

# INFORMATION

Quality Management



EIP-1035-USA-AL

Revision 4

Date of issue: 04.09.2023

1/2

## Elemental Impurities

Ph. Eur. General Text 5.20; USP-NF General Chapter (232) and (233); ICH Guideline Q3D(R2)

MEGGLE USA Product:

Anhydrous Lactose: DuraLac® H

In the production process of the above mentioned MEGGLE USA Product, the elements classified in Class 1, 2A, 2B and 3 are not intentionally added in form of metal catalysts, metal reagents etc.

Permitted concentrations limits were calculated using the Permitted Daily Exposures and assuming a daily intake of the excipients of 10 g (ICH Q3D(R2), No 7 Option 1, stated in table A.2.2). Acceptance levels were defined as 30 % of the permitted concentrations.

Testing was conducted for the elements categorized as Class 1 and 2A relevant for oral route of administration according to the ICH Guideline Q3D(R2). Several representative lots of the product were tested using ICP-MS method in conformance to USP-NF (233). Testing method has been validated for the matrix of the products.

Results are mainly < LoQ and shown on the table below which is valid the mentioned product. All results are below 30 % of the acceptance levels for oral application. In consequence, additional controls are not required.

MEGGLE USA has implemented an ongoing monitoring program for elemental impurities in accordance to the regime of the initial study performed. The program is conducted at MEGGLE GmbH & Co. KG.

*This MEGGLE USA Information was electronically released and is valid without signature.*

# INFORMATION

Quality Management



EIP-1035-USA-AL | Revision 4 | Date of issue: 04.09.2023 | 2/2

## Elemental impurities – Summary Results

Ph. Eur. General Text 5.20;  
USP-NF General Chapter (232) and (233);  
ICH Guideline Q3D(R2)

Material Name: Anhydrous Lactose; trade name see above  
Production and Release Site: Agropur Inc., 719 North Main Street, Le Sueur, MN 56058-1404, USA  
Source/Type of Excipient: Lactose: Animal derived (Milk of bovine origin)  
Route of administration (RoA): Oral

Class	Elements	Elements to be considered		Oral PDE µg/day	Perm. Conc. µg/g	Accept. Level µg/g	Results* µg/g	Method	Comments	
		Added	Based on RoA							
1	Cadmium	Cd	No	Yes	5	0.5	0.15	< 0.0009	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
1	Lead	Pb	No	Yes	5	0.5	0.15	< 0.0006	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
1	Arsenic (inorg.)	As	No	Yes	15	1.5	0.45	< 0.0081	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
1	Mercury (inorg.)	Hg	No	Yes	30	3	0.9	< 0.0005	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
2A	Cobalt	Co	No	Yes	50	5	1.5	0.0003	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
2A	Vanadium	V	No	Yes	100	10	3	< 0.0093	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
2A	Nickel	Ni	No	Yes	200	20	6	< 0.0084	ICP-MS; USP-NF (233)	7 batches tested. Monitoring installed (1 / year)
2B	Thallium	Tl	No	No	n/a					
2B	Gold	Au	No	No	n/a					
2B	Palladium	Pd	No	No	n/a					
2B	Iridium	Ir	No	No	n/a					
2B	Osmium	Os	No	No	n/a					
2B	Rhodium	Rh	No	No	n/a					
2B	Ruthenium	Ru	No	No	n/a					
2B	Selenium	Se	No	No	n/a					
2B	Silver	Ag	No	No	n/a					
2B	Platinum	Pt	No	No	n/a					
3	Lithium	Li	No	No	n/a					
3	Antimony	Sb	No	No	n/a					
3	Barium	Ba	No	No	n/a					
3	Molybdenum	Mo	No	No	n/a					
3	Copper	Cu	No	No	n/a					
3	Tin	Sn	No	No	n/a					
3	Chromium	Cr	No	No	n/a					

\* "< X": < LoQ (Limit of Quantification)

This MEGGLE USA Information was electronically released and is valid without signature.