

Ref: VIG-07-2017-01

Lisses, on 19 July 2017

To the attention of: Directors of Health Establishments
Persons in charge of Laboratories
Local Correspondents of Vigilance

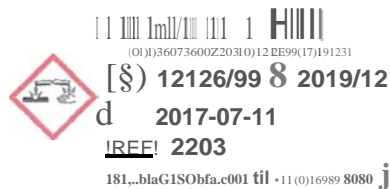
RE: **INFORMATION/ RECOMMENDATION**
MINICAP PROTIEN(E) 6 (product number 2203)
Batch number : 05096/02
Expiry date : 2019/09

Dear Sir/ Madam,

Users have demonstrated that in few kits the batch of buffer is not the expected one: lot 12126/01 instead of 05096/01. Our investigations revealed a human error in the packaging of these kits. This inversion does not affect the quality or functionality of these kits.

Our traceability indicates that you are a user of this batch number, therefore, we kindly ask you to:

1. Check that in the kits of lot number 05096/02 which are in your possession, the lot number of the buffer vials is 05096/01.
2. Fill-in the enclosed form and indicate precisely the number of non compliant kits in your possession. At your request we will send you the label to correctly identify the kits concerned.



The French Health Products Safety Agency (ANSM) has been informed about this communication.

Please do not hesitate to call your local SEBIA contact for further information.

We apologize for any inconvenience caused and we thank you for your confidence in Sebia.

Yours sincerely,

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INFORMATION CERTIFICATE
MINICAP PROTEIN(E) 6 (product number 2203)
Batch Number: 05096/02
Expiry date: 2019109

Please fill out this document and return it to us upon reception

Laboratory Stamp (mandatory)

We certify, Madam, Mister

To have taken knowledge of the mail « VIG-07-2017-01 ».

To have checked the concerned kit(s) at receipt of this mail.

Number of non compliant kits:

Request the identification label for the kits concerned YES NO

(Place) _____, (Date) _____

Signature:

SEBIA Pare technologique Leonard de Vinci CP 8010 91008 EVRY CEDEX FRANCE	Tél.: + 33 1 69 89 80 80 Fax: + 3316989 78 78 E-mail: sebia@sebia.com
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