

Déclaration de Conformité Declaration of Conformity

(No. dc200066)

Selon
According to ISO 17050-1

NOUS LE FABRICANT WE THE MANUFACTURER

Nom <i>Name</i>	HORIBA ABX SAS
Adresse <i>Address</i>	Parc Euromédecine Rue du Caducée – BP7290 34184 MONTPELLIER Cedex 4 - FRANCE

**ETABLISSEMENTS SOUS NOTRE SEULE RESPONSABILITE LA DECLARATION SUIVANTE ET
DECLARONS QUE LE PRODUIT
ESTABLISH UNDER OUR ONLY RESPONSIBILITY THE FOLLOWING DECLARATION AND
DECLARE THAT THE PRODUCT**

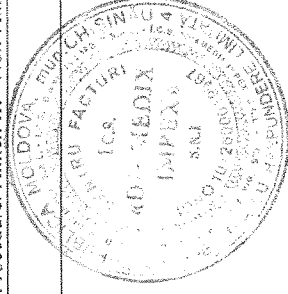
Catégorie du dispositif <i>Category device</i>	HEMATOLOGY REAGENT
Nom du produit <i>Product name</i>	ABX DILUENT (20L) / ABX DILUENT (10L) For application on : ABX Pentra 60, ABX Pentra 60C+, Pentra ES60, Pentra MS60, Pentra MS CRP, ABX Pentra 80, ABX Pentra XL80, Pentra XLR, ABX Pentra 120, ABX Pentra 120 Reite, ABX Pentra DX120, ABX Pentra DF120, Pentra DX Nexus, Pentra DF Nexus, Yumizen H500 OT, Yumizen H500 CT, Yumizen H1500, Yumizen H2500, Yumizen H550
Modèles <i>Models</i>	0901020 / 0901010
Pays d'origine <i>Country of origin</i>	France

**EST CONFORME AUX DIRECTIVES ET NORMES
CONFORMS TO DIRECTIVES AND STANDARDS**

Directives <i>Directives</i>	98/79/EC – IVD Medical Devices Classification : General IVD (others) - Outside Annex II and not for self-testing Conformity Assessment Procedure: Annex III / Non-Annex II
Normes <i>Standards</i>	N/A

Montpellier, France
06 Juin 2017
June 06th, 2017

Christian DUBUC
Directeur
Senior Director



TEMP-4018 Rev.12

HORIBA

M E D I C A L

HORIBA ABX SAS

Parc Euromédecine

Rue du Caducée - BP7290

34184 MONTPELLIER Cedex 4 - France

Web : www.horiba.com

Tél. : 33 (0) 4 67 14 15 16

Déclaration de Conformité Declaration of Conformity

(No. dc20014/s)

DECLARATION

Selon

According to ISO 17050-1

**NOUS LE FABRICANT
WE THE MANUFACTURER**

Nom <i>Name</i>	HORIBA ABX SAS
Adresse <i>Address</i>	Parc Euromédecine Rue du Caducée - BP7290 34184 MONTPELLIER Cedex 4 - FRANCE

**DECLARONS QUE LE PRODUIT
DECLARE THAT THE PRODUCT**

Nom du produit <i>Product name</i>	ABX LYSEBIO (0.4L) For application on : ABX Pentra 60, ABX Pentra 60 C+, Pentra ES60, Pentra MS60, Pentra MS CRP, ABX Pentra 80, ABX Pentra XL 80, Pentra XLR, Microsemi CRP ABX LYSEBIO (1L), For application on : ABX Pentra 80, ABX Pentra XL 80, ABX Pentra 120, ABX Pentra 120 RETIC, ABX Pentra DX 120, ABX Pentra DF 120, Microsemi CRP, Pentra DX Nexus, Pentra DF Nexus, Pentra XLR, Yumizen III 500, Yumizen H2500
Modèles <i>Models</i>	0906013 0906012
Pays d'origine <i>Country of origin</i>	France

**EST CONFORME AUX DIRECTIVES ET NORMES
CONFORMS TO DIRECTIVES AND STANDARDS**

Directives <i>Directives</i>	98/79/EC - IVD Medical Devices Classification : General IVD (others) - Outside Annex II and not for self-testing Conformity Assessment Procedure Annex III / Non-Annex II
Normes <i>Standards</i>	N/A

Montpellier, France

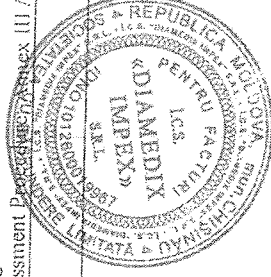
05 Avril 2017

April 05th, 2017

Christian DUBUC

Directeur

Senior Director



TE-MP-4018 Rev.9

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Déclaration de Conformité Declaration of Conformity

(No.de20017k)

Selon
According to ISO 17050-1

**NOUS LE FABRICANT
WE THE MANUFACTURER**

Nom Name	HORIBA ABX SAS
Adresse Address	Parc Euromédecine Rue du Caducée – BP7290 34184 MONTPELLIER Cedex 4 - FRANCE

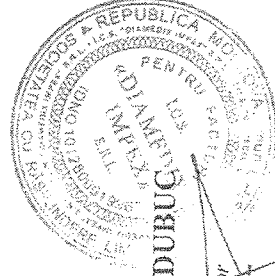
**DECLARONS QUE LE PRODUIT
DECLARE THAT THE PRODUCT**

Nom du produit Product name	ABX EOSINOFIX (IL) For application on : ABX Pentra 60, ABX Pentra 60 C+, Pentra ES60, Pentra MS60, Pentra MS CRP, ABX Pentra 80, ABX Pentra XL 80, Pentra XLR, ABX Pentra 120, ABX Pentra 120 Rette
Modèles Models	0206010
Pays d'origine Country of origin	France

**EST CONFORME AUX DIRECTIVES ET NORMES
CONFORMS TO DIRECTIVES AND STANDARDS**

Directives Directives	98/79/EC – IVD Medical Devices Classification : General IVD (others) - Outside Annex II and not for self-testing Conformity Assessment Procedure: Annex III / Non-Annex II
Normes Standards	N/A

Montpellier, France
03 Février 2016
February 03rd, 2016



Christian DUBUC
Directeur
Senior Director

TEMP-4018 Rev.9

Déclaration de Conformité Declaration of Conformity

(No.de200161)

Selon
 According to ISO 17050-1

**NOUS LE FABRICANT
 WE THE MANUFACTURER**

Nom <i>Name</i>	HORIBA ABX SAS
Adresse <i>Address</i>	Parc Euromédecine Rue du Caducée – BP7290 34184 MONTPELLIER Cedex 4 - FRANCE

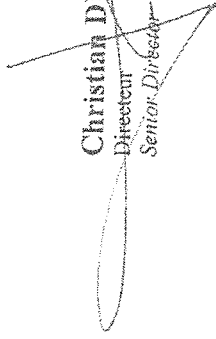
**DECLARONS QUE LE PRODUIT
 DECLARE THAT THE PRODUCT**

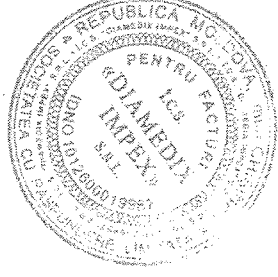
Nom du produit <i>Product name</i>	ABX BASOLYSE II (IL) For application on : ABX Pentra 60, ABX Pentra 69C [®] , Pentra ES 60, Pentra MS 60, Pentra MS CRP, ABX Pentra 80, ABX Pentra XL 80, Pentra XLK
Modèles <i>Models</i>	0906003
Pays d'origine <i>Country of origin</i>	France

**EST CONFORME AUX DIRECTIVES ET NORMES
 CONFORMS TO DIRECTIVES AND STANDARDS**

Directives <i>Directives</i>	98/79/EC – IVD Medical Devices Classification : General IVD (others) - Outside Annex II and not for self-testing Conformity Assessment Procedure: Annex III / Not-Annex II
Normes <i>Standards</i>	N/A

Montpellier, France
 03 Février 2016
 February 03rd, 2016


Christian DURIEZ
 Directeur
 Senior Director



TEMP-4018 Rev.9

HORIBA

Medical

HORIBA ABX SAS

Parc Euromédecine

Rue du Caducée – BP7290

34184 MONTPELLIER Cedex 4 – France

Web : www.horiba.com

Tél. : 33 (0) 4 67 14 15 16

Déclaration de Conformité Declaration of Conformity

(No.de20025o)

Selon
According to ISO 17050-1

**NOUS LE FABRICANT
WE THE MANUFACTURER**

Nom <i>Name</i>	HORIBA ABX SAS
Adresse <i>Address</i>	Parc Euromédecine Rue du Caducée – BP7290 34184 MONTPELLIER Cedex 4 - FRANCE

**ETABLISSEONS SOUS NOTRE SEULE RESPONSABILITE LA DECLARATION SUIVANTE ET
DECLARONS QUE LE PRODUIT
ESTABLISH UNDER OUR ONLY RESPONSIBILITY THE FOLLOWING DECLARATION AND
DECLARE THAT THE PRODUCT**

Catégorie du dispositif <i>Device category</i>	HEMATOLOGY REAGENT ABX CLEANER (IL) For application on : ABX Pentra 60, ABX Pentra 60 C+, Pentra ES 60, Pentra MS 60, Pentra MS CRP, ABX Pentra 80, ABX Pentra XL 80, Pentra XLR, ABX Pentra 120, ABX Pentra 120 Retic, ABX Pentra DX 120, ABX Pentra DF 120, Pentra DX Nexus, Pentra DF Nexus, Yumizen H500 OT, Yumizen H500 CT, Yumizen H1500, Yumizen H2500, Yumizen H550 ABX CLEANER (0.5L) For application on : ABX Micros ES 60, ABX Micros CRP, ABX Micros CRP 200
Modèles <i>Models</i>	0903010, 0903011
Pays d'origine <i>Country of origin</i>	France

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Directives <i>Directives</i>	98/79/EC – IVD Medical Devices Classification : General IVD (others) - Outside Annex II and not for self-testing Conformity Assessment Procedure: Annex III / Non-Annex II
Normes <i>Standards</i>	N/A

Montpellier, France

06 Juin 2017

June 06th, 2017

Christian DUBUC

Directeur
Senior Director

TEMP-4018 Rev.12

