Teriparatide prefilled pens and cartridges:

multidose formulations as a source of medication errors

TO. RECRPV

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INTRODUCTION

Teriparatide: recombinant human parathyroid hormone analog (rhPTH (1-34)) approved for the treatment of osteoporosis in patients at high risk of fracture

the only injectable drug (subcutaneous route) administered once daily in this indication currently

 multidose formulations (28 doses of 20 μg)





Prefilled pens

 ${\tt FORSTEO}^{\circledast}, {\tt LIVOGIVA}^{\circledast}, {\tt SONDELBAY}^{\$}, {\tt TERIPARATIDE\,BIOGARAN}$

Cartridges for reusable pens MOVYMIA®, TERROSA®

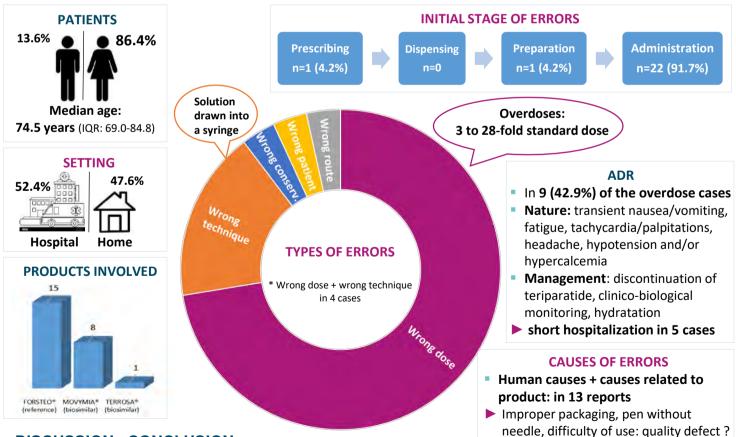
- Risk factors for medication errors (MEs)
 National safety alert 2021¹: dosing errors = injection of the entire solution (28 doses) as a single dose
- ► Objective: to characterize MEs involving teriparatide formulations reported in the French Pharmacovigilance Database (FPVD)

MATERIAL AND METHODS

Inclusion of all MEs (risk, intercepted and actual MEs with or without adverse drug reaction (ADR)) involving teriparatide (substance name) identified in the FPVD until December 2022.

RESULTS

▶ 24 MEs cases identified: 24 actual MEs, including 4 cases occurred after the ANSM safety alert of 2021



DISCUSSION - CONCLUSION

- MEs associated with multidose formulations of teriparatide mainly consist of overdoses without serious clinical consequences but may require hospitalizations for surveillance with costs to the health system.
- Risk minimization and prevention measures (patients' education...) still need to be strengthened.