

EPEAT Program

Continuous Monitoring Outcomes Report



Mobile Phones
MP-2024-01
July 8, 2024

1.0 Background

EPEAT® is a comprehensive voluntary sustainability Type 1 ecolabel that helps purchasers identify sustainable technology products and services. Central to EPEAT are conformity assurance activities that meet the technical rigor and credibility needs of the institutional purchasers who rely upon EPEAT. The EPEAT Program ensures the ongoing conformance of EPEAT-registered products through an ongoing surveillance process known as Continuous Monitoring. Continuous Monitoring activities occur throughout the year and test the ability of Participating Manufacturers to prove conformance with EPEAT Criteria on an ongoing basis.

Some Continuous Monitoring activities require that Investigations be conducted in discrete timeframes called Rounds. The EPEAT Program develops an individual plan for each Continuous Monitoring Round, which specifies the EPEAT Criteria to be investigated, the method of investigation that GEC-approved Conformity Assurance Bodies (CABs) must use and the specific dates when the Investigation activities must be completed. The EPEAT Program also selects the Participating Manufacturers and EPEAT-registered products and assigns Investigations to CABs, which must fully participate in and are responsible for implementing Continuous Monitoring Round activities with their Participating Manufacturer clients. Participating Manufacturers are required to cooperate fully with their GEC-approved CAB during Round activities.

To maintain the level of transparency relied on by purchasers, the EPEAT Program publishes an Outcomes Report at the conclusion of each Round to summarize the activities conducted and to identify the products and Participating Manufacturers that received nonconformances and the actions taken to restore accuracy of the EPEAT Registry.

This document summarizes the activities and results of Continuous Monitoring Round MP-2024-01 conducted for the Mobile Phones category.

2.0 Overview of Continuous Monitoring Round MP-2024-01

2.1 Investigation Activities

As per the published [Round Plan](#), Continuous Monitoring Round MP-2024-01 used Level 2 Investigations (laboratory evaluation of products to determine the products' conformance with specific EPEAT Criteria). GEC-approved CABs obtained the products, as identified by the EPEAT Program, from the open market without involvement of the Participating Manufacturers, where possible, and sent them for laboratory evaluation. The laboratories evaluated the products against the specified criteria and produced reports summarizing the activities conducted and the results. GEC-approved CABs reviewed the reports, made recommendations on conformity, and sent the reports to the EPEAT Program. The EPEAT Program made the final decisions on conformity for the Investigations.

2.2 Criteria Investigated

Continuous Monitoring Round MP-2024-01 focused on sustainable use of resources. The unsustainable use of resources has triggered raw material scarcities, contributed to climate change, and caused widespread environmental degradation with implications for, and negative impacts on, human health and our environment. Globally, electronic waste is the fastest growing waste stream. The United Nations attributes this growth in e-waste to technological and product proliferation, along with shorter lifecycles and fewer repair options.

Sustainable use of resources to enable a circular economy is a priority for government policy, institutional purchasers, and manufacturers worldwide. A circular economy is paramount for the electronics industry to become more sustainable and resilient. Circularity seeks to keep products in use for as long as possible, emphasizing durability, repairability, reuse, and the importance of recycling. To this end, criteria which focus on circularity and sustainable use of resources were selected for investigation in this Round. When products or components fail, the ability to repair and refurbish the product is essential to keeping it in service, and the product and packaging design should facilitate reuse and recycling.

Products are selected randomly using a random number generator from a list of Participating Manufacturers. Each product is investigated for the criteria identified in the table below, however if a product has not selected a criterion, that criterion is not investigated.

Criteria Number	Criterion Title
11.3.1	Battery removability/replacement by qualified repair service providers or authorized repair providers
11.3.3	Battery removability/replacement without use of tools
11.4.1	Ease of disassembling mobile phone
12.1.1	Use of recyclable fiber-based packaging materials
12.2.1	Separability and labeling of plastics in packaging
12.9.1	Improve packaging efficiency

3.0 Summary of Investigations and Final Decisions on Conformity for MP-2024-01

Highlights from this Continuous Monitoring Round are:

- 6 investigations completed
- 6 decisions of Conformance

All investigations in Continuous Monitoring Round MP-2024-01 were conformant. No nonconformances were identified.

4.0 Further Details on Nonconformances for MP-2024-01

All nonconformances must be categorized as either a minor error, nonconformance, or nonconformance due to CAB inaction or delay not attributable to the Participating Manufacturer. For Level 2 Investigations, nonconformances may be categorized as minor errors if a GEC-approved CAB is unable to obtain a product from the market and the Participating Manufacturer indicated the product has reached end-of-life and is no

longer available on the market. All nonconformances that do not meet the definition of minor errors are categorized as nonconformances (unless they are due to CAB inaction or delay).

No nonconformances or minor errors were identified in this Round.

5.0 Actions to Restore Conformance

Where the final conformity decision is nonconformance (including minor errors and those due to CAB inaction or delay), Participating Manufacturers must make corrections to restore the accuracy of the EPEAT Registry during the Corrective Action Phase. These activities may include providing additional evidence to demonstrate conformance with the criterion or unselecting the criteria in the EPEAT Registry. Where the product was found nonconformant and is no longer available in the marketplace, the product must be archived.

During the Corrective Action Phase, Participating Manufacturers must also develop Corrective Action Plans for other EPEAT-registered products that may be affected by the same underlying issue causing the nonconformance but were not the subject of investigation (called “similarly affected products”).

Since no nonconformances or minor errors were identified in this Round, no corrective actions were taken.

6.0 Key Findings

6.1 Scope of Criterion 12.2.1: Separability and labeling of plastics in packaging

The first part of required criterion 12.2.1 – Separability and labeling of plastics in packaging, applies to all packaging components ≥ 25 g. Any packaging component over this weight threshold must be identified unless exempted by the criterion (tape or labels affixed to plastic bags or wraps, staples, and top sheet adhered to chipboard, corrugate or other paperboard, or films, coatings or adhesives).

6.2 Criterion 11.4.1 Ease of disassembling mobile phone

Participating Manufacturers are reminded that 11.4.1 requires the criterion to be met without causing functional damage that would preclude re-use or refurbishment of the mobile phone.

Document Control and Change History						
Issue	Revision	Owner	Approver	Description	Approval Date	Effective Date
1	0	EPEAT Conformity Assurance Manager	Director, EPEAT Program	Initial release		
1	1	EPEAT Conformity Assurance Manager	Director, EPEAT Program		2018 Dec 11	2018 Dec 11
2	0	Senior Manager, Ecolabels and Resources	Senior Director, Ecolabels and Manufacturer Resources	Reformatting of document. Addition of standardized text.	2021 Mar 25	2021 Mar 30
2	1	Senior Manager, Ecolabels and Resources	Vice President, Ecolabels and Manufacturer Resources	Updated terminology for nonconformances to include "nonconformances" and "minor errors", in alignment with revisions to P66.	2022 Sep 15	2022 Sep 30
2	2	Senior Manager, Ecolabels and Resources	Vice President, Ecolabels and Manufacturer Resources	Updated to reflect new nonconformance category for CAB inaction or delay	2023 Mar 24	2023 Mar 24