



Executive Summary

This report details the novel technology for post-consumer mechanical HDPE recycling developed by Veolia ES Plastics UK Limited at the Dagenham facility. The process, based on a recycling system developed by WRAP, focuses on recycling HDPE bottles back into food-grade materials, specifically for milk bottle applications. The technology has demonstrated consistent output quality and high decontamination efficiency, with ongoing monitoring to ensure compliance with EC 2022/1616 regulations.

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1. **Background**

a. **Purpose of the report**

The process now owned by Veolia ES Plastics UK Limited for the Dagenham facility is based on a recycling system developed by WRAP (2005) ISBN: 1-84405-225-7.

This was followed by a large-scale trial by Wrap (2007) ISBN: 1-84405-308-3.

“The development of world leading UK recycling technology allows post-consumer milk bottles to be recycled back into food contact milk bottles. Milk bottles with 30% recycled content perform identically as virgin resin bottles, have been extensively tested and have passed all EU, UK and consumer tests and are currently in production within UK dairies. The novel technology represents the first time post-consumer HDPE Milk bottles have been recycled back into Milk Bottles with full food contact status.”The process was submitted to the United States Food and Drug Administration for approval. Subsequently a No Objection Letter 108 was issued (2007).

Based on this the Dagenham facility was built (2009). The process contains the following key steps: Sorting of natural HDPE milk bottles, grinding and washing of the bottles then decontamination. Quality management of the feedstock, combined with improvements in pre-processing and decontamination reduce the risk of incidental contamination.

Since 2010 the Dagenham facility has supplied 10,000 tonnes / year to the UK dairy industry with no incidents and each batch of output material is tested for its suitability to come into contact with food.

The process was also submitted to EFSA (RECYC063) with a final safety assessment still pending on further additional data (2015).

The regulation was updated and the process was deemed a Novel Technology requiring self monitoring data and reporting based on EC 2022/1616 Article 13

The controlled version will be available online and previously superseded versions archived offline.

b. **Description of the Novel Technology**

Polymer type	Short description of the recycling technology	Specification of plastic input	Specification of output	Subject to the authorisation of individual processes	Recycling scheme applies
HDPE	Mechanical recycling	Only HDPE PCW containing maximum 1% of materials and articles that were used in contact with non-food materials or substances	Decontaminated HDPE, final materials and articles intended use for milk bottle applications	Yes	No

This a Novel Technology and there were no deviations from the requirements set out in Articles 6, 7 and 8. No recycling scheme was used. This data is consistent with the [Technical dossier](#) page 21 / 57. Confirmation of input source and traceability through the process has been independently assessed using RecyClass - Certification compliant with EN 15343:2007 Audit Report and Certificate Code: RP261-VEO-07-25-CIR-PC)

Figure 1

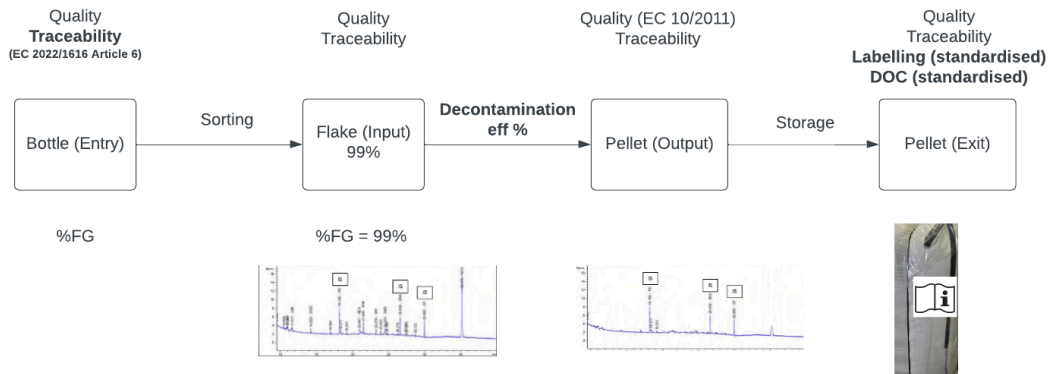
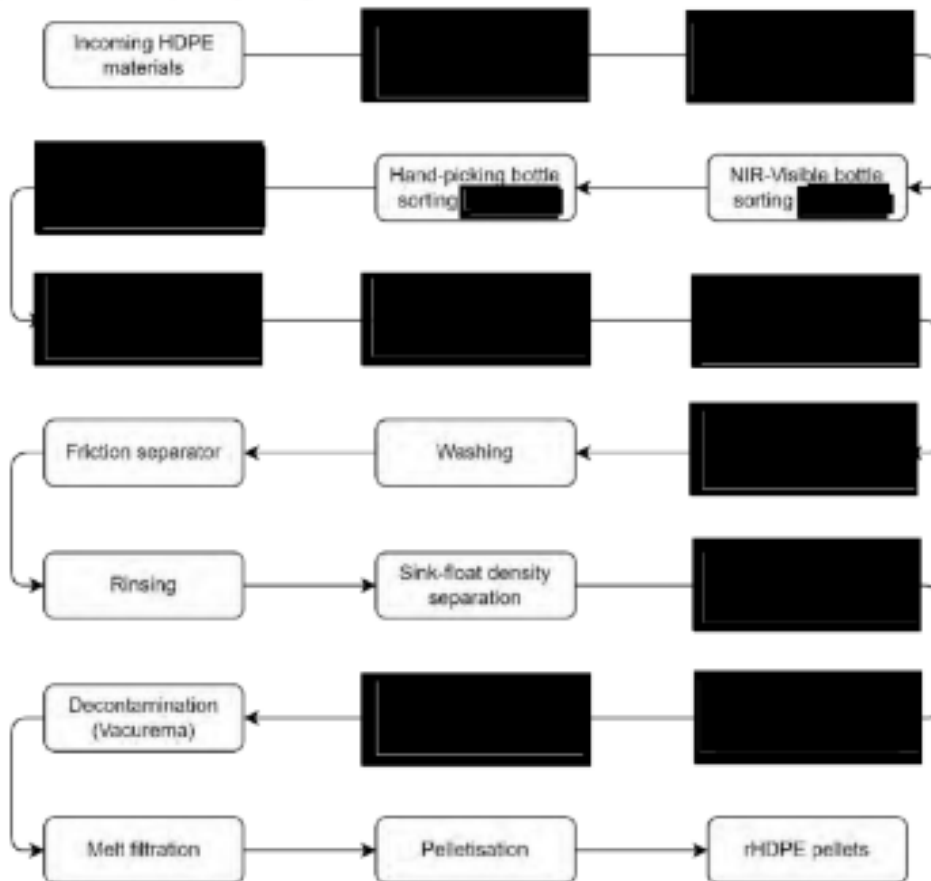


Figure 2: Veolia Plastics process flow diagram



2. Sampling strategy

The strategy is designed to ensure consistent quality and safety at each stage of the recycling process. Here's a detailed breakdown of the sampling and testing procedures:

- I. Entry Stage (Every batch):
 - A. Visual inspection of all bales
 - B. Random testing of one bale per load using ISO 15344:2008
 - C. Traceability verification through supplier pre-assessments, declarations, regulated waste transfer notes and supplier audits
 - D. Individual batch identification based on ISO 22095:2020
- II. Post-sorting (after the hand-picking bottle sorting stage) and pre-wash
 - A. Bottle count every 2 hours to confirm previous food use purity
- III. Input Stage (Every batch - random spot sample during production of that batch):
 - A. Physical characteristics:
 1. Presence of non-HDPE plastics
 2. Glue content
 - B. Chemical analysis:
 1. Quantitative analysis for volatiles (Limonene) using GC-MS(Headspace)
 2. Semi-quantitative analysis using THF extraction (ASTM D7210-21)
 3. Migration study using 50% ethanol (EC 10/2011)
- IV. Output Stage (Composite sample, minimum 1 kg/hr at 1t/hr production rate):
 - A. Physical characteristics:
 1. Melt Flow Rate at 2.16 kg and 21.6kg, density, shape and colour
 - B. Chemical analysis:
 1. Semi-quantitative analysis using THF extraction (ASTM D7210-21)
 2. Migration study using 50% ethanol (EC 10/2011)
- V. Exit Stage:
 - A. Declaration of Compliance
 - B. Compliant labelling

Sampling every batch with full traceability for at least a full year is designed to factor in seasonal, suppliers and process variations. This will provide a larger dataset for trending and reduce the overall uncertainty.

3. **Monitoring**

a. **Entry quality**

For commercial reasons results for incoming material are not discussed in this report although the records are readily available for the competent authorities to inspect at any time.

REC-001-01 Suppliers Register

REC-002-01 Incoming HDPE Load Inspections - LAB

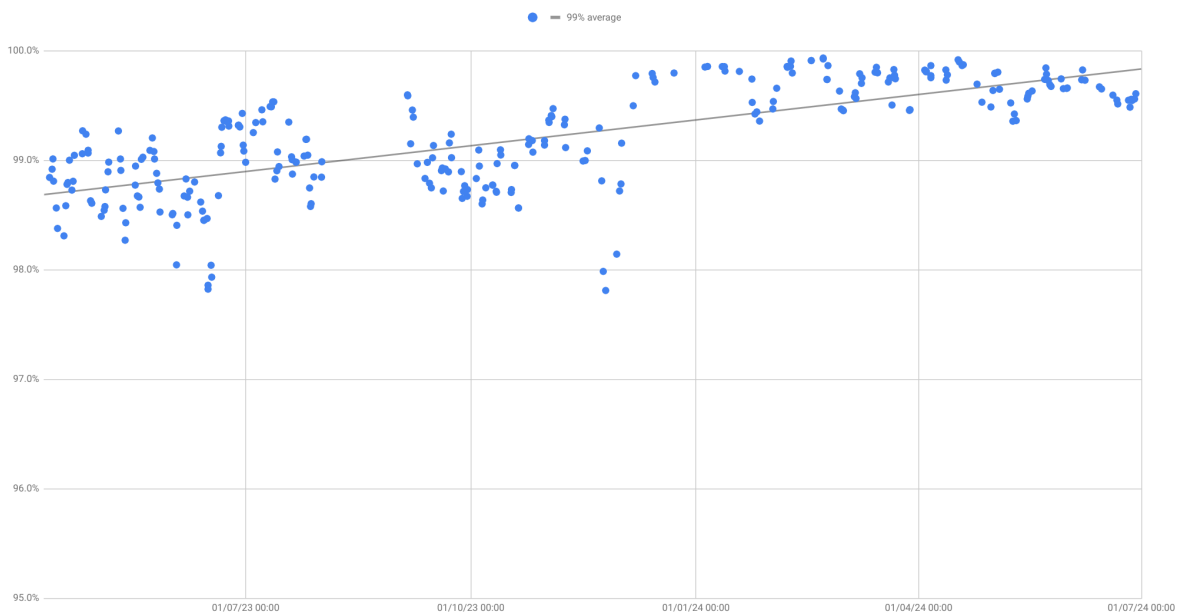
REC-018-02 Supplier and Storage Audit Log

Traceability of the process has been independently assessed using PRE RecyClass - Certification compliant with EN 15343:2007 audit report and certificate code: RP261-VEO-07-25-CIR-PC)

b. **Pre-sorted Quality**

Average 99% after the initial sorting but before granulation and further sorting. This data is consistent with the [Technical dossier](#) page 26 / 57

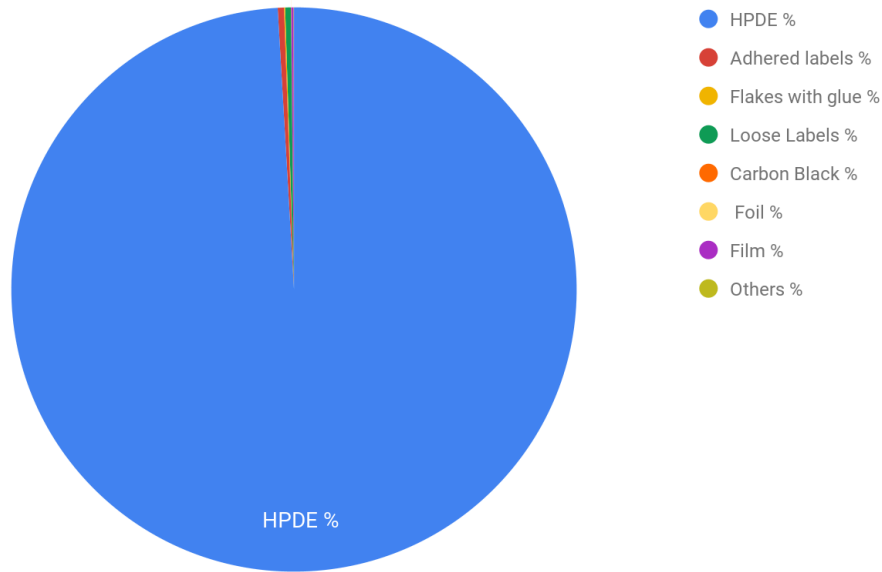
Batch Average Food Use Purity After Sorting by Counting Method



c. Input Quality
i. Contaminating Materials in Plastic Input

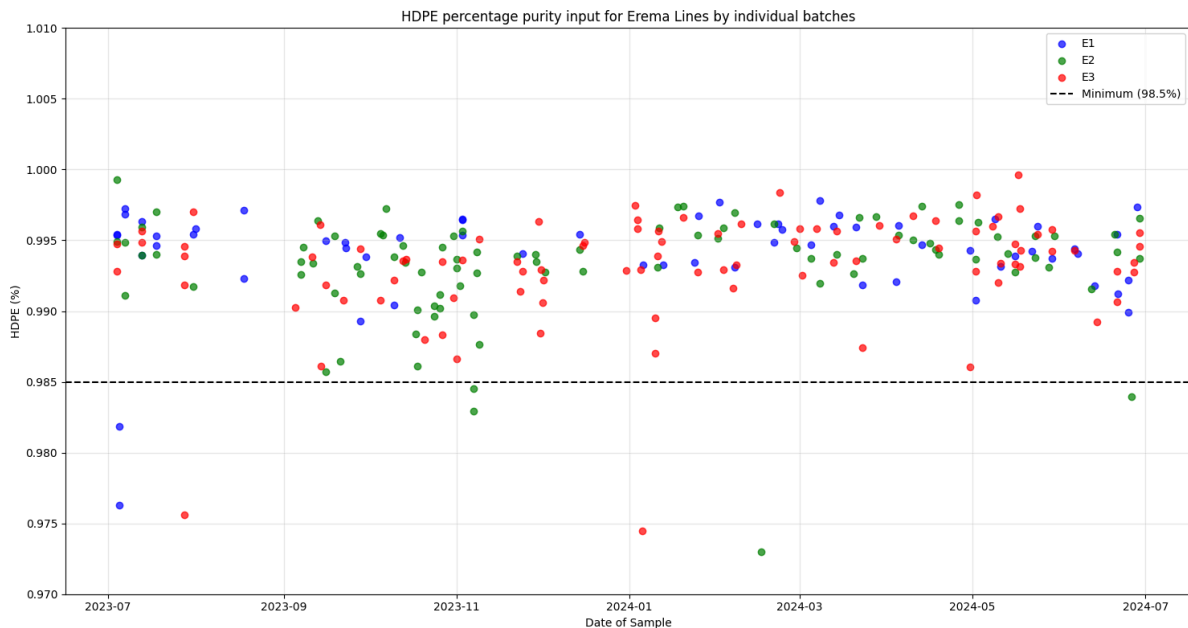
Average of input quality by mass percentage (raw data: REC-061 HDPE Flake)

From	06/06/2023	TO	30/6/2024	Number of samples		284	
HPDE %	Adhered labels %	Flakes with glue %	Loose Labels %	Carbon Black %	Foil %	Film %	Others %
99.34	0.35	0.06	0.29	0.01	0.00	0.16	0.02



Carbon black is typically from the middle layer in long life milk bottles.
 Foil is typically from the milk bottle foil seal at the top of the bottle.

99% HDPE input is consistent with the data in the [Technical dossier](#) page 27 / 57



Samples below 98.5% HDPE:

All samples below 98.5 were retested and passed based on the average result; this is consistent with the high purity at the initial sorting stage.

ii. Substances found in plastic inputs

Semi-quantitative analysis using THF extraction (ASTM D7210-21) on the flakes which was analysed on a GC-MS (raw data: REC-062 THF screening on flakes).

All Input samples were checked between 10 June 2023 and 10 June 2024 this covered all the batches produced during that time, screening down to 1.5 ppm

Library/ID	Total	Washed Flake Mean Conc	Washed Flake Max Conc	Washed Flake RT	CAS#	Qual	Source
1-Docosene	320	6.96	13.31	23.39	001599-67-3	64	Oligomer
1-Hexacosene	303	4.88	14.34	26.56	018835-33-1	64	Oligomer
Cetene	270	7.48	20.16	16.85	000629-73-2	64	Oligomer
Cyclotetracosane	204	7.12	11.51	24.77	000297-03-0	62	Oligomer
5-Eicosene, (E)-	200	6.5	16.41	22.04	074685-30-6	64	Oligomer
Tris(2,4-di-tert-butylphenyl) phosphate	196	14.24	46.3	32.65	95906-11-9	60	Irgafos 168
Hexadecane	165	4.33	41.87	16.62	000544-76-3	64	Oligomer
1-Tetracosene	162	6.69	17.77	24.75	010192-32-2	64	Oligomer
1-Octadecene	151	6.64	19.61	19.03	000112-88-9	64	Oligomer
Isopropyl myristate	139	4.85	45.89	18.95	000110-27-0	62	Cosmetics
Cyclooctacosane	132	4.2	10.53	27.48	000297-24-5	64	Oligomer
Tetradecane	124	4.27	41.89	14.11	000629-59-4	64	Oligomer
3-Eicosene, (E)-	116	7.64	18.23	21.09	074685-33-9	64	Oligomer
Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	116	3.88	6.97	32.97	002082-79-3	60	Irganox 1076
Eicosane	112	2.63	11.39	20.59	000112-95-8	72	Oligomer
Octadecane	104	3.33	9.57	18.77	000593-45-3	70	Oligomer
Octanal, 2-(phenylmethylene)-	98	2.84	8.73	18.14	000101-86-0	64	Cosmetics
E-15-Heptadecenal	95	8.04	19.26	19.36	1000130-97-9	72	TBC
Isopropyl palmitate	95	2.71	10.69	21.02	000142-91-6	62	Cosmetics
D-Limonene	89	7.22	49.8	8.86	005989-27-5	78	Flavour
1-Tricosene	73	5.96	16.33	25.56	018835-32-0	70	Oligomer

* Multiply the concentration by 10 to convert the concentration in solution to concentration in the polymer as mg/kg

d. Output Quality

i. Substances found in plastic output

Semi-quantitative analysis using THF extraction (ASTM D7210-21) analysed on GC-MS

Raw data: REC-062 THF screening on pellets

All samples were checked between 10 June 2023 and 10 June 2024 this covered all of the batches produced during that time, screening down to 1.5 ppm.

Library/ID	Total	Pellet Mean Conc	Pellet Max Conc	Pellet RT	CAS#	Qual	Source
1-Docosene	284	5.38	11.64	23.15	001599-67-3	72	Oligomer
Cyclotetracosane	225	4.85	10.73	24.69	000297-03-0	62	Oligomer
1-Hexacosene	208	3.58	12.37	26.11	018835-33-1	64	Oligomer
5-Eicosene, (E)-	157	5.89	16.63	21.3	074685-30-6	64	Oligomer
Tris(2,4-di-tert-butylphenyl) phosphate	139	9.45	25.52	32.59	95906-11-9	60	Irgafos 168
1-Tetracosene	135	5.02	13.06	24.67	010192-32-2	64	Oligomer
1-Octadecene	133	5.4	14.62	19.44	000112-88-9	78	Oligomer
Isopropyl myristate	129	3.16	5.35	18.89	000110-27-0	62	Cosmetics
Cetene	128	5.28	13.21	17.68	000629-73-2	70	Oligomer
3-Eicosene, (E)-	112	6.29	14.65	21.44	074685-33-9	68	Oligomer
Cyclooctacosane	105	2.96	7.07	27.14	000297-24-5	62	Oligomer
Hexadecane	95	4.92	39.78	16.57	000544-76-3	64	Oligomer
9-Eicosene, (E)-	89	6.45	19.69	20.25	074685-29-3	64	Oligomer
E-15-Heptadecenal	86	6	16.48	19.99	1000130-97-9	64	TBC
Octadecane	80	2.48	6.53	18.64	000593-45-3	72	Oligomer
Isopropyl palmitate	76	2.14	3.65	20.98	000142-91-6	60	Cosmetics
17-Pentatriacontene	72	3.55	8.66	27.65	006971-40-0	62	Oligomer
1-Tricosene	51	6.08	15.23	24.17	018835-32-0	64	Oligomer
Nonacos-1-ene	50	3.82	8.03	25.9	018835-35-3	76	Oligomer
5-Octadecene, (E)-	48	9.4	19.48	18.89	007206-21-5	72	Oligomer

1-Nonadecene	46	6.49	14.66	23.04	018435-45-5	64	Oligomer
Eicosane	44	1.89	3.62	20.4	000112-95-8	68	Oligomer
Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy- , octadecyl ester	43	2.66	5.04	32.83	002082-79-3	60	Irganox 1076

* Multiply the concentration by 10 to convert the concentration in solution to concentration in the polymer as mg/kg

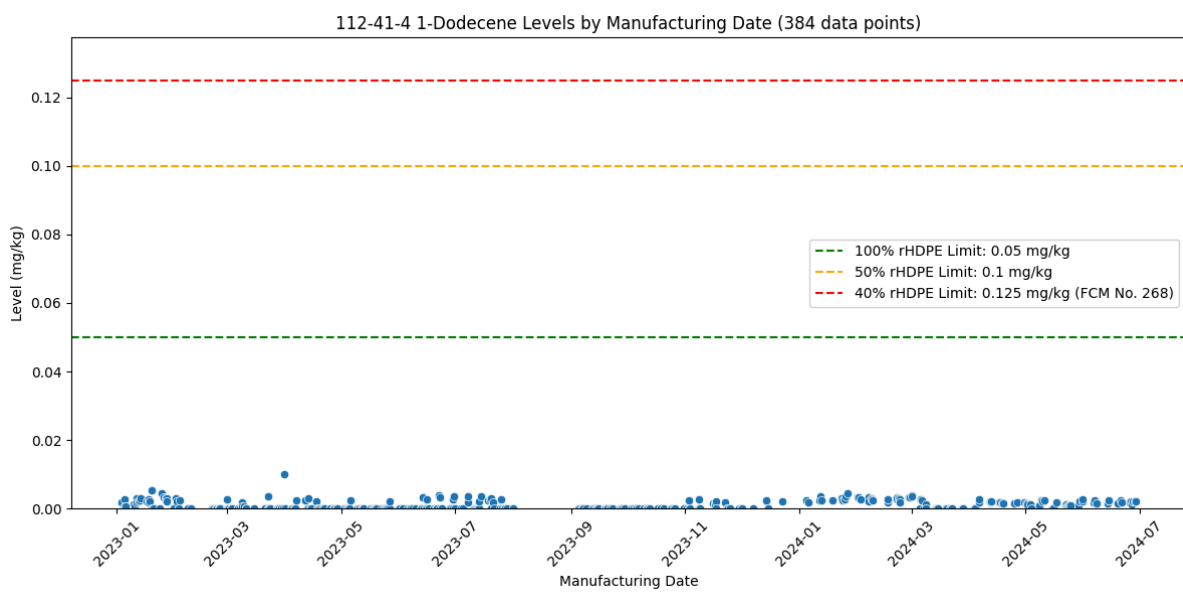
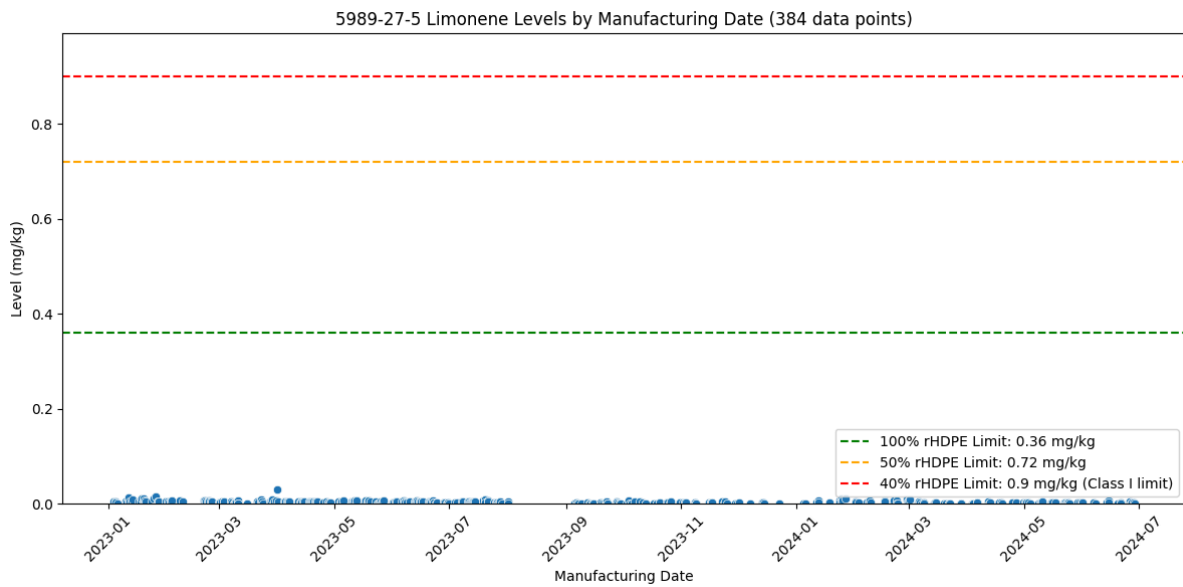
Qual = Minimum match score using NIST spectral search algorithm on GC-MS

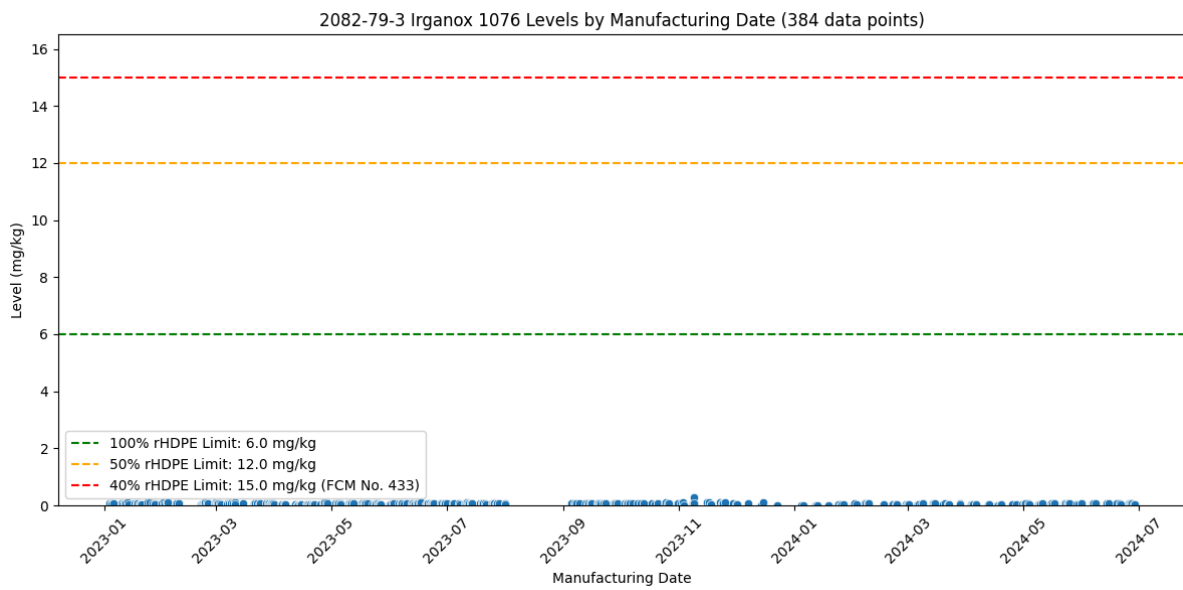
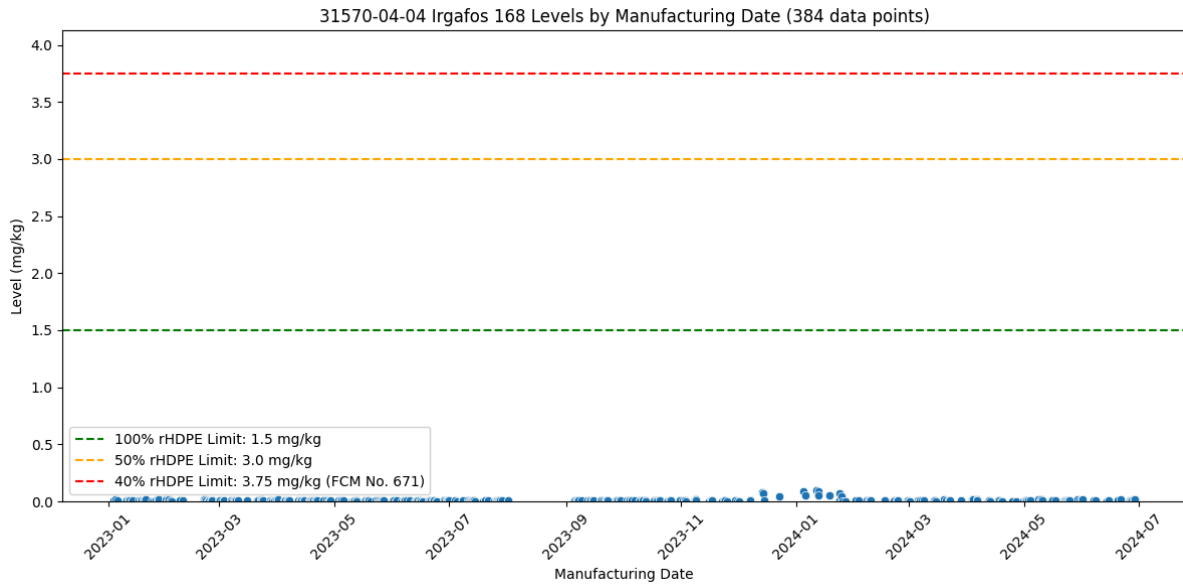
- ii. **Migration levels of contaminants** (Article 13(5)(f) a measurement or estimation of the migration levels);

Every batch produced complied with the limits stated on the declaration of compliance when blended with virgin at 40%. This can be found on the website

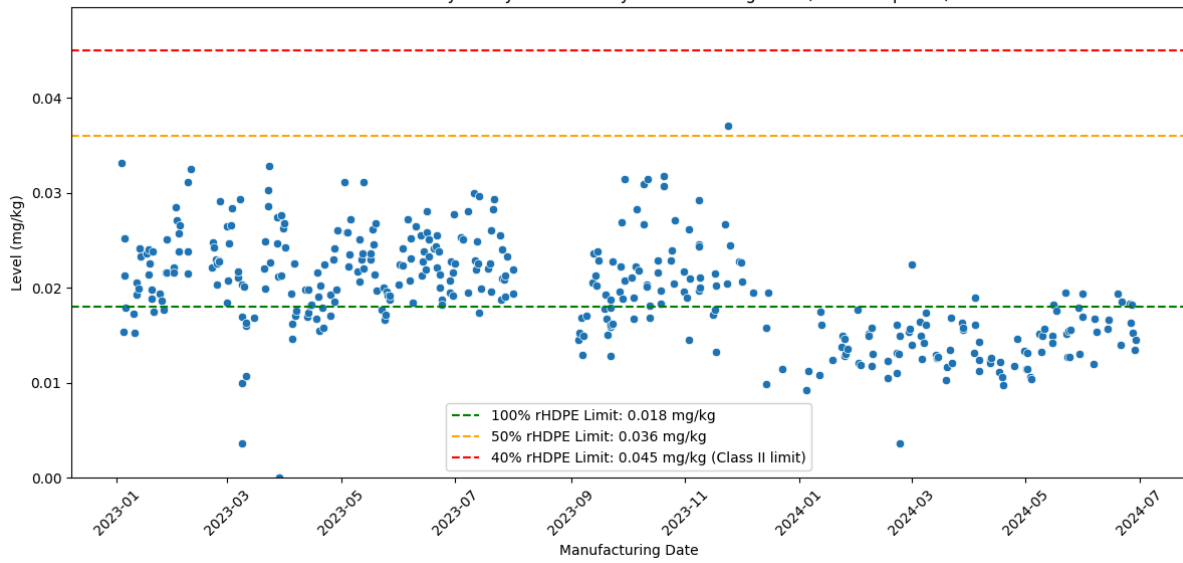
<https://dagenhamplastics.veolia.co.uk/>

CAS	Name	mg/kg	FCM
5989-27-5	Limonene	<0.017	(Class I)
112-41-4	1-Dodecene	<0.008	268
31570-04-04	Irgafos 168	<0.040	671
2082-79-3	Irganox 1076	<0.181	433
6259-76-3	n-Hexyl salicylate	<0.045	(Class II)
5444-75-7	2-Ethylhexyl benzoate	<0.049	(Class I)
101-86-0	α -Hexyl cinnamaldehyde	<0.045	(Class II)
110-27-0	Isopropyl myristate	<0.150	(Class I)
142-91-6	Isopropyl palmitate	<0.040	(Class I)
109-43-3	Dibutyl sebacate	<0.181	242
-----	Unknown	<0.010	

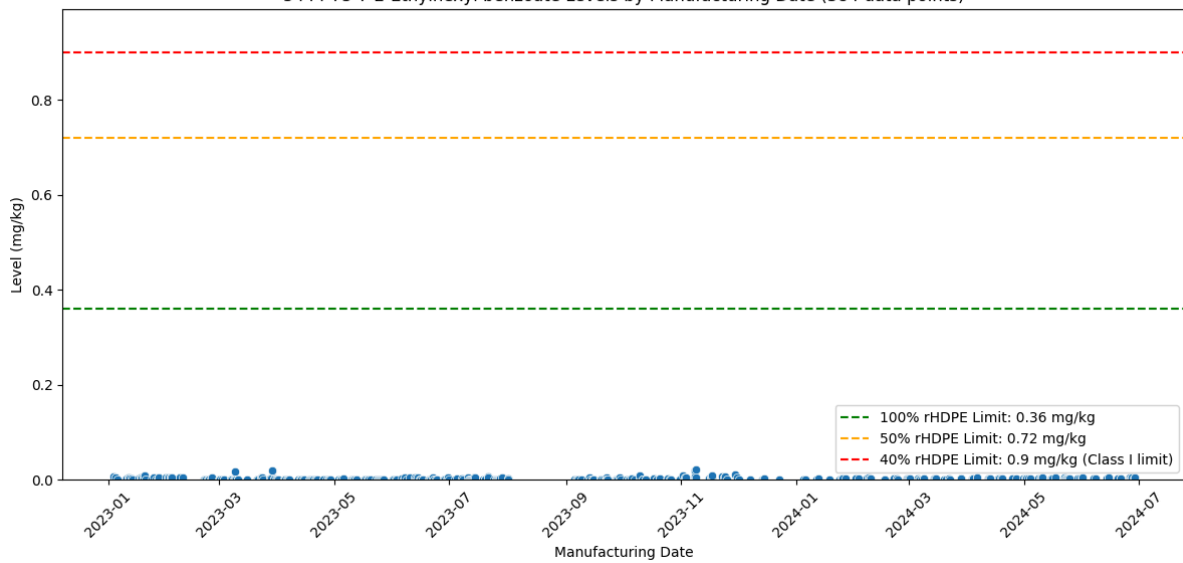




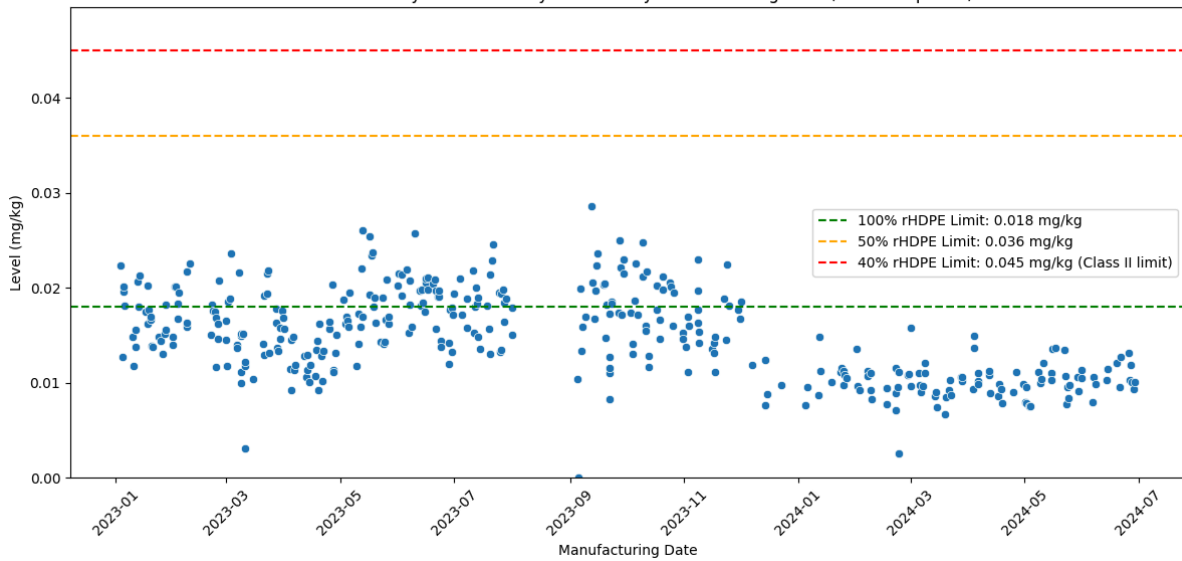
6259-76-3 n-Hexyl salicylate Levels by Manufacturing Date (384 data points)



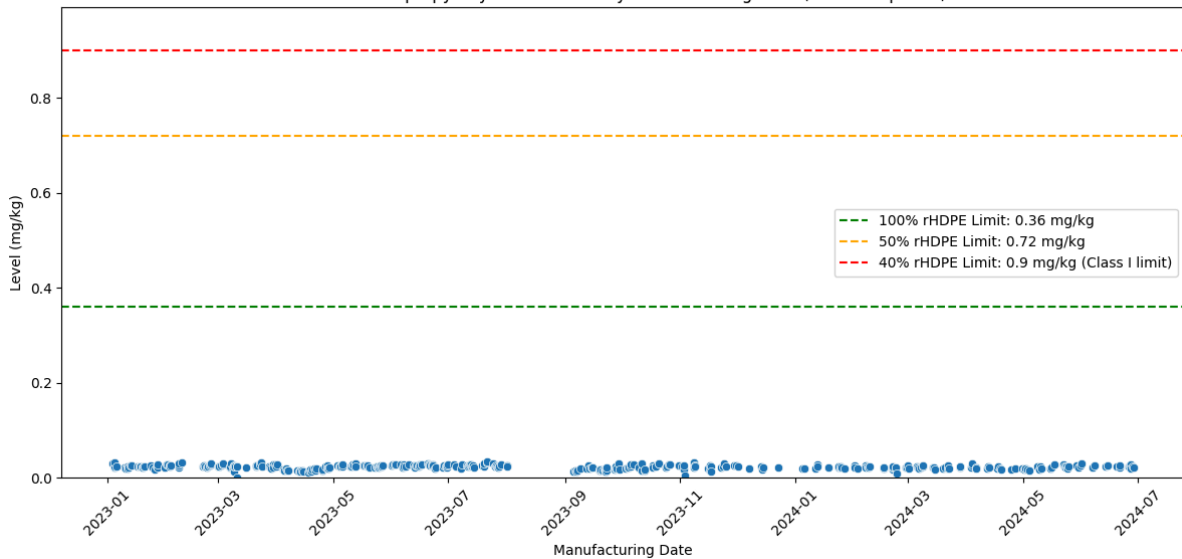
5444-75-7 2-Ethylhexyl benzoate Levels by Manufacturing Date (384 data points)



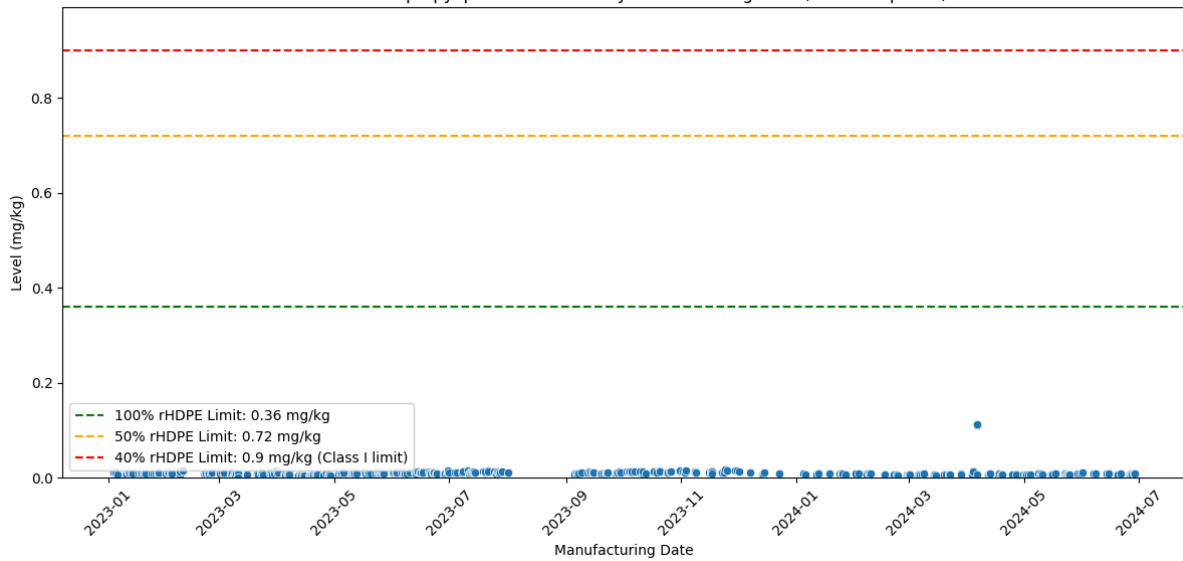
101-86-0 α -Hexyl cinnamaldehyde Levels by Manufacturing Date (384 data points)



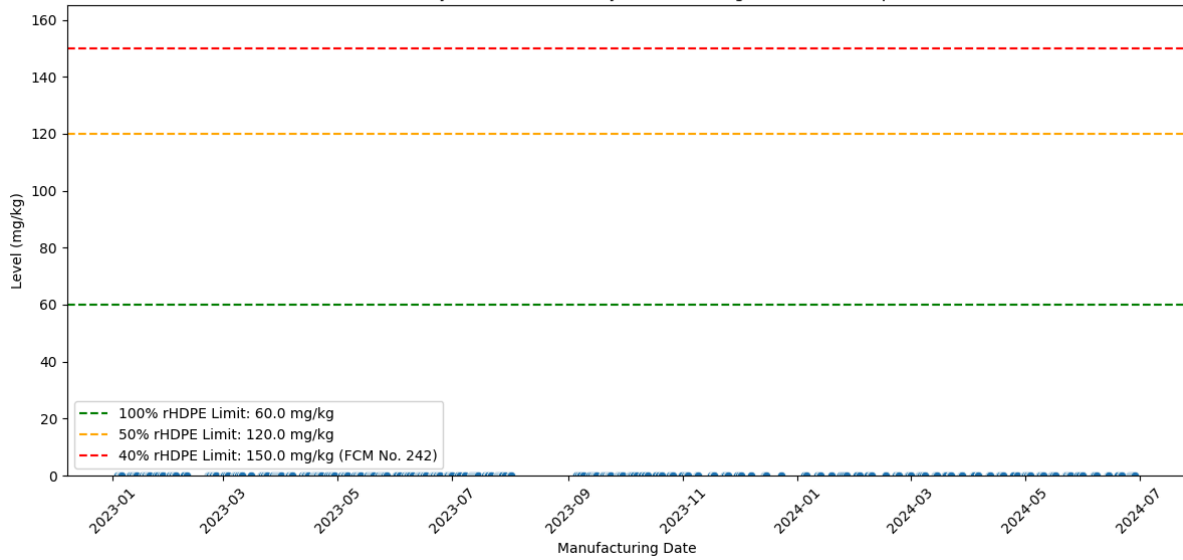
110-27-0 Isopropyl myristate Levels by Manufacturing Date (384 data points)

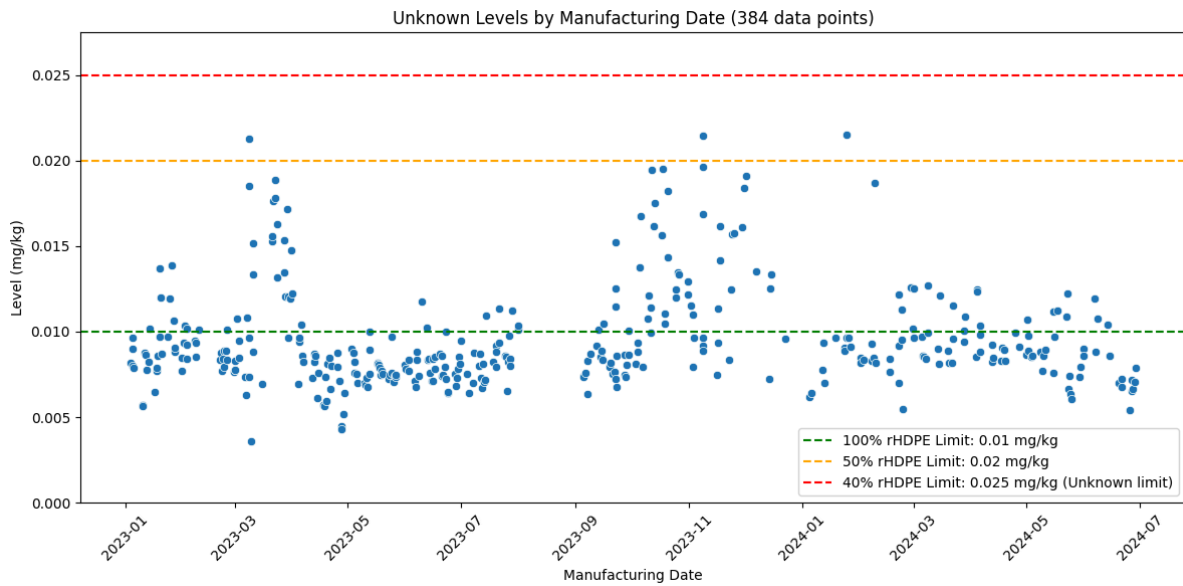


142-91-6 Isopropyl palmitate Levels by Manufacturing Date (384 data points)



109-43-3 Dibutyl sebacate Levels by Manufacturing Date (384 data points)

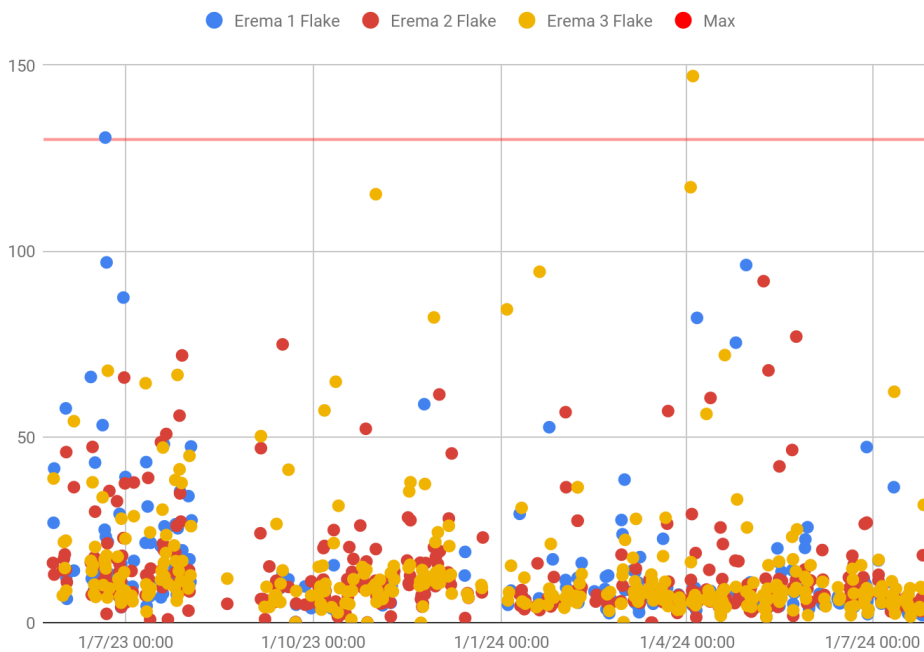




Whilst it is possible to meet the limits of 50% in most cases further work is ongoing to identify unknowns and sources of Class II compounds to maintain that margin of safety.

e. Decontamination efficiency

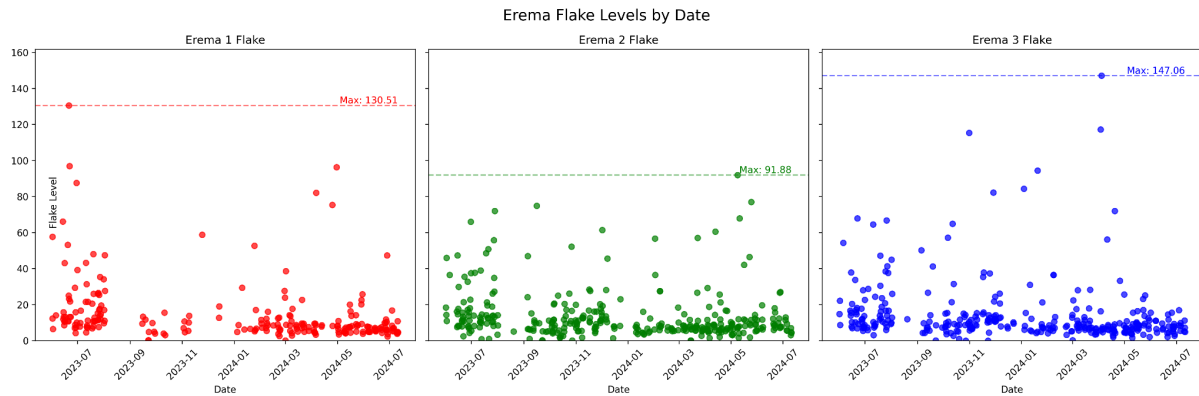
Input flake limonene concentrations (ppm) by headspace method (overlaid)



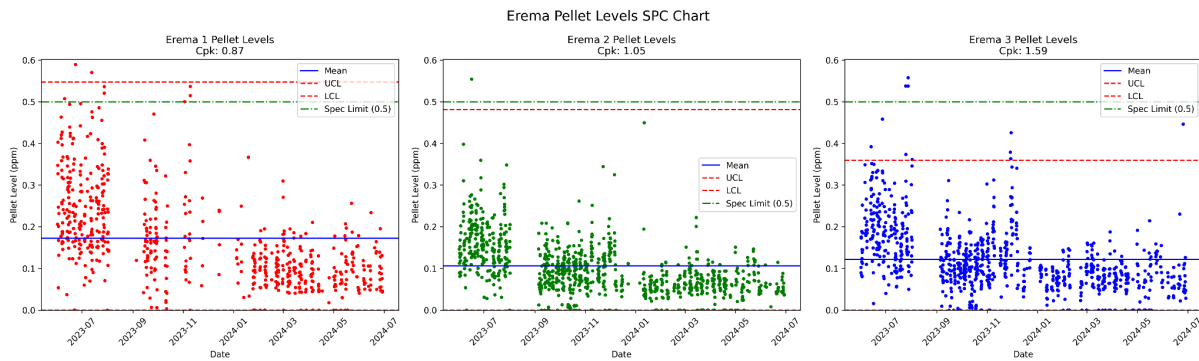


(Raw data: REC-009-01 FG rHDPE Pellet Results)

Input limonene concentrations (ppm) by headspace method (individuals plots)



Output limonene concentrations (ppm)

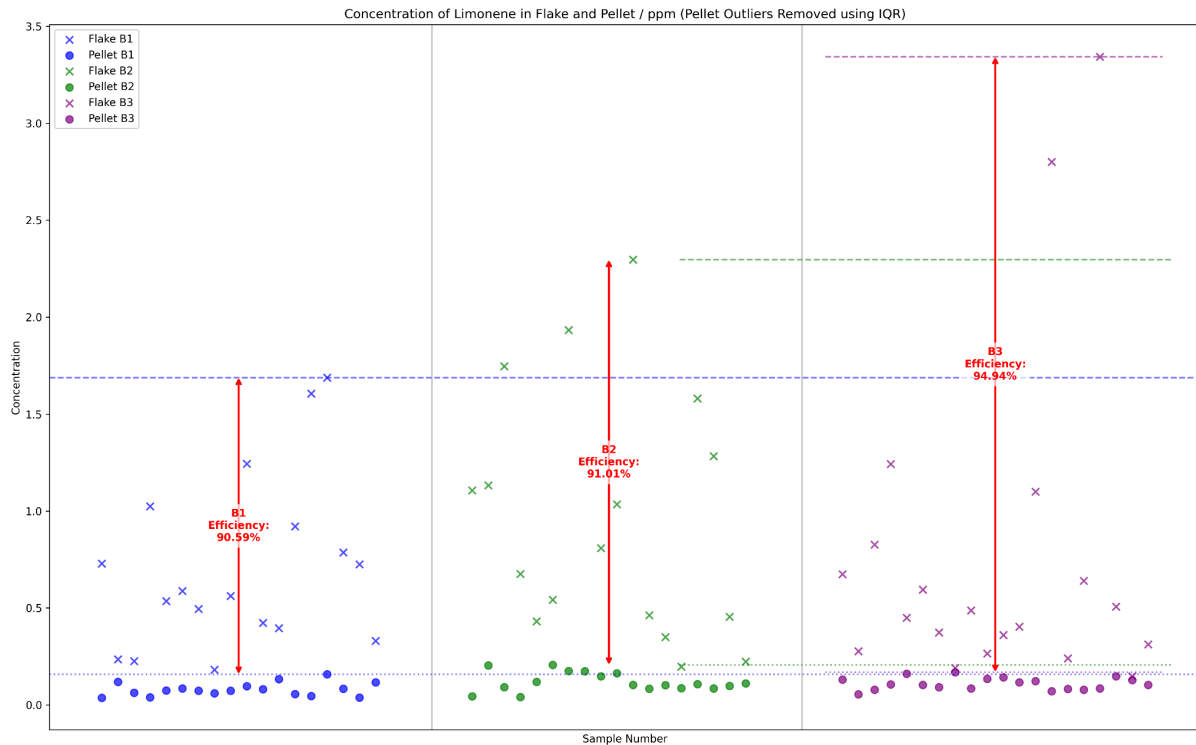


(Raw data: REC-009-01 FG rHDPE Pellet Results)

June 2023 - June 2024

Erema	Input Date Time	Output Date Time	Input Limonene	Output Limonene	Decontamination Efficiency (%)
1	2023-06-21 03:00:00	2023-06-21 12:00:00	130.51	0.25	99.81
2	2024-05-08 15:00:00	2024-05-09 00:00:00	91.88	0.05	99.94
3	2024-04-04 00:00:00	2024-04-04 06:00:00	147.06	0.06	99.96

Decontamination efficiency was also calculated for limonene using the THF extraction results

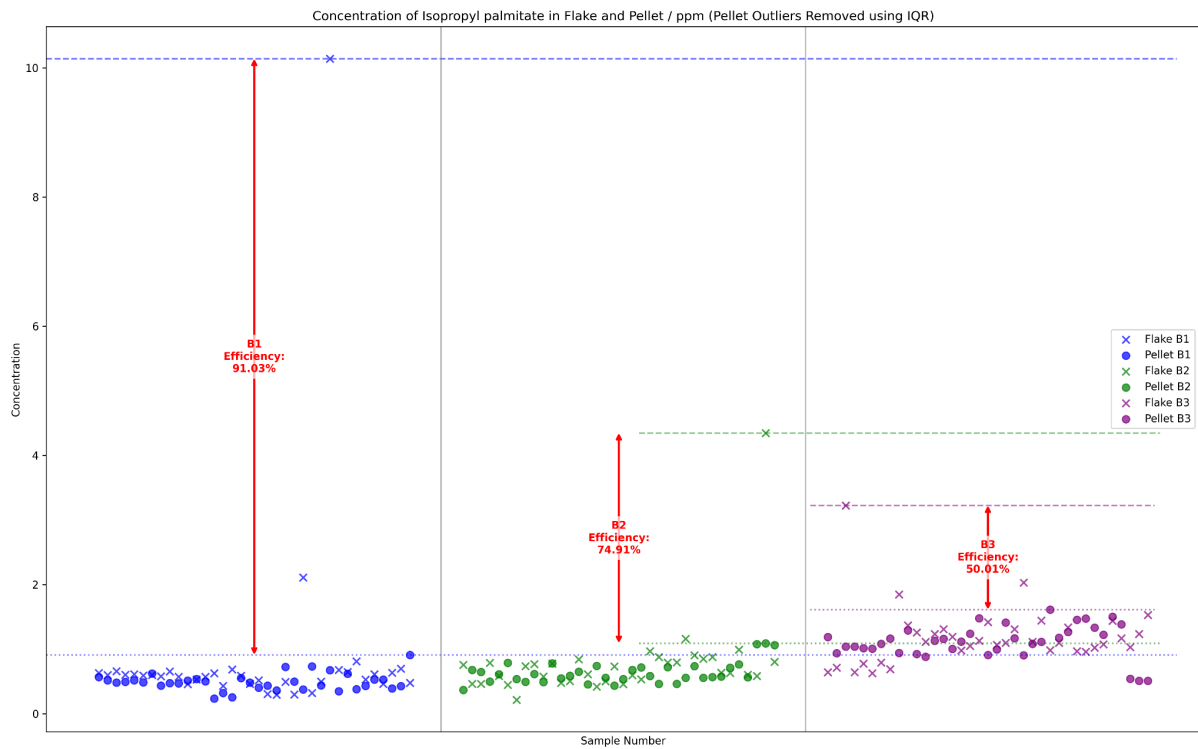
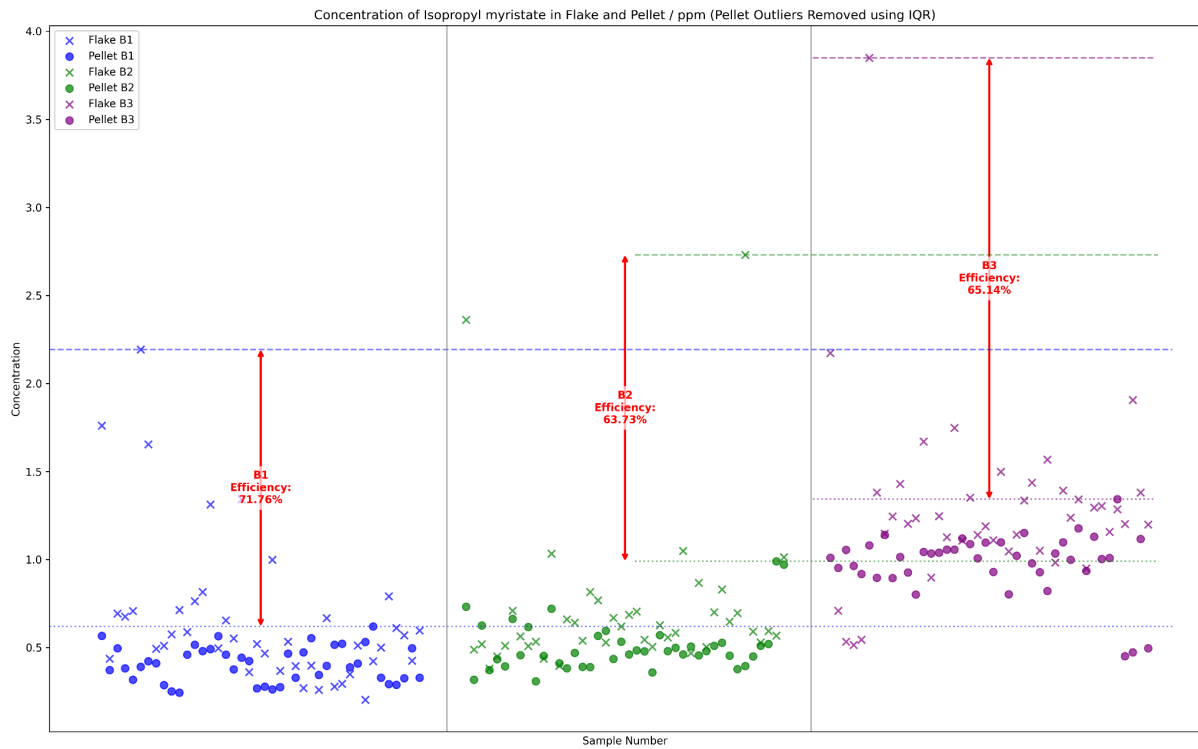


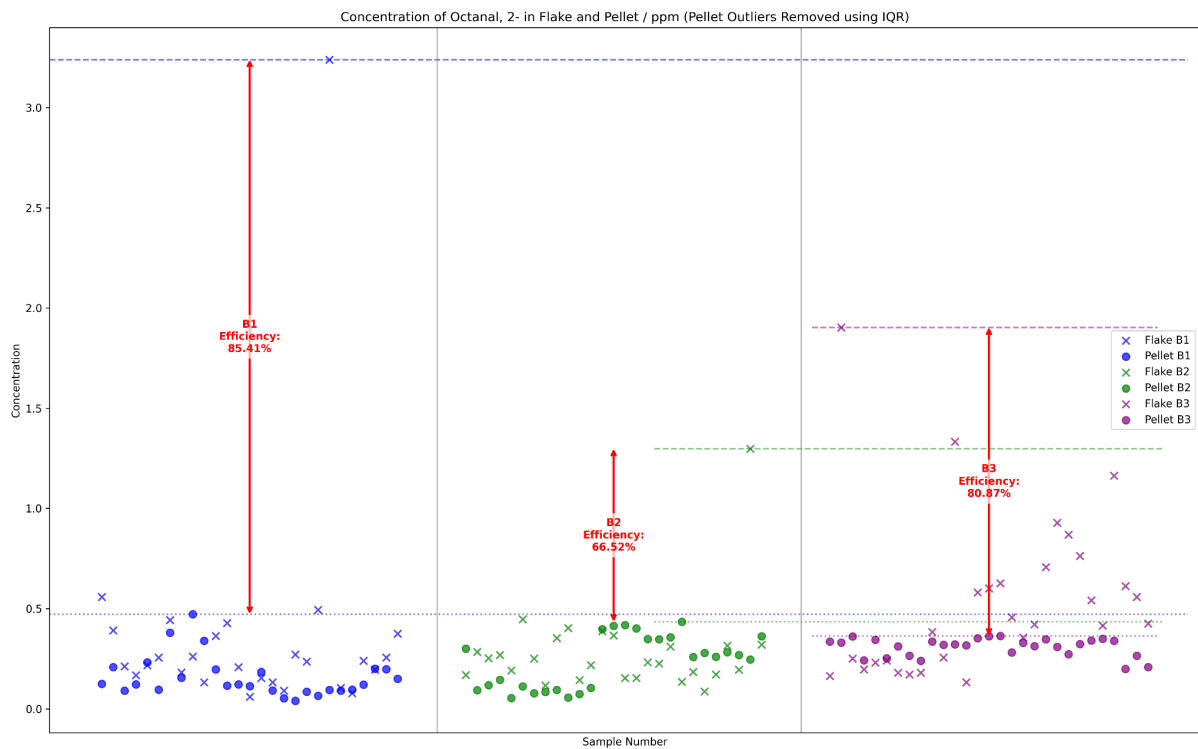
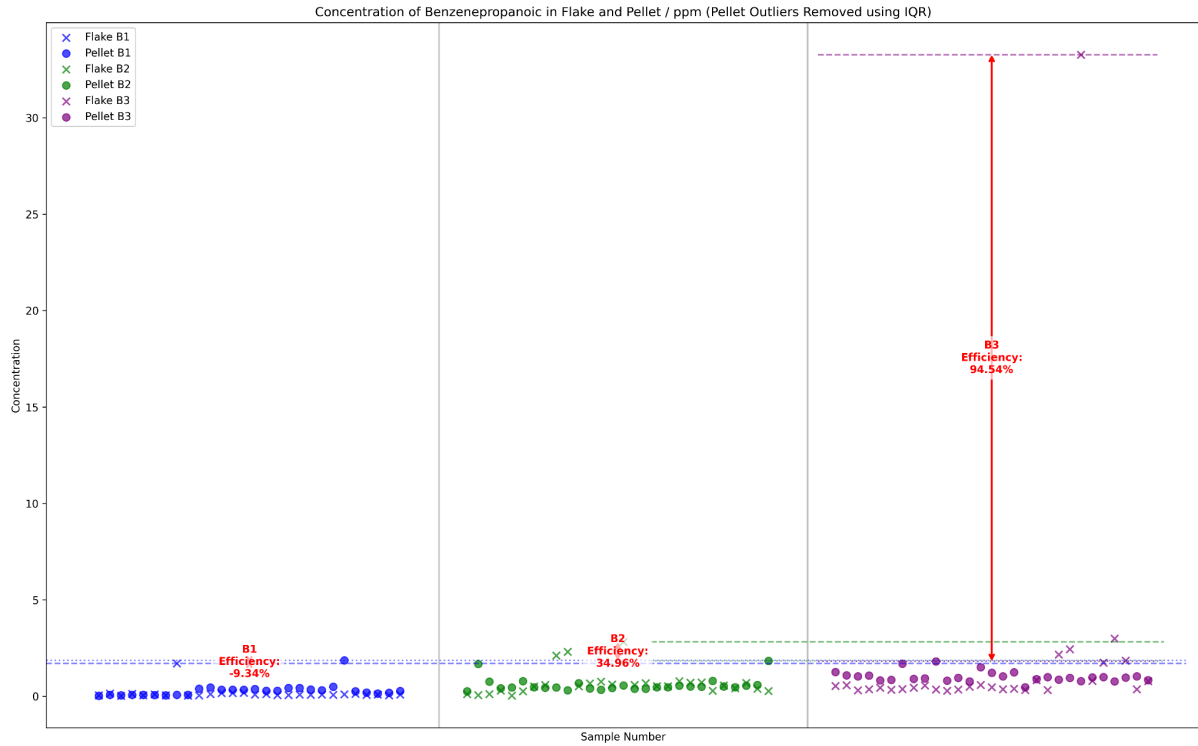
Decontamination efficiencies for a similar set of samples between June 2023 and December 2023 calculated using THF extraction were slightly lower than the headspace extraction.

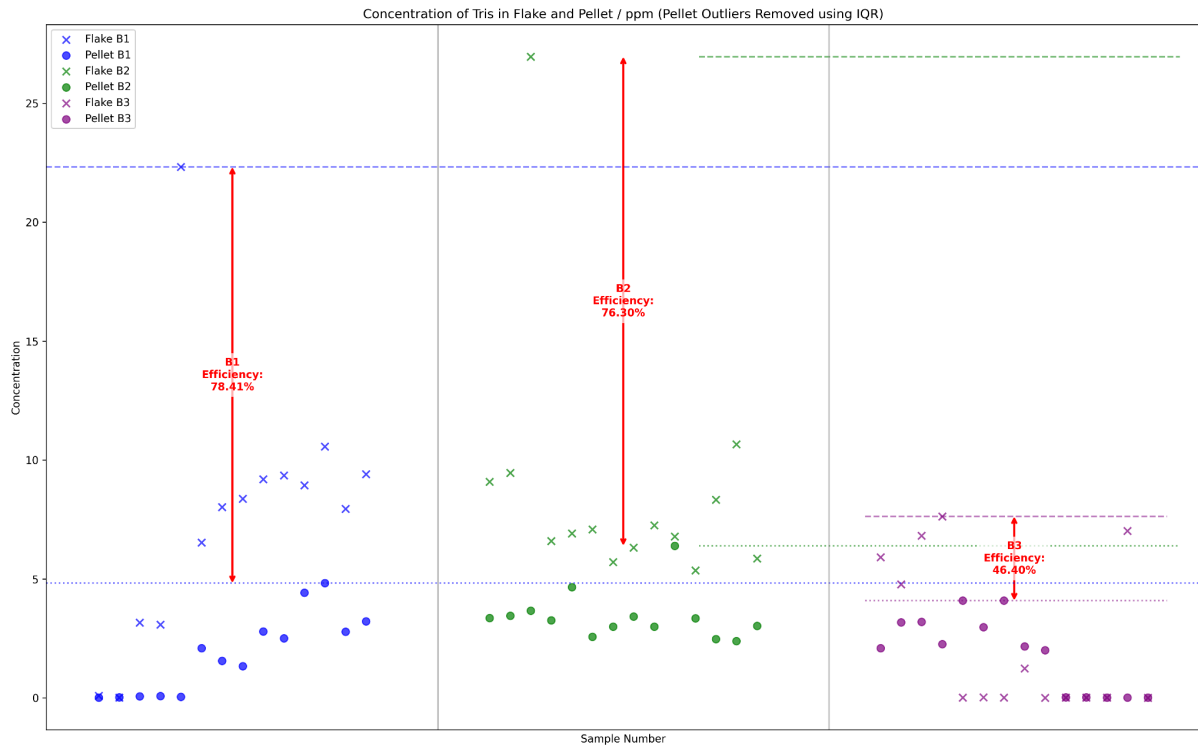
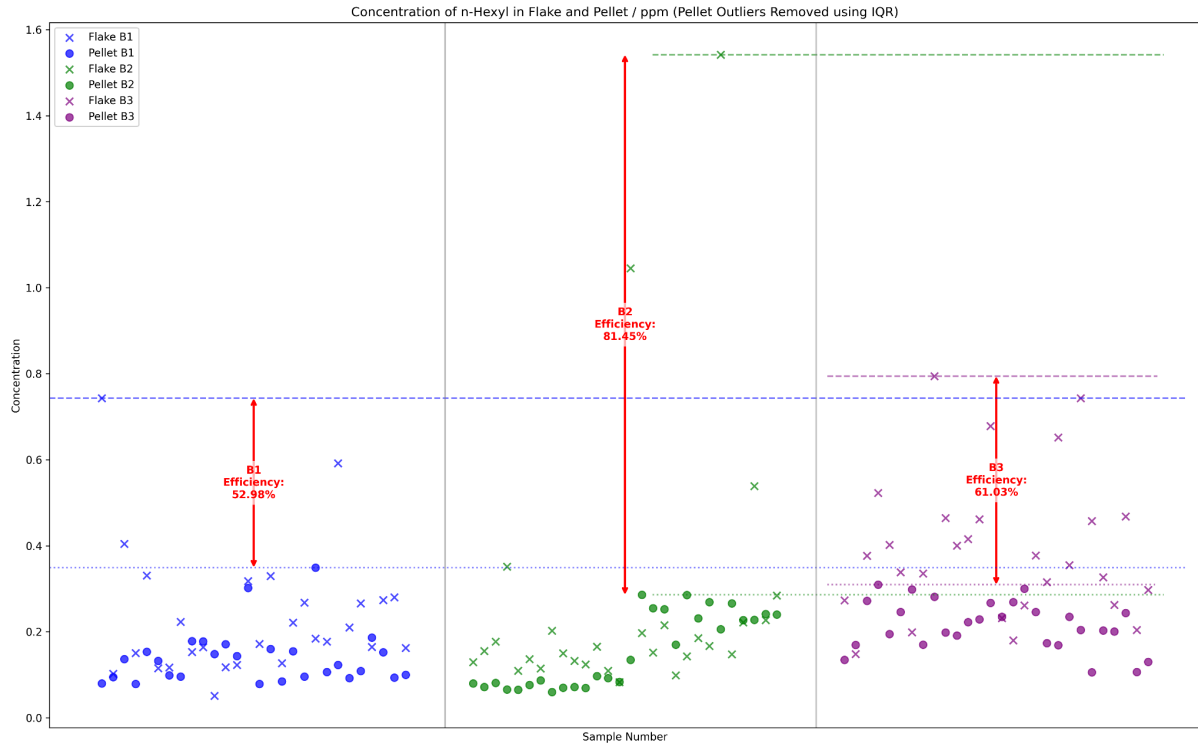
Other THF extractions were studied as they represented compounds found in the output migrations

Compound	Batch	Max Flake Sample ID	Max Flake Concentration	Max Pellet Sample ID	Max Pellet Concentration	Decontamination Efficiency (%)
Limonene	B1	D-Limonene	1.69	D-Limonene	0.16	90.59
Limonene	B2	D-Limonene	2.30	D-Limonene	0.21	91.01
Limonene	B3	D-Limonene	3.34	D-Limonene	0.17	94.94
Isopropyl myristate	B1	Isopropyl myristate	2.19	Isopropyl myristate	0.62	71.76
Isopropyl myristate	B2	Isopropyl myristate	2.73	Isopropyl myristate	0.99	63.73
Isopropyl myristate	B3	Isopropyl myristate	3.85	Isopropyl myristate	1.34	65.14
Isopropyl palmitate	B1	Isopropyl palmitate	10.14	Isopropyl palmitate	0.91	91.03
Isopropyl palmitate	B2	Isopropyl palmitate	4.35	Isopropyl palmitate	1.09	74.91
Isopropyl palmitate	B3	Isopropyl palmitate	3.22	Isopropyl palmitate	1.61	50.01

Benzenepropanoic	B1	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, methyl ester	1.70	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	1.86	-9.34
Benzenepropanoic	B2	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	2.82	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	1.84	34.96
Benzenepropanoic	B3	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	33.28	Benzenepropanoic acid, 3,5-bis(1,1-dimethylethyl)-4-hydroxy-, octadecyl ester	1.82	94.54
Octanal, 2-	B1	Octanal, 2-(phenylmethylene)-	3.24	Octanal, 2-(phenylmethylene)-	0.47	85.41
Octanal, 2-	B2	Octanal, 2-(phenylmethylene)-	1.30	Octanal, 2-(phenylmethylene)-	0.43	66.52
Octanal, 2-	B3	Octanal, 2-(phenylmethylene)-	1.90	Octanal, 2-(phenylmethylene)-	0.36	80.87
n-Hexyl	B1	n-Hexyl salicylate	0.74	n-Hexyl salicylate	0.35	52.98
n-Hexyl	B2	n-Hexyl salicylate	1.54	n-Hexyl salicylate	0.29	81.45
n-Hexyl	B3	n-Hexyl salicylate	0.79	n-Hexyl salicylate	0.31	61.03
Tris	B1	Tris(2,4-di-tert-butylphenyl) phosphate	22.33	Tris(2,4-di-tert-butylphenyl) phosphate	4.82	78.41
Tris	B2	Tris(2,4-di-tert-butylphenyl) phosphate	26.96	Tris(2,4-di-tert-butylphenyl) phosphate	6.39	76.30
Tris	B3	Tris(2,4-di-tert-butylphenyl) phosphate	7.63	Tris(2,4-di-tert-butylphenyl) phosphate	4.09	46.40







4. Analytical Procedures

- a. **Appendix 1: Migration: Quantitative Evaluation of Purity of Recycled HDPE by rapid migration test.** This method was originally developed by (Pira International Research) reference: 08A11J0703 Development of an accelerated migration test for recycled rHDPE (Pira International Research) September 2008. The method has been updated to include calibrations for known substances commonly occurring.
- b. **Appendix 2: Headspace: Quantitative Determination of Limonene Concentration in HDPE by Headspace GC-MS.** This method was originally developed by (Pira International Research) reference: 08A11J0703
- c. **Appendix 3: Extraction: Analysis of Non-Intentionally Added Substances (NIAS) in Recycled HDPE.** Based on ASTM D7210-21

5. Summary

a. Review

Summary of the reasoning on the capability of the technology to produce safe recycled materials (Article 13.5 b):

The Veolia Plastics rHDPE recycling process demonstrates a robust capability to produce safe recycled materials for food contact applications, particularly milk bottles. This conclusion is supported by several key factors: First, the process consistently achieves a high decontamination efficiency, with worst-case scenarios showing 99.8% removal of surrogate contaminants like limonene. This level of decontamination is comparable to that demonstrated in challenge tests, indicating the process's effectiveness in removing potential contaminants. Secondly, the output quality remains consistently high, regardless of input variability. Migration testing on every batch confirms compliance with EU Regulation 10/2011, with all monitored substances falling below specified limits. The process's ability to maintain this quality, even with varying input contamination levels, demonstrates its reliability in producing safe recycled materials. Additionally, the comprehensive monitoring strategy, which includes analysis of both known contaminants and potential NIAS, provides ongoing assurance of the technology's capability to meet food safety requirements.

Analysis of whether contaminant origins could lead to other undetected substances of concern (Article 13.5 e):

The majority of contaminants identified in the input material can be traced to three primary sources: residual product contents (e.g., milk components), packaging additives, and environmental contamination. While the current analytical methods effectively detect and quantify a wide range of substances, there is a possibility of undetected compounds of concern.

For instance, residual milk components could potentially harbor unidentified metabolites or breakdown products. Packaging additives, while generally well-characterized, may include proprietary formulations with undisclosed components. Environmental contaminants present perhaps the most significant unknown, as they can vary widely based on the post-consumer waste's origin and handling. To address these potential unknowns, ongoing research is focused on expanding the range of analyzed compounds and lowering detection limits. The



sorting process, which achieves at least 99% food-use HDPE packaging, significantly reduces the risk of non-authorized additives. Additionally, the process's high decontamination efficiency provides a safety margin against potential undetected substances, assuming they would behave similarly to known contaminants during processing.

b. Discussion of the difference of previous report

Compared to previous reports, this dataset demonstrates a high degree of consistency in both input quality and process performance. The decontamination efficiency remains stable, with no significant deviations from earlier findings. However, one notable difference is the increased focus on identifying and quantifying frequently occurring compounds. This shift has led to the identification of a new substance E-15-Heptadecenal, which was not prominently featured in previous reports.

The previous technical reports used decontamination efficiencies based on challenge test data. Monitoring day-to-day decontamination efficiencies are only indicative as a large number of tests need to be performed to find occasional input with elevated levels.

Even on these rare occasions the input is still approximately 90% lower than the levels used in the challenge test.

c. Analysis of discrepancies in decontamination efficiency

Three methods could have been used to monitor the decontamination efficiency.

Each method has advantages and disadvantages. The headspace method is quantitative and fast.

This allows for more measurements throughout the batch for every batch. Limonene can be monitored as an indicator of decontamination efficiency for volatile compounds. The disadvantage of this method is that there is less confidence in detection of semi-volatile compounds. The extraction method is performed on every batch using an aggressive solvent to give more confidence in extracting all chemicals present. This technique is ideal for NIAS and satisfactory for the decontamination efficiency; it should be noted that numbers are semi-quantitative. The migration is much better suited to quantitative measurements of the output. All three methods provide valuable information about decontamination performance and efficiency. **This larger dataset covering a year highlights the correlation between bottle purity post-sorting and the output quality in terms of migration for Class II (n-Hexyl salicylate & α -Hexyl cinnamaldehyde) and unknown compounds.**

d. Other considerations

The monitoring data focuses on decontamination efficiency

e. Recommendations

i. Developer

1. Continue to monitor data for trends

ii. Recycler

1. Continue to identify frequently occurring unknowns and quantify them using calibrated standards
2. Find the source of E-15-Heptadecenal and confirm any migration levels using a calibrated standard

iii. Reprocessors

1. Feedback on both positive and negative issues
2. Share external test reports

iv. Competent Authority Assessment

Article 13(5)(i) an analysis and explanation of any discrepancies observed between contaminant levels expected in the input plastic and in the output of the installation and its decontamination efficiency based on the reasoning provided under point (b) and the actual results under point (c).

This dataset remains similar to the initial technical report and shows batch batch consistency irrespective of any variation in the input material quality.

The data showed that routinely both input and output contamination levels were low. The main reason is due to a high quality of input material and also a consistently good decontamination stage. The low level of input contamination made it difficult to compare the maximum decontamination efficiency relative to the previous challenge test. Although the data gave an insight into what real contamination was found in the input material before decontamination and actual decontamination levels achieved. Irrespective of the input contamination levels a baseline decontamination was always achieved.

The following evaluation criteria are proposed for the Authority's future assessment of recycling processes applying this novel technology: (Article 10.3.f)

1. Input Material Characterization:
 - a. Detailed analysis of the post-consumer HDPE input, including:
 - b. Polymer composition (minimum 98.5% HDPE), plastics other than HDPE, Glue
 - c. Presence and quantification of potential migrants
 - d. Contamination levels (maximum 1% non-food contact materials for a defined blend rate with virgin material)
 - e. Composition and bulk density of the flake (range - average)
 - f. Flake dimensions, such as range and average of thickness, size distribution
2. Process Parameters Monitoring:
 - a. Continuous monitoring and recording of critical process parameters: a) Washing temperature and detergent concentration
 - b. Drying temperature and duration
 - c. Extrusion temperature profile
 - d. Vacuum levels in decontamination reactors
 - e. Residence time in each critical step of the process
3. Decontamination Efficiency:
 - a. Assessment of decontamination efficiency using challenge tests:
 - i. Challenge tests which are based on a standard methodology or matching previous challenge tests by the developer
 - ii. Requirement for the decontamination efficiency to match or be better than previous results
 - b. Regular assessment of decontamination efficiency by monitoring each batch
 - i. Using compounds consistently identified in the input e.g. limonene
 - ii. Day to day input material may show lower calculated decontamination efficiencies than the challenge test when the input quality is good because the Erema is not challenged. The focus should be on a consistent output level
 - iii. A large dataset approximately 100 results or more are needed to spot outliers



- iv. Accelerated migration results for input and output can be used to calculate decontamination efficiency. This is because most compounds identified in THF extractions are also identified in 50% migration results quantitatively
4. Output Quality Control:
 - a. Characterisation of the output HDPE, output including:
 - i. Melt flow index
 - ii. Colour
 - iii. Density
 - iv. Visual appearance
 - v. Form e.g. flakes, pellets, sheets
5. Migration Testing:
 - a. Compliance with EU Regulation 10/2011 on plastic materials and articles intended to come into contact with food
 - b. Specific migration limits (SML) for identified substances on each batch initially
 - c. Overall migration limit (OML) testing
6. Non-intentionally Added Substances (NIAS) Evaluation:
 - a. Screening for NIAS using advanced analytical extraction techniques (THF/GC-MS or ethanol 95% LC-MS)
 - b. Risk assessment of identified NIAS and likely sources of contamination
7. Instruction and labelling for reprocessors
 - a. Specify safe blend rate with virgin, intended application
8. Traceability System:
 - a. Implementation of a robust traceability system compliant with EU Regulation 1935/2004
 - b. Ability to trace each batch of recycled material back to its input source
9. Quality Management System:
 - a. Implementation of a comprehensive quality management system
 - b. Regular internal audits and third-party certifications
10. Long-term Performance Monitoring:
 - a. Ongoing monitoring of recycled material performance in final applications
 - b. Collection and analysis of data from end-users (e.g., dairy industry)
11. Continuous Improvement Plan:
 - a. Unknown compound retention times or mass fragments and concentrations should be trended to identify frequently occurring unknowns as they can significantly affect the blend rate.
12. Completion of a CMSS (ANNEX II) template provided by the developer highlighting critical control points, quality assessment stages and corrective actions.

6. References

- a. 08A11J0703 Development of an accelerated migration test for recycled rHDPE (Pira International Research) September 2008
- b. 08A11J0798 Studies on Recycled HDPE used for milk packaging (Pira International Research) November 2008
- c. <https://efsa.onlinelibrary.wiley.com/doi/epdf/10.2903/j.efsa.2015.4016>

7. Amendments log

- a. Section 3: Monitoring data has been updated
- b. Section 5: Review updated
- c. Section Appendices are

8. Appendices

- a. Appendix 1: Quantitative migration of contaminants in recycled HDPE by GC-FID and GC-MS



- b. Appendix 2: Quantitative Determination of Limonene Concentration in HDPE by Headspace GC-MS
- c. Appendix 3: Analysis of Non-Intentionally Added Substances (NIAS) in Recycled HDPE

Abbreviations:

ASTM: American Society for Testing and Materials
AVG: Average
CAS: Chemical Abstracts Service
CIR: Circular Economy Certification
CMSS: Compliance Monitoring Summary Sheet
CONC: Concentration
EC: European Commission
EFSA: European Food Safety Authority
EN: European Standard
EU: European Union
FCM: Food Contact Material
FDA: Food and Drug Administration
FID: Flame Ionization Detector
GC-MS: Gas Chromatography-Mass Spectrometry
HDPE: High-Density Polyethylene
ID: Identification
ISO: International Organization for Standardization
LAB: Laboratory
MAX: Maximum
MIN: Minimum
NIAS: Non-Intentionally Added Substances
NOL: No Objection Letter
OML: Overall Migration Limit
PCW: Post-Consumer Waste
ppm: Parts Per Million
PRE: Plastics Recyclers Europe
QUAL: Quality
QC: Quality Control
REC: Record (as in REC-001-01, REC-002-01, etc.)
rHDPE: Recycled High-Density Polyethylene
RP: Recycled Plastics
RT: Retention Time in minutes
SML: Specific Migration Limit
TBC: To Be Confirmed
THF: Tetrahydrofuran
WRAP: Waste and Resources Action Programme