# SUITABILITY OF ELASTOMERIC PUMPS FOR DRUG STORAGE

N Ott<sup>1</sup>, W Bello<sup>2</sup>, C Lanfranchi<sup>3</sup>, M Czernek<sup>1</sup>, G Kiefer<sup>1</sup>, B Thomas<sup>1</sup>, M Senn<sup>1</sup>, J Pezzatti<sup>2</sup>, U Lösch<sup>1</sup>

<sup>1</sup>Hospital Pharmacy, University Hospital Basel, <sup>2</sup>Pharmacy Department, Lausanne University Hospital, <sup>3</sup>Pharmacie interjurassienne, Hôpitaux du Jura et du Jura bernois

## **Background and objectives**

#### **Elastomeric pumps (EPs):**

- autonomous application system in outpatient settings (e.g. oncology, infectiology)
- continuous intravenous drug administration
- no electronic pumps needed
- stability data for > 130 active pharmaceutical ingredients for up to 60 days promote storage
- of data about leachables from various polymers and plastic additives

**Examination of storage of hydrophilic** solutions in elastomeric pumps over 180 days.

### Materials and methods

#### Examined pump devices (supplier / manufacturer)

ON-Q 400 mL (Avanos) Surefuser+ SFS-2-30P (Nipro) AutoSelector 550 mL (Oncomedical) Accufuser Selectus C0005M (WYM-Theramed) Folfusor LV system 1.5 mL/h and 5 mL/h, Intermate LV 250 mL (Baxter) PANT Easypump II LT 500-12.5-S (B.Braun) Dosifuser 500 D2 (PlusMedica)

Device were filled with ad hoc produced NaCl 0.9% (avoid leaching from plastic materials, simulate hydrophilic solutions).

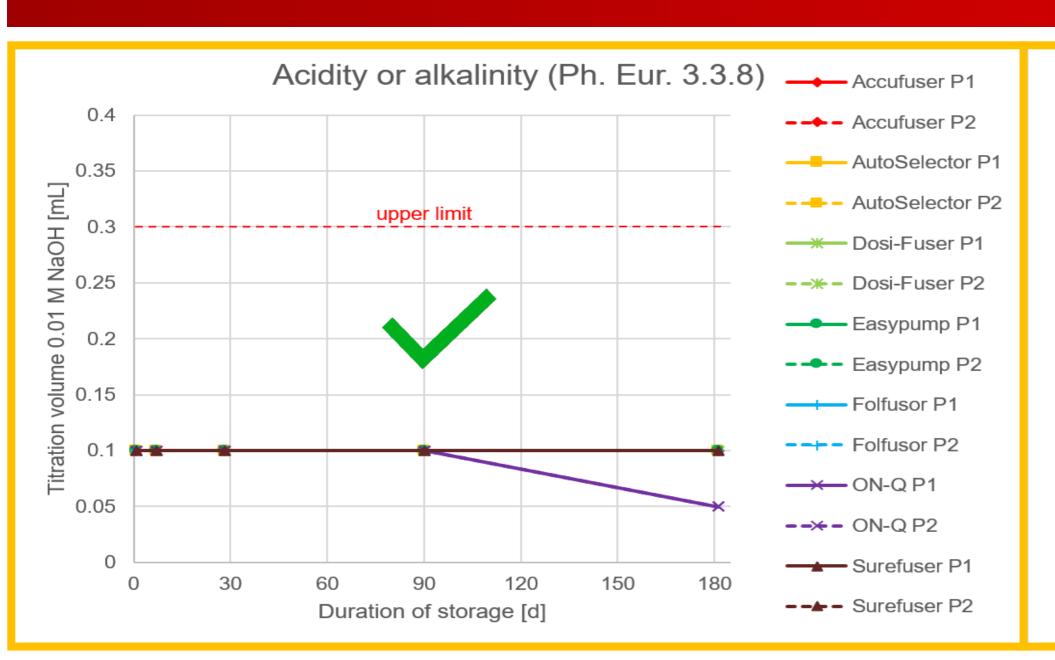
#### Measurements (days 1, 7, 28, 90 and 180):

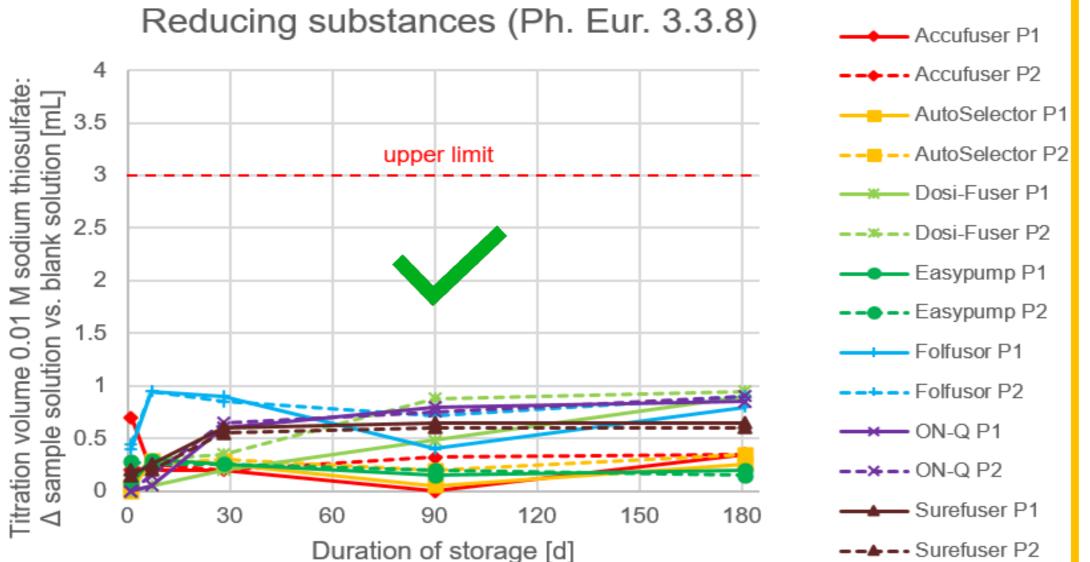
- 1. According to European Pharmacopoeia (Ph. Eur.) 3.3.8\* "Sterile single-use plastic syringes" [1,2]
  - Absorption
  - Acidity or alkalinity
  - Reducing Substances

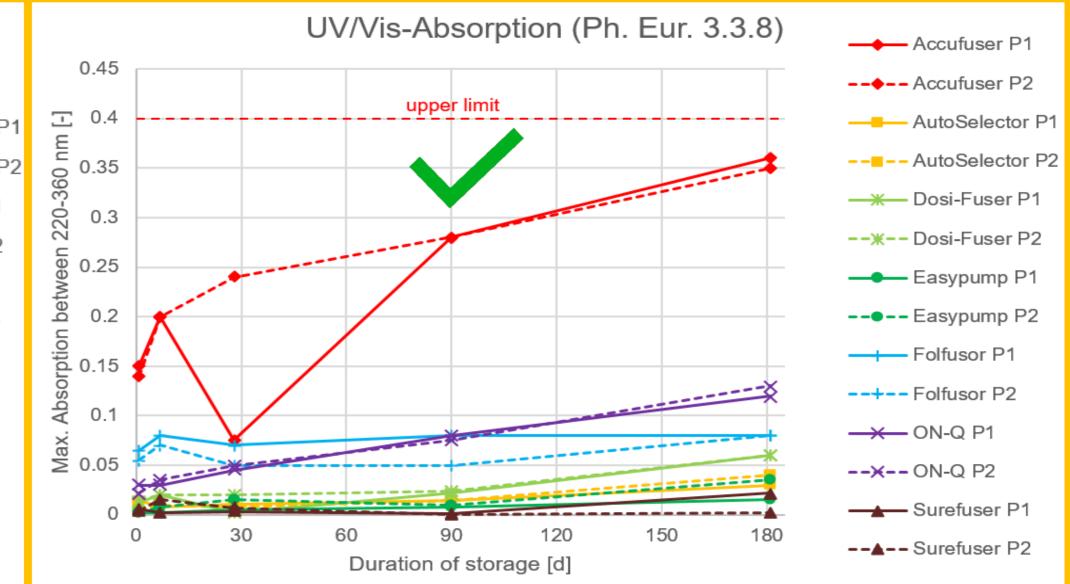
\*Ph. Eur. 3.2.2.1 "Plastic container for aqueous solutions for infusion" requires autoclaving – not possible with EPs [1,2].

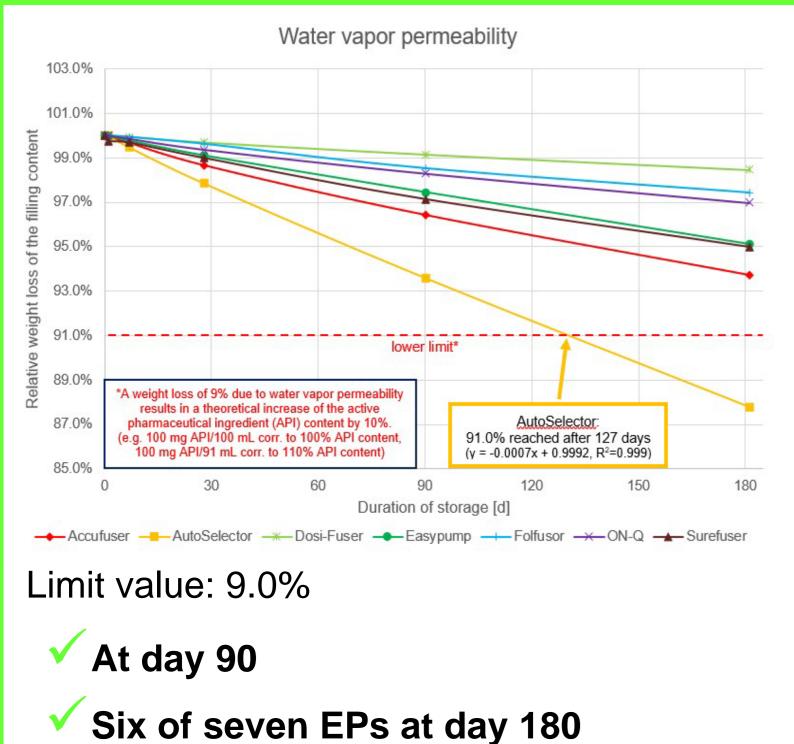
- 2. Water vapor permeability quantifying weight loss over time
- 3. HPLC-MS identifying leachables from plastic additives and recording semi-quantitatively [3]

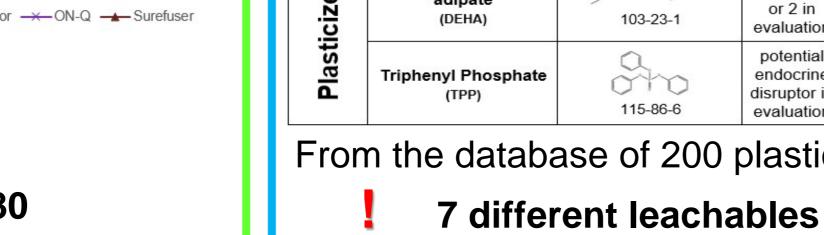
#### Results

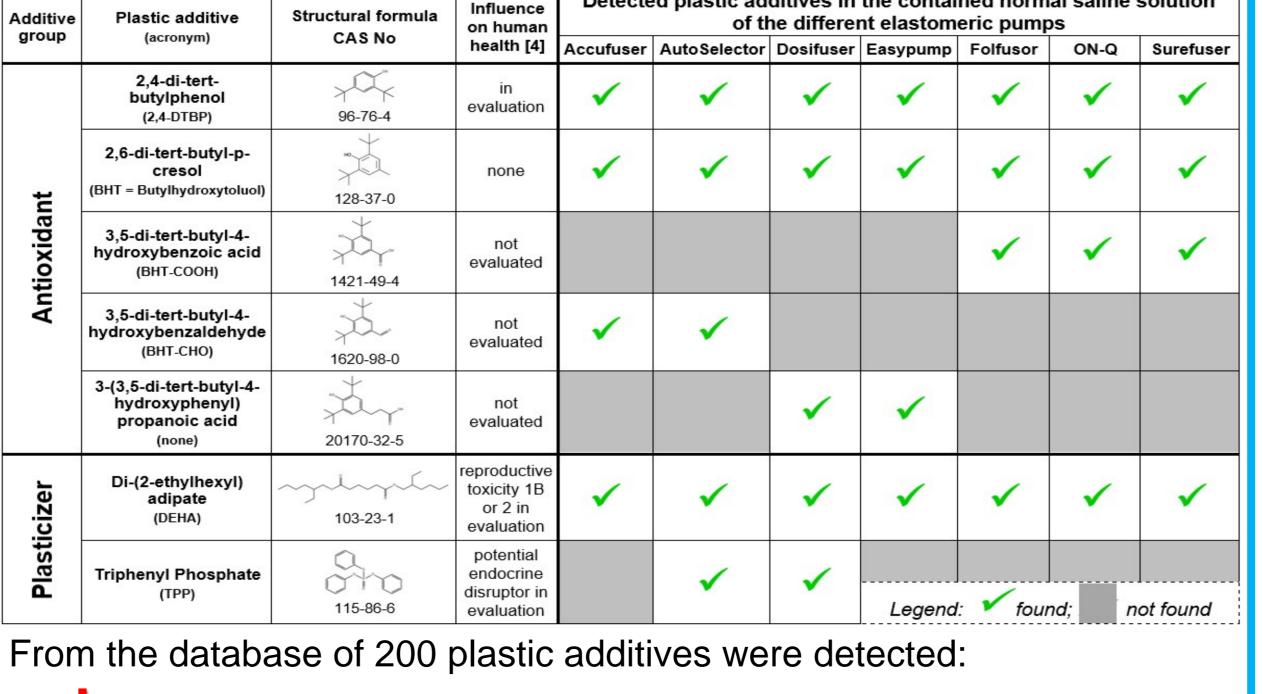




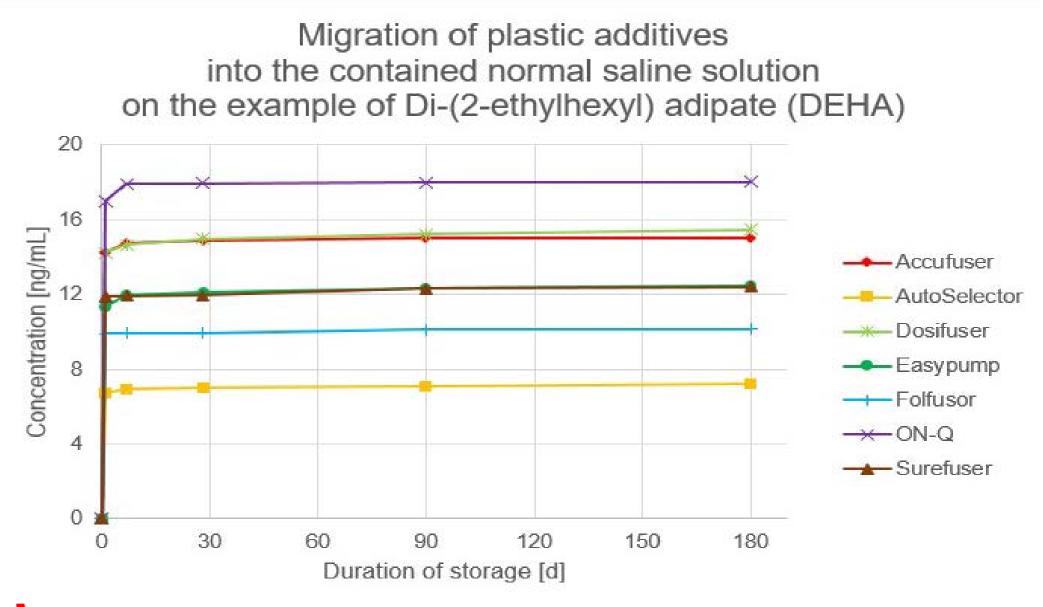








- 7 different leachables (5 antioxidants, 2 plasticizers)
- 2,4-DTBP, BHT and DEHA from each EP



- Example DEHA: > 90% migration within first 24 hours
- Overall migration in hydrophilic solutions:

most migration (43-97%, median: 80%) Day 1-180: minor migration

# Conclusions

No transfer of impurities in unacceptable quantities for the period of 180 days [1,2]

X AutoSelector out of limit at day 127

- **Continuous evaporation**
- X Limits the storage time (increasing concentration of ingredients)
- X Promotes precipitation of ingredients (solubility limit)
- Migration of antioxidants and plasticizers from every EP
- Not validated HPLC-MS method → only identification & semiquantitative detection (comparison of dimension (ng/mL)) with
- estimated NOAEL (No Observed Adverse Effect Level) and
- PDE (Permitted Daily Exposure) limits of the individual plastic additives [3]
- 2,4-DTBP, BHT derivatives, DEHA and TPP: incomplete data on the toxicology and long-term effects → unknown consequences of exposure [4].

Recommendation for use of the examined EPs:

Patient group	Duration of therapy	
	Days to weeks	Long-term
Adults		
ediatrics	! Prior: risk-benefit assessment*	X

Hydrophilic solutions can be stored for 127 days (AutoSelector) resp. 180 days (6 other EPs), if the removable volume of parenterals (Ph. Eur. 2.9.17) is observed.

**Presentation:** 

20 to 22 March 2024

#### Literature:

- [1] European Pharmacopoeia. In: European Pharmacopoeia Commission, editor. European Pharmacopoeia. 11.0. European Directorate for the Quality of Medicines and HealthCare; 2022.
- [2] Bracher, F., Heisig, P. & G. Schriba et al.: Kommentar zum Europäischen Arzneibuch - Band 2. In: Wissenschaftliche Verlagsgesellschaft mbH (Hrsg.), Allgemeiner Teil, Methoden 3-5, 72. Aktualisierungslieferung, Stuttgart, Deutschland: Govi-Verlag, 2023
- [3] Bello W, Pezzatti J, Berger-Gryllakia M, Rudaz S, Sadeghipour S. J. Pharm. Biomed. Anal. 236 (2023) 115640

action-plan/corap-table/ [cited: 04.09.2023]

[4] https://echa.europa.eu/information-on-chemicals/evaluation/community-rolling-

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#### **Corresponding author:**

Norman Ott Universitätsspital Basel Spital-Pharmazie Spitalstrasse 26, CH-4031 Basel norman.ott@usb.ch





