Section B CHED-P (for products referred to in point (b) of Article 47(1) of Regulation (EU) 2017/625)

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Common Health Entry Document for Products							PART I – DESCRIPTION OF CONSIGNMENT						
	I.2 CHED reference				I.1 Consignor/Exporter								
		I.3 Loc		Name									
	QR CODE	I.4 Boi		Address									
		I.5 Boi	code	de									
						Country ISO country code					/ code		
I.6	Consignee/Importer Name Address					1.7					Registration/Approval No		
	Country I	SO country co	de				Country ISO			ISO country code			
1.8	Operator respon	nsible for the	consi	gnment		1.9	, , , ,						
	Name Address					Type Code Country							
	Country	ISO count				Commercial document references							
I.10	Prior notification	n	D	ate				Time					
1.13	Means of transp	ort					I.11 Country of origin ISO count				ISO country code		
	□Airplane □	Vessel						I.12	Region of or	igin	Code		
		□Railway □Road Identification											
1.14	Country of dispa	atch		l.15 Es	tablishm	nent of a	origin						
	Country			Na	me Registration/Approval No					al No			
	ISO country code)		Ac	dress Country			Country		ISO country code			
I.16	Transport cond	itions	□Ar	nbient		□Chilled			□Frozen				
1.17	Container numb	er/Seal Numl	ber										
	Container No		Seal	No						Of	ficial Seal		
I.18	Certified as or fe	or:				I.19 Conformity of the goods				f the goods			
	Human consumption DPharmaceutical			□Trade sample □Other									
	□Feedstuff use □Technical use			□Further process			Conforming Non-conforming						
1.20					[Details of controlled destinations for I.20 to I.22 and I.24							
1.22	□For transit to:	:											
1.24	□For non-conforming goods												
	□Specially approved customs warehouse												
	□Free zone												
	□Vessel												
1.23	I.23 DFor Internal market				1.25	□For i	re-entry						
I.27 Means of transport after BCP/storage													
DAirplane DR						□Ra	ilway						
□Vessel					□Road vehicle								
Identification:													

1.29	Date of dep	arture		Date		Time					
1.31	Description of consignment										
	CN code Species		Batch Number Quantity		No of packages	Net weight(kg)	IAS Permit	Final consumer			
1.32	Total number of packages I.33			tal quantity	1.34	Total net weight	Total net weight/gross weight				
1.35	Declaration: I, the undersigned operator responsible for the consignment detailed above, certify that to the best of my knowledge and belief the statements made in Part I of this document are true and complete, and I agree to comply with the requirements of Regulation (EU) 2017/625 on official controls, including payment for official controls, as well as for re-dispatching of consignments, for quarantine or isolation of animals, or costs of euthanasia and disposal where necessary.										
	Date of declaration			ame of signator	у	Signature					

In processing the personal data included in the CHEDs, Member States shall comply with Regulation (EU) 2016/679 and Directive (EU) 2016/680 and the Commission with Regulation (EU) 2018/1725.

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PART II - CONTROLS **Common Health Entry Document for Products** II.1 **Previous CHED** 11.2 CHED reference 11.24 Subsequent CHED Identity check 11.3 Documentary 11.4 □Yes □No check EU requirements □Satisfactory □Not satisfactory DSeal check □Not □Satisfactory National satisfactory □Satisfactory □Not satisfactory Even Full check requirements Laboratory □No □No □Yes 11.5 Physical check □Yes 11.6 test Test: □Intensified controls Required □Random □ Reduced check Emergency measures □Satisfactory □Not satisfactory Suspicion □Not □Others Test result: □Pending □Satisfactory satisfactory Acceptable for (II.9 to II.16): II.13 □Monitoring 11.9 □Transhipment to II.11 □Transit to: □Entry monitoring □Re-entry monitoring □Non-conforming II.12 II.14 II.16 □Internal market □Not acceptable goods □Trade □Human □Specially approved Destruction By (date) consumption sample customs □Feedstuff □Other □Re-dispatch warehouse □ Pharmaceutical Local use □Free zone □Special treatment use □Further □Use for other □Technical use Vessel process purposes II.18 Details of controlled destinations for II.9 to II.16 II.17 Reason for refusal □Physical Documentary □ldentity □Origin □Laboratory □IAS □Other II.19 New seal number □Consignment resealed 11.20 Identification of BCP 11.21 Certifying officer BCP Stamp I, the undersigned official veterinarian, certify that the checks on the consignment have been carried out in accordance with the Union requirements and where applicable in accordance with the national Control Unit code requirements of the Member State of destination 11.22 Inspection fees Name (in capital letters) Date Signature 11.23 Customs document reference

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Common Health Entry Document for Products

PART III - FOLLOW-UP

III.1	Previous CHED	III.2	CHED reference		III.3	Subsequent CHED
III.4	Details on re-dispatch					
	Country of destination		ISO country Code			
	Exit BCP		Control Unit code			
	Means of transport					
	□Airplane □Road □Vessel □Other □Railway	Vehicle	Identification			
	Date of re-dispatch					
III.5	Follow up by					
	□Exit		Arrival of consignment:	□Yes		No
	□Final destination □Local competent authority	BCP	Compliance of consignment:	□Yes		No
			Further destination:		R	easons
III.6	Certifying officer					
	Name (in capital letters)				U	nit name
	Address				C	ontrol Unit code
	Date		Stamp	Si	gnature	