
Anorganic growth in Pharma and regulations

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Teva Pharmaceuticals

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Teva Pharmaceuticals

60+
markets

Over the counter
medicines
& active
pharmaceutical
ingredients

2016 revenues:
\$21.9
billion

The world's largest
'medicine cabinet'

Serving 200 million
people every day



A strong
specialty
medicines
portfolio

The leading
global generic
company

50,000+
employees



Teva's history



1901 - 1940

A new pharmaceutical industry is founded

- 1901: Established in Jerusalem by Chaim Salomon, Moshe Levin and Yitschak Elstein.

1960 - 1980

Consolidation of the local pharmaceutical industry

- 1976: Eli Hurvitz forms Teva Pharmaceutical Industries Ltd.

1980 - 1990

Global expansion

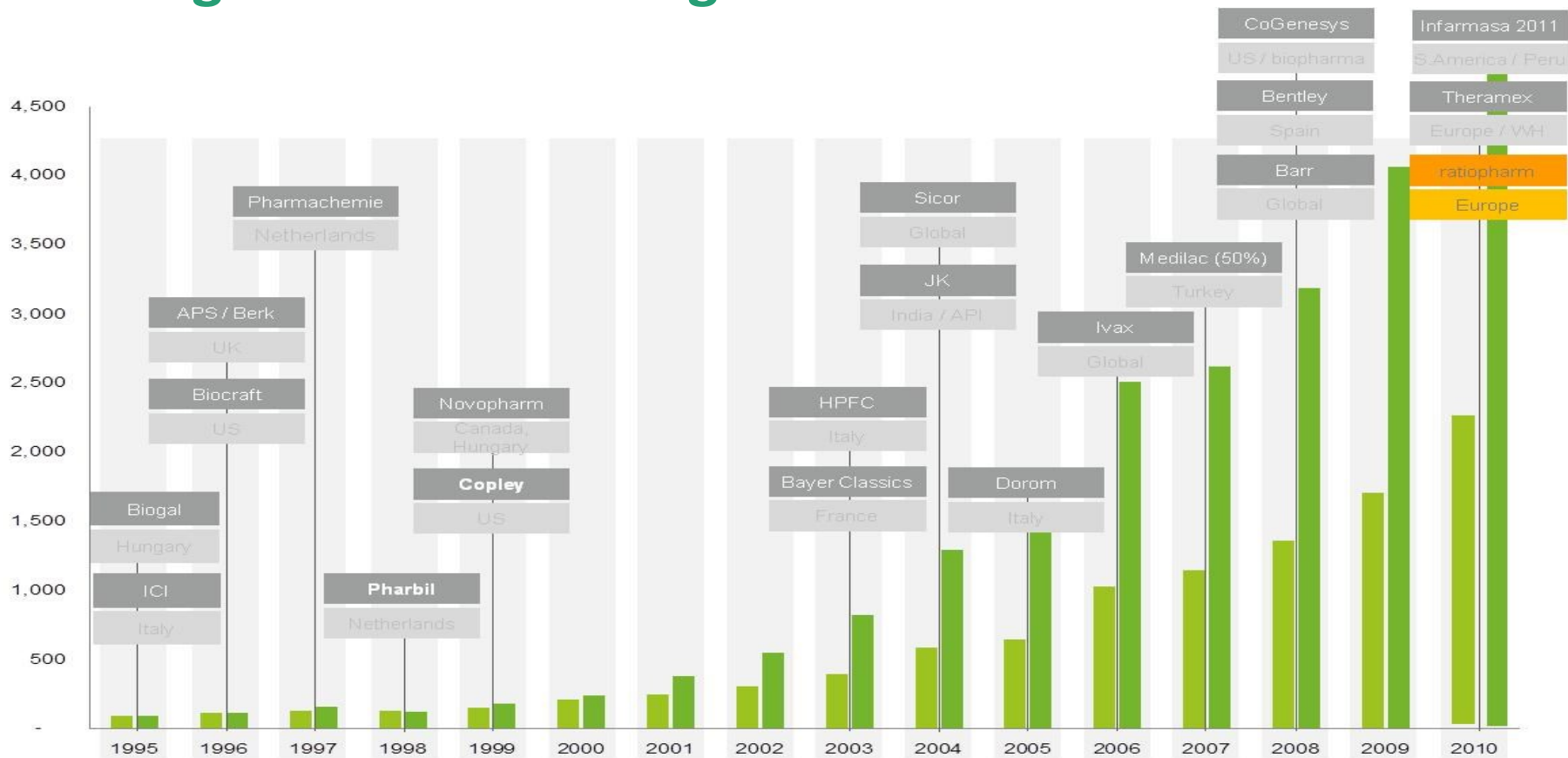
- 1984: Hatch-Waxman Act paves way for U.S. generic entry
- 1987: Teva begins trading on NASDAQ

1990 - present

A global leader in generics / establishes specialty franchise

- 1996: Teva launches Copaxone® in the US.
- 2006: Teva acquires Ivax
- 2016: Teva acquires Actavis Generics

Teva grew and become global leader via M&As



*Index 1995 = 100

Are there still „so many“ deals in Pharma?

- Yes indeed, although the number is decreasing in the last year or two
- Lower number of deals in 2016 – possibly Brexit impact, Donald Trump election and pressure on pharma industry in general, tax inversion deals under higher scrutiny, six M+As deals over \$5bn in 2016 compared to nine in 2015

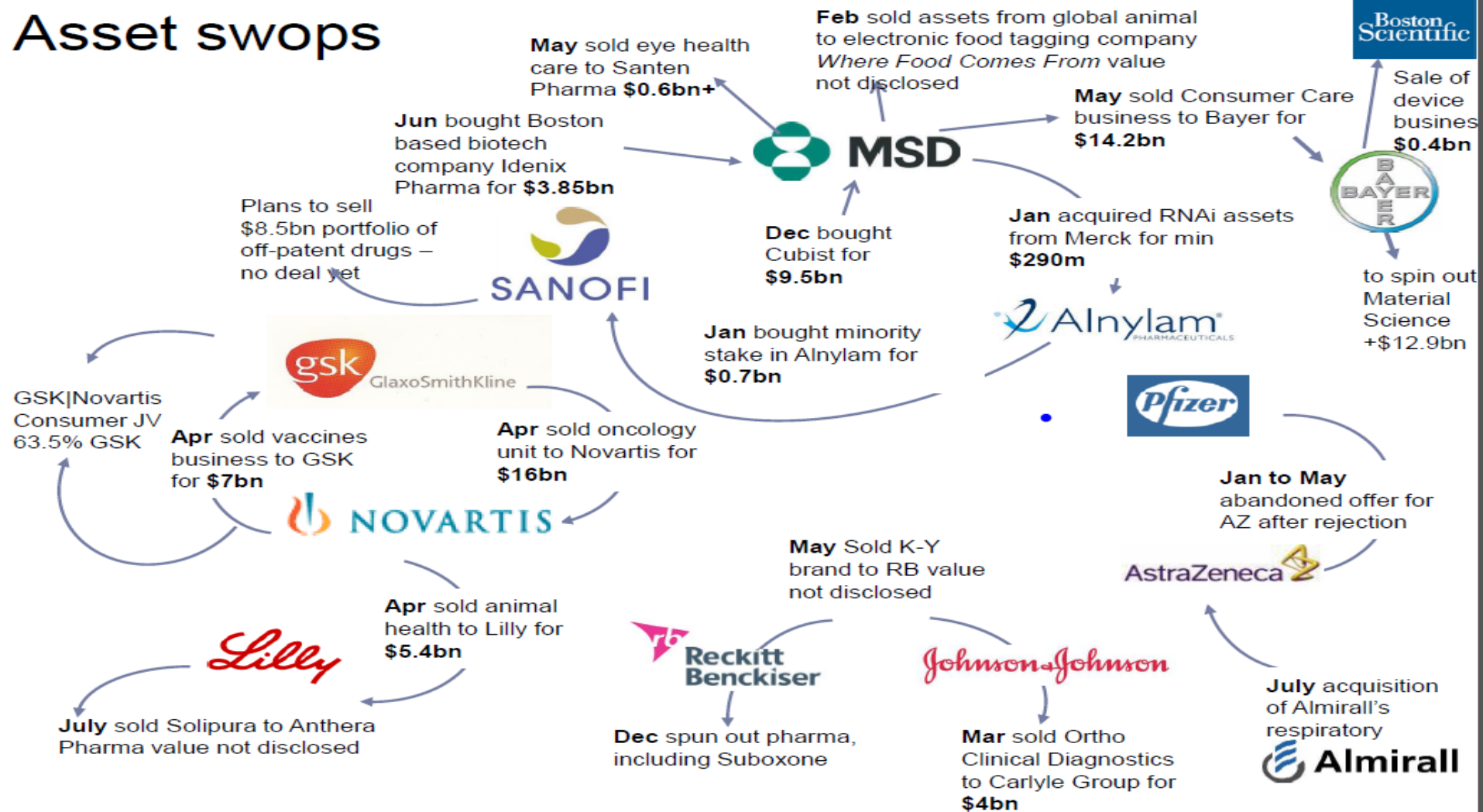
Acquirer	Target	Price \$bn	Sales multiple	Net profit multiple	Share price prem.
Teva	Allergan generics	40.5	6x	15x EBITDA	NA
Novartis	Alcon	51.6	7.2x	23.5x	NA
Sanofi	Genzyme	20.1+	5x	>500	40%
Teva	Cephalon	6.8	2.3x	5.5x	44%
Teva	ratiopharm	5.0	2.25x	10x	NA
GSK	HGH	3.6	27x	n/a	99%

Are there still „so many“ deals in Pharma?

- New deals are not necessarily classical company M&As but more and more creative
 - JVs,
 - Product Acquisitions / Licensing
 - Portfolio Swaps
 - Asset deals
 - Product divestments / switches by big pharma continue at a low level
- New R+D activities continue full speed

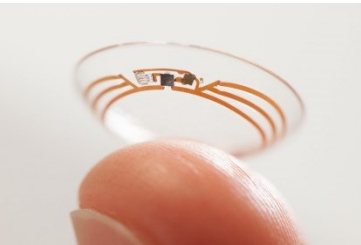
Top 25 companies	2016 number of deals*	% growth 2016	% growth 2015
Partnering-in	296	-15%	29%
Partnering-out	146	-19%	50%
M&A	53	0%	39%

Asset swaps



Are there still „so many“ deals in Pharma?

Digital health is a major area for company deal investments (e.g smart inhalers, big data deployment, smart eye lenses, smart phones in treatment ...)

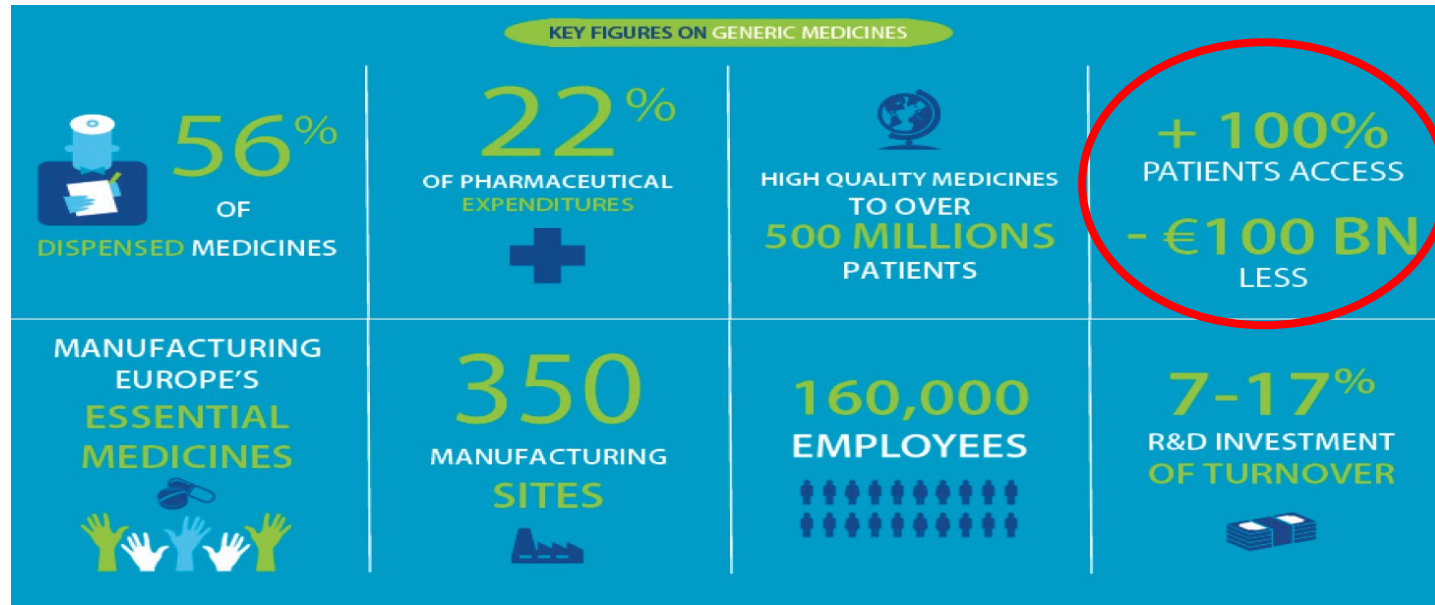


Why deals in Pharma?

- **Digital** era forces pharma sector to look into this area as well as a part of the necessary innovation
- Critical mass, R&D pipeline, Cost Savings, Geography, Synergy, Cheap Money, Re-Focus
- **R&D costs very high** (innovative products up to 2B per product), savings to finance R&D; sharing costs is now common approach, start-ups can tolerate higher development risk, cheaper to buy than built,
- Huge pressure from the payers – **austerity measures** are driving costs of pharmaceuticals down – economy of scale needs to help to cover price erosion and to free up resources for further R&D activities in this highly innovative sector

What it brings to payors and patients?

- Innovation – **new medication** for unmet medical needs, increased costs
- Affordability – **affordable medicines** save billions for healthcare systems worldwide



Position on current merger control thresholds

- Introduction of value-based threshold designed to fill an « enforcement gap »
 - allow the review of potentially problematic transactions that are not caught by revenue based thresholds
 - In the pharma sector: acquisition of pipelines that do not generate revenues but may still be competitively significant
- Added value of new thresholds is questionable, in particular in the pharma space
 - No experience of a significant deal that would have stayed below the radar (very few examples given by the Commission)
 - Increased burden and costs for the companies: need to file many deals with no competitive impact, complicates the initial assessment of whether the deal is reportable
 - Ultimately a question of proportionality as all thresholds are „imperfect“: filling a tiny gap by creating a huge burden is not appropriate
 - Burden and complication also for the authority (relevant market; specific market conditions, etc.)

Thank
you

Enabling
people to
live better,
healthier
lives

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This presentation contains forward-looking statements, which are based on management's current beliefs and expectations and involve a number of known and unknown risks and uncertainties that could cause our future results, performance or achievements to differ significantly from the results, performance or achievements expressed or implied by such forward-looking statements. Important factors that could cause or contribute to such differences include risks relating to: our ability to develop and commercialize additional pharmaceutical products; competition for our specialty products, especially Copaxone® (which faces competition from orally-administered alternatives and a generic version); our ability to integrate the acquisition of Allergan plc's worldwide generic pharmaceuticals business ("Actavis Generics") and to realize the anticipated benefits of such acquisition (and the timing of realizing such benefits); the fact that following the consummation of the Actavis Generics acquisition, we are dependent to a much larger extent than previously on our generic pharmaceutical business; potential restrictions on our ability to engage in additional transactions or incur additional indebtedness as a result of the substantial amount of debt incurred to finance the Actavis Generics acquisition; the fact that for a period of time following the consummation of the Actavis Generics acquisition, we will have significantly less cash on hand than previously, which could adversely affect our ability to grow; ; the possibility of material fines, penalties and other sanctions and other adverse consequences arising out of our ongoing FCPA investigations and related matters; our ability to achieve expected results from investments in our pipeline of specialty and other products; our ability to identify and successfully bid for suitable acquisition targets or licensing opportunities, or to consummate and integrate acquisitions; the extent to which any manufacturing or quality control problems damage our reputation for quality production and require costly remediation; increased government scrutiny in both the U.S. and Europe of our patent settlement agreements; our exposure to currency fluctuations and restrictions as well as credit risks; the effectiveness of our patents, confidentiality agreements and other measures to protect the intellectual property rights of our specialty medicines; the effects of reforms in healthcare regulation and pharmaceutical pricing, reimbursement and coverage; competition for our generic products, both from other pharmaceutical companies and as a result of increased governmental pricing pressures; governmental investigations into sales and marketing practices, particularly for our specialty pharmaceutical products; adverse effects of political or economic instability, major hostilities or acts of terrorism on our significant worldwide operations; interruptions in our supply chain or problems with internal or third-party information technology systems that adversely affect our complex manufacturing processes; significant disruptions of our information technology systems or breaches of our data security; competition for our specialty pharmaceutical businesses from companies with greater resources and capabilities; the impact of continuing consolidation of our distributors and customers; decreased opportunities to obtain U.S. market exclusivity for significant new generic products; potential liability in the U.S., Europe and other markets for sales of generic products prior to a final resolution of outstanding patent litigation; our potential exposure to product liability claims that are not covered by insurance; any failure to recruit or retain key personnel, or to attract additional executive and managerial talent; any failures to comply with complex Medicare and Medicaid reporting and payment obligations; significant impairment charges relating to intangible assets, goodwill and property, plant and equipment; the effects of increased leverage and our resulting reliance on access to the capital markets; potentially significant increases in tax liabilities; the effect on our overall effective tax rate of the termination or expiration of governmental programs or tax benefits, or of a change in our business; variations in patent laws that may adversely affect our ability to manufacture our products in the most efficient manner; environmental risks; and other factors that are discussed in our Annual Report on Form 20-F for the year ended December 31, 2015 and in our other filings with the U.S. Securities and Exchange Commission (the "SEC"). Forward-looking statements speak only as of the date on which they are made and we assume no obligation to update or revise any forward-looking statements or other information, whether as a result of new information, future events or otherwise.