



Brussels, 15.5.2024  
C(2024) 3082 final

**COMMISSION DECISION**

**of 15.5.2024**

**appointing members of the Committee for Orphan Medicinal Products to represent  
patients' organisations**

# COMMISSION DECISION

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## appointing members of the Committee for Orphan Medicinal Products to represent patients' organisations

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to [Regulation (EC) No 141/2000 of the European Parliament and of the Council of 16 December 1999 on orphan medicinal products]<sup>1</sup>, and in particular Article 4(3) thereof,

Whereas:

- (1) Article 4(3) of the Regulation (EC) No 141/2000 requires that the Commission appoints representatives of patients' organisations for a mandate of three years.
- (2) The term of office of the three members representing patients' organisations expires on 30 June 2024.
- (3) The members of the Committee should be appointed for a period of three years starting 1 July 2024.
- (4) In accordance with Article 4(3) of Regulation (EC) No 141/2000, a new public call for expression of interest has been undertaken by the Commission.
- (5) Where a member appointed by this decision is no longer capable of contributing efficiently to the work of the Committee, or resigns, the Commission may replace this member from the reserve list, for the duration of the member's mandate.

HAS DECIDED AS FOLLOWS:

### *Article 1*

The following are hereby appointed members of the Committee for a term of three years, from 1 July 2024:

- ISLA GOMEZ Julian, representing EURORDIS-Rare Diseases Europe,
- ALVEZ Ines, representing ANDO Portugal – The National Association for Skeletal Dysplasias,
- DRIESENS Mariette, representing EURORDIS-Rare Diseases Europe.

### *Article 2*

The following are hereby placed on a reserve list:

- NGUYEN Gerard, representing RSE – Rett Syndrome Europe and AFSR – Association Française du Syndrome de Rett,

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<sup>1</sup> OJ L 18, 22.1.2000, p. 1.

- CAVALLER Maria, representing EURORDIS-Rare Diseases Europe,
- TODOROVA-YANAKIEVA Yoana, representing RARE DISEASES BULGARIA.

Done at Brussels, 15.5.2024

*For the Commission*  
*Stella KYRIAKIDES*  
*Member of the Commission*