



## Declaration of Conformity

**Beckman Coulter Inc. hereby ensures and declares that the product(s) listed below comply with the requirement of the European Union In-vitro Diagnostics Medical Device Directive 98/79/EC.**

*Beckman Coulter Inc. assure et déclare par la présente que le(s) produit(s) listé(s) ci-dessous sont conformes aux exigences de la directive européenne 98/79/CE relative aux dispositifs médicaux de diagnostic in vitro.*

*Beckman Coulter Inc. dichiara ed assicura che i prodotti qui elencati sono conformi ai requisiti della direttiva comunitaria 98/79/CE relative ai dispositivi medico-diagnostici in vitro.*

*Beckman Coulter Inc. versichert und erklärt hiermit, daß die im Folgenden aufgeführten Produkte den Auflagen der IVD-Richtlinie für In-vitro-Diagnostika der Europäischen Union (98/79/EC) entsprechen.*

*Beckman Coulter Inc. asegura y declara que los productos listados a continuación cumplen con los requisitos establecidos en la directiva 98/79/EC de la Comunidad Europea para dispositivos médicos de diagnóstico in vitro.*

**Product(s) /Produkt(e) /Prodotto(i) / Produit(s) / Producto(s):**

*MicroScan Pos ID Panel Type 3, B1017-221*

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**Conformity Assessment Procedure**  
*Annex III, Self-Declared*

**Classification**  
*General*

2015-03-06  
Date

*Robert Eusebio*  
*Director Regulatory Affairs*



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