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VOC EMISSION TEST REPORT

Indoor Air Comfort GOLD

8 July 2026

1 Sample Information

Sample name	Crilux
Batch no.	26PR001434
Stated production date	31/03/2026
Product type	Primer, water based
Sample reception	09/04/2026

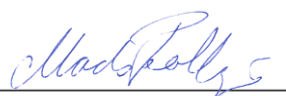
2 Brief Evaluation of the Results

Regulation or protocol	Conclusion	Version of regulation or protocol
Indoor Air Comfort GOLD	Pass	Indoor Air Comfort GOLD v2026 of January 2026
Indoor Air Comfort	Pass	Indoor Air Comfort v2026 of January 2026
ABG/AgBB	Pass	Ausschuss zur gesundheitlichen Bewertung von Bauprodukten (September 2024)
Belgian Regulation	Pass	Royal decree of May 2014 (C-2014/24239)
EU Taxonomy	Pass	Regulation (EU) 2020/852 of the European Parliament and of the Council
EU Ecolabel	Pass	Commission Decision 2025/2607 of 17 December 2025
Blue Angel (DE-UZ 12a)	Pass	January 2019 - Version 12
Italian CAM Edilizia	Premium Score	Decreto Ministeriale 24 novembre 2025 – Criteri Ambientali Minimi per l'edilizia
French VOC Regulation		Decree of March 2011 (DEVL1101903D) and Arrêté of April 2011 (DEVL1104875A) modified in February 2012 (DEVL1133129A)
French CMR components	Pass	Regulation of April and May 2009 (DEVP0908633A and DEVP0910046A)

Full details based on the testing and direct comparison with limit values are available in the following pages
Regarding pass/fail decision rule please see appendix



Rasmus Verdier
Analytical Service Manager



Mads Folkjær
Analytical Chemist

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3 Applied Test Methods

3.1 General Test References

Regulation, protocol or standard	Version	Reporting limit VOC [µg/m³]	Calculation of TVOC	Combined uncertainty ^a [RSD(%)]
EN 16516	2017 + A1:2020	5	Toluene equivalents	22%
ISO 16000 -3 -6 -9 -11	2021-2024 depending on part	2	Toluene equivalents	22%
EN 16402	2019	2	Toluene equivalents	22%
ASTM D5116	2025	-	-	-
Specifications Indoor Air Comfort Gold	v2026 of January 2026	5	Toluene equivalents	22%
AgBB (MVV TB/ABG)	September 2024 (2024/2025)	5	Compound Specific	22%
Belgian VOC	Royal decree of May 2014 (C - 2014 / 24239)	5	Toluene equivalents	22%
EU Taxonomy	Regulation (EU) 2020/852; Annex 1/2 section 7.1 and 7.2	-	-	22%
EU Ecolabel	Commission Decision 2025/2607 of 17 December 2025	5	Toluene equivalents	22%
Blue Angel (DE-UZ 12a)	"Low-emission and low-pollutant paints and varnishes", January 2019 - Version 12	5	Compound Specific	22%
BREEAM NOR	BREEAM NOR v6.1.1 New Construction (2024)	5	Toluene equivalents	22%
BREEAM International	BREEAM International New Construction v7.0 (2025)	5	Toluene equivalents	22%
LEED v4.1 BETA (outside U.S.)	November 2025	5	Compound Specific	22%
WELL V2	Q2 2025	5	Compound Specific	22%
Italian CAM Edilizia	Decreto Ministeriale 24 novembre 2025 – Criteri Ambientali Minimi per l'edilizia	2	Toluene equivalents	22%
French VOC Classes	Decree of 03/2011 (DEVL1101903D) and arrêté of 02/2012 (DEVL1133129A)	2	Toluene equivalents	22%
French CMR	Regulation of April and May 2009 (DEVP0908633A and DEVP0910046A)	1	Toluene equivalents	22%

3.2 Specific Laboratory Sampling and Analyses

Procedure	External Method	Internal SOP	Quantification limit / sampling volume	Analytical principle	Uncertainty ^a [RSD(%)]
Sample preparation	ISO 16000-11:2024, EN 16516:2017+A1:2020, AgBB:2024, EMICODE:2025	71M549810	-	-	-
Emission chamber testing	ISO 16000-9:2024, EN 16516:2017+A1:2020	71M549811	-	Chamber and air control	-
Sampling of VOC	ISO 16000-6:2021, EN 16516:2017+A1:2020	71M549812	5 L	Tenax TA	-
Analysis of VOC	ISO 16000-6:2021, EN 16516:2017+A1:2020	71M542808B	1 µg/m³	ATD-GC/MS	10%
Sampling of aldehydes	ISO 16000-3:2022, EN 16516:2017+A1:2020	71M549812	35 L	DNPH	-
Analysis of aldehydes	ISO 16000-3:2022, EN 16516:2017+A1:2020	71M548400	3-6 µg/m³	HPLC-UV	10%
Sampling of phthalates*	ISO 16000-33:2017, MEL-09:2003	71M549812	60 L	Florisil	-
Analysis of phthalates*	ISO 16000-33:2017	71M546060	0.6 µg/m³	GC/MS	10%

The analysis are carried out on the sample(s) as received and the result(s) are only valid for the tested sample(s).

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4 Test Parameters, Sample Preparation and Deviations

4.1 VOC Emission Chamber Test Parameters

Parameters	Value	Sample Conditions	Value
Chamber volume, V[L]	119	Date and time of unpacking and start of sample preparation	25/05/2026 - 10:37
Air change rate, n[h ⁻¹]	0.5	Preconditioning period	-
Air Velocity [m/s]	0.1	Chamber test period	25/05/2026 - 22/06/2026
Area specific ventilation rate, q [m/h or m ³ /m ² /h]	0.5	Analytical test period	25/05/2026 - 01/07/2026
Relative humidity of supply air, RH [%]	50 ± 3	Exposed sample area [m ²]	0.120
Temperature of supply air, T [°C]	23 ± 1	Loading factor [m ² /m ³]	1.0
Background concentration of individual VOCs [µg/m ³]	< 2	Test scenario	Wall
Background concentration of TVOC [µg/m ³]	< 20		

4.2 Preparation of the Test Specimen

The sample was homogenised and applied onto a glass plate.

Number of Layers	Application amount per layer, g/m ²	Drying time, h
1	151	-

4.3 Picture of Sample



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4.4 Deviations from Referenced Protocols and Regulations

No deviations from the referenced test methods were observed.

4.5 Air Samplings from the Test Chamber

Sampling media	Day (yyyy-mm-dd)	Time (hh:mm)	Volume [L]
3 Day, DNPH silicagel	2026-05-28	09:54 - 11:43	35
3 Day-Res, DNPH silicagel	2026-05-28	09:54 - 11:44	33
3 Day, Tenax TA	2026-05-28	09:55 - 10:54	5.3
3 Day-Res, Tenax TA	2026-05-28	10:54 - 11:44	2.4
3 Day, Tenax TA	2026-05-28	07:17 - 07:27	0.47
28 Day, Florisil	2026-06-22	07:21 - 09:51	60
28 Day-Res, Florisil	2026-06-22	07:22 - 09:52	61
28 Day, DNPH silicagel	2026-06-22	11:43 - 13:33	36
28 Day-Res, DNPH silicagel	2026-06-22	11:43 - 13:33	36
28 Day, Tenax TA	2026-06-22	11:44 - 12:43	5.1
28 Day-Res, Tenax TA	2026-06-22	12:43 - 13:33	2.3

5 Results

5.1 VOC Emission Test Results after 3 Days

	CAS No.	Retention time [min]	ID-Cat	Specific Conc. [µg/m³]	Toluene eq. [µg/m³]	Specific SER [µg/(m²·h)]	R _D	R _B
VOC with NIK/LCI								
1,2-Propandiol (Propylene glycol)	57-55-6	3.53	1	110	22	55	0.052	0.052
2-methyl-4-isothiazolin-3-on	2682-20-4	10.86	1	15	< 5	7.5	0.15	0.15
5-Chlor-2-methyl-4-isothiazolin-3-on (CIT)	26172-55-4	11.43	1	11	< 5	5.5	11	11
VOC without NIK/LCI								
None determined								
Sum of VOC without NIK/LCI				< 5	< 5	< 3		
VVOC compounds								
None determined								
TVOC				< 5	< 5	< 3		
SVOC compounds								
None determined								
TSVOC				< 5	< 5	< 3		
Carcinogens								
Total carcinogens				< 1	< 1	< 1		
Aldehydes								
Formaldehyde	50-00-0		1	< 3		< 2		
Acetaldehyde	75-07-0		1	< 3		< 2		
Propionaldehyde	123-38-6		1	< 3		< 2		
Butyraldehyde	123-72-8		1	< 3		< 2		
Acrolein *	107-02-8		1	< 5		< 3		
2-Butenal *	123-73-9		1	< 5		< 3		
Glutaraldehyde *	111-30-8		1	< 3		< 2		
R-values							11	11
TVOC				140	22	68		

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5.2 VOC Emission Test Results after 28 Days

	CAS No.	Retention time [min]	ID-Cat	Specific Conc. [µg/m³]	Toluene eq. [µg/m³]	Specific SER [µg/(m²·h)]	R _D	R _B
VOC with NIK/LCI								
None determined								
VOC without NIK/LCI								
None determined								
Sum of VOC without NIK/LCI				< 5	< 5	< 3		
VVOC compounds								
None determined								
TVOC				< 5	< 5	< 3		
SVOC compounds								
None determined								
TSVOC				< 5	< 5	< 3		
Carcinogens								
Total carcinogens				< 1	< 1	< 1		
CMR (French reg.)								
Benzene	71-43-2		1	< 1		< 1		
Trichloroethylene	79-01-6		1	< 1		< 1		
Dibutylphthalate (DBP)*	84-74-2		1	< 1		< 1		
Diethylhexylphthalate (DEHP)*	117-81-7		1	< 1		< 1		
Aldehydes								
Formaldehyde	50-00-0		1	< 3		< 2		
Acetaldehyde	75-07-0		1	< 3		< 2		
Propionaldehyde	123-38-6		1	< 3		< 2		
Butyraldehyde	123-72-8		1	< 3		< 2		
Acrolein *	107-02-8		1	< 5		< 3		
2-Butenal *	123-73-9		1	< 5		< 3		
Glutaraldehyde *	111-30-8		1	< 3		< 2		
R-values							0	0
TVOC				< 5	< 5	< 3		
TVOC (French label)					< 2			
Toluene	108-88-3			< 2	< 2	< 1		
Tetrachloroethylene *	127-18-4			< 2	< 2	< 1		

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	CAS No.	Retention time [min]	ID- Cat	Specific Conc. [µg/m³]	Toluene eq. [µg/m³]	Specific SER [µg/(m²·h)]	R _D	R _B
Ethylbenzene	100-41-4			< 2	< 2	< 1		
Xylene *	1330-20-7			< 2	< 2	< 1		
Styrene	100-42-5			< 2	< 2	< 1		
2-Butoxyethanol	111-76-2			< 2	< 2	< 1		
1,2,4-Trimethylbenzene	95-63-6			< 2	< 2	< 1		
1,4-Dichlorobenzene	106-46-7			< 2	< 2	< 1		

6 Summary and Evaluation of the Results

6.1 Comparison with Limit Values of Indoor Air Comfort Gold

	Test after 3 days		Test after 28 days	
	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$
TVOC (EN 16516)	22	≤ 1000	< 5	≤ 100
TVOC (AgBB)	140	≤ 3000	< 5	≤ 300
TSVOC	< 5	-	< 5	≤ 50
R _D -value (NIK) (dimensionless)	11	-	0	≤ 1.0
R _B -value (LCI) (dimensionless)	11	-	0	≤ 1.0
Sum of VOC without NIK/LCI	< 5	-	< 5	≤ 40
Total carcinogens	< 1	< 10	-	-
Any individual carcinogens	-	-	< 1	< 1
CMR (French reg.)	-	-	< 1	< 1
Formaldehyde	< 3	≤ 50	< 3	< 10
Acetaldehyde	< 3	≤ 50	< 3	≤ 50
French A+ and Italian CAM Premium Score Criteria	-	-	Complies	

6.2 Comparison with Limit Values of Indoor Air Comfort

	Test after 3 days		Test after 28 days	
	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$
TVOC (EN 16516)	22	≤ 10000	< 5	≤ 1000
TVOC (AgBB)	140	≤ 10000	< 5	≤ 1000
TSVOC	< 5	-	< 5	≤ 100
R _D -value (NIK) (dimensionless)	11	-	0	≤ 1
R _B -value (LCI) (dimensionless)	11	-	0	≤ 1
Sum of VOC without NIK/LCI	< 5	-	< 5	≤ 100
Total carcinogens	< 1	≤ 10	-	-
Any individual carcinogens	-	-	< 1	< 1
CMR (French reg.)	-	-	< 1	< 1
Formaldehyde	< 3	-	< 3	< 60
Acetaldehyde	< 3	-	< 3	< 200
French A+/A	-	-	Complies	

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6.3 Comparison with Limit Values of AgBB/ABG

Parameters	Test after 3 days		Test after 28 days	
	Concentration mg/m ³	Limit Value mg/m ³	Concentration mg/m ³	Limit Value mg/m ³
TVOC	0.14	≤ 10	< 0.005	≤ 1.0
TSVOC	< 0.005	-	< 0.005	≤ 0.1
R-value (dimensionless)	11	-	0	≤ 1
Sum of VOC without NIK/LCI	< 0.005	-	< 0.005	≤ 0.1
Formaldehyde	-	-	< 0.003	≤ 0.1
Any individual carcinogens	< 0.001	≤ 0.01	< 0.001	≤ 0.001

Compliance with the limits alone does not replace an approval or voluntary documentation by a Technical Assessment Body according to the Construction Product Regulation. This requires an application and approval.

6.4 Comparison with Limit Values of the Belgian Regulation

Parameters	Test after 28 days	
	Concentration µg/m ³	Limit Value µg/m ³
TVOC (EN 16516)	< 5	≤ 1000
TSVOC	< 5	≤ 100
R-value (dimensionless)	0	≤ 1
Total carcinogens	< 1	≤ 1
Toluene	< 5	≤ 300
Formaldehyde	< 3	≤ 100
Acetaldehyde	< 3	≤ 200

6.5 Comparison with Limit Values of EU Taxonomy

Parameters	Test after 28 days	
	Concentration mg/m ³	Limit Value mg/m ³
Formaldehyde	< 0.003	≤ 0.06
Any individual carcinogens	< 0.001	< 0.001

The formaldehyde emission is tested according to EN 16516 with air change rate 0.5 /h and relative humidity of 50 ± 3 %.

6.6 Comparison with Limit Values of EU Ecolabel

Parameters	Test after 3 days		Test after 28 days	
	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$	Concentration $\mu\text{g}/\text{m}^3$	Limit Value $\mu\text{g}/\text{m}^3$
TVOC	22	≤ 3000	< 5	≤ 300
R-value (dimensionless)	11	-	0	≤ 1.0
Formaldehyde	-	-	< 3	≤ 10
Any individual carcinogens	< 1	≤ 10	< 1	≤ 1

6.7 Comparison with Limit Values of Blue Angel (DE-UZ 12a)

	Test after 3 days		Test after 28 days	
	Concentration mg/m^3	Limit Value mg/m^3	Concentration mg/m^3	Limit Value mg/m^3
TVOC	0.14	< 3	< 0.005	< 0.3
TSVOC	< 0.005	-	< 0.005	< 0.1
R-value (dimensionless)	11	-	0	≤ 1
TVOC without LCI	< 0.005	-	< 0.005	< 0.04
Total carcinogens	< 0.001	< 0.01	-	-
Any individual carcinogens	-	-	< 0.001	< 0.001
Formaldehyde	-	-	< 0.003	< 0.02

6.8 Comparison with Limit Values of BREEAM International

Parameters	Concentration mg/m ³	Basic Level mg/m ³	Exemplary Level mg/m ³
Formaldehyde 28 days	< 0.003	≤ 0.06	≤ 0.01
TVOC (EN 16516) 28 days	< 0.005	≤ 1.0	≤ 0.3
TSVOC 28 days	< 0.005	-	≤ 0.1
Total carcinogens 28 days	< 0.001	≤ 0.001	≤ 0.001

This evaluation only concerns the emission requirements of BREEAM International. In order to satisfy the credit on "Health and wellbeing", according to the requirements of BREEAM International New Construction 2025, the product must also show compliance with the VOC content requirements.

6.9 Comparison with Limit Values of BREEAM NOR

Parameters	Concentration mg/m ³	Basic Level mg/m ³	Exemplary Level mg/m ³
Formaldehyde 28 days	< 0.003	≤ 0.06	≤ 0.01
TVOC (EN 16516) 28 days	< 0.005	≤ 0.3	≤ 0.3
TSVOC 28 days	< 0.005	-	≤ 0.1
Total carcinogens 28 days	< 0.001	≤ 0.001	≤ 0.001

This evaluation only concerns the emission requirements of BREEAM NOR. In order to satisfy the credit on "Health and wellbeing", according to the requirements of BREEAM NOR v6.1.1 New Construction, the product must also show compliance with the VOC content requirements.

6.10 Comparison with Limit Values of LEED v4.1 BETA and WELL V2

Parameters	Test after 28 days	
	Concentration µg/m ³	Limit Value µg/m ³
TVOC	< 5	≤ 1000
Sum of VOC without NIK/LCI	< 5	< 100
Formaldehyde	< 3	≤ 10
R-value (dimensionless)	0	≤ 1
AgBB Compliancy	Pass	

This evaluation only concerns the emission requirements of LEED v4.1 BETA. In order to satisfy the credit on "Low-Emitting Material" according to the requirements of LEED v4.1 BETA (November 2025), the product must also show compliance with the VOC content requirements.

6.11 Comparison with Limit Values of the Italian CAM Regulation

	CAS No.	Conc. 28 days $\mu\text{g}/\text{m}^3$	Limit value 28 days $\mu\text{g}/\text{m}^3$	Limit Value Premium Score 28 days $\mu\text{g}/\text{m}^3$
TVOC	-	< 2	≤ 1000	≤ 500
Formaldehyde	50-00-0	< 3	< 60	< 10
Acetaldehyde	75-07-0	< 3	< 200	< 100
Toluene	108-88-3	< 2	< 300	< 150
Tetrachloroethylene *	127-18-4	< 2	< 250	< 120
Ethylbenzene	100-41-4	< 2	< 750	< 350
Xylene *	1330-20-7	< 2	< 300	< 100
Styrene	100-42-5	< 2	< 250	< 120
2-Butoxyethanol	111-76-2	< 2	< 1000	< 500
1,2,4-Trimethylbenzene	95-63-6	< 2	< 1000	< 500
1,4-Dichlorobenzene	106-46-7	< 2	< 60	< 30
Benzene	71-43-2	< 1	< 1	
Trichloroethylene	79-01-6	< 1	< 1	
Dibutylphthalate (DBP)*	84-74-2	< 1	< 1	
Diethylhexylphthalate (DEHP)*	117-81-7	< 1	< 1	

The product was assigned a VOC emission class without taking into account the measurement uncertainty associated with the result. Awarding of the 'premium score' is at the sole discretion of the Contracting Authority (Art. 57(2) of the Code). Eurofins provides CAM compliant test results and does not assign points.

6.12 Comparison with Limit Values of the French VOC Regulation

	CAS No.	Conc. 28 days $\mu\text{g}/\text{m}^3$	 $\mu\text{g}/\text{m}^3$	 $\mu\text{g}/\text{m}^3$	 $\mu\text{g}/\text{m}^3$	 $\mu\text{g}/\text{m}^3$
TVOC	-	< 2	> 2000	< 2000	< 1500	< 1000
Formaldehyde	50-00-0	< 3	> 120	< 120	< 60	< 10
Acetaldehyde	75-07-0	< 3	> 400	< 400	< 300	< 200
Toluene	108-88-3	< 2	> 600	< 600	< 450	< 300
Tetrachloroethylene *	127-18-4	< 2	> 500	< 500	< 350	< 250
Ethylbenzene	100-41-4	< 2	> 1500	< 1500	< 1000	< 750
Xylene *	1330-20-7	< 2	> 400	< 400	< 300	< 200
Styrene	100-42-5	< 2	> 500	< 500	< 350	< 250
2-Butoxyethanol	111-76-2	< 2	> 2000	< 2000	< 1500	< 1000
1,2,4-Trimethylbenzene	95-63-6	< 2	> 2000	< 2000	< 1500	< 1000
1,4-Dichlorobenzene	106-46-7	< 2	> 120	< 120	< 90	< 60

The product was assigned a VOC emission class without taking into account the measurement uncertainty associated with the result. As specified in French Decree no. 2011-321 of March 23 2011, correct assignment of the VOC emission class is the sole responsibility of the party responsible for distribution of the product in the French market.

6.13 Comparison with Limit Values of the CMR Components

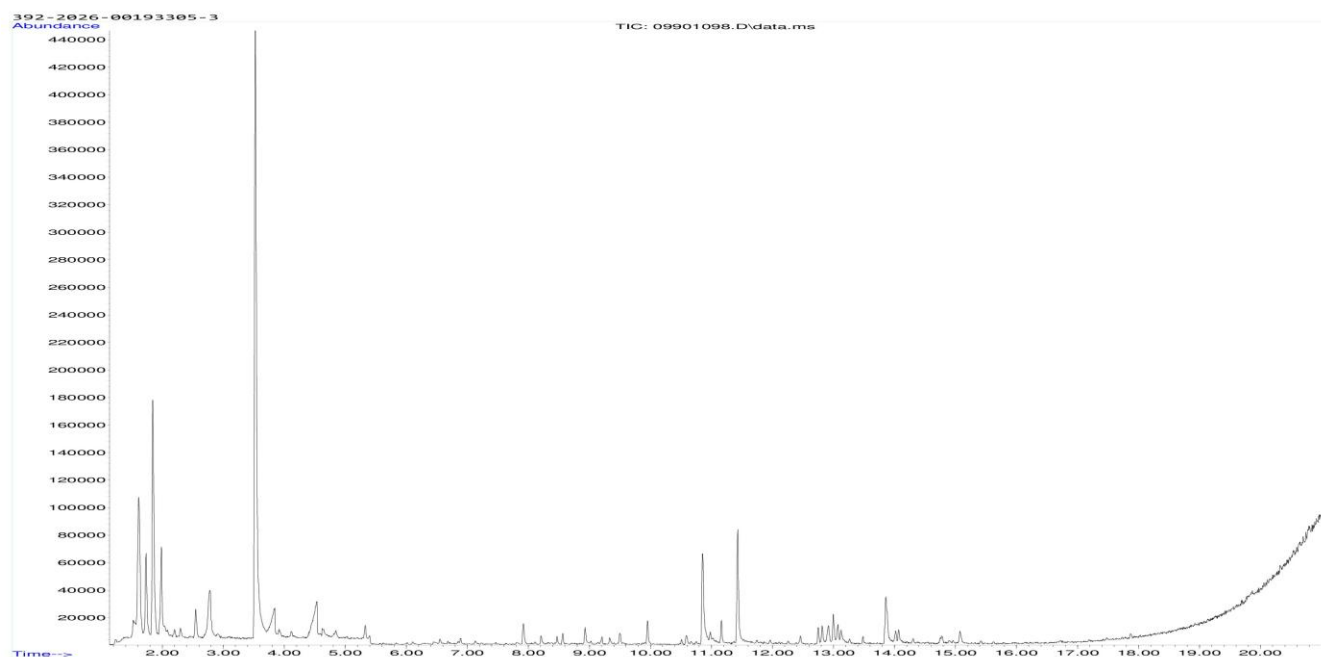
CMR (French reg.)	CAS No.	Conc. 28 days $\mu\text{g}/\text{m}^3$	Max. allowed air concentration $\mu\text{g}/\text{m}^3$
Benzene	71-43-2	< 1	< 1
Trichloroethylene	79-01-6	< 1	< 1
Dibutylphthalate (DBP)*	84-74-2	< 1	< 1
Diethylhexylphthalate (DEHP)*	117-81-7	< 1	< 1

The analysis are carried out on the sample(s) as received and the result(s) are only valid for the tested sample(s).

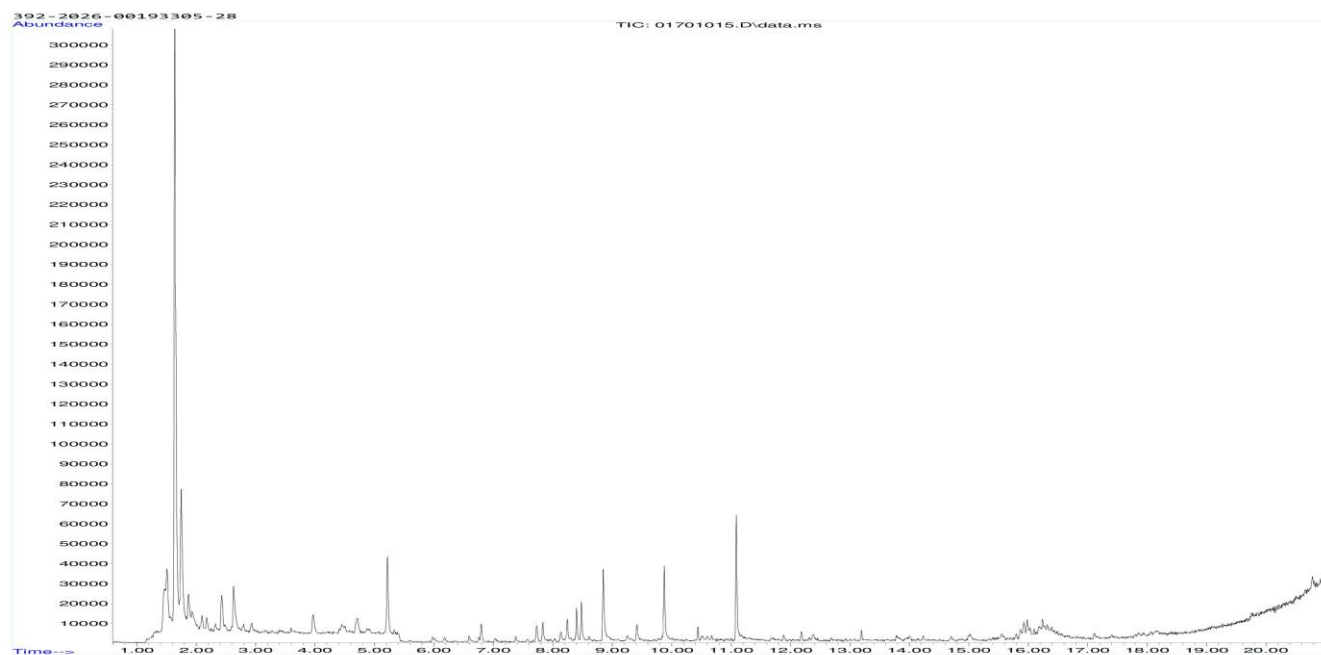
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7 Appendices

7.1 Chromatogram of VOC Emissions after 3 Days



7.2 Chromatogram of VOC Emissions after 28 Days



Please consider the different scales.

The analysis are carried out on the sample(s) as received and the result(s) are only valid for the tested sample(s).

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7.3 Sampling Report


Product Testing

*This document must be
printed and shipped with
the sample.*

Combined Sampling Report and Chain of Custody	
Name of applicant: Francesca Perazzolo, OIKOS PAINT S.R.L., Via Cherubini, 2, 47043 Gatteo (name, company, phone) (FC), ITALY, +390547681460	
Product information	
Name of the product: CRILUX	Product type: Paint, water based
Batch N°: 26PR001434	Article N°:
Model / Program / Series:	Manufacture: OIKOSPAINT S.R.L., via Cherubini 2, 47043 Gatteo ITALY (Company, Address, Stamp)
Production & Sampling information	
Production Date : 31/03/2026 Time: 17:30:00	Sampling Date: 01/04/2026 Time:
Place of sampling: WAREHOUSE (if deviating from the manufacture)	Sample is taken from: ☐ ongoing production ☒ stocks ☐ retained sample
Person in charge of sampling: Francesca Perazzolo, OIKOS PAINT SRL. (Name, company, telephone) +393923096564	Signature of sample collector: 
Where has the product been stored prior to sampling? ☐ production ☒ store ☐ miscellaneous	How has the product been stored prior to sampling? ☐ open ☒ in the stack ☐ wrapped up
Place of storage:	Packing material:
Specifics (possible negative influences by air contamination where sample was taken, by petrol emissions, by solvent emissions from production; any other uncertainties, questions, etc).	
Cut edges (identification of cut edges when present and identification of new surfaces and surface to be exposed in the emission test):	
Confirmation from the applicant	
Herewith the signer confirms the correctness of the data given above. The sample was selected, drawn and packed personally in accordance with the instructions for the taking of samples.	
Date:	Signature:
03/04/2026	(Stamp)  Oikos Paint S.r.l. sede legale in Gatteo (FC) Via Cherubini n. 2 C.F. e P. IVA 04969870270

The analysis are carried out on the sample(s) as received and the result(s) are only valid for the tested sample(s).

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Chain of Custody			
What is a Chain of custody?			
Whenever the sample is handed over, please fill out the below information			
Handed over between:	Initials + Signature	Date + Time	Condition
Handed over by			
Handed over to			
Handed over by			
Handed over to			
Handed over by			
Handed over to			
Laboratory receiving details (date, condition of package and sample, assigned lab no.): 9/4-26, OK, 392-2026-00193305			
Receptionist, Eurofins Product Testing A/S:		Signature of receptionist:	
91X2		Andrew Jamison	

7.4 How to Understand the Results

7.4.1 Acronyms Used in the Report

<	Means less than
>	Means bigger than
*	Not a part of our accreditation
α	Please see section regarding uncertainty in the Appendices
§	Deviation from method. Please see deviation section
a	The method is not optimal for very volatile compounds. For these substances smaller results and a higher measurement uncertainty cannot be ruled out
b	The component originates from the substrate and is thus removed
c	The results have been corrected by the emission from the substrate
d	Very polar organic compounds are not suitable for reliable quantification using Tenax TA adsorbent and HP-5ms GC column. A high degree of uncertainty must be expected
e	The component may be overestimated due to contribution from the system
SER	Specific Emission Rate

7.4.2 Explanation of ID Category

Categories of Identity:

- 1: Identified by comparison with a mass spectrum obtained from library and supported by other information and quantified through specific calibration.
- 2: Identified by comparison with a mass spectrum obtained from library and supported by other information. Quantified as toluene equivalent.
- 3: Identified with a lower match by comparison with a mass spectrum obtained from a library. Quantified as toluene equivalent.
- 4: Not identified, quantified as toluene equivalent.

7.5 Applied LCI and NIK Values

7.5.1 LCI/NIK Values for Compounds Found After 3 Day Measurements

Compound	CAS No.	AgBB 2024 NIK [µg/m³]	Belgian NIK [µg/m³]
1,2-Propandiol (Propylene glycol)	57-55-6	2100	2100
2-methyl-4-isothiazolin-3-on	2682-20-4	100	100
5-Chlor-2-methyl-4-isothiazolin-3-on (CIT)	26172-55-4	1	1

7.5.2 LCI/NIK Values for Compounds found after 28 Day Measurements

Compound	CAS No.	AgBB 2024 NIK [µg/m³]	Belgian NIK [µg/m³]
None determined	-	-	-

7.6 Description of VOC Emission Test

7.6.1 Test Chamber

The test chamber is made of stainless steel. A multi-step air clean-up is performed before loading the chamber, and a blank check of the empty chamber is performed.

The chamber operation parameters are as described in the test method section. (EN 16516, ISO 16000-9, internal method no.: 71M549811).

The recovery rates in the climate test chamber have been investigated using toluene and n-dodecane. The mean recovery rates of toluene and n-dodecane were concluded to be between 95 % and 100 % depending on the chamber size. These values comply with the criteria of a minimum mean recovery rate of 80 % stated in the 16000-9 test method.

Air sampling from the test chamber is carried out in a clean test chamber room at ambient air pressure and 23 ± 1 °C.

7.6.2 Expression of the Test Results

All test results are calculated as specific emission rate, and as extrapolated air concentration in the European Reference Room (EN 16516, AgBB, EMICODE, M1 and Indoor Air Comfort).

7.6.3 Testing of Carcinogenic VOCs

The emission of carcinogens (EU Categories C1A and C1B, as per European law) is tested by drawing sample air from the test chamber outlet through Tenax TA tubes after the specified duration of storage in the ventilated test chamber. Analysis is performed by ATD-GC/MS (automated thermal desorption coupled with gas chromatography and mass spectroscopy using 30 m HP-5 (slightly polar) column with 0.25 mm ID and 0.25 µm film, Agilent) (EN 16516, ISO 16000-6, internal methods no.: 71M549812 / 71M542808B).

All identified carcinogenic VOCs are listed; if a carcinogenic VOC is not listed then it has not been detected. Quantification is performed using the TIC signal and authentic response factors, or the relative response factors relative to toluene for the individual compounds.

This test only covers substances that can be adsorbed on Tenax TA and can be thermally desorbed. If other emissions occur, then these substances cannot be detected (or with limited reliability only).

7.6.4 Testing of VOC, SVOC and VVOC

The emissions of volatile organic compounds are tested by drawing sample air from the test chamber outlet through Tenax TA tubes after the specified duration of storage in the ventilated test chamber. Analysis is performed by ATD-GC/MS using HP-5 column (30 m, 0.25mm ID, 0.25µm film) (EN 16516, ISO 16000-6, internal methods no.: 71M549812 / 71M542808B).

All single substances that are listed with an LCI/NIK value in the latest publications (hereafter referred to as target compounds) are identified if present. All other appearing VOCs are identified as far as possible. Quantification of target compounds is done using the TIC signal and authentic response factors, or the relative response factors relative to toluene. For certain compound groups, which differ significantly in chemistry from toluene, quantification is performed relative to a representative member of the group for more accurate and precise results. This can include quantification of for example glycols and acids. In addition to that, all results are also expressed in toluene equivalents. All non-target compounds, as well as all non-identified substances, are quantified in toluene equivalents.

The results of the individual substances are calculated in three groups depending on their retention time when analyzing using a non-polar column (HP-1):

- Volatile Organic Compounds (VOC) are defined as: All substances eluting between and including n-hexane (n-C6) and n-hexadecane (n-C16)

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- Semi-Volatile Organic Compounds (SVOC) are defined as: All substances eluting after n-hexadecane (n-C16) and before and including n-docosane (n-C22)
- Very Volatile Organic Compounds (VVOC) are defined as: All substances eluting before n-hexane (n-C6).

Total Volatile Organic Compounds (TVOC) is calculated by summation of all individual VOCs with a concentration $\geq 5 \mu\text{g}/\text{m}^3$. The TVOC can be expressed either in toluene equivalents as defined in EN 16516 and similar to ISO 16000-6, or as the sum of concentrations using specific or relative response factors. In the case of summation of concentrations using authentic or relative response factors, the toluene equivalent is applied to all non-target and non-identified VOCs before summing up. Compounds regarded as VOC in line with the above definition but elute before n-C6 or after n-C16 on the HP-5 column are treated as VOC, and are thus added to the TVOC.

Total Semi-Volatile Organic Compounds (TSVOC) is calculated by the summation of all individual SVOCs expressed in toluene equivalents with a concentration $\geq 5 \mu\text{g}/\text{m}^3$, as defined in EN 16516. VOCs that are regarded as VOC in line with the above definition, but elute after n-C16 in this test, are not added to the TSVOC.

Total Very Volatile Organic Compounds (TVVOC) is calculated by the summation of all individual VVOCs with a concentration $\geq 5 \mu\text{g}/\text{m}^3$ and expressed in toluene equivalents. VOCs that are regarded as VOC in line with the above definition, but elute before n-C6 in this test, are not added to the TVVOC.

This test only covers substances which can be adsorbed on Tenax TA and can be thermally desorbed. If emissions of substances outside these specifications occur then these substances cannot be detected (or with limited reliability only). The relative standard deviation of the analysis of VOCs can vary greatly depending on the identified compound. For most apolar to slightly polar compounds it is around 10%. For most polar compounds the relative standard deviation will be much higher. The specific relative standard deviation for the reported compounds can be reported upon request.

7.6.5 Calculation of R Values with LCI Lists

The concentrations of detected compounds $\geq 5 \mu\text{g}/\text{m}^3$ are divided by their respective LCI/NIK value (if defined in the given publication). The sum of the quotients gives the R value, which can be mathematically expressed:

$$R = \sum_1^n \left(\frac{c_i}{\text{NIK}_i} + \dots + \frac{c_n}{\text{NIK}_n} \right)$$

This R value is calculated, depending on the purpose of this test, for the European LCI list (published November 2023), for the German LCI/NIK list (R_D) (published September 2024), and/or for the Belgian LCI list (R_B) (updated September 2024). The relative standard deviation of the R-value varies depending on the identified compounds, but will usually be around 22%. The specific relative standard deviation for the reported results can be reported upon request.

All VOCs without published LCI/NIK value and concentration $\geq 5 \mu\text{g}/\text{m}^3$ are summed up as sum of VOCs without LCI/NIK if required by the standard or protocol.

7.6.6 Testing of Aldehydes

The presence of aldehydes is tested by drawing air samples from the test chamber outlet through DNPH-coated silicagel tubes after the specified duration of storage in the ventilated test chamber. Analysis is performed by solvent desorption and subsequently by HPLC and UV-/diode array detection.

The absence of formaldehyde and other aldehydes is stated if UV detector response at the specific wavelength is lacking at the specific retention time in the chromatogram. Otherwise it is checked whether the

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reporting limit is exceeded. In this case the identity is finally checked by comparing full scan sample UV spectra with full scan standard UV spectra.

Conversions of specific aldehydes from $\mu\text{g}/\text{m}^3$ to ppm are done by the ideal gas law using a temperature of 23 degree Celsius and standard atmospheric pressure.

7.6.7 Testing of Phthalates

The presence of phthalates is tested by drawing air samples from the test chamber outlet through tube with Florisil adsorbent after the specified duration of storage in the ventilated test chamber. Analysis is performed by solvent desorption and subsequently by GC/MS. Analysis of phthalates is not currently covered by the accreditation (Internal methods no.: 71M549812 / 71M546060).

7.7 Quality Assurance

Before loading the test chamber, a blank check of the empty chamber is performed and compliance with background concentrations in accordance with EN 16516 / ISO 16000-9 is determined.

Air sampling at the chamber outlet and subsequent analysis is performed in duplicate. Relative humidity, temperature and air change rate in the chambers is logged every 5 minutes and checked daily. A double determination is performed on random samples at a regular interval and results are registered in a control chart to ensure the uncertainty and reproducibility of the method.

The stability of the analytical system is checked by a general function test of device and column, and by use of control charts for monitoring the response of individual substances prior to each analytical sequence.

7.8 Accreditation

The testing methods described above are accredited online with EN ISO/IEC 17025 by DANAK (no. 522). This accreditation is valid worldwide due to mutual approvals of the national accreditation bodies (ILAC/IAF, see also www.eurofins.com/galten.aspx#accreditation).

Eurofins Product Testing Denmark A/S is notified body for the construction products regulation (EU) No 305/2011 with number NB 2657 under system 3.

Not all parameters are covered by this accreditation. The accreditation does not cover parameters marked with an asterisk (*), however analysis of these parameters is conducted at the same level of quality as for the accredited parameters.

The customer is solely responsible for the accuracy and completeness of the sampling report and all associated information. Eurofins accepts no liability for any errors or consequences arising from customer-provided data. Results are based solely on the samples as received and the information provided by the customer.

7.9 Uncertainty of the Test Method

The relative standard deviation of the overall analysis is 22%. The expanded uncertainty U_m equals 2 x RSD. For further information please visit www.eurofins.dk/product-testing/uncertainty/.

7.10 Decision Rules

Eurofins Product Testing A/S, declare statement of conformity based on the "Binary Statement for Simple Acceptance Rule" described in ILAC's "Guidelines on decision Rules and Statements of Conformity" ILAC-G8:09/2019.

This means that results above the detection limit are always reported with two significant digits. Results are evaluated with the same number of significant digits as the corresponding limit values, and conformity is

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based on results being less than or equal to limit values.

For limit values with more than two significant digits, the third digit will be used to confirm whether a result is below or equal to the limit value. It will always be indicated in the evaluation table if this expanded evaluation is performed.

For further information, please visit www.eurofins.dk/product-testing/om-os/beslutningsregler/

7.11 Version History

Report date	Report number	Modification
08/07/2026	392-2026-00193305_A_EN	Current version