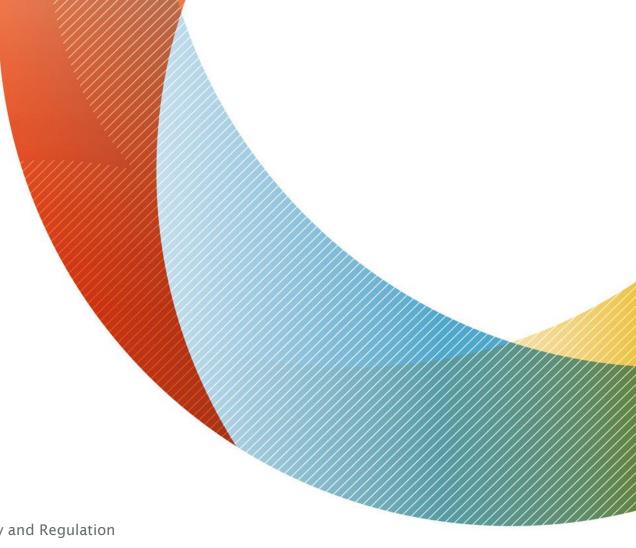
An analysis of price developments of biological medicines after patent expiration

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Gesundheit Österreich

Background | Product life cycle of a medicine

Research

- Basic research
- Applied research

Development

- Pre-Clinical research / Non-Clinical research
- Clinical Trials

Commercialisation

- Regulatory Approval
- Pricing
- Reimbursement

On-Patent-Phase

Off-Patent-Phase

• Dynamics during both phases are determined through different forces

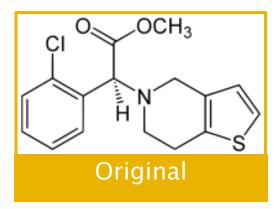
Background | Generics ≠ Biosimilar

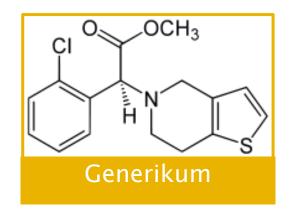
Generic

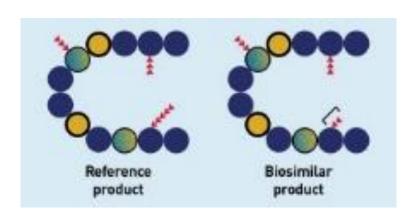
A generic medicine is a pharmaceutical product which has the **same** qualitative and quantititative composition in active [**chemical**] substances and the same pharmaceutical form as the original product

Biosimilar

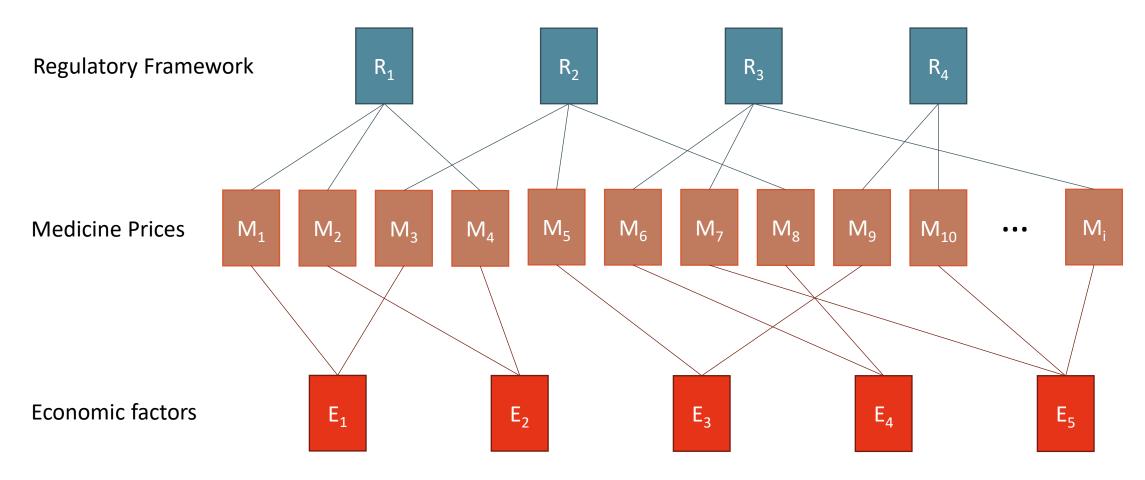
A **biological** medicine that is developed to be **similar** to an existing biological mecine (the ,reference medicine')

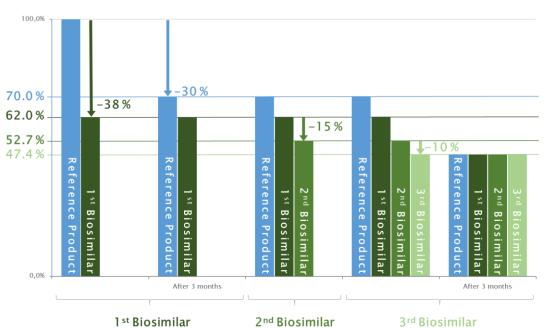






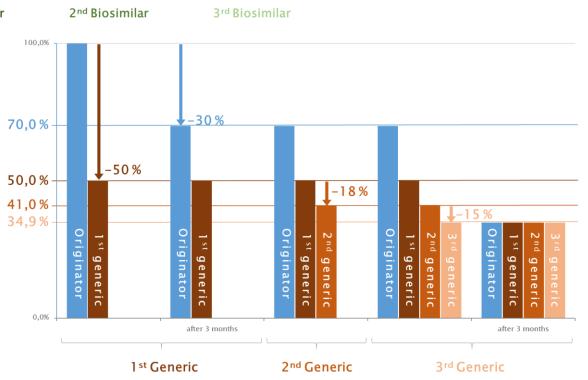
Methods | Two level nested Structure





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Results | Supply Side Measures

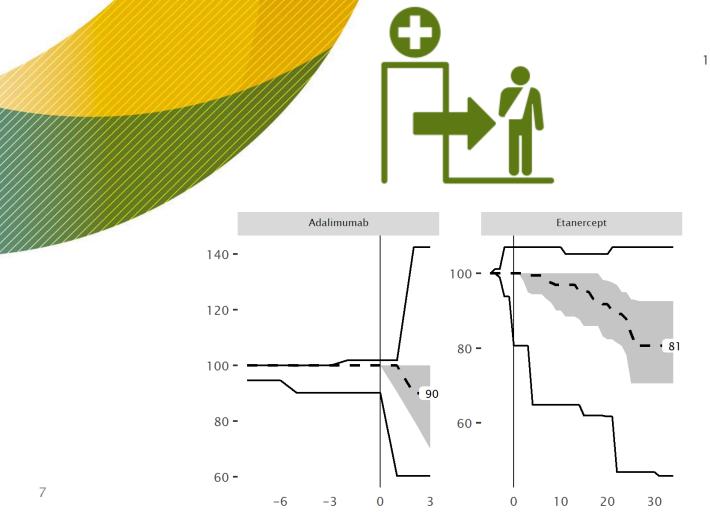


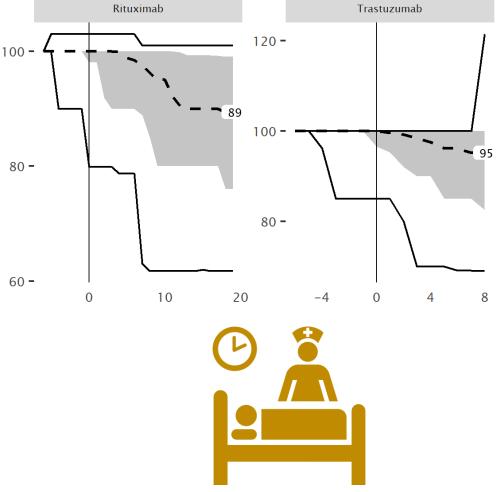


Results | Demand Side Measures



Results | Point of Administration





Results | Conjectured Factors

Pharmaceutical Expenditures are a major driving factor in pharmaceutical prices at launch, but are they also a driving factor for price developments?

Does the market size measured in population size (large, medium and small markets) have an impact on price developments

Conclusions

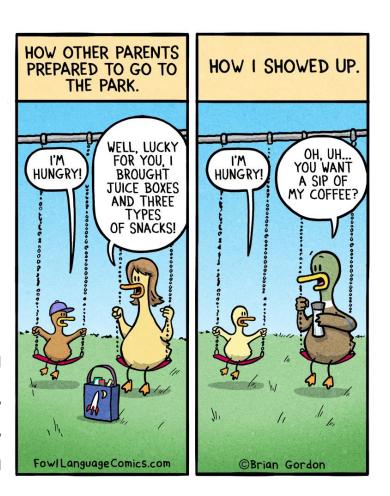


Lack of consistency

Available regulations for offpatent medicines are not consistently applied to biological medicines / biosimilars

Lack of competition

country-specific factors hamper price decreases due to a lack of competition



Thank you for your attention

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