

| Client Claim Form | | | | |
|--|-----------------------------|--|-----------|---------------------|
| Date: | | | Number: | |
| ТҮРЕ | completed by the Client | Client: Address: | | |
| | | Country and Department N°: | Re | ference: |
| | | Series: Distribution product concerned: | | duct concerned: |
| | | Batch Number: | Quantity: | |
| DESCRIPTIO | Client Claim Description | | | |
| INVESTIGATION | | Investigation description: | | |
| CURATIVE MEDICAL DEVICE VIGILANCE INVEST | reserved for BIOTECH DENTAL | CC type: | | |
| | | CC occurrence: Identified risk code : | | |
| | | Serious or not serious : Materiovigilance Classification: | | |
| | | Medical device vigilance report (yes/no) : | | |
| | | Person responsible for the statement: Date of the statement: | | |
| | | Product quantity (if medical device vigilance report) : | | |
| | | Reasons for the decision : | | |
| | | Person responsible for medical device vigilance | : | Authorisation : |
| | | Description of the CURATIVE ACTION | | |
| | : rese | Client Information fulfilled: Exchange : | | DS number: |
| | Part | Sales Manager: | | Date of discussion: |
| CAPA | | Opening of a corrective/preventive action yes/ne Justification: | o : | |
| | | Person responsible for CAPA: | | |
| CLOSING | | Closing date of customer complaint : | | |
| D CLO | | Name of person in charge of managing the com | plaint : | Authorisation : |