



## DECLARATION OF CONFORMITY

**Immunotech a.s. hereby ensures and declares that the product(s) listed below comply with the requirements of the European Union In Vitro Diagnostic Medical Device Directive 98/79/EC.**

Immunotech a.s. 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.

Immunotech a.s. dichiara ed assicura che i prodotti qui elencati sono conformi ai requisiti della direttiva comunitaria 98/79/CE relativa ai dispositivi medico-diagnostici in vitro.

Immunotech a.s. 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.

Immunotech a.s. 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):**

**Total Thyroxine RIA, Part Number IM1447  
Total Thyroxine RIA, Part Number IM3286**

Petr Šmídl  
Quality Assurance and  
Regulatory Affairs Manager

**Date:** September 15, 2004

**Conformity Assessment Procedure**  
Annex III



Immunotech a.s.  
Radiová 1  
102 27 Praha 10, Czech Republic  
Tel.: +420 267 008 444 Fax: +420 267 008 385

**Document Control**  
Issue Date: May 2, 2004  
Revision Lev: 3.0  
Revision Date: September 15, 2004  
File Name: IM1447(3286) DoC-TT4 RIA-rev3.0