

## DECLARATION OF CONFORMITY

Diamond Diagnostics, Inc. hereby ensures and declares that the product(s) listed below comply with the requirement of In Vitro Diagnostics Medical Devices Regulation EU 2017/746 and Directive 2011/65/EU.

Diamond Diagnostics Inc. ezúton biztosítja és kijelenti, hogy az alább felsorolt termék(ek) megfelelnek az In Vitro Diagnostics Medical Devices EU 2017/46 rendelet és a 2011/65/EU irányelv követelményeinek.

Diamond Diagnostics Inc. versichert und erklärt hiermit, dass das/die unten aufgeführte(n) Produkt(e) den Anforderungen der In-vitro-Diagnostik-Medizinprodukte-Verordnung EU 2017/46 und der Richtlinie 2011/65/EU entsprechen.

Diamond Diagnostics Inc. garantit et déclare par la présente que le ou les produits répertoriés ci-dessous sont conformes aux exigences du règlement sur les dispositifs médicaux de diagnostic in vitro UE 2017/46 et de la directive 2011/65/EU.

Diamond Diagnostics Inc. por la presente garantiza y declara que los productos enumerados a continuación cumplen con los requisitos del Reglamento de dispositivos médicos de diagnóstico in vitro EU 2017/46 y la Directiva 2011/65/EU.

Diamond Diagnostics Inc. 特此确保并声明下列产品符合体外诊断医疗器械法规 EU 2017/46 和指令 2011/65/EU 的要求。

Diamond Diagnostics Inc. garante e declara que o(s) produto(s) listado(s) abaixo cumpre(m) o requisito do Regulamento de Dispositivos Médicos de Diagnóstico In Vitro EU 2017/46 e Diretiva 2011/65/EU.

Diamond Diagnostics Inc. гарантирует и заявляет, что продукты, перечисленные ниже, соответствуют требованиям Регламента ЕС 2017/46 о медицинских устройствах для диагностики in vitro и Директивы 2011/65/EU.

Vitro Diagnostica Medical Regulation 2017/746 and Directive 2011/65/EU المنتجات المذكورة أدناه تتوافق مع متطلبات الاتحاد الأوروبي المدرجة في ن شركة دايموند داياغنوستكس تصرح و تؤكد أن التعلیمة

Diamond Diagnostics Inc. garantisce e dichiara che i prodotti elencati di seguito sono conformi ai requisiti del Regolamento UE 2017/46 sui dispositivi medici per la diagnostica in vitro e della Direttiva 2011/65/EU.

Diamond Diagnostics, Inc. işbu belge ile aşağıda listelenen ürün(ler)in In Vitro Diagnostics Tıbbi Cihazlar Yönetmeliği EU 2017/746 ve Direktif 2011/65/EU gerekliliklerine uygun olduğunu garanti ve beyan eder.

**Product(s) / Produkt(e) / Produit(s) / Producto(s) / 产品 (S) / Produto(s) / Продукт (ы) / المنتج (ق) / Prodott(i) / Ürün :**

**Model: Horiba ABX Micros, Horiba ABX Pentra, Siemens Advia 60, Beckman Coulter ACT 5 Diff**

### Electrodes & Accessories

Part Number	Device Name:	Basic UDI-DI	Intended Use
HM-XEA328ASD	Maintenance Kit	081140301HMSEA328ASDHN	For use with Horiba ABX Micros, Horiba ABX Pentra, Siemens Advia 60, Beckman Coulter ACT 5 Diff
HM-XEA486ORD	O-Ring Kit	081140301HMSEA486ORDMS	For use with Horiba ABX Micros, Horiba ABX Pentra, Siemens Advia 60, Beckman Coulter ACT 5 Diff

**Manufacturer's Name:**

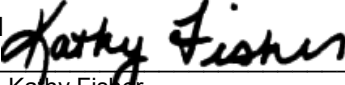
Diamond Diagnostics Inc.

**Manufacturer's Address:**

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1044 Budapest, Hungary  
Tel: + 3617872222

**(AR) Authorized Representative:** Diamond Diagnostics Inc., Hungary Branch Office  
**(SRN) Single Registration Number:** US-MF-000035435  
**Risk Class:** Class A, Rule 5  
**Conformity Route:** Annex IV, Self-Declared  
**Notified Body:** N/A  
**Quality Systems Registration:** ISO 9001:2015  
ISO 13485:2016  
EN ISO 9001:2015  
EN ISO 13485:2016

**Authorized Officer:**  **Date:** 6 May, 2024  
Kathy Fisher  
Director, Quality Assurance

