

Trade Mark Cluttering - Evidence from EU Enlargement

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Agenda

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Infinite Name Spaces?

- ▶ Landes and Posner (1987) argue that the supply of names for trademarks is unlimited.
- ▶ Regulation of names for prescription drugs by FDA and EMA introduces scarcity.
- ▶ Competition for viable names is leading to multiple applications per drug - some parallels to the patent thickets problem. (Shapiro, 2001; Hall and Ziedonis, 2001)
- ▶ Only one name will be used, most of the remainder “clutter” the register.
- ▶ Not much is known about the importance of cluttering.
- ▶ **Research question:** Did the 2004 expansion of the european trade mark system induce a jump in simultaneous applications?



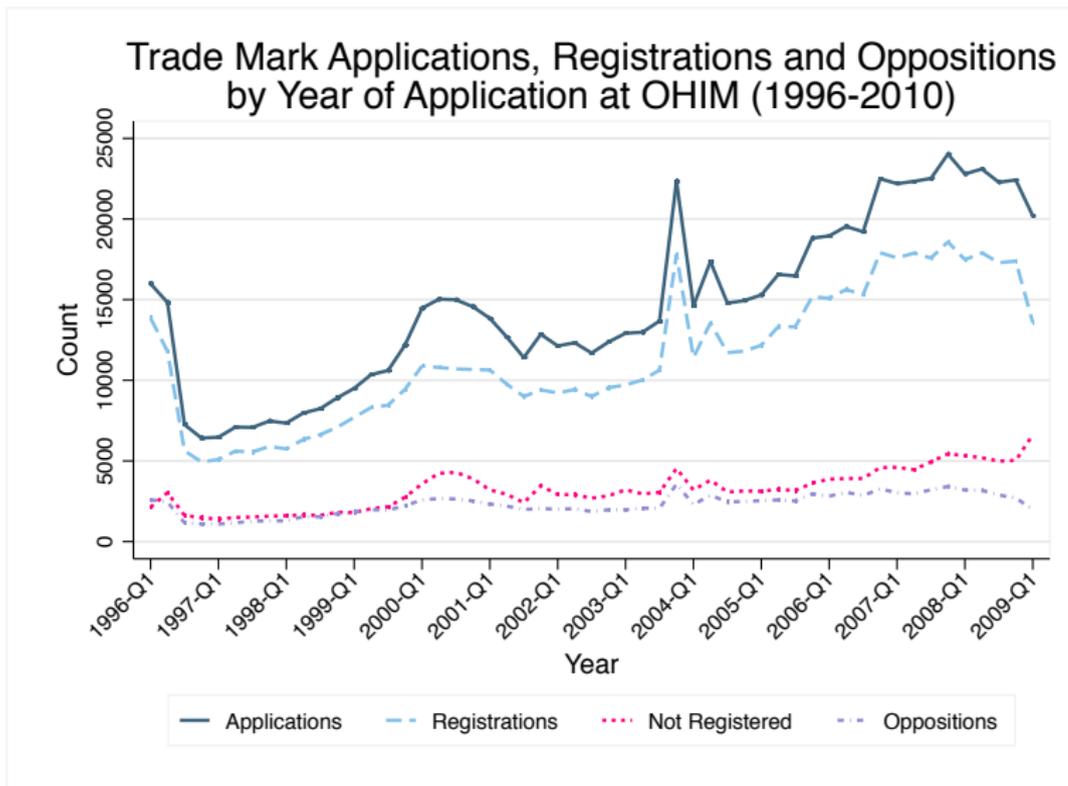


Figure : Demand for the Community Trade Mark

Simultaneous applications and Extension of TM System

- ▶ Are there simultaneous applications leading to unused trade marks?

A recent survey suggests the problem exists.

(Kur et al., 2011).

- ▶ “..pharmaceutical manufacturers are routinely filing five to ten different trademarks for each trial drug in their pipelines.”

(Lallemand, 2011)

- ▶ Enlargement:

On the 1.5.2004 Czech Republic, Estonia, Cyprus, Latvia, Lithuania, Hungary, Malta, Poland, Slovenia and Slovakia joined EU and thereby OHIM and EMA.

- ▶ As trade marks accumulate on the register it becomes harder to ensure a new mark is not opposed or rejected.



The Naming Committee at EMA

- ▶ Losec (proton pump inhibitor) and Lasix (hypertension)
- ▶ FDA forces AstraZeneca to change Losec to Prilosec
- ▶ Prilosec was subsequently mistaken for Prozac by a pharmacist. The patient had a gastric ulcer!!

- ▶ The naming committee exists to prevent such mistakes.
- ▶ The naming committee works on the premise that mistakes can happen and will often be dangerous - it is very restrictive.

- ! Interestingly the opposition chambers at OHIM operate on the premise that in pharmaceuticals everybody is extremely attentive, due to high costs of error.



What drives simultaneous applications?

- ▶ Trade mark approval rates at EMA:

	2000	2001	2002	2003	2004	2005	2006	2007	2008
# Reviewed	177	132	104	109	146	204	315	311	455
% Accepted	63%	54%	70%	67%	63%	51%	52%	53%	57%

Note the “break” after 2004.

- ▶ Ideally a pharmaceutical company would like to secure the same trademark worldwide.
- ▶ It faces at least USPTO, OHIM, FDA and EMA.
- ▶ Rejection of trademarks means a product launch is delayed → big reductions in share prices.



Cost of simultaneous applications?

- ▶ Creating a set of names for a pharmaceutical product costs US \$ 100,000 - US \$ 700,000 (Kenagy and Stein, 2001) and sometimes even US \$ 2.25 million (Wick, 2011).
- ▶ This is for: prescription simulation exercises, tests of name similarity, tests of implied claims conveyed by a name, tests of visual and verbal similarity, linguistic analysis.
- ▶ Important question that remains unanswered here: which proportion of these costs is due to cluttering?
- ▶ Creating names for other products costs US \$ 9400 in 2004 prices (Kohli and LaBahn, 1997).



Experiment?

- ▶ Medical name regulation imposes costs.
- ▶ The European Union is enlarged in 2004.
- ▶ Pharmaceutical firms face a large increase in costs of getting names approved after 2004.
- ▶ Can try to use this to estimate how much cluttering is going on.

But, need to answer additional questions:

- ▶ Is this an experiment?
- ▶ What are the potential outcomes?
- ▶ How does assignment work here?



A model

- ▶ Firms make long run, uncertain R&D investments and short run marketing and regulatory compliance investments.
- ▶ R&D investments in period t pay off in $t + 2$, marketing and compliance investments are made in $t + 1$.
- ▶ Firms invest to maximize profits.

- ▶ **Result:** A regulatory change in period T that is announced in period $T - 1$ leads to increased trade mark applications per product for the cohort of R&D investments made in period $T - 2$.
- ▶ **Result:** R&D investments made in period $T - 1$ will adjust for the higher cost of compliance - i.e. fewer projects.



Counterfactuals and Outcomes

- ▶ What would have been pharmaceutical firms' simultaneous application numbers, absent EU enlargement?
- ▶ Use Difference-in-Differences to construct counterfactual using firms not affected by pharmaceuticals.

$$A_i = \alpha + \beta T_i + \sum_{g=1}^4 \gamma_g \cdot 1[G_i = g] + \sum_{g=1}^3 \tau_g \cdot 1[G_i = g, T_i = 1] + \mathbf{X}'\boldsymbol{\theta} + \epsilon_i, \quad (1)$$

A_i – Applications $T_i (= 1)$ EU enlargement indicator

$G_i (= g)$ group indicator \mathbf{X} covariates

Assn. Common trends for both types of firms - how likely?

- ▶ Pharmaceutical firms tend to be large - concentrated markets.
- ▶ Size is imperfectly observed here - have no matched firm data!



Alternative: Apply the Rubin Causal Model

- ▶ Potential outcomes: number of names simultaneously applied for given level of regulation.
 - ▶ Assignment not random, as firms do not randomly choose to become or stay in pharmaceuticals in 1996-1999.
 - ▶ Is assignment unconfounded?
- ⇒ Imbens (2004): if treatment participation (being a pharmaceutical firm) is a separate decision from the outcome (how many names to apply for) then it may be.
- ⇒ It will be if treatment uptake depends on variables that do not affect outcomes, even if these variables are not observable!
- ▶ Note here that treatment uptake and outcomes are based on decisions taken several years apart.



Who is Treated?

Trade marks may be registered in multiple Nice classes simultaneously:

- ▶ define *related* to Pharmaceuticals if more than 24% of applications in the class are also registered in class 5.
- ▶ define *unrelated* to Pharmaceuticals if less than 5% of applications in the class are also registered in class 5.

Table : Groups in Sample

Applications in Classes Related to Pharmaceuticals	Applications in Classes un-related to Pharmaceuticals		
	No	Yes	Total
No	71,226	395,991	467,217
Yes	76,201	54,806	131,007
Total	147,427	450,797	598,224

Note: EU Enlargement in 2004 did not have statistically significant effects on proportion of application events in **pharmaceuticals** and **artifacts**.



Research Design

- ▷ I compare the number of simultaneous trade marks applied for in four sets of trade mark classes before and after 2004:

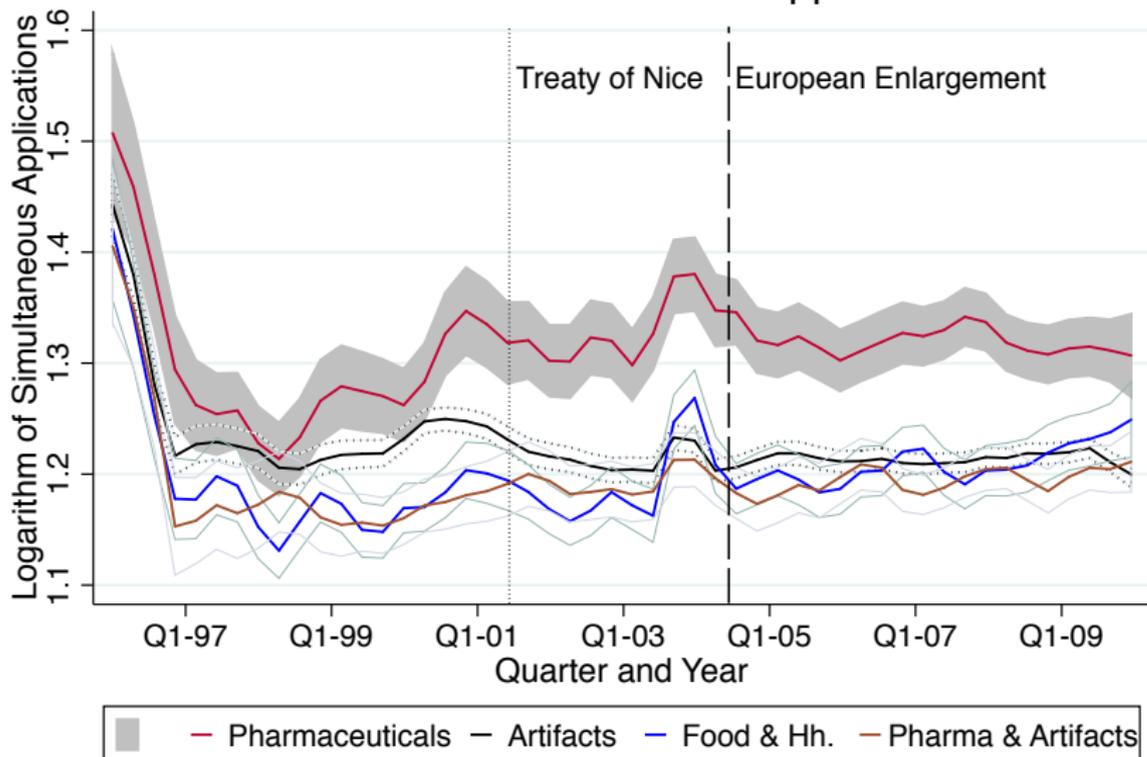
Table : Distribution of Trade Mark Applications

	Type of Industry		
	Artifacts	Food & Household	Pharmaceuticals
Nice Classes	6, 7, 9, 12, 16, 18 19, 25, 28, 33, 35, 36, 37, 38, 39, 41, 43 ,45	11, 20, 21, 29, 30, 31, 32, 42	1, 3, 5, 10, 13, 44
Des- crip- tion	Metals, Machines, Scientific Apparatus, Vehicles, Leather, Building, Clothing, Games, ...	Lighting, Furniture, Household utensils Food, Coffee, Produce, Beer,	Chemicals, Laundry, Pharmaceuticals, Medical Apparatus, Firearms, Medical Services
N	395,991	71,226	76,201



Graphical Analysis - I

Incidence of Simultaneous Applications



Data are 597,450 application events - these are units of analysis.

Descriptive Statistics

Variable	Mean	S.D.	O.C. Mean	Min.	Max.
Simultaneous Applications Count	1.200	1.317	1.194	1	634
Fee Dummy	0.426	-	0.426	0	1
Anticipation Dummy	0.197	-	0.197	0	1
Expansion Dummy	0.549	-	0.549	0	1
Pharmaceutical Dummy	0.127	-	0.127	0	1
Food & Household Dummy	0.092	-	0.092	0	1
Pharmaceuticals & Artifacts	0.119	-	0.119	0	1
Seniority Dummy	0.074	-	0.074	0	1
Breadth	2.738	2.400	2.738	0	45
Opposition rate	0.170	0.134	0.170	0	1
Opposition rate × Food & Hh.	0.026	0.076	0.026	0	1
Opposition rate × Pharma.	0.019	0.100	0.019	0	1
Opposition rt. × Pharma & Atfs.	0.019	0.058	0.019	0	1
Pre-existing TM Mark stock	2.980	16.945	2.977	0	349

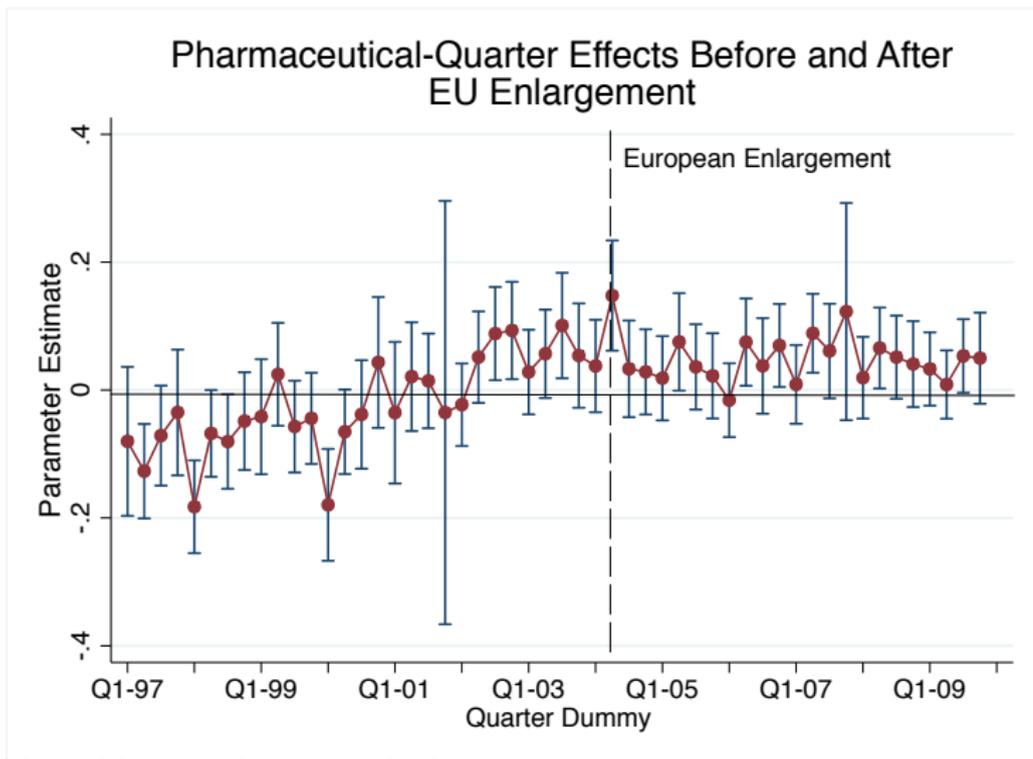


Table 3: Results from Difference-in-Differences Models

	Base	Outliers	Quarter dummies	Time Trend
Expansion dummy	-0.014 (0.029)	-0.040** (0.013)	-0.028* (0.012)	-0.026* (0.013)
Pharma dummy	0.004 (0.018)	0.009 (0.016)	0.005 (0.018)	-0.042 (0.131)
Food & Hh dummy	-0.033* (0.015)	0.016 (0.016)	-0.034* (0.015)	-0.217* (0.091)
Pharma & Artifacts dummy	0.024 [†] (0.013)	0.018 [†] (0.011)	0.024 [†] (0.013)	0.097 (0.097)
Anticipation × Pharma	0.059*** (0.014)	0.061*** (0.013)	0.058*** (0.014)	0.054** (0.020)
Expansion × Pharma	0.043*** (0.013)	0.043*** (0.012)	0.043*** (0.013)	0.034 (0.028)
Expansion × Food & Household	0.018* (0.008)	0.020** (0.007)	0.018* (0.008)	-0.017 (0.019)
Expansion × Pharma & Artifact	0.016* (0.008)	0.021** (0.007)	0.016* (0.008)	0.030 (0.021)
R-squared	0.220	0.114	0.220	0.220
N	597450	597339	597450	597450

[†] p<0.10, * p<0.05, ** p<0.01, *** p<0.001. Robust standard errors in parentheses

Robustness - Stability over time



Note: Vertical lines mark two standard errors.

What do the estimated effects imply?

- ▷ Note that I observe:

$$\bar{n}_j = \frac{Om_j + NM_j}{N + O} = \frac{\lambda Am_j + (1 - \lambda)AM_j}{A} = \lambda m_j + (1 - \lambda)M_j \quad (2)$$

- ▷ I assume that the probability of passing name review in each country (ρ) is identical and use the table provided by Lallemand (2011) to calculate this probability.
- ▷ I assume that λ, ρ do not change 2004.
- ▷ Combining this with sample values of n_l, n_h, m_l, m_h and the estimate of τ_p and a cost of US \$ 25,000 per name it is possible to work out that between 2001 and 2004 per year US \$ 17.7 million was spent on inventing surplus names.
- ▷ This is likely a very conservative lower bound on the true costs.



Are these effects reliable?

- ▶ Covariate balance (Imbens and Wooldridge, 2009)?
- ▶ Alternative is to use matching and bias adjusted matching (Abadie et al., 2004).
- ▶ To mimic DID I match on covariates and the average of the dependent variable before expansion (Abadie and Imbens, 2011).
- ▶ PATT or PATE? - DID estimates PATT.

Table 4: Covariate Balance Before EU Enlargement

Variable	Treated	(sd)	Controls	(sd)	Norm. Diff.
	Pharmaceuticals		Artifacts		
Breadth	1.74	1.00	2.68	1.73	-0.489
Opposition rate	0.20	0.09	0.14	0.08	0.776
Registration rate	0.78	0.14	0.71	0.21	0.395
Past applications	1.00	1.01	0.74	2.48	0.195
New combination dummy	0.50	0.50	0.69	0.46	-0.377
Seniorities	8.56	31.46	2.20	13.64	0.275
Age (days)	543.99	2507.11	346.78	1801.60	0.072
No seniorities dummy	0.93	0.26	0.95	0.23	-0.075

Normalized differences calculated using pstest in STATA.

Note: Imbens and Wooldridge (2009) and Imbens and Rubin (2011) suggest that the normalized differences should not be greater than a quarter.



Matching conditioning on 2004

Table 5: Results from Matching Estimators

Number of matches	PATT		PATE	
	Robust, Bias adjusted		Robust, Bias adjusted	
1	0.1806*** (0.0441)	0.1879*** (0.0429)	0.0862*** (0.0289)	0.0242 (0.0258)
	53.21%		57.60%	
4	0.1417*** (0.0378)	0.1675*** (0.0352)	0.1002*** (0.0268)	0.0356 (0.0233)
	45.00%		50.72%	

N=21162

† $p < 0.10$, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$. Standard errors in parentheses



Matching conditioning on 2001 & 2004

Table 6: Results from Matching Estimators

Number of matches	PATT		PATE	
	Robust, Bias adjusted		Robust, Bias adjusted	
1	0.1676*** (0.0444)	0.1796*** (0.0404)	0.1059*** (0.0303)	0.0340 (0.0269)
	52.58%		56.96%	
4	0.1383*** (0.0379)	0.1529*** (0.0363)	0.1138*** (0.0286)	0.0754*** (0.0261)
	43.90%		49.20%	

N=21162

† $p < 0.10$, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$. Standard errors in parentheses



What do I find?

- ▶ Simple graphical evidence is not favourable to a big jump in cluttering.
- ▶ Simple estimation results (DiD) suggest there was a substantial jump in cluttering.
- ▶ Estimating time period specific treatment effects also suggests a jump, also some evidence of a jump before 2004, anticipating expansion.
- ▶ Once a time trend is included the (DiD) treatment effect falls and is no longer significant, the trend is also not significant. Results are ambivalent here.
- ▶ Results from a matching estimator suggest significant effects of enlargement on simultaneous applications.
These effects are **four** times larger than those obtained from DID.



Conclusion

- ▶ If group membership is exogenous to expansion then results show either (i) that 2004 there was a significant increase in simultaneous applications, or (ii) that there is a positive time trend in simultaneous applications.
Both of these results suggest something is going on.
- ▶ The matching estimator shows that effects are quite significant.
- ▶ Extension to comparison with US or Germany - DDD?
- ▶ Other control groups - different types of pharmaceutical firms
- ▶ Control for rate of conversion (applications to registrations)
- ▶ Control for reassignments

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