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VOC EMISSION TEST REPORT

AgBB

31 August 2018

Sample Information 1

Sample name Batch no. Production date Product type Sample reception

06/07/2018 Wood based panel 11/07/2018

Fimaplast E-Z

2 Brief Evaluation of the Results

Regulation or protocol	Conclusion	Version of regulation or protocol
AgBB	Pass	AgBB of February 2015. DIBt of October 2010
Full data is in a sector of the test from an distance		

Full details based on the testing and direct comparison with limit values are available in the following pages

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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 [*] [RSD(%)]
EN 16516	October 2017	5	Toluene equivalents	22%
ISO 16000 -3 -6 -9 -11	2006-2011 depending on part	2	Toluene equivalents	22%
ASTM D5116-10	2010	-	-	-
AgBB/DIBt	February 2015/October 2010	5	Compound Specific	22%

3.2 Specific Laboratory Sampling and Analyses

Procedure	External Method	Internal SOP	Quantification limit / sampling volume	Analytical principle	Uncertainty [¤] [RSD(%)]
Sample preparation	ISO 16000-11:2006, EN16402:2013, CDPH, AgBB/DIBt, EMICODE	71M549810	-	-	-
Emission chamber testing	ISO 16000-9:2006, EN 16516:2017	71M549811	-	Chamber and air control	-
Sampling of VOC	ISO 16000-6:2011, EN 16516:2017	71M549812	5 L	Tenax TA	-
Analysis of VOC	ISO 16000-6:2011, EN 16516:2017	71M542808B	1 µg/m³	ATD-GC/MS	10%
Sampling of aldehydes	ISO 16000-3:2011, EN 16516:2017	71M549812	35 L	DNPH	-
Analysis of aldehydes	ISO 16000-3:2011, EN 717-1, EN 16516:2017	71M548400	3-6 µg/m³	HPLC-UV	10%





4 Test Parameters, Sample Preparation and Deviations

4.1 VOC Emission Chamber Test Parameters

Parameter	Value	Parameter	Value
Chamber volume, V[L]	238	Preconditioning period	-
Air Change rate, n[h ⁻¹]	0.5	Test period	27/07/2018 - 24/08/2018
Relative humidity of supply air, RH [%]	50 ± 3	Area specific ventilation rate, q [m/h or m³/m²/h]	0.36
Temperature of supply air, T [°C]	23 ± 1	Loading factor [m ² /m ³]	1.4
		Test scenario	Wall and ceiling

4.2 Preparation of the Test Specimen

Edges and back were covered with aluminium foil.

4.3 Picture of Sample



4.4 Deviations from Referenced Protocols and Regulations

No deviations from the referenced test methods were observed.





5 Results

5.1 VOC Emission Test Results after 3 Days

	CAS No.	Retention time	ID- Cat	Specific Conc.	Toluene eq.	Specific SER	R _D
		[min]	eat	[µg/m³]	[µg/m³]	[µg/(m²·h)]	
VOC with NIK							
Acetic acid * ^a	64-19-7	1.88	1	95	11	34	0.076
Toluene	108-88-3	3.81	1	2.7	2.7	0.95	
Hexanal	66-25-1	4.53	1	73	36	26	0.081
Cyclohexanone *	108-94-1	6.51	1	3.9	2.0	1.4	
Heptanal	111-71-7	6.69	1	3.7	2.1	1.3	
α-Pinene *	80-56-8	7.30	1	290	280	100	0.12
Camphene *	79-92-5	7.56	1	6.2	6.8	2.2	0.0041
β-Pinene *	127-91-3	8.05	1	340	360	120	0.24
3-Caren *	13466-78-9	8.57	1	12	12	4.3	0.0080
Other Alkylbenzene *	1-1-1	8.78	2	5.4	5.4	1.9	0.012
Limonene *	138-86-3	8.85	1	41	42	15	0.0082
Fenchol *	2217-02-9	10.04	1	4.2	3.3	1.5	
Borneol *	507-70-0	10.70	1	3.1	3.0	1.1	
1-Terpinen-4-ol *	562-74-3	10.83	1	2.8	2.9	0.98	
α-Terpineol *	98-55-5	10.98	1	8.8	8.6	3.2	0.0059
Caryophyllene *	87-44-5	13.49	1	7.1	7.5	2.5	0.0047
α-Caryophyllen *	6753-98-6	13.82	2	1.5	1.5	0.53	
VOC without NIK							
Not identified *		7.66	4	1.2	1.2	0.44	
Not identified *		8.19	4	1.1	1.1	0.38	
Not identified *		8.26	4	7.3	7.3	2.6	
Not identified *		9.70	4	3.4	3.4	1.2	
Not identified *		10.19	4	1.4	1.4	0.49	
Not identified *		10.38	4	4.1	4.1	1.5	
(+)-Campher *	464-49-3	10.45	2	2.8	2.8	1.0	
Not identified *		10.50	4	1.2	1.2	0.44	
Not identified *		10.67	4	6.1	6.1	2.2	
Not identified *		11.22	4	1.8	1.8	0.65	
Not identified *		11.32	4	1.5	1.5	0.55	
Not identified *		11.90	4	1.1	1.1	0.40	
(-)-Bornylacetat *	5655-61-8	12.05	2	1.2	1.2	0.43	
Not identified *		12.20	4	1.3	1.3	0.45	
Not identified *		12.80	4	2.9	2.9	1.0	

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

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392-2018-00272301_D_EN





	CAS No.	Retention time	ID- Cat	Specific Conc.	Toluene eq.	Specific SER	R _D
		[min]		[µg/m³]	[µg/m³]	[µg/(m²·h)]	
Not identified *		13.03	4	2.5	2.5	0.88	
Not identified *		13.21	4	1.4	1.4	0.50	
Not identified *		13.39	4	15	15	5.3	
Not identified *		13.68	4	1.6	1.6	0.57	
Not identified *		14.61	4	1.0	1.0	0.36	
Not identified *		15.76	4	1.0	1.0	0.37	
Sum of VOC without NIK				28	28	10	
VVOC compounds							
Acetone * a	67-64-1		1	160		58	0.14
TVVOC				160		58	
SVOC compounds							
None determined							
TSVOC				< 5	< 5	< 2	
Carcinogens							
Total carcinogens				< 1	< 1	< 1	
Aldehydes							
Formaldehyde	50-00-0		1	30		11	0.30
Acetaldehyde	75-07-0		1	31		11	0.026
Propionaldehyde	123-38-6		1	11		3.9	
Butyraldehyde	123-72-8		1	< 3		< 2	
2-butenal	123-73-9		1	< 5		< 2	
Glutaraldehyde	111-30-8		1	< 5		< 2	
R-values							1.0
TVOC				910	800	320	





5.2 VOC Emission Test Results after 28 Days

	CAS No.	Retention	ID-	Specific	Toluene	Specific	R _D
		time	Cat	Conc.	eq.	SER	
		[min]		[µg/m³]	[µg/m³]	[µg/(m²·h)]	
VOC with NIK							
Toluene	108-88-3	3.81	1	2.0	2.0	0.70	
Hexanal	66-25-1	4.52	1	53	27	19	0.059
α-Pinene *	80-56-8	7.29	1	46	46	16	0.018
Camphene *	79-92-5	7.55	1	1.5	1.7	0.54	
β-Pinene *	127-91-3	8.03	1	54	57	19	0.039
3-Caren *	13466-78-9	8.56	1	2.3	2.4	0.84	
Limonene *	138-86-3	8.84	1	12	12	4.3	0.0024
Fenchol *	2217-02-9	10.03	1	2.4	1.8	0.84	
Borneol *	507-70-0	10.69	1	2.2	2.1	0.78	
1-Terpinen-4-ol *	562-74-3	10.82	1	1.9	2.0	0.68	
α-Terpineol *	98-55-5	10.97	1	5.0	4.9	1.8	
Longifolene *	475-20-7	13.38	1	10	12	3.6	0.0067
Caryophyllene *	87-44-5	13.48	1	2.8	3.0	1.0	
Other alkylbenzenes *		8.7-8.8	2	1.8	1.8	0.64	
VOC without NIK							
2-Pentylfurane *	3777-69-3	8.26	2	3.3	3.3	1.2	
Not identified *		10.37	4	2.3	2.3	0.81	
Not identified *		10.66	4	3.1	3.1	1.1	
Not identified *		11.06	4	1.8	1.8	0.65	
Not identified *		13.02	4	1.6	1.6	0.56	
Not identified *		13.81	4	1.6	1.6	0.56	
Sum of VOC without NIK				< 5	< 5	< 2	
VVOC compounds							
Acetone * a	67-64-1		1	100		37	0.086
TVVOC				100		37	
SVOC compounds							
None determined							
TSVOC				< 5	< 5	< 2	
Carcinogens							
Total carcinogens				< 1	< 1	< 1	

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

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	CAS No.	Retention time	ID- Cat	Specific Conc.	Toluene eq.	Specific SER	R _D
		[min]		[µg/m³]	[µg/m³]	[µg/(m²·h)]	
Aldehydes							
Formaldehyde	50-00-0		1	29		10	0.29
Acetaldehyde	75-07-0		1	19		6.8	0.016
Propionaldehyde	123-38-6		1	6.9		2.5	
Butyraldehyde	123-72-8		1	< 3		< 2	
2-butenal	123-73-9		1	< 5		< 2	
Glutaraldehyde	111-30-8		1	< 5		< 2	
R-values							0.52
TVOC				180	150	63	





6 Summary and Evaluation of the Results

6.1 Comparison with Limit Values of AgBB

Parameter	Test afte	er 3 days	Test after 28 days		
	Concentration mg/m ³	Limit Value mg/m³	Concentration mg/m ³	Limit Value mg/m³	
TVOC	0.91	≤ 10	0.18	≤ 1.0	
TSVOC	< 0.005	-	< 0.005	≤ 0.1	
R-value (dimensionless)	1.0	-	0.52	≤ 1	
Sum without NIK	0.028	-	< 0.005	≤ 0.1	
Formaldehyde	-	-	0.029	≤ 0.1	
Total carcinogens	< 0.001	≤ 0.01	< 0.001	≤ 0.001	

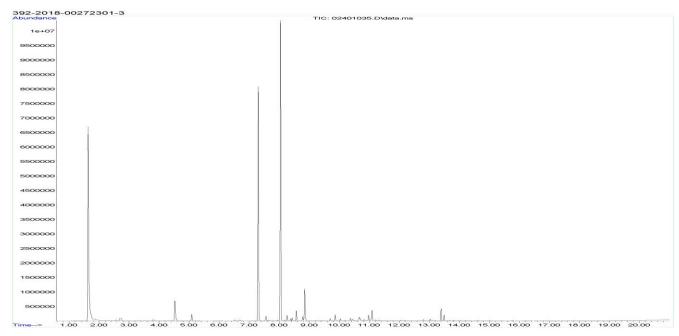
Compliance with the limits alone does not entitle to use the AgBB requirements in conjunction with approval by DIBt. This requires an application, site inspection, and approval. See www.eurofins.com/dibt-procedures.



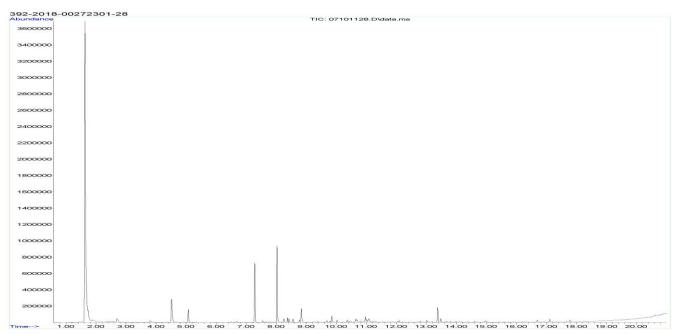


7 Appendices

7.1 Chromatogram of VOC Emissions after 3 Days



7.2 Chromatogram of VOC Emissions after 28 Days



Please consider the different scales.





7.3 Sampling Report

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EN 16516 Sampling Report

(name, company, phone): FINANCIERA MADERERA, S.A FINSA. (if different from company's name at place of sampling): *34 981050000 *34 981050000 Sampler * (Please mark): Production plant, where sampling takes place FINSA SANTIAGO Ctra. N-550, km.57 (Please mark): 15707 - Santiago de Compostela, Spain Spain (name, company, phone): Name of the product: Fimaplast E-Z Type of product Melamine faced wood based particleboard Model / Program / Series: Batch N*: Sampler * Article N*: Date of batch production: 06/07/2018 Sample was taken retained sample production: Date of sampling: 12:30 Where had the production stocks retained sample Time of sampling: 12:30 Where had the production store armic miscellaneous Production: Sampler very and the stock sampling? Place of storage: Factory warehouse 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). Packing material: Confirmation Hore sample was selected, drawn and packed personally in accordance with the instructions for the taking or samples return the sample was selected, drawn and packed personally in accordance with the instruc	Name of applicant:	RAFAEL FORT	Producer	
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Date: 31/07/2018 Signature: (Stamp)	Herewith the signer co packed personally in a	onfirms the correctness of the d accordance with the instructions	ata given above. The samples	e was selected, drawn and
OT SANTIAGO		Signature:	or the termination	*

* Please fill in an additional sampling description form per product! Sampling instruction has to be followed correctly!

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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
- ¤ 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 wooden panels and is thus removed.
- c The results have been corrected by the emission from wooden panels.
- d Very polar organic compounds are not suitable for reliable quantification using tenax TA adsorbent and HP-5 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 2015 NIK [µg/m³]
Acetic acid * ^a	64-19-7	1250
Toluene	108-88-3	2900
Hexanal	66-25-1	900
Cyclohexanone *	108-94-1	410
Heptanal	111-71-7	900
α-Pinene *	80-56-8	2500
Camphene *	79-92-5	1500
β-Pinene *	127-91-3	1400
3-Caren *	13466-78-9	1500
Other Alkylbenzene *	1-1-1	450
Limonene *	138-86-3	5000
Fenchol *	2217-02-9	1500
Borneol *	507-70-0	1500
1-Terpinen-4-ol *	562-74-3	1500
α-Terpineol *	98-55-5	1500
Caryophyllene *	87-44-5	1500
alpha-Caryophyllen *	6753-98-6	1500
Formaldehyde	50-00-0	100
Acetaldehyde	75-07-0	1200
Acetone * ^a	67-64-1	1200

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

Compound	CAS No.	AgBB 2015 NIK [µg/m³]
Toluene	108-88-3	2900
Hexanal	66-25-1	900
α-Pinene *	80-56-8	2500
Camphene *	79-92-5	1500
β-Pinene *	127-91-3	1400
3-Caren *	13466-78-9	1500
Limonene *	138-86-3	5000
Fenchol *	2217-02-9	1500

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

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Compound	CAS No.	AgBB 2015 NIK [µg/m³]
Borneol *	507-70-0	1500
1-Terpinen-4-ol *	562-74-3	1500
α-Terpineol *	98-55-5	1500
Longifolene *	475-20-7	1500
Caryophyllene *	87-44-5	1500
Other alkylbenzenes *		450
Formaldehyde	50-00-0	100
Acetaldehyde	75-07-0	1200
Acetone * ^a	67-64-1	1200





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).

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 μ m 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 a 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-5):

- Volatile Organic Compounds (VOC) are defined as: All substances eluting between and including
- n-hexane (n-C6) and n-hexadecane (n-C16)
- 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 $\ge 5 \ \mu g/m^3$. 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 $\geq 5 \ \mu g/m^3$, 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 µg/m³ 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).

7.6.5 Calculation of R Values with LCI Lists

The concentrations of detected compounds $\geq 5 \ \mu g/m^3$ 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:

$$R = \sum_{i}^{n} \left(\frac{c_{i}}{NIK_{i}} + \dots + \frac{c_{n}}{NIK_{n}} \right)$$

This R value is calculated, depending on the purpose of this test, for the European LCI list, for the German LCI/NIK list (R_D), and/or for the Belgian LCI list (R_B).

All VOCs without published LCI/NIK value and concentration $\geq 5 \ \mu g/m^3$ 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 DNPHcoated 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 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.

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 on line 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).

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.

7.9 Uncertainty of the Test Method

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