



**Client Claim Form**

<b>Date:</b>		<b>Number:</b>	
<b>TYPE</b>	Part to be completed by the Client	Client: _____ Address: _____	Reference: _____
		Country and Department N°: _____	Distribution product concerned: _____
<b>DESCRIPTION</b>	Part reserved for BIOTECH DENTAL	Series: _____	Quantity: _____
		Batch Number: _____	
<b>INVESTIGATION</b>	Part reserved for BIOTECH DENTAL	Client Claim Description	
		Investigation description: _____	
<b>MEDICAL DEVICE VIGILANCE REPORT</b>	Part reserved for BIOTECH DENTAL	CC type: _____	CC occurrence: _____
		Identified risk code : _____	
<b>CURATIVE ACTION</b>	Part reserved for BIOTECH DENTAL	Serious or not serious : _____	Materiovigilance Classification: _____
		Medical device vigilance report (yes/no) : _____	
<b>CAPA</b>	Part reserved for BIOTECH DENTAL	Person responsible for the statement: _____	Date of the statement: _____
		Product quantity (if medical device vigilance report) : _____	
<b>CLOSING DATE</b>	Part reserved for BIOTECH DENTAL	Reasons for the decision : _____	
		Person responsible for medical device vigilance : _____	Authorisation : _____
		Description of the CURATIVE ACTION	
		Client Information fulfilled: _____	Exchange : _____ DS number: _____
		Sales Manager: _____	Date of discussion: _____
		Opening of a corrective/preventive action yes/no : _____	
		Justification: _____	
		Person responsible for CAPA: _____	
		Closing date of customer complaint : _____	
		Name of person in charge of managing the complaint : _____	Authorisation : _____