

Colombia

Implementing a National Rare Disease Plan starting with National Registry

Germán Escobar Morales, MD. MPH .MSc



RARE DISEASE DAY®



@GERMANESCOBARM



german.escobar.morales@gmail.com



Ministry of Health and Social Protection

Health=fundamental right

Patients with RD=subjects of special protection

Public/Private Insurers

Package of Benefits



Hospitals/Clinics (Providers)

Universal coverage

Out of pocket exp. < 15%

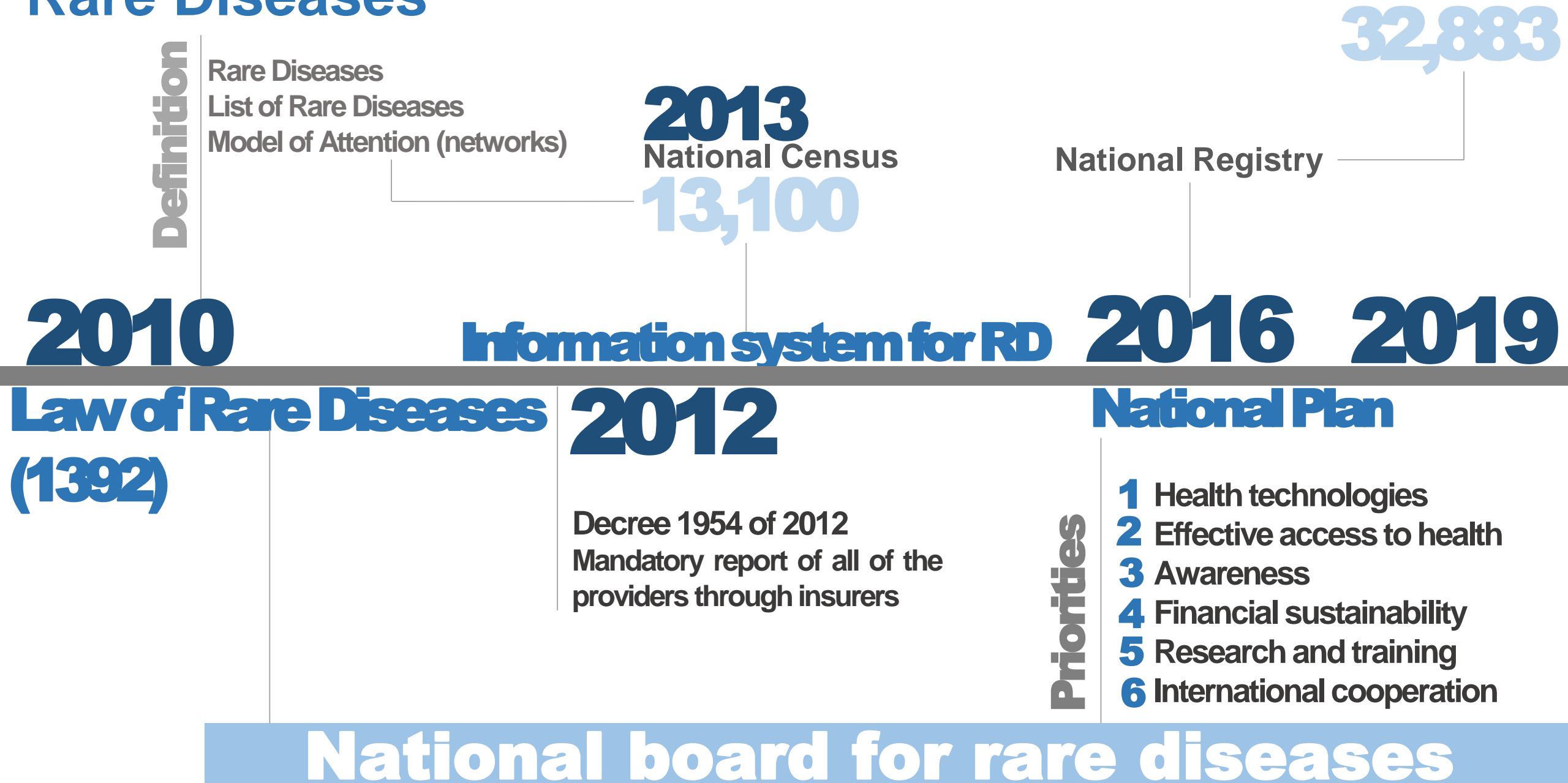


@GERMANESCOBARM



german.escobar.morales@gmail.com

Timeline of the Public Policy Rare Diseases



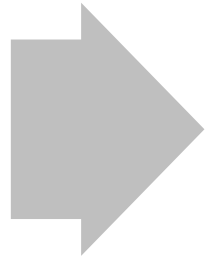
National Registry of Rare Diseases

Rare Diseases

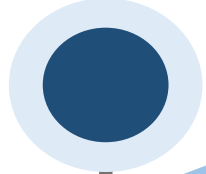
Minsalud

32,883

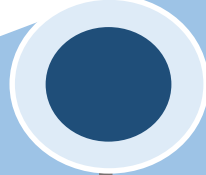
Provision records
Payment records



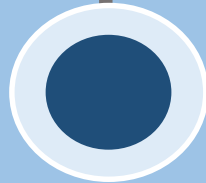
NHI



Insurers



Certified Providers



Public Health surveillance system

Mandatory

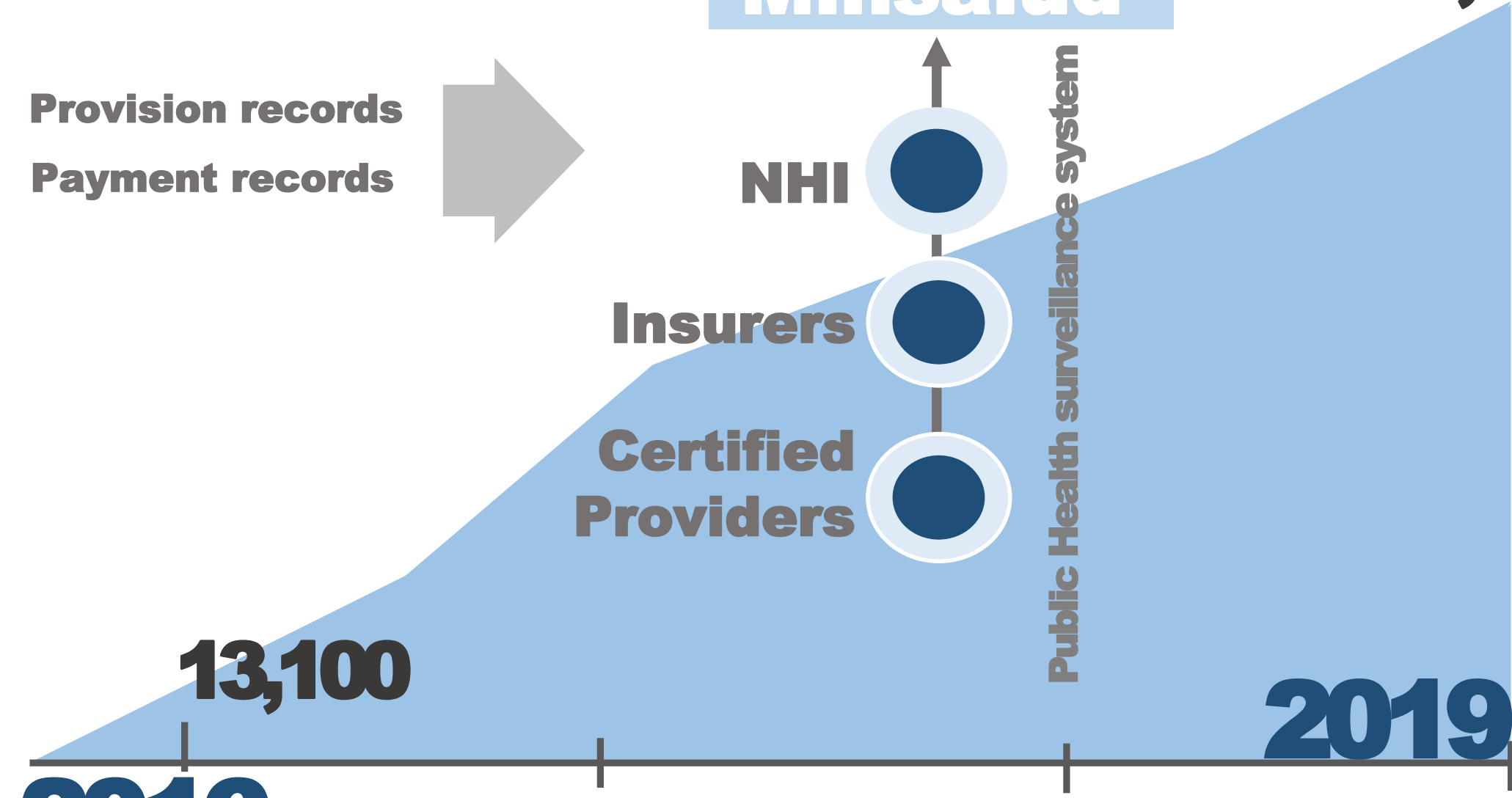
Necessary for payments of some specialized technologies

Instrument for patients

13,100

2019

2016



National Plan for Rare Diseases

Health technologies	Value-based pricing Special criteria for introducing HT into BP
Awareness	National Board for rare diseases Patients associations (FECOER/ACOPEL, etc)
Access/Financial S.	Reference centers New financial resources Health expenditure control
Research/int. coop	4.2 USD million for research 2016 WFH (Memorandum of Agreement)

Challenges

Assure financial resources
Improve effective access
Multi-sector policy



[@GERMANESCOBARM](https://twitter.com/GERMANESCOBARM)



german.escobar.morales@gmail.com

Thank You!

 [@GERMANESCOBARM](https://twitter.com/@GERMANESCOBARM)

 german.escobar.morales@gmail.com



RARE DISEASE DAY®