CE marking	Class I medical device in accordance with EU 2017/745 Personal protective equipment category III in accordance with Regulation (EU) 2016/425
CH REP	Swiss AR Services AG, Industriestrasse 47, 6300 Zug, Switzerland
EN ISO 21420:2020	Standard: Protective gloves - General requirements and test methods
EN 455-1:2022	Medical gloves for single use - Part 1: Requirements and testing for freedom from holes
EN 455-3:2015	Medical gloves for single use - Part 3: Requirements and testing for biological evaluation
EN 455-4:2009	Medical gloves for single use - Part 4: Requirements and testing for shelf life determination
EN ISO 374-1:2016 + A1:2018 TYPE B	Standard: Protective gloves against dangerous chemicals and micro-organisms - Part 1: Terminology and performance requirements for chemical risks.
EN ISO 374-2:2019	Protective gloves against dangerous chemicals and microorganisms - Part 2: Determination of resistance to penetration
EN ISO 374-4:2019	Protective gloves against dangerous chemicals and microorganisms - Part 4: Determination of resistance to degradation by
EN ISO 374-5:2016 VIRUS	Standard: Protective gloves against dangerous chemicals and micro-organisms - Part 5: Terminology and performance requirements for micro-organism risks
VO (EU) 2023/988	Regulation (EU) Nr. 2023/988 on general product safety.
VO (EG) 1935/2004	Regulation regarding plastic materials and articles intended to come into contact with food.
VO (EU) 10/2011	Regulation on plastic materials and articles intended to come into contact with food

P		2
T	Formaldehyde 37%	2