CERTIFICATE of EXAMINATION

Certificate Validation

NOTIFIED BODY EU-TYPE EXAMINATION CERTIFICATE

AGCC3904-EU / *17 Mar 2025* / Rev A

Electromagnetic Compatibility Directive 2014/30/EU

MiCOM Labs Inc., Notified Body Number 2280 declares, on the basis of the assessment of the tests and the technical documentation provided by the applicant that the following product complies with the essential requirements of the above noted Directive.

Product Name:
High Fidelity Speaker

Approval Holder Name:
Zhuhai HiVi Technology Co., Ltd.

MICOM Labs, Inc. INCORPORATED JAN. 25, 2002 CALIFORNIA

CE

Gordon Hurst, Product Certifier

This Certificate is Issued under the Authority of:
MiCOM Labs Inc., 575 Boulder Court, Pleasanton, California 94566, USA
Notified Body Number: 2280

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Notified Body EU-Type Examination Certificate

AGCC3904-EU / *17 Mar 2025* / Rev A

for Electromagnetic Compatibility Directive 2014/30/EU

Product Name:

High Fidelity Speaker

Product Model Numbers: **Oasis 140**

Approval Holder: **Zhuhai HiVi Technology Co., Ltd.**, NO.1, South Dongcheng Road, Liangang Industrial Zone, Zhuhai, Guangdong, P.R. China

Product Manufacturer: **Zhuhai HiVi Technology Co., Ltd.**, NO.1, South Dongcheng Road, Liangang Industrial Zone, Zhuhai, Guangdong, P.R. China

Standards

Group	Name
EMC Directive: Disturbance(Annex 1.1.a)	EN 55032:2015/A1:2020
EMC Directive: Immunity(Annex 1.1.b)	EN 55035:2017/A11:2020

MiCOM Labs

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Annex 1 to EU-Type Examination

EU-Type examination on the essential requirements
Article 3

Annex 1.1.a - EMC Directive: Disturbance	Assessed
Annex 1.1.b - EMC Directive: Immunity	Assessed

Note 1: Article 3.4 Dates of Application
This aspect of the EU-Type Examination Certificate does not become legally applicable until the dates indicated below;-

28th December 2024 – Date of Applicability for handheld mobile phones, tablets, digital cameras, headphones, headsets, handheld videogame consoles,portable speakers, e-readers, keyboards, mice, portable navigation systems, earbuds.

28th April 2026 – Date of Applicability for laptops.

Description of Apparatus

Company Name	**Zhuhai HiVi Technology Co., Ltd.**
Certification No.	**AGCC3904-EU**
Issue Date / Rev	**17 Mar 2025 / Rev A**
Equipment Description	**High Fidelity Speaker**

Technical Construction File Details: (Documents Reviewed)

Technical Report(s):
EMC Directive: Disturbance(Annex 1.1.a):
AGC06182250223EE01
EMC Directive: Immunity(Annex 1.1.b):
AGC06182250223EE01

Supporting Documentation:
Service Agreement
Letter of Authorization
EU Application
EU Declaration of Conformity
Block Diagram
BOM or Parts List
External Photographs
Internal Photographs
Label and its Location
Operational Description
Risk Assessment Report
Schematics
Test Setup - EU
User Manual

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Notes

This device also satisfies standard EN IEC 62368-1: 2020+A11:2020, Report No.: AGC06182250223ES01

Scope

This EU-Type Examination Certificate is given in with respect to the Electromagnetic Compatibility Directive 2014/30/EU. EU Type Examination was performed according to Module B: EUtype examination procedure per Annex III the Directive on the essential requirements in Annex I, for the specific product and Certificate Number referenced above.

This EU Type Examination Certificate is based upon the review of the Technical Documentation and supporting evidence for the adequacy of the technical design solution, it is only valid in conjunction with the attached Annexes. The scope of this statement relates to a single sample of the apparatus identified above and of the submitted documents only.

Annex 2 to EU-Type Examination
Obligations of the Applicant

Ref EMCD 2014/30/EU Article 7
Obligations of manufacturers

1. When placing their apparatus on the market, manufacturers shall ensure that they have been designed and manufactured in accordance with the essential requirements set out in Annex I.

2. Manufacturers shall draw up the technical documentation referred to in Annex II or Annex III and carry out the relevant conformity assessment procedure referred to in Article 14 or have it carried out.

Where compliance of apparatus with the applicable requirements has been demonstrated by that procedure, manufacturers shall draw up an EU declaration of conformity and affix the CE marking.

3. Manufacturers shall keep the technical documentation and the EU declaration of conformity for 10 years after the apparatus has been placed on the market.

4. Manufacturers shall ensure that procedures are in place for series production to remain in conformity with this Directive. Changes in apparatus design or characteristics and changes in the harmonised standards or in other technical specifications by reference to which conformity of apparatus is declared shall be adequately taken into account.

5. Manufacturers shall ensure that apparatus which they have placed on the market bear a type, batch or serial number or other element allowing their identification, or, where the size or nature of the apparatus does not allow it, that the required information is provided on the packaging or in a document accompanying the apparatus.

6. Manufacturers shall indicate, on the apparatus, their name, registered trade name or registered trade mark and the postal address at which they can be contacted or, where that is not possible, on its packaging or in a document accompanying the apparatus. The address shall indicate a single point at which the manufacturer can be contacted. The contact details shall be in a language easily understood by end-users and market surveillance authorities.

7. Manufacturers shall ensure that the apparatus is accompanied by instructions and the information referred to in Article 18 in a language which can be easily understood by consumers and other end-users, as determined by the Member State concerned. Such instructions and information, as well as any labelling, shall be clear, understandable and intelligible.

8. Manufacturers who consider or have reason to believe that an apparatus which they have placed on the market is not in conformity with this Directive shall immediately take the corrective measures necessary to bring that apparatus into conformity, to withdraw it or recall it, if appropriate. Furthermore, where the apparatus presents a risk, manufacturers shall immediately inform the competent national authorities of the Member States in which they made the apparatus available on the market to that effect, giving details, in particular, of the non-compliance and of any corrective measures taken.

9. Manufacturers shall, further to a reasoned request from a competent national authority, provide it with all the information and documentation in paper or electronic form necessary to demonstrate the conformity of the apparatus with this Directive, in a language which can be easily understood by that authority. They shall cooperate with that authority, at its request, on any action taken to eliminate the risks posed by apparatus which they have placed on the market.

Ref EMCD 2014/30/EU Article 8
Authorised representatives

1. A manufacturer may, by a written mandate, appoint an authorised representative.

The obligations laid down in Article 7(1) and the obligation to draw up technical documentation referred to in Article 7(2) shall not form part of the authorised representative's mandate.

2. An authorised representative shall perform the tasks specified in the mandate received from the manufacturer. The mandate shall allow the authorised representative to do at least the following:

(a) keep the EU declaration of conformity and the technical documentation at the disposal of national market surveillance authorities for 10 years after the apparatus has been placed on the market;

(b) further to a reasoned request from a competent national authority, provide that authority with all the information and documentation necessary to demonstrate the conformity of the apparatus;

(c) cooperate with the competent national authorities, at their request, on any action taken to eliminate the risks posed by the apparatus covered by the authorised representative's mandate.

Ref EMCD 2014/30/EU Article 16
General Principles of the CE Marking

The CE marking shall be subject to the general principles set out in Article 30 of Regulation (EC) No 765/2008.

Ref EMCD 2014/30/EU Article 17
Rules and Conditions for affixing the CE Marking

1. The CE marking shall be affixed visibly, legibly and indelibly to the apparatus or to its data plate. Where that is not possible or not warranted on account of the nature of the apparatus, it shall be affixed to the packaging and to the accompanying documents.

2. The CE marking shall be affixed before the apparatus is placed on the market.

3. Member States shall build upon existing mechanisms to ensure correct application of the regime governing the CE marking and shall take appropriate action in the event of improper use of that marking.