11 November 2021
EMA/629749/2021
Human Medicines Division

Statement indicating compliance with the agreed completed paediatric investigation plan

Medicinal product	
ADCETRIS (brentuximab vedotin)	
Pharmaceutical form:	See Annex A of the CHMP Opinion
Strength(s):	See Annex A
Route(s) of administration:	See Annex A
Packaging and package size(s):	See Annex A
Number(s)in the Community Register of Medicinal Products:	See Annex A

Marketing Authorisation Holder (MAH):	
Name and address of the MAH:	Takeda Pharma A/S Delta Park 45 2665 Vallensbaek Strand DENMARK

Procedure	
Procedure number:	EMEA/H/C/002455/II/0093

Further to the compliance check performed under Article 23 of Regulation (EC) No 1901/2006, and in accordance with Article 28(3) and Article 45(3) of the said Regulation it is concluded that:

- the development of this product has complied with all measures in the agreed paediatric investigation plan P/0013/2021. All studies in the agreed paediatric investigation plan P/0013/2021 were conducted after the entry into force of that Regulation and

- the Summary of Product Characteristics reflects the results of studies conducted in compliance with this agreed paediatric investigation plan.