ICORD 2008



Access to Information & to Orphan Products: Industry's View

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25 years of US Orphan Drug Act: How Do We Continue?

- CONGRATULATIONS for the pioneers : NORD, NIH and FDA
- Other regions have followed the USA in creating a system to help patients by fostering orphan drug development
- After 8 Years of EU Orphan Medicines Regulation: similar annual number of designations and approvals in EU and USA
- Stakeholders, including industry, are delivering on their promise: more therapies for patients with rare disorders!

Hope

- For patients and their care takers looking forward to the prospect of a treatment
- For children born with treatable rare diseases: the potential to live a normal life









Fear

- For payers afraid of the impact of orphan drug costs on health care budgets
 - High prices?
 - A Tsunami of orphan drugs coming?
 - Salami-slicing? Off-label use?
 - Misuse of the legislations?







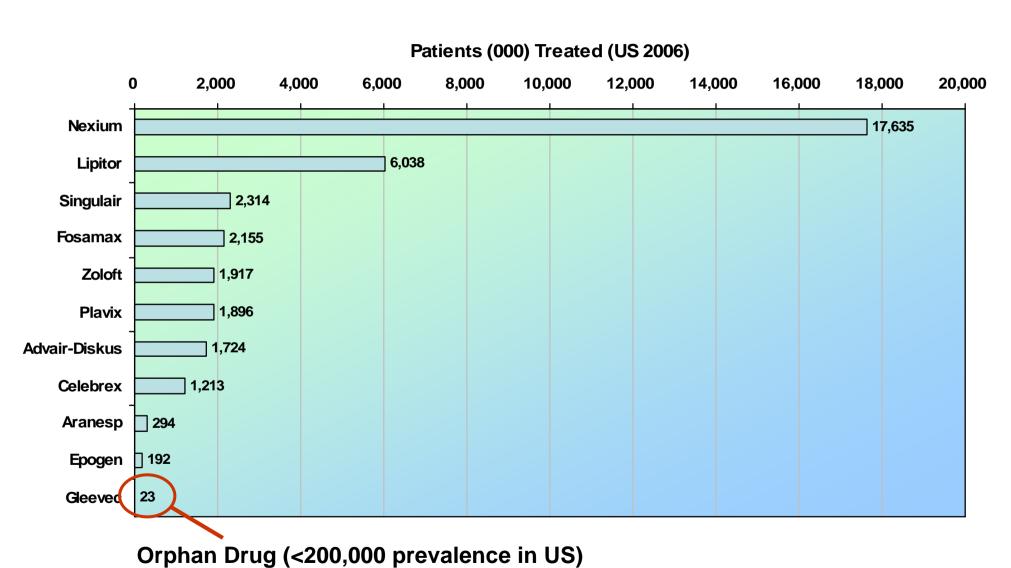




Department of Health and Ageing

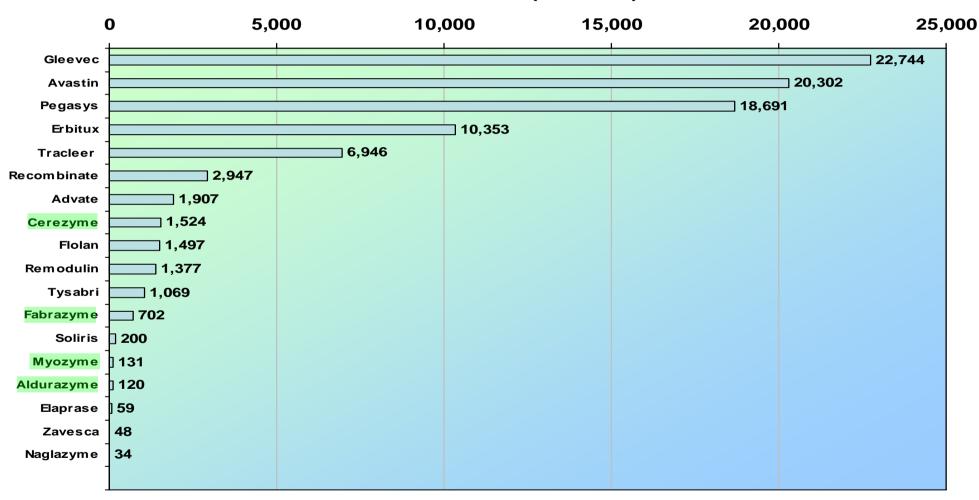


Understanding Rarity



Understanding Rarity

Patients Treated (US 2006)



Address misconceptions with clearer messages from us all

- Orphan medicines are treatments for lifethreatening or serious and chronic diseases
- Estimated prevalence is not the same as treatable number of patients
- Common, rare (orphan), ultra-rare (ultra-orphan) diseases: the reality is a continuum with research and treatment complexity increasing with rarity
- There is no avalanche, but rather a steady increase of 10-12 new products annually (EPPOSI workshop Copenhagen, 2007)

Industry makes substantial contributions

- Innovation for unmet medical needs
- A major source of healthcare products, including orphan drugs
- Expanded Access/compassionate use programs
- Wealth & employment creation

But needs to communicate better

Policy needs: in the US and EU: from R+R → A+A

The focus needs shifting
from "only"
Research and Regulatory issues
to
Awareness and Access

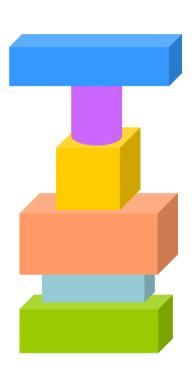
Policy needs in other parts of the world without OD system

Include Awareness and Access issues into the discussions on new legislation regarding orphan drugs?

A broader Healthcare System Building Access & Optimal Patient Care

- Diagnostic infrastructure
- Treatment guidelines
- Centres of excellence
- Registries
- Compassionate use
- Dedicated and "centralized" funding
- Patient network







THANK YOU