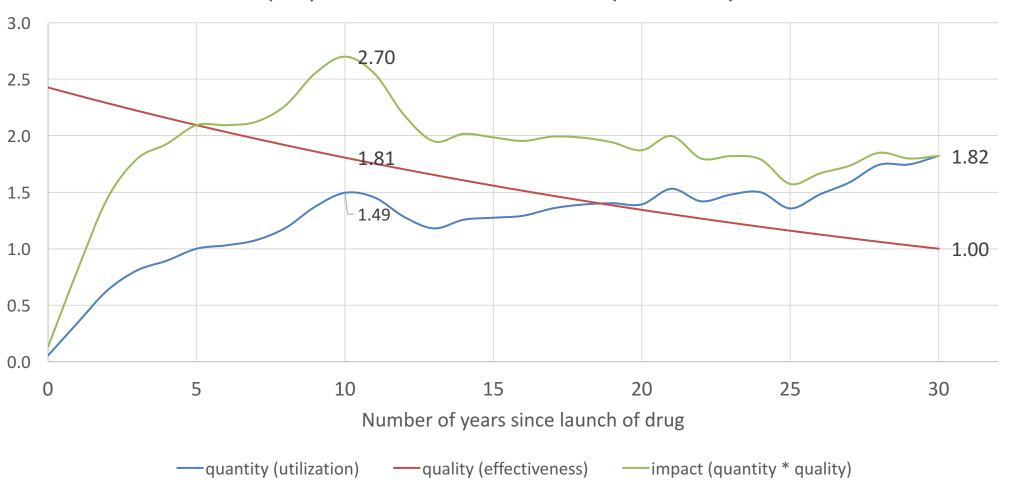
Mean utilization of a drug is about twice as high 5-9 years after launch as it was 0-4 years after launch

years since launch	mean utilization (relative to utilization 5 years after launch)
0-4	0.55
5-9	1.13
10-14	1.33
15-31	1.49



Figure 6

Hypothetical quantity, quality, and impact (= quantity * quality) of a drug when quality increases at a 3% annual rate with respect to launch year





Data sources

- Mortality: WHO Global Health Estimates 2015: Disease burden by Cause and Deaths by Cause databases
- Cancer incidence: GLOBOCAN 2002 and 2012
- Drug indications: Thériaque database (Centre National Hospitalier d'Information sur le Médicament (2017))
- Drug launch years: IMS Health New Product Focus database
- Drug utilization: IMS Health MIDAS database



Table 1							
Summary statistics, 19 major cancer	sites in 36 coun	trios					
Summary statistics, 19 major cancer	sites iii 30 couii	tries					
	2005	2015	% change				
disability-adjusted life years (DALYs)	68,179,003	76,596,299	12%				
years of life lost, as measured in the WHO Global Burden of							
Disease Estimates (YLL)	65,246,858	72,439,899	11%				
years lost due to disability (YLD)	2,932,144	4,156,401	42%				
years of life lost before age 75 (YLL75)	23,398,525	25,137,974	7%				
years of life lost before age 65 (YLL65)	11,163,603	11,545,184	3%				
number diagnosed 3 years earlier (CASES)	4,474,445	5,716,879	28%				

Source: Author's calculations based on WHO Global Health Estimates 2015: Disease burden by Cause database (World Health Organization (2016a)); WHO Global Health Estimates 2015: Deaths by Cause database (World Health Organization (2016b)); GLOBOCAN 2002 (Ferlay et al (2004)); and GLOBOCAN 2012 (International Agency for Research on Cancer (2017b)).



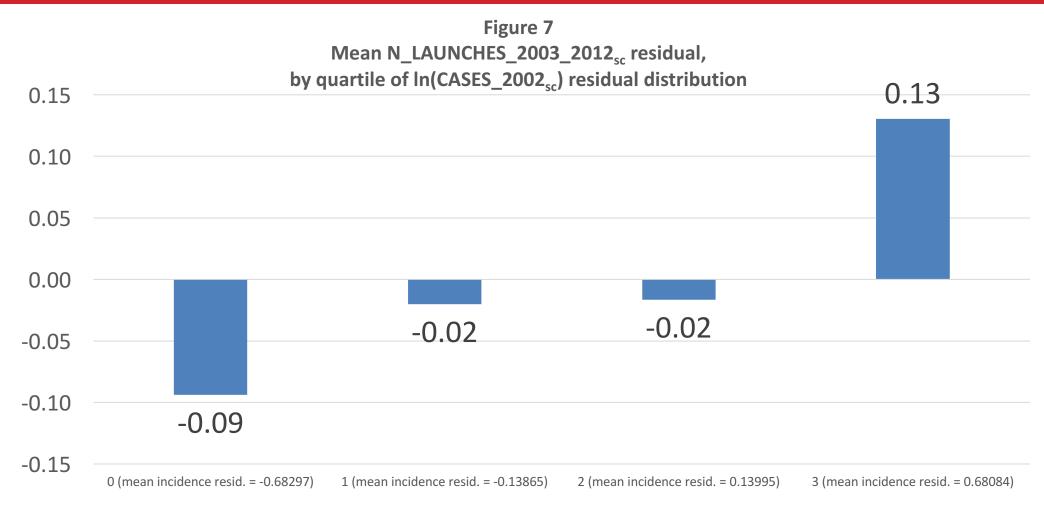
The effect of incidence on the number of new drug launches

- Previous studies have shown that both innovation (the number of drugs developed) and diffusion (the number of drugs launched in a country) depend on market size.
- N_LAUNCHES_2003_2012_{sc} = σ ln(CASES_2002_{sc}) + α_s + δ_c + ϵ_{sc} (3)

N_LAUNCHES_2003_2012 _{sc}	= the number of drugs to treat cancer at site s launched in
	country c during 2003-2012
CASES_2002 _{sc}	= the number of patients diagnosed with cancer at site s in
	country c in 2002

The estimate of σ is positive and significant: estimate = 0.1872; standard error = .0662; Z = 2.83; p-value = .0047. This signifies that larger relative market size (number of patients diagnosed) increases the relative number of drugs launched.

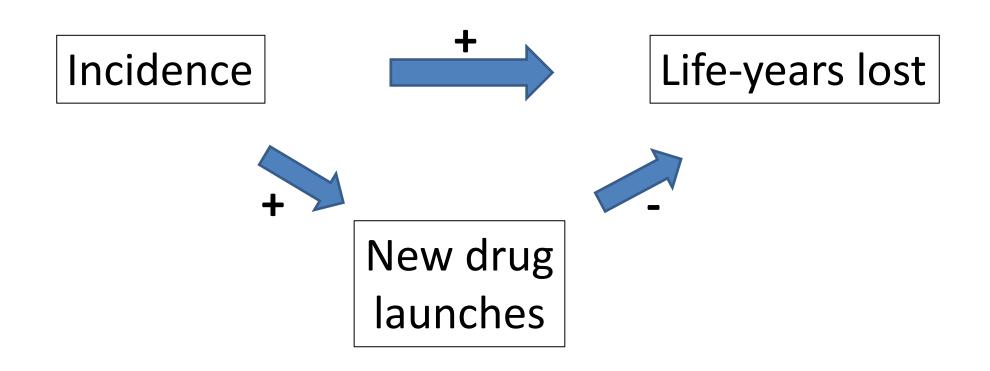




Quartile of In(CASES_2002_{sc}) residual distribution



Direct and indirect effects of incidence on life-years lost



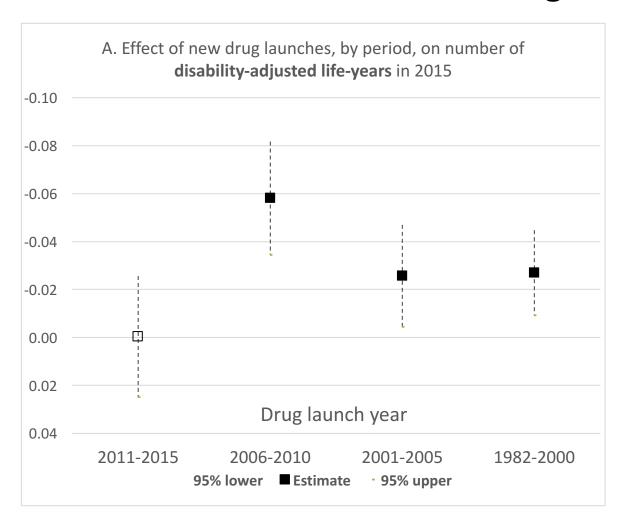


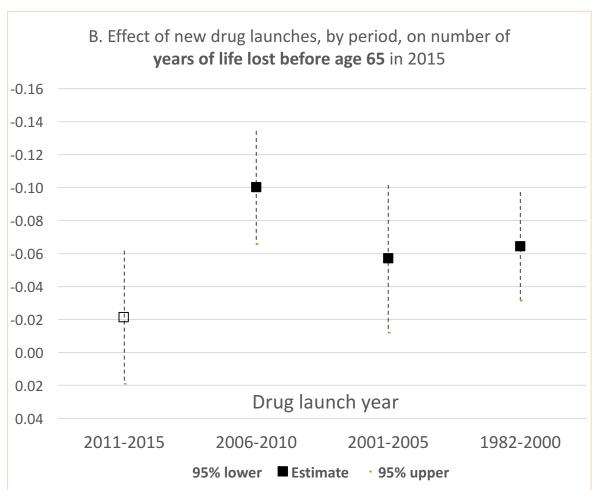
Estimates of difference-in-differences model of life-years lost (eq. (1))

Parameter	Regressor	Estimate	Std. Err.	Z	Pr > Z	
Dependent variable = DALYS_2015						
β0-4	LAUNCHES_2011_2015	0.000	0.013	-0.03	0.977	
β5-9	LAUNCHES_2006_2010	-0.058	0.012	-4.81	<.0001	
β10-14	LAUNCHES_2001_2005	-0.026	0.011	-2.35	0.019	
β15-33	LAUNCHES_1982_2000	-0.027	0.009	-2.96	0.003	
γ	In(CASES_2012)	0.849	0.032	26.27	<.0001	
	Dependent varia	ble = YLL65_2	2015			
β0-4	LAUNCHES_2011_2015	-0.021	0.021	-1.03	0.303	
β5-9	LAUNCHES_2006_2010	-0.100	0.018	-5.69	<.0001	
β10-14	LAUNCHES_2001_2005	-0.057	0.023	-2.48	0.013	
β15-33	LAUNCHES_1982_2000	-0.064	0.017	-3.81	1E-04	
γ	In(CASES_2012)	0.833	0.063	13.16	<.0001	



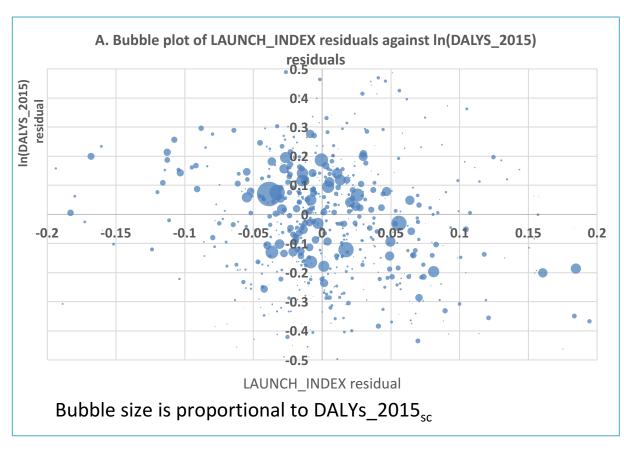
Estimated effects of new drug launches on DALYs and YLL65 in 2015

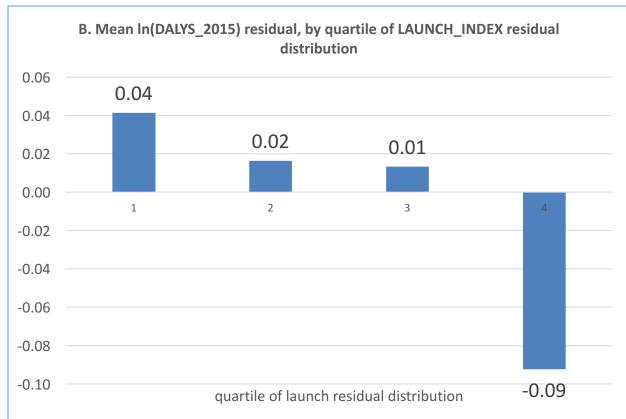






Relationship between LAUNCH_INDEX residual and ln(DALYS_2015) residual







Effect on mortality of one drug launch

- DALYs and life-years lost are unrelated to drug launches 0-4 years earlier.
- However, mortality is significantly inversely related to the number of drug launches at least 5 years earlier, especially to drug launches 5-9 years earlier.
 - One additional drug for a cancer site launched during 2006-2010 is estimated to have reduced the number of 2015 DALYs due to cancer at that site by 5.8%
 - One additional drug launched during 1982-2005 is estimated to have reduced the number of 2015 DALYs by about 2.6%.



Life-years gained calculation

Column		1	2	3	4	
			Disease burd	len measure)	
Row		DALY	YLL	YLL75	YLL65	Basis
		Life-y	ears gained	calculation		
1	b ₅₋₉	-0.058	-0.064	-0.091	-0.100	Table 2
2	weighted mean(LAUNCHES_2006_2010)	1.505	1.487	1.532	1.607	Author's calculations based on IMS New Product Focus and Theriaque databases
3	log-change in 2015 life-years lost due to LAUNCHES_2006_2010	-0.087	-0.095	-0.139	-0.161	(1) * (2)
4	life-years lost due to all types of cancer in 36 countries in 2015	88,108,225	83,467,085	30,255,229	114 451 (191	World Health Organization (2016a, 2016b).
5	reduction in 2015 life-years lost due to LAUNCHES_2006_2010	8,035,792	8,280,097	4,509,546	2,520,071	(exp(-(3))-1) * (4)



Reduction in YLL

- The estimates implied that drugs launched during 2006-2010 reduced the number of cancer DALYs in 2015 by about 8.7%
- In the absence of new drug launches during 2006-2010, there would have been 8.04 million additional DALYs due to cancer in the 36 countries.
- The estimates also implied that, in the absence of new drug launches during 2006-2010, there would have been 4.51 million additional years of life lost before age 75, and 2.52 million additional years of life lost before age 65.



Reduction in DALYs

- We also estimated that drugs launched during the entire 1982-2010 period reduced the number of cancer DALYs in 2015 by about 23.0%
- In the absence of new drug launches during 1982-2010, there would have been 26.3 million additional DALYs in 2015.
- The 9 countries with the largest number of drug launches during 1982-2010 are estimated to have had 14% fewer cancer DALYs (controlling for incidence) in 2015 than the 9 countries with the smallest number of drug launches during 1982-2010.

Column		1	2	3	4	
		Disease burden measure				
Row		DALY	YLL	YLL75	YLL65	Basis
		Pharmac	eutical expen	diture calculat	ion	
6	global cost (in millions) of oncology therapeutics and supportive care drugs in 2015, measured at invoice price levels		• ,		IMS Institute for Healthcare Informatics (2016, p. 20)	
7	36-country share of total pharmaceutical expenditure in 2014		78	3%		International Federation of Pharmaceutical Manufacturers & Associations (2017, Annex 2)
8	36-country cost (in millions) of oncology therapeutics and supportive care drugs in 2015		\$83,076			(6) * (7)
9	fraction of 2010 pharma expend. on drugs launched in country during 2001-2005, 31 countries	16%		Author's calculations based on IMS MIDAS data		
10	estimated 36-country expenditure (in millions) in 2015 on cancer drugs launched during 2006-2010		\$13	,539		(8) * (9)
	age group	All ages	All ages	Below 75	Below 65	
11	estimated age group share of cancer drug expenditure	100%	100%	76%	52%	International Agency for Research on Cancer (2017b)
12	estimated 2015 36-country expenditure (in millions) by age group on cancer drugs launched during 2006-2010	\$13,539	\$13,539	\$10,264	\$7,106	(10) * (11)
	Phari	maceutical exp	penditure per	life-year gaine	ed calculation	
13	pharmaceutical expenditure per life-year gained	\$1,685	\$1,635	\$2,276	\$2,820	(12) / (5)



Estimated cost per life-year gained

- Estimates of the cost per life-year gained in 2015 from drugs launched during 2006-2010 ranged between \$1635 (life-years gained at all ages) and \$2820 (life-years gained before age 65).
- These estimates are similar to those obtained in previous country-specific studies of Belgium, Canada, and Mexico, and are well below the estimate obtained in one study of Switzerland.



Forecast

- Mortality in 2015 is strongly inversely related to the number of drug launches in 2006-2010.
- If the relationship between mortality in 2020 and the number of drug launches in 2011-2015 is similar, drug launches 5-9 years earlier will reduce mortality even more (by 9.9%) between 2015 and 2020 than they did (by 8.4%) between 2010 and 2015.

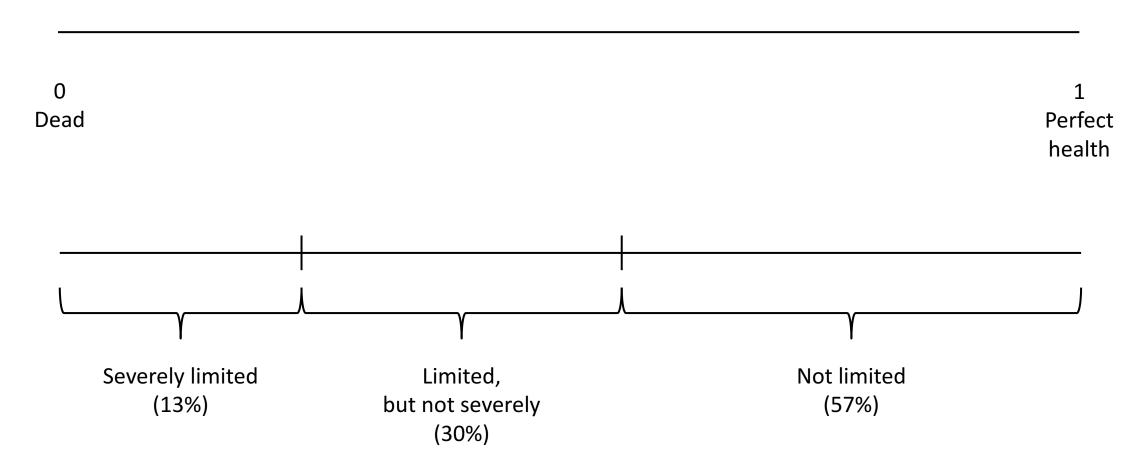


4 studies of impact of prescription drugs on health

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Quality of life





Clinical studies have shown that the use of certain drugs can reduce disability

- Post-menopausal women treated for osteoporosis with alendronate sodium for three years reported 63 per cent fewer days of disability requiring bed-rest for back pain related to those fractures.
- Etanercept, adalimumab, and infliximab reduced disability in rheumatoid arthritis patients, even those with a longstanding history and highly-active form of the disease.
- Multiple sclerosis patients given alemtuzumab (first launched in 2001) were almost twice as likely to achieve an improvement in physical disabilities as those given interferon beta-1a (first launched in 1995).

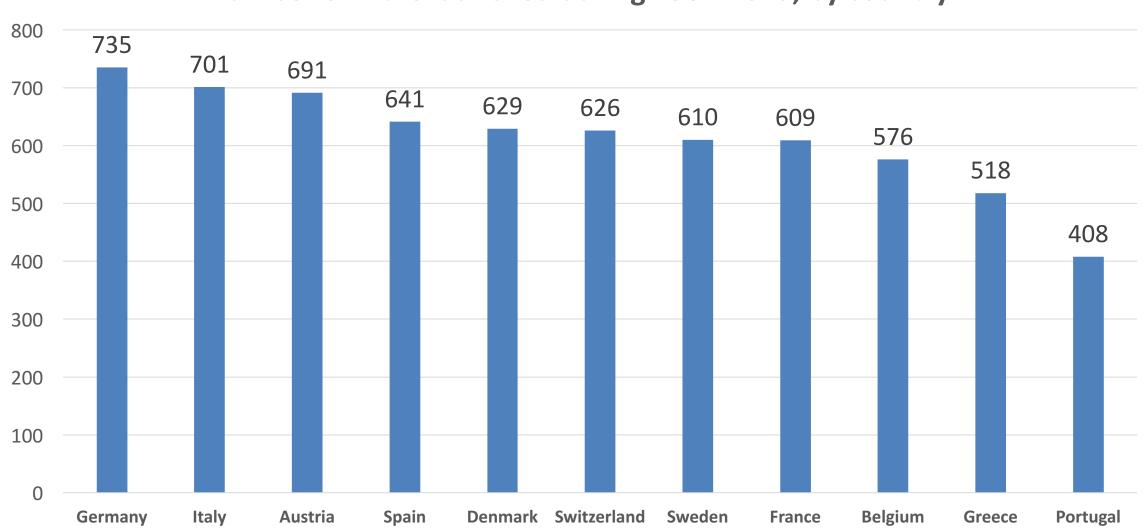


Access to prescription drugs varies across countries

- The average number of new chemical entities (NCEs) launched during 1982-2015 in Germany, Italy, and Austria (709) was 42% greater than the average number of NCEs launched in Belgium, Greece, and Portugal (501).
- Even when the total number of drugs launched in two countries is similar, the specific drugs that were launched, and the diseases those drugs are used to treat, may differ.

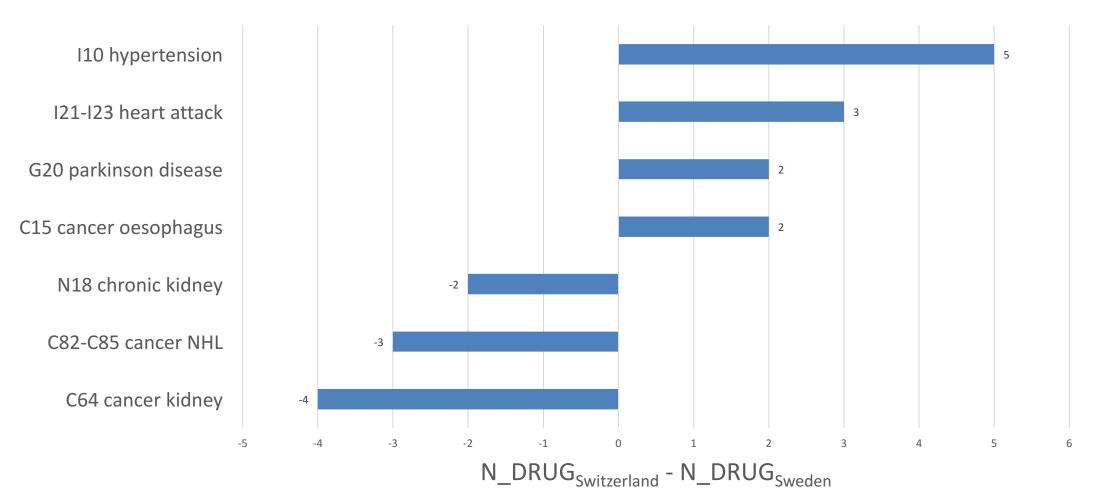


Number of NCEs launched during 1982-2015, by country





Difference between number of drugs launched during 1982-2015 for 7 diseases in Switzerland and Sweden





Main hypothesis

- I will test the hypothesis that the larger the relative number of drugs for a disease that have been launched in a country, the lower the relative disability from that disease in that country, controlling for the average level of disability (and the average number of drug launches) in each country and for each disease, and for the number of patients with the disease and their mean age.
- The hypothesis will be tested using data about 31 diseases collected from over 45,000 people aged 50 and over in eleven European countries, partially derived from the <u>Survey of Health, Ageing and Retirement in</u> <u>Europe</u>.



Impact of a drug launch on disability in 2015 is likely to depend on launch year

- Impact is likely to depend on the interaction between drug quantity (utilization) and drug quality (effectiveness)
- Drugs launched more recently tend to be used less than drugs launched many years before: utilization increases with time since launch
- But drugs launched more recently tend to be more effective than drugs launched many years before: quality decreases with time since launch
- Impact (which depends on quantity * quality) will increase with time since launch if rate of quantity increase is greater than rate of quality decrease; otherwise, impact will decrease with time since launch



Number of drug launches, by period

- 2011-2015
- 2006-2010
- 2001-2005
- 1982-2000

Disease prevalence

Mean age

Disease fixed effects

Country fixed effects











Disability, by disease and country

- have severe limitations?
- have any limitations?
- the mean number of ADL limitations
- the mean number of IADL limitations
- the mean CASP index for quality of life and well-being



Diff-in-diff model of the effect of the number of 1982-2015 drug launches on disability in 2015 (eq. (2))

$$\begin{split} & Y_{dc} = \beta_{0\text{-}4} \text{ LAUNCHES}_2011_2015_{dc} + \beta_{5\text{-}9} \text{ LAUNCHES}_2006_2010_{dc} \\ & + \beta_{10\text{-}14} \text{ LAUNCHES}_2001_2005_{dc} + \beta_{15\text{-}33} \text{ LAUNCHES}_1982_2000_{dc} \\ & + \gamma \ln(\text{PREV}_{dc}) + \rho \text{ AGE_MEAN}_{dc} + \alpha_{d} + \pi_{c} + \epsilon_{dc} \end{split}$$

where Y_{dc} is one of the following variables:

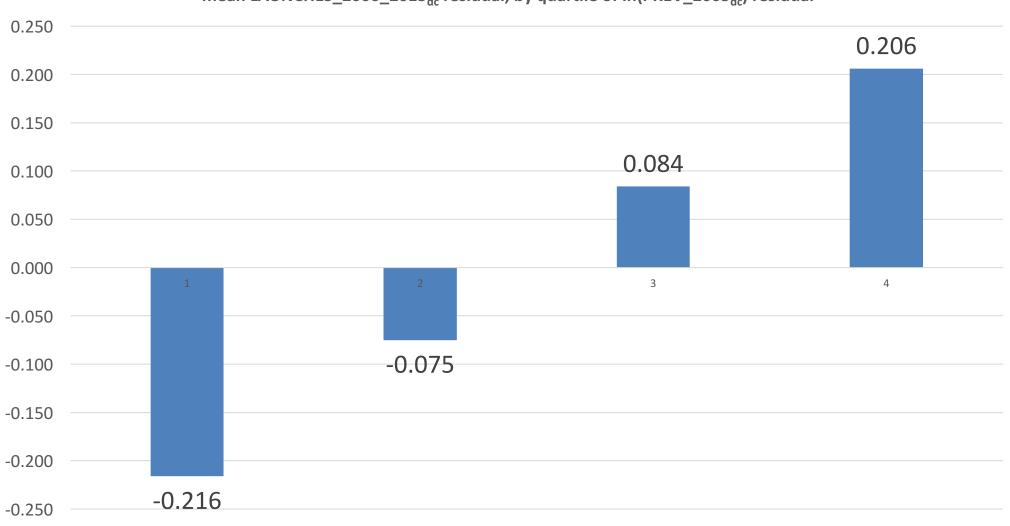
- the log-odds that individuals with disease d in country c have severe limitations
- the log-odds that individuals with disease d in country c have any limitations
- the mean number of limitations with activities of daily living of individuals with disease d in country c
- the **mean number of limitations with instrumental activities of daily living** of individuals with disease d in country c
- the mean CASP index for quality of life and well-being of individuals with disease d in country c



LAUNCHES_2011_2015 _{dc}	= the number of post-1981 new chemical entities used to treat disease d
	launched in country c during 2011-2015
LAUNCHES_2006_2010 _{dc}	= the number of post-1981 new chemical entities used to treat disease d
	launched in country c during 2006-2010
LAUNCHES_2001_2005 _{dc}	= the number of post-1981 new chemical entities used to treat disease d
	launched in country c during 2001-2005
LAUNCHES_1982_2000 _{dc}	= the number of post-1981 new chemical entities used to treat disease d
	launched in country c during 1982-2000
PREV _{dc}	= the number of people in country c who said that a doctor ever told
	them that they had disease d
AGE_{dc}	= the mean age of people in country c who said that a doctor ever told
	them that they had disease d
$\alpha_{\sf d}$	= a fixed effect for disease d
π_{c}	= a fixed effect for country c



Figure 5 mean LAUNCHES_2006_2015_{dc} residual, by quartile of ln(PREV_2005_{dc}) residual





Data sources

- Data on disability were obtained from Wave 6 of the <u>Survey of Health</u>,
 <u>Ageing and Retirement in Europe (SHARE)</u>, a multidisciplinary and cross national panel database of micro data on health, socio-economic status and
 social and family networks of more than 120,000 individuals aged 50 or
 older.
- Data on drug launch years, by molecule and country, were obtained from the IMS Health *New Product Focus database*.
- Data on the indications of each drug were obtained from the *Thériaque* database (Centre National Hospitalier d'Information sur le Médicament (2017)).

Summary statistics, by country

Country	number of persons in sample	mean age	mean number of medical conditions	% severely limited	% with any limitation	mean number of ADL limitations	mean number of IADL limitations	mean CASP index of quality of life and well-being
11 countries combined	45,592	67.8	1.6	13%	43%	0.26	0.54	37.4
Austria	3402	69.1	1.5	17%	50%	0.27	0.64	39.8
Belgium	5823	66.4	1.7	16%	48%	0.31	0.62	38.3
Denmark	3733	65.6	1.3	9%	38%	0.17	0.37	41.4
France	3948	68.0	1.6	16%	46%	0.28	0.54	37.9
Germany	4412	66.3	1.7	18%	55%	0.23	0.40	39.2
Greece	4937	66.8	1.6	8%	30%	0.18	0.50	31.8
Italy	5313	67.2	1.4	14%	40%	0.27	0.53	34.8
Portugal	1676	67.7	2.3	23%	60%	0.52	0.83	33.3
Spain	5636	70.0	1.7	7%	40%	0.37	0.82	36.1
Sweden	3906	70.4	1.3	13%	44%	0.17	0.36	39.5
Switzerland	2806	68.6	1.1	9%	35%	0.12	0.26	40.8

Summary statistics, by medical condition

(10 most prevalent conditions)

medical condition	number of conditions in sample	mean age	% severely limited	% with any limitation	mean number of ADL limitations	mean number of IADL limitations	mean CASP index of quality of life and well-being
31 medical conditions combined	62,424	69.7	17%	54%	0.36	0.73	36.4
I10 hypertension	17,438	69.2	10%	41%	0.21	0.44	37.3
E78 high cholesterol	11,080	68.0	10%	39%	0.18	0.39	37.1
M15-M19 osteoarthritis	8,858	68.9	17%	63%	0.30	0.58	36.8
E10-E14 diabetes	5,717	70.2	16%	53%	0.34	0.68	36.2
I21-I23 heart attack	4,581	73.0	24%	68%	0.42	0.92	35.8
M05-M06 rheumatoid arthritis	3,603	70.4	20%	68%	0.41	0.83	35.1
J40–J47 chronic lung	2,883	68.9	25%	70%	0.39	0.82	35.7
K25-K27 ulcer	1,626	66.6	17%	54%	0.35	0.62	34.6
I63-I64 stroke	1,571	73.1	43%	78%	1.24	2.34	34.6
G30 alzheimer	1,057	81.3	60%	89%	2.30	5.23	32.1

Number of drugs launched during 1982-2015, by country and medical condition

(10 conditions with highest mean number of launches)

medical condition	Austria	Belgium	Denmark	France	Germany	Greece	Italy	Portugal	Spain	Sweden	Switzerland	Mean
I10 hypertension	31	27	29	37	37	32	33	31	32	26	31	31.5
E10-E14 diabetes	33	29	32	33	33	32	30	24	32	32	32	31.1
C50 cancer breast	24	22	22	25	24	21	24	13	23	22	23	22.1
C91-C95 cancer leukaemia	22	17	21	21	25	14	20	7	17	20	19	18.5
M05-M06 rheumatoid arthritis	19	18	19	18	18	18	19	12	20	18	18	17.9
I21-I23 heart attack	17	15	16	15	18	16	17	11	15	14	17	15.5
C34 cancer lung	16	13	16	17	16	11	11	9	13	15	14	13.7
C82-C85 cancer NHL	15	12	14	12	15	9	0	5	11	14	11	10.7
C61 cancer prostate	11	11	12	12	11	10	9	8	11	11	11	10.6
N18 chronic kidney	10	10	11	10	11	11	11	8	10	11	9	10.2

Estimates of model of the effect of the number of drug launches on disability (eq. (2))

Column		1	2	3	4	5		
			Dependent variable					
		log-odds of	log-odds of	number of	number of	CASP index		
		severe limitation	any limitation	ADL	IADL			
Regressor				limitations	limitations			
	Estimate	-0.025	-0.015	-0.005	0.000	0.008		
LAUNCHES_2011_2015	Std. err.	0.026	0.018	0.006	0.009	0.045		
	Pr > Z	0.340	0.405	0.355	0.987	0.866		
	Estimate	-0.013	-0.006	-0.017	-0.006	0.036		
LAUNCHES 2006 2010	Std. err.	0.024	0.026	0.007	0.013	0.045		
	Pr > Z	0.595	0.809	0.020	0.638	0.423		
	Estimate	-0.022	-0.023	-0.011	-0.013	0.006		
LAUNCHES 2001 2005	Std. err.	0.006	0.010	0.006	0.010	0.033		
	Pr > Z	0.001	0.023	0.055	0.198	0.864		
	Estimate	-0.015	-0.017	-0.005	0.000	0.031		
LAUNCHES 1982 2000	Std. err.	0.005	0.006	0.004	0.006	0.016		
	Pr > Z	0.004	0.007	0.203	0.958	0.052		
	Estimate	-0.282	-0.297	-0.119	-0.039	0.478		
IMPACT	Std. err.	0.103	0.108	0.052	0.084	0.230		
	Pr > Z	0.006	0.006	0.022	0.642	0.037		
						0.110		
	Estimate	-0.232	-0.213	-0.045	-0.100	0.119		
In(PREV)	Std. err.	0.060	0.061	0.018	0.030	0.082		
	Pr > Z	0.000	0.000	0.014	0.001	0.146		
	Estimate	0.038	0.010	0.028	0.078	-0.173		
AGE_MEAN	Std. err.	0.013	0.013	0.004	0.009	0.052		
	Pr > Z	0.003	0.441	<.0001	<.0001	0.001		



Estimated effects of 1982-2015 drug launches on mean 2015 disability of people with at least one medical condition

Column	1	2	3
		counterfactual	
		(no 1982-2015	
		drug launches)	effect of 1982-2015 drug
Disability measure	actual mean	mean	launches
probability of severe limitation	16.9%	21.2%	-4.3%
probability of any limitation	53.4%	60.7%	-7.3%
mean number of ADL limitations	0.34	0.46	-0.12
mean CASP index	36.58	36.10	0.48



Cost-effectiveness

- I estimate that mean pharmaceutical expenditure in 2014 by people 45 and over in the eleven European countries was \$1184 (= 1.77 * \$667).
- Expenditure of \$1184 reduced the probability of being severely limited by 4.3 percentage points.
- If people would have been willing to pay at least \$27,318 (= \$1184 / 4.3%) to avoid being severely limited, drugs launched during 1982-2015 would have been cost-effective, even if they did not provide any other benefits, e.g. increased longevity and reduced hospitalization.
- However my previous research has demonstrated that new drug launches have also provided those benefits.



The impact of access to prescription drugs on disability in eleven European countries

- Access to prescription drugs varies across countries: the average number of new chemical entities (NCEs) launched during 1982-2015 in Germany, Italy, and Austria (709) was 42% greater than the average number of NCEs launched in Belgium, Greece, and Portugal (501).
- Even when the total number of drugs launched in two countries is similar, the specific drugs that were launched, and the diseases those drugs are used to treat, may differ.
- I test the hypothesis that the larger the relative number of drugs for a disease that have been launched in a country, the lower the relative disability from that disease in that country, controlling for the average level of disability (and the average number of drug launches) in each country and for each disease, and for the number of patients with the disease and their mean age.
- The hypothesis is tested (and confirmed) using data about 31 diseases collected from over 45,000 people aged 50 and over in eleven European countries, partially derived from the Survey of Health, Ageing and Retirement in Europe.
- The estimates imply that drug launches during 1982-2011:
 - reduced the probability of severe limitation in 2015 by 4.3 percentage points, from 21.2% to 16.9%
 - reduced the probability of any limitation by 7.3 percentage points, from 60.7% to 53.4%
 - reduced the mean number of Activities of Daily Living limitations by about 26%
- Drug launches also yielded a small but significant increase in an index of quality of life and well-being.
- The population age 50 and over of these 11 countries in 2015 was 128 million, so we estimate that drug launches during 1982-2011 reduced the number of severely limited people in these countries by 5.5 million (= 4.3% * 128 million).
- Disability could have been reduced even more if there had been greater access to prescription drugs.



4 studies of impact of prescription drugs on health

study	health outcome	drug measure	region(s)	data
				19 cancer sites,
1	mortality	number of drugs	36 countries	36 countries
				31 diseases, 11
2	disability	number of drugs	11 European countries	countries
				X diseases, 15
3	hospitalization	vintage of drugs	USA	years
				500,000
4	mortality	vintage of drugs	Puerto Rico	patients

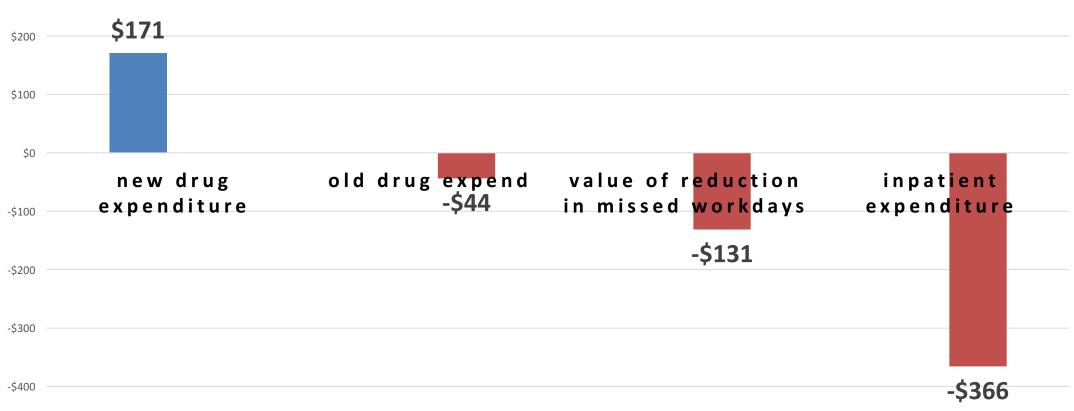


Pharmaceutical innovation is often cost-saving, not just cost-effective

- In a recent study (Lichtenberg (2014)), I investigated whether diseases subject to more rapid pharmaceutical innovation experienced greater declines in Americans' disability days and use of medical services during the period 1997–2010, controlling for several other factors, using data from a U.S. government household survey.
- The mean number of work loss days, school loss days, and hospital admissions
 declined more rapidly among medical conditions with larger increases in the mean
 number of new (post-1990) prescription drugs consumed.
- The value of reductions in work loss days and hospital admissions attributable to pharmaceutical innovation was estimated to be three times as large as the cost of new drugs consumed.
- Lichtenberg FR (2014), "The impact of pharmaceutical innovation on disability days and the use of medical services in the United States, 1997-2010," Journal of Human Capital 8(4): 432-480.



Impact of pharmaceutical innovation on per capita drug expenditure, work-loss days, and inpatient expenditure, USA, 2010



Lichtenberg FR (2014), "The impact of pharmaceutical innovation on disability days and the use of medical services in the United States, 1997-2010," Journal of Human Capital 8(4): 432-480.



4 studies of impact of prescription drugs on health

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The Effect of Drug Vintage on Survival: Micro Evidence From Puerto Rico's Medicaid Program

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THE EFFECT OF DRUG VINTAGE ON SURVIVAL: MICRO EVIDENCE FROM PUERTO RICO'S MEDICAID PROGRAM

Frank R. Lichtenberg

ABSTRACT

Using micro data on virtually all of the drugs and diseases of over 500,000 people enrolled in Puerto Rico's Medicaid program, the impact of the vintage (original FDA approval year) of drugs used to treat a patient on the patient's three-year probability of survival, controlling for demographic characteristics (age, sex, and region), utilization of medical services, and the nature and complexity of illness are examined. It is found that people using newer drugs during January—June, 2000, were less likely to die by the end of 2002, conditional on the covariates. The estimated mortality rates are strictly declining with respect to drug vintage. For pre-1970 drugs, the estimated mortality rate is 4.4%. The mortality rates for 1970s, 1980s, and 1990s drugs are 3.6%, 3.0%, and 2.5%, respectively. The actual mortality rate is about 16% (3.7% vs. 4.4%) lower than it would have been if all of the drugs utilized in 2000 had been pre-1970 drugs. Estimates for subgroups of people with specific diseases display the same general pattern.

Pharmaceutical Markets and Insurance Worldwide

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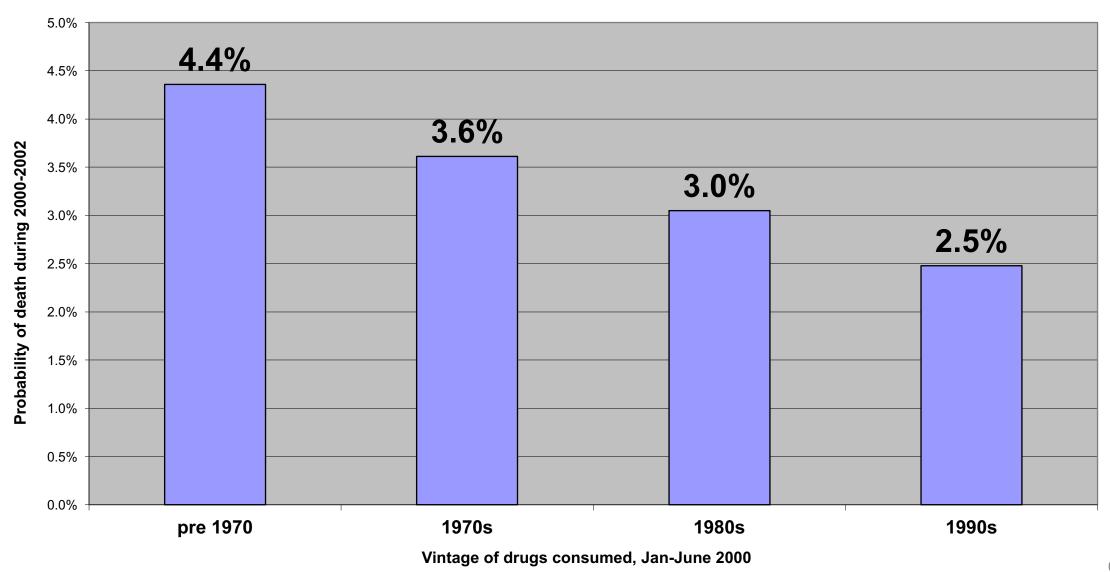
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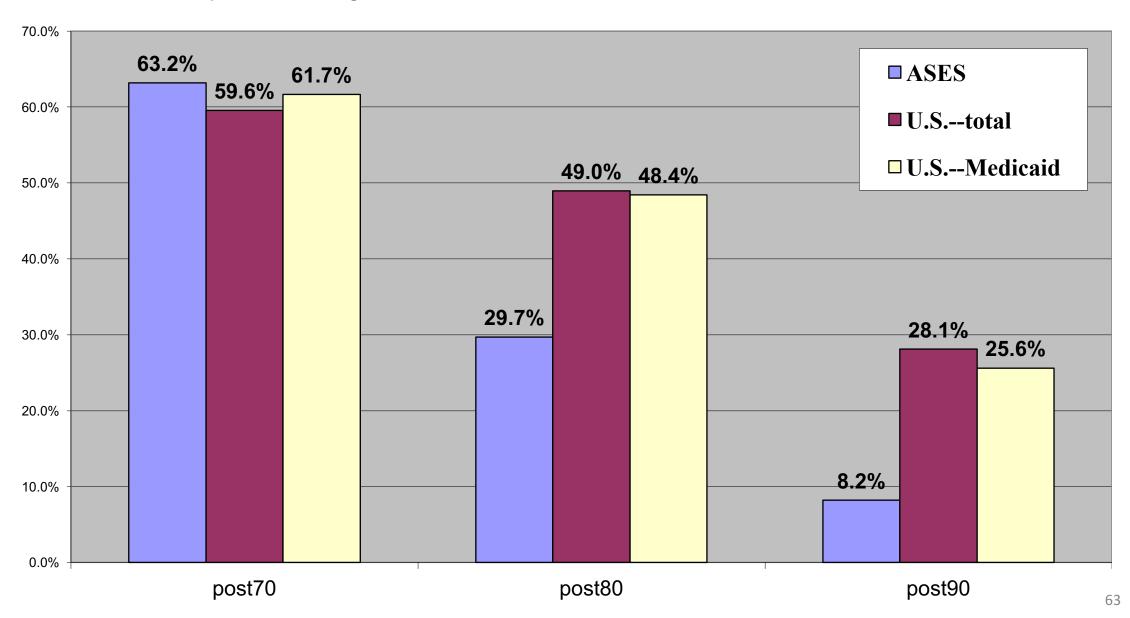


- Using micro data on virtually all of the drugs and diseases of over 500,000 people enrolled in Puerto Rico's Medicaid program, I examined the impact of the vintage (original FDA approval year) of drugs used to treat a patient on the patient's three-year probability of survival, controlling for:
 - demographic characteristics (age, sex, and region)
 - utilization of medical services
 - the nature and complexity of illness
- I found that people using newer drugs during January—June 2000, were less likely to die by the end of 2002, conditional on the covariates.
- The actual mortality rate was about 16% (3.7% vs. 4.4%) lower than it would have been if all of the drugs utilized in 2000 had been pre-1970 drugs.
- Estimates for subgroups of people with specific diseases displayed the same general pattern.

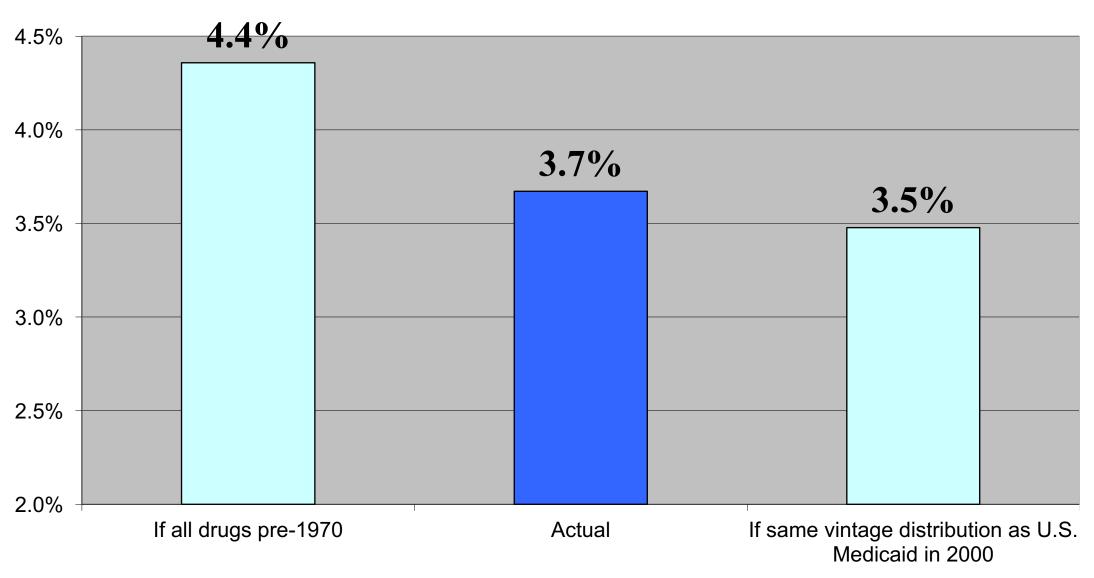
Estimated vintage-specific mortality rates, entire ASES population



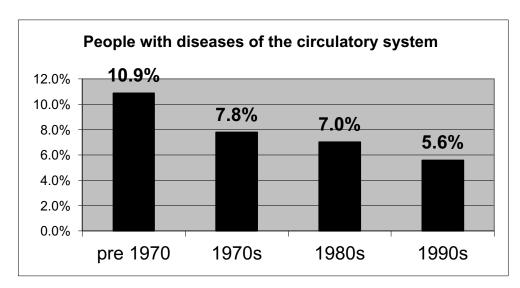
Comparison of vintage distributions of ASES Rx's, all U.S. Rx's, and U.S. Medicaid Rx's

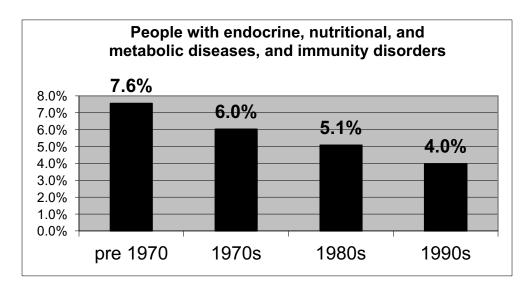


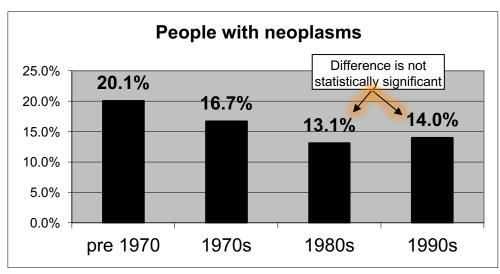
Actual vs. hypothetical ASES mortality rates













Pharmaceutical innovation has increased longevity and reduced disability and hospitalization

Pharmaceutical innovation

(development and use of new drugs)







Longevity

Disability

(work- and school-loss days, quality of life

Hospitalization

and nursing home use



Thank You

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