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EMICODE Test Report

1 Sample Information

Sample identification	CRAFT MSP Crystal Clear Seal
Batch no.	-
Production date	-
Product type	Sealant
Date when sample was received	02/01/2014
Testing (start - end)	27/01/2014-23/02/2014

2 Evaluation of the Results

The tested product complies with the requirements of GEV and the results correspond to the EMICODE emission class EC 1^{PLUS}

Parameter	Concentration µg/m³	EC 2 µg/m³	EC 1 μg/m³	EC 1^{PLUS} μg/m³			
TVOC _{3 days}	270	≤ 3000 ≤ 1000		≤ 750			
TVOC _{28 days}	41	≤ 300	≤ 100	≤ 60			
TSVOC _{28 days}	< 5	≤ 100	≤ 100 ≤ 50				
Total VOC without NIK 28 days	23	> 40	≤ 40				
R-value (dimensionless)	< 1	> 1	≤ 1				
Formaldehyde _{3 days}	< 3	≤ 50					
Acetaldehyde _{3 days}	< 3	≤ 50					
Sum formaldehyde and acetaldehyde after 3 days	< 0.002 ppm	≤ 0.05 ppm					
Total Carcinogens _{3 days}	< 1	≤ 10					
Total Carcinogens _{28 days}	< 1	≤ 1					

This test report does not alone entitle to use the protected trademark label EMICODE. For the use of an EMICODE label a license has to be applied for at the GEV, Düsseldorf, Germany. A license can only be granted for ready-to use products, if some additional requirements on contents of certain chemicals (e.g. solvent-free) are fulfilled.

Note: The label is supplemented with a final letter R (e.g. EMICODE EC 1 R) for installation products that fulfill the specification in clause 3.1.2 sentence 2 of GEV classification criteria and that therefore may require measures for ensuring occupational safety during application.

The results are only valid for the tested sample(s).

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3 Test Method

Method		Principle	Parameter	Quantification limit	Uncertainty			
GEV method in its current	version of	f 23.02.2011.						
Internal method numbers: 9811, 9812, 2808, 8400	9810,	GC/MS	VVOC, VOC, SVOC	5 μg/m³	22% (RSD)			
	HPLC Volatile Aldehydes				U _m = 2 x RSD= 45 %			
Test chamber parameter	r							
Chamber volume, I	119	Temperature, °C	23±1	Relative humidity	, % 50±5			
Air exchange rate, 1/h	0.5	Loading ratio, m ²	/m³ 0.007					
Sample preparation								
The sample was applied of layer with a broadness of		s plate and drawr	n off over a model	giving a 3 mm thic	k and uniform			
Deviations from the test method: None								

For detailed method description see: 5.1 Description of the applied test method on page 7





4 Results

4.1 Emissions Test after 3 Days

	CAS No.	Retention time	ID- Cat	Concen- tration	NIK- value	R- value	Emission rate	Toluene equivalent
		Min		μg/m³	µg/m³		μg/(m²*h)	μg/m³
TVOC (C ₆ -C ₁₆)				270	-	-	19000	230
VOC with NIK								
n-Hexane	110-54-3	2.10	1	2.7	72	(<5)	190	1.3
1-Methoxy-2-propanol *	107-98-2	2.71	1	16	3700	0.0043	1100	4.7
2-Ethyl-1-hexanol	104-76-7	9.16	1	3.5	540	(<5)	250	2.6
(S)-(+)-6-Methyl-1-octanol *	110453-78-6	10.66	1	76	500	0.15	5400	51
R-value = Σ Conc _i /NIK _i						0.16		
VOC without NIK								
Methylcyclopentane *	96-37-7	2.32	2	1.6	-	-	110	1.6
Not identified *	-	2.61	4	1.1	-	-	80	1.1
Not identified *	-	5.00	4	1.2	-	-	89	1.2
Not identified *	-	8.54	4	1.1	-	-	82	1.1
Not identified *	-	9.74	4	1.6	-	-	120	1.6
Not identified *	-	9.97	4	2.5	-	-	180	2.5
Not identified *	-	10.01	4	4.0	-	-	290	4.0
Not identified *	-	10.13	4	25	-	-	1800	25
Not identified *	-	10.25	4	1.5	-	-	110	1.5
Not identified *	-	10.34	4	18	-	-	1300	18
Not identified *	-	10.50	4	56	-	-	4000	56
Not identified *	-	10.54	4	63	-	-	4500	63
Not identified *	-	10.82	4	2.3	-	-	170	2.3
1-Nonanol *	143-08-8	10.99	2	14	-	-	1000	14
Total VOC without NIK				180	-	-	13000	180
Total VVOC (< C ₆)				< 5	-	-	< 400	< 5
n.d.	-	-	-	< 5	-	-	< 400	< 5
Total SVOC (> C ₁₆)				< 5	-	-	< 400	< 5
n.d.	-	-	-	< 5	-	-	< 400	< 5
Total Carcinogens				< 1	-	-	< 80	< 1
n.d.	-	-	-	< 1	1	-	< 80	< 1





	CAS No.	Retention time Min	ID- Cat	Concentration	NIK- value µg/m³	R- value	Emission rate	Toluene equivalent µg/m³	
Volatile Aldehydes C₁-C₂ measured with DNPH-Method (see 5.1.4)									
Formaldehyde	50-00-0	-	-	< 3	-	-	< 300	-	
Acetaldehyde	75-07-0	-	-	< 3	-	-	< 300	-	

n.d. Not detected

< Means less than

^{*} Not a part of our accreditation. See 5.1.6 Accreditation





4.2 Emissions Test after 28 Days

	CAS No.	Retention time	ID- Cat	Concen- tration	NIK- value	R- value	Emission rate	Toluene equivalent
		min		μg/m³	μg/m³		μg/(m²*h)	μg/m³
TVOC (C ₆ -C ₁₆)				41	-	-	2900	35
VOC with NIK								
(S)-(+)-6-Methyl-1-octanol *	110453-78-6	10.65	1	19	500	0.037	1300	13
R-value = Σ Conc _i /NIK _i						0.037		
VOC without NIK								
Not identified *	-	10.13	4	2.4	-	-	170	2.4
Not identified *	-	10.34	4	2.6	-	-	190	2.6
Not identified *	-	10.49	4	8.4	-	-	600	8.4
Not identified *	-	10.53	4	14	-	-	1000	14
Not identified *	-	10.98	4	3.6	-	-	260	3.6
Total VOC without NIK				23	-	-	1600	23
Total VVOC (< C ₆)				< 5	-	-	< 400	< 5
n.d.	-	-	-	< 5	-	-	< 400	< 5
Total SVOC (> C ₁₆)				< 5	-	-	< 400	< 5
n.d.	-	-	-	< 5	-	-	< 400	< 5
Total Carcinogens				< 1	-	-	< 80	< 1
n.d.	-	-	-	< 1	-	-	< 80	< 1

n.d. Not detected

Categories of Identity:

- 1: Identified and specifically calibrated
- 2: Identified by comparison with a mass spectrum obtained from library and supported by other information. Calibrated as toluene equivalent
- 3: Identified by comparison with a mass spectrum obtained from a library. Calibrated as toluene equivalent
- 4: Not identified, calibrated as toluene equivalent

Pascal Ge

Analytical Service Manager

< Means less than

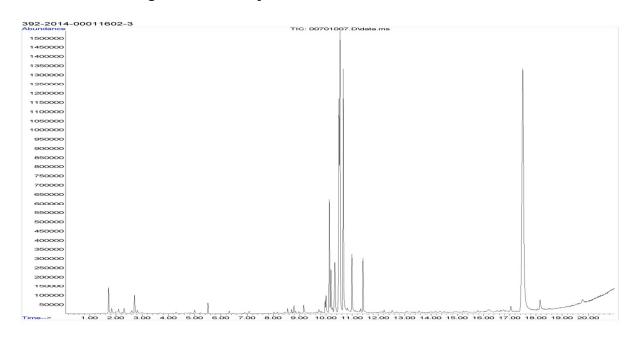
^{*} Not a part of our accreditation. See 5.1.6 Accreditation

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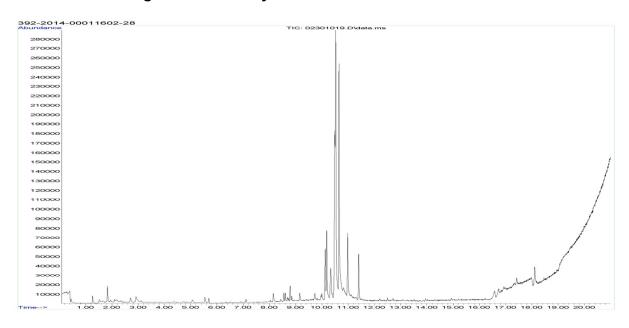




4.3.1 Chromatogram after 3 days



4.3.2 Chromatogram after 28 days



Please consider the different scales

The results are only valid for the tested sample(s).

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5 Appendices

5.1 Description of the applied test method

5.1.1 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 operation parameters are 23 °C, 50 % relative air humidity in the supply air.

5.1.2 Sampling, Desorption, Analysis

Testing for Carcinogens

The presence of carcinogens (EU Categories C1 and C2, as per the latest publication on the homepage of German BGIA Institute) was tested by drawing sample air from the chamber outlet through 2 Tenax TA tubes (main tube and backup tube). Analysis was performed by thermal desorption (Perkin Elmer) and gas chromatography / mass spectroscopy (30 m column, 0.25 mm ID, 0.25 μ m HP-1 film, Agilent) (internal methods no.: 9812 / 2808). The absence of a listed carcinogen was stated if the specific combination of fragment ions was absent at the specific retention time in the chromatogram. If no listed carcinogens were found but the required detection limit was exceeded, the identity was checked by comparing full scan sample mass spectra with full scan standard mass spectra.

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

VOC Emissions Testing

The emissions of organic compounds were tested by drawing sample air from the chamber outlet through 2 Tenax TA tubes (main tube and backup tube). Analysis was performed by thermal desorption (Perkin Elmer) and gas chromatography / mass spectroscopy (30 m column, 0.25 mm ID, 0.25 μ m HP-1 film, Agilent) (internal methods no.: 9812 / 2808).

All single substances that are listed with a NIK value in the latest AgBB publication were identified. Quantification was done with the respective response factor and the TIC signal or in case of overlapping peaks by calculating with fragment ions. All other single substances, as well as all non-identified substances, were quantified as toluene equivalent.

The results of the individual substances were calculated in three groups depending on their appearance in a gas chromatogram when analysing with a non-polar column (HP-1):

- Volatile organic compounds VOC: All substances appearing between these limits.
- Semi-volatile organic compounds SVOC: All substances appearing after n-hexadecane (n-C16).
- Very volatile organic compounds VVOC: All substances appearing before n-hexane (n-C6).

Calculation of the TVOC (Total Volatile Organic Compounds) was performed according to the AgBB/DIBt test method, by addition of the results of all individual substances with concentrations $\geq 5 \,\mu\text{g/m}^3$ in the retention time interval C6-C16. Furthermore the TVOC was calculated as the toluene equivalent, as defined in ISO 16000-6.

Calculation of the TSVOC (Total Semi-Volatile Organic Compounds) was performed by addition of the results of all substances with concentrations \geq 5 $\mu g/m^3$ between C16 and C22 as toluene equivalent, as defined in ISO 16000-6.





Calculation of the TVVOC (Total Very Volatile Organic Compounds) was performed by addition of the results of all substances with concentrations \geq 5 µg/m³ appearing before C6 as toluene equivalent, as defined in ISO 16000-6.

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

5.1.3 Calculation of R Values with the German NIK List

The concentrations of all substances $\geq 5 \,\mu\text{g/m}^3$ in the interval between n-C6 and n-C16 were divided by their respective NIK value (if given). The sum of the quotients gives the R value:

$$R = \sum_{i}^{n} \left({c_i \atop NIK_I} + ... + {c_n \atop NIK_n} \right)$$

In addition, all results were summed up for the substances without published NIK value, but in the interval be-tween n-C6 and n-C16, when concentrations were $\geq 5 \,\mu\text{g/m}^3$

5.1.4 Testing of Aldehydes after 28 Days

The presence of aldehydes was tested by drawing air samples from the chamber outlet through DNPH-coated silicagel tubes. Analysis was done by solvent desorption, HPLC and UV-/diode array detection (ISO 16000-3, internal methods no.: 9812 / 8400).

The absence of formaldehyde was stated if the specific wavelength UV detector response was lacking at the specific retention time in the chromatogram. Otherwise it was checked whether the detection limit was exceeded. In this case the identity was finally checked by comparing full scan sample UV spectra with full scan standard UV spectra.

5.1.5 Quality assurance

Before loading the chamber a blank check of the empty chamber was performed and compliance with background concentrations in accordance with ISO 16000-9 was determined. Sampling at the chamber outlet and subsequent analysis was performed in duplicate. For monitoring any breakthrough or overloading of the tubes, two Tenax TA tubes were used in series.

In each sequence stability of GC system was checked by a general function test of device and column, and by use of control charts for monitoring mean values and standard deviations for individual VOCs. Reproducibility of the method was monitored for two selected VOCs per sequence.

5.1.6 Accreditation

The testing methods described above are accredited to EN ISO/IEC 17025:2005 by DANAK (no. 522). Not all parameters are covered by this accreditation. At present the accreditation does not cover the parameters marked with a note *, however analysis was performed for these parameters at the same level of quality as for the accredited parameters.

5.1.7 Uncertainty of the test method

The relative standard deviation of the test method amounts to 22% (RSD). The expanded uncertainty U_m is 45% and equals 2 x RSD%. For further information please visit www.eurofins.dk/uncertainty.