



# SMETA Corrective Action Plan Report (CAPR)

Version 6.0



Audit Details				
Sedex Company Reference: <i>(only available on Sedex System)</i>	ZC: N/A	Sedex Site Reference: <i>(only available on Sedex System)</i>	ZS: N/A	
Business name (Company name):	BIONET WEST HELLAS SA			
Site name:	Gastouni Ilias, Thesi Agkinara			
Site address: <i>(Please include full address)</i>	Gastouni, Thesi Agkinara, 27300, Iliia	Country:	Greece	
Site contact and job title:	Diamantis Milinis – Plant Manager			
Site phone:	+30 2623033324	Site e-mail:	<a href="mailto:palmina00@gmail.com">palmina00@gmail.com</a>	
SMETA Audit Type:	<input checked="" type="checkbox"/> Labour Standards	<input checked="" type="checkbox"/> Health & Safety	<input type="checkbox"/> Environment	<input checked="" type="checkbox"/> Labour Standards
Date of Audit:	10/1/2019 (2-pillar)			



<b>Audit Company Name &amp; Logo:</b>  EUROCERT SA	<b>Report Owner (payee):</b> <i>(If paid for by the customer of the site please remove for Sedex upload)</i>
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Audit Conducted By					
Commercial	<input checked="" type="checkbox"/> (EUROCERT SA)	Purchaser	<input type="checkbox"/>	Retailer	<input type="checkbox"/>
Brand owner	<input type="checkbox"/>	NGO	<input type="checkbox"/>	Trade Union	<input type="checkbox"/>
Multi-stakeholder	<input type="checkbox"/>	Combined Audit (select all that apply)			

## Audit Content:

(1) A SMETA audit was conducted which included some or all of Labour Standards, Health & Safety, Environment and Business Ethics. The SMETA Best Practice Version 6.0 April 2017 was applied. The scope of workers included all types at the site e.g. direct employees, agency workers, workers employed by service providers and workers provided by other contractors. Any deviations from the SMETA Methodology are stated (with reasons for deviation) in the SMETA Declaration.

(2) The audit scope was against the following reference documents

### 2-Pillar SMETA Audit

- ETI Base Code
- SMETA Additions
  - Universal rights covering UNGP
  - Management systems and code implementation,
  - Responsible Recruitment
  - Entitlement to Work & Immigration,
  - Sub-Contracting and Home working,

### 4-Pillar SMETA

- 2-Pillar requirements plus
- Additional Pillar assessment of Environment
- Additional Pillar assessment of Business Ethics
- The Customer's Supplier Code (Appendix 1)

(3) Where appropriate non-compliances were raised against the ETI code / SMETA Additions & local law and recorded as non compliances on both the audit report, CAPR and on Sedex.

(4) Any Non-Compliance against customer code shall not be uploaded to Sedex. However, in the CAPR these 'Variances in compliance between ETI code / SMETA Additions/ local law and customer code' shall be noted in the observations section of the CAPR.

## SMETA Declaration

I declare that the audit underpinning the following report was conducted in accordance with SMETA Best Practice Guidance and SMETA Measurement Criteria.

- (1) Where appropriate non-compliances were raised against the ETI code / SMETA Additions & local law and recorded as non-compliances on both the audit report, CAPR and on Sedex.
- (2) Any Non-Compliance against customer code alone shall not be uploaded to Sedex. However, in the CAPR these 'Variances in compliance between ETI code / SMETA Additions/ local law and customer code' shall be noted in the observations section of the CAPR.

Any exceptions to this must be recorded here (e.g. different sample size):

Auditor Team (s) (please list all including all interviewers):

Lead auditor: Konstantinos Karvounakis

Team auditor:

Trainee auditor: Panagiotis Tzinieris

Interviewers: Konstantinos Karvounakis / Panagiotis Tzinieris

Report writer: Konstantinos Karvounakis

Report reviewer: Nikolaos Koulas

Date of declaration: 1/2/2019

*Note: The focus of this ethical audit is on the ETI Base Code and local law. The additional elements will not be audited in such depth or scope, but the audit process will still highlight any specific issues.*

*This report provides a summary of the findings and other applicable information found/gathered during the social audit conducted on the above date only and does not officially confirm or certify compliance with any legal regulations or industry standards. The social audit process requires that information be gathered and considered from records review, worker interviews, management interviews and visual observation. More information is gathered during the social audit process than is provided here. The audit process is a sampling exercise only and does not guarantee that the audited site prior, during or post-audit, are in full compliance with the Code being audited against. The provisions of this Code constitute minimum and not maximum standards and this Code should not be used to prevent companies from exceeding these standards. Companies applying this Code are expected to comply with national and other applicable laws and where the provisions of law and this Code address the same subject, to apply that provision which affords the greater protection. The ownership of this report remains with the party who has paid for the audit. Release permission must be provided by the owner prior to release to any third parties.*

## Audit Parameters

Audit Parameters			
A: Time in and time out	Day 1 Time in: 10:00 Day 1 Time out: 18:00	Day 2 Time in: Day 2 Time out:	Day 3 Time in: Day 3 Time out:
B: Number of Auditor Days Used:	1		
C: Audit type:	<input checked="" type="checkbox"/> Full Initial <input type="checkbox"/> Periodic <input type="checkbox"/> Full Follow-up <input type="checkbox"/> Partial Follow-Up <input type="checkbox"/> Partial Other – Define		
D: Was the audit announced?	<input checked="" type="checkbox"/> Announced <input type="checkbox"/> Semi – announced: Window detail:    weeks <input type="checkbox"/> Unannounced		
E: Was the Sedex SAQ available for review?	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No		
F: Any conflicting information SAQ/Pre-Audit Info to Audit findings?	<input type="checkbox"/> Yes <input type="checkbox"/> No If <b>Yes</b> , please capture detail in appropriate audit by clause  N/A		
G: Who signed and agreed CAPR (Name and job title)	Asimina Palaiologou / Quality Manager		
H: Is further information available (if Y please contact audit company for details)	<input type="checkbox"/> Yes <input checked="" type="checkbox"/> No		
I: Previous audit date:	N/A		
J: Previous audit type:	N/A		
K: Was any previous audit reviewed during this audit	<input type="checkbox"/> Yes <input type="checkbox"/> No  <input checked="" type="checkbox"/> N/A		

Audit attendance	Management	Worker Representatives	
	Senior management	Worker Committee representatives	Union representatives
A: Present at the opening meeting?	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No

B: Present at the audit?	X Yes <input type="checkbox"/> No	X Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
C: Present at the closing meeting?	X Yes <input type="checkbox"/> No	X Yes <input type="checkbox"/> No	<input type="checkbox"/> Yes <input type="checkbox"/> No
D: If Worker Representatives were not present please explain reasons why (only complete if no worker reps present)	N/A		
E: If Union Representatives were not present please explain reasons why: (only complete if no union reps present)	N/A		

**Guidance:**

The Corrective Action Plan Report summarises the site audit findings and a corrective, and preventative action plan that both the auditor and the site manager believe is reasonable to ensure conformity with the ETI Base Code, Local Laws and additional audited requirements. After the initial audit, the form is used to re-record actions taken and to categorise the status of the non-compliances.

N.B. observations and good practice examples should be pointed out at the closing meeting as well as discussing non-compliances and corrective actions.

To ensure that good practice examples are highlighted to the supplier and to give a more 'balanced' audit a section to record these has been provided on the CAPR document (see following pages) which will remain with the supplier. They will be further confirmed on receipt of the audit report.

**Root cause (see column 4)**

*Note: it is not mandatory to complete this column at this time.*

**Root cause refers to the specific procedure or lack of procedure which caused the issue to arise. Before a corrective action can sustainably rectify the situation it is important to find out the real cause of the non-compliance and whether a system change is necessary to ensure the issue will not arise again in the future.**

**See SMETA BPG Chapter 7 'Audit Execution' for more explanation of "root cause".**

**Next Steps:**

1. The site shall request, via Sedex, that the audit body upload the audit report, non-compliances, observations and good examples. If you have not already received instructions on how to do this then please visit the web site [www.sedexglobal.com](http://www.sedexglobal.com).
2. Sites shall action its non-compliances and document its progress via Sedex.
3. Once the site has effectively progressed through its actions then it shall request via Sedex that the audit body verify its actions. Please visit [www.sedexglobal.com](http://www.sedexglobal.com) web site for information on how to do this.
4. The audit body shall verify corrective actions taken by the site by either a "Desk-Top" review process via Sedex or by Follow-up Audit (see point 5).
5. Some non-compliances that cannot be closed off by "Desk-Top" review may need to be closed off via a "1 Day Follow Up Audit" charged at normal fee rates. If this is the case then the site will be notified after its submission of documentary evidence relating to that non-compliance. Any follow-up audit must take place within twelve months of the initial audit and the information from the initial audit must be available for sign off of corrective action.
6. For changes to wages and hours to be correctly verified it will normally require a follow up site visit. Auditors will generally require to see a minimum of two months wages and hours records, showing

new rates in order to confirm changes (note some clients may ask for a longer period, if in doubt please check with the client).

## Corrective Action Plan

Corrective Action Plan – non-compliances									
Non-Compliance Number <i>The reference number of the non-compliance from the Audit Report, for example, Discrimination No.7</i>	New or Carried Over <i>Is this a new non-compliance identified at the follow-up or one carried over (C) that is still outstanding</i>	Details of Non-Compliance <i>Details of Non-Compliance</i>	Root cause <i>(completed by the site)</i>	Preventative and Corrective Actions <i>Details of actions to be taken to clear non-compliance, and the system change to prevent re-occurrence (agreed between site and auditor)</i>	Timescale <i>(Immediate, 30, 60, 90, 180, 365)</i>	Verification Method <i>Desktop / Follow-Up [D/F]</i>	Agreed by Management and Name of Responsible Person: <i>Note if management agree to the non-compliance, and document name of responsible person</i>	Verification Evidence and Comments <i>Details on corrective action evidence</i>	Status <i>Open/Closed or comment</i>
Safety and Hygienic Conditions No.1	New	Company does not ensure the minimisation of the H&S hazards, so far as reasonable practicable: a) There is no available maintenance report for the burner, b) There is no valid certificate for the forklifts, c) Risk assessment for health & safety is not approved by Safety Officer, d) There is no documented definition of required PPE usage per workplace	<input type="checkbox"/> Training <input checked="" type="checkbox"/> Systems <input type="checkbox"/> Costs <input type="checkbox"/> lack of workers <input type="checkbox"/> Other – please give details:  Ineffective monitoring of health & safety requirements	Closure of all pending issues Improvement in process and frequency of monitoring health & safety requirements	60	D	Asimina Palaiologou	-Maintenance report of the burner by a licensed technician -Approval regarding the forklift control -Amendment of the risk assessment (form E5) -Documentation of PPE usage per workplace (form E2)  -Introduction of a dedicated checklist, form E606 for monitoring – to be completed along with periodic internal audits	Closed



Environment 2-Pillar No 1	New	The company has not registered for inclusion of Environmental Conditions (PPD) in its permit	<input type="checkbox"/> Training <input checked="" type="checkbox"/> Systems <input type="checkbox"/> Costs <input type="checkbox"/> lack of workers <input type="checkbox"/> Other – please give details:  Ineffective monitoring of legislative requirements	Registration at the competent authority Improvement in monitoring	30	D	Asimina Palaiologou	Registration at Prefecture of Peloponnese, P.E. Ilias, No. PDE/DAH/27522/437/28-1-2019 Amendment of monitoring methodology – introduction of a new checklist	Closed
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**Corrective Action Plan – Observations**

<b>Observation Number</b> <i>The reference number of the observation from the Audit Report, for example, Discrimination No.7</i>	<b>New or Carried Over</b> <i>Is this a new observation identified at the follow-up or one carried over (C) that is still outstanding</i>	<b>Details of Observation</b> <i>Details of Observation</i>	<b>Root cause</b> <i>(completed by the site)</i>	<b>Any improvement actions discussed</b> <i>(Not uploaded on to SEDEX)</i>
Safety and Hygienic Conditions No 1	New	a) There is no record of a recent evacuation drill, b) Signage in the plant needs to be improved regarding use of PPE and evacuation routes	Risk assessment and evacuation plans require amendment	Risk assessment updated Evacuation plans and signage to be posted – drill planned within 2019
Management system and Code Implementation No 1	New	The company has not completed relevant notification for applying ETI Base code to all suppliers of raw material (farmers – packing materials suppliers)	Initial audit / not clear what to communicate	Company planning to complete Code notification to all suppliers within next months. Communication initiated, verified e-mail on 30/1/2019
Harsh or Inhumane Treatment	New	Grievance mechanism does not assure unanimous reporting of a complaint	It was not deemed as necessary (existing mechanism)	Grievance procedure and relevant forms to be amended

			and worker rep were considered satisfactory)	
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Good examples		
Good example Number <i>The reference number of the non-compliance from the Audit Report, for example, Discrimination No.7</i>	Details of good example noted	Any relevant Evidence and Comments
	Excellent working environment and rapport between workers	

## Confirmation

<p><b>Please sign this document confirming that the above findings have been discussed with and understood by you:</b> (site management)  <i>If actual signatures are not possible in electronic versions, please state the name of the signatory in applicable boxes, as indicating the signature.</i></p>		
A: Site Representative Signature:	Asimina Palaiologou	Title – Quality Manager Date 1/2/2019
B: Auditor Signature:	Konstantinos Karvounakis 	Title – Lead Auditor Date 1/2/2019
C: Please indicate below if you, the site management, dispute any of the findings. No need to complete D-E, if no disputes.		
D: I dispute the following numbered non-compliances:		
E: Signed: (If <u>any</u> entry in box D, please complete a signature on this line)		Title Date
F: Any other site Comments:		

## Guidance on Root Cause

### Explanation of the Root Cause Column

If a non-compliance is to be rectified by a corrective action which will also prevent the non-compliance re-occurring, it is necessary to consider whether a system change is required.

Understanding the root cause of the non-compliance is essential if a site is to prevent the issue re-occurring.

The root cause refers to the specific activity/ procedure or lack of activity /procedure which caused the non-compliance to arise. Before a corrective action can rectify the situation it is important to find out the real cause of the non-compliance and whether a system change is necessary to ensure the issue will not arise again in the future.

Since this is a new addition, it is not a mandatory requirement to complete this column at this time. We hope to encourage auditors and sites to think about Root Causes and where they are able to agree, this column may be used to describe their discussion.

### Some examples of finding a "root cause"

#### Example 1

Where excessive hours have been noted the real reason for these needs to be understood, whether due to production planning, bottle necks in the operation, insufficient training of operators, delays in receiving trims, etc.

#### Example 2

A non-compliance may be found where workers are not using PPE that has been provided to them. This could be the result of insufficient training for workers to understand the need for its use; a lack of follow-up by supervisors aligned to a proper set of factory rules or the fact that workers feel their productivity (and thus potential earnings) is affected by use of items such as metal gloves.

#### Example 3

A site uses fines to control unacceptable behaviour of workers.

International standards (and often local laws) may require that workers should not be fined for disciplinary reasons.

It may be difficult to stop fines immediately as the site rules may have been in place for some time, but to prevent the non-compliance re-occurring it will be necessary to make a system change.

The symptom is fines, but the root cause is a management system which may break the law. To prevent the problem re-occurring it will be necessary to make a system change for example the site could consider a system which rewards for good behaviour

Only by understanding the underlying cause can effective corrective actions be taken to ensure continuous compliance.

The site is encouraged to complete this section so as to indicate their understanding of the issues raised and the actions to be taken.



For more information visit: [Sedexglobal.com](http://Sedexglobal.com)

Your feedback on your experience of the SMETA audit you have observed is extremely valuable. It will help to make improvements to future versions.

You can leave feedback by following the appropriate link to our questionnaire:

**[Click here for Buyer \(A\) & Buyer/Supplier \(A/B\) members:](#)**

[http://www.surveymonkey.com/s.aspx?sm=riPsbE0PQ52ehCo3lnq5lw\\_3d\\_3d](http://www.surveymonkey.com/s.aspx?sm=riPsbE0PQ52ehCo3lnq5lw_3d_3d)

**[Click here for Supplier \(B\) members:](#)**

[http://www.surveymonkey.com/s.aspx?sm=d3vYsCe48fre69DRgLY\\_2brg\\_3d\\_3d](http://www.surveymonkey.com/s.aspx?sm=d3vYsCe48fre69DRgLY_2brg_3d_3d)