

INTERNATIONAL
STANDARD

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**Implants for surgery — Femoral and tibial
components for partial and total knee joint
prostheses —**

Part 1:

Classification, definitions and designation of
dimensions

*Implants chirurgicaux — Éléments fémoral et tibial de prothèses partielle
et totale de l'articulation du genou —*

Partie 1: Classification, définitions et désignation des dimensions



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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

International Standard ISO 7207-1 was prepared by Technical Committee ISO/TC 150, *Implants for surgery*, Subcommittee SC 4, *Bone and joint replacements*.

This second edition cancels and replaces the first edition (ISO 7207-1:1985), of which it constitutes a technical revision.

ISO 7207 consists of the following parts, under the general title *Implants for surgery — Femoral and tibial components for partial and total knee joint prostheses*:

- Part 1: *Classification, definitions and designation of dimensions*
- Part 2: *Bearing surfaces made of metallic and plastics materials*

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Introduction

Partial and total knee joint prostheses are designed to transmit load and allow movement under high stress conditions. The task of preparing International Standards to cover all requirements is complicated by the limited range of biologically suitable materials.

The purpose of this part of ISO 7207 and of other International Standards relating to joint prostheses is to provide direction in the control of manufacture and standard specifications for the different components of prostheses.

Attention is drawn to ISO 5839, *Implants for surgery — Orthopaedic joint prostheses — Basic requirements*, in which the requirements for orthopaedic joint prostheses are specified.

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Implants for surgery — Femoral and tibial components for partial and total knee joint prostheses —

Part 1:

Classification, definitions and designation of dimensions

1 Scope

This part of ISO 7207 classifies femoral and tibial components for knee joint prostheses in which one or both bearing surfaces of at least one compartment of the knee are replaced, and gives definitions of components and the designation of dimensions.

Patellar components and prostheses that include an interposed floating component are not included in this part of ISO 7207.

2 Definitions

For the purposes of this part of ISO 7207, the following definitions apply.

2.1 partial unicompartamental knee joint component: Prosthesis designed to replace the bearing surface of one condyle of either the femur or the tibia (see figure 1, position A, B, C or D).

2.2 partial bicompartamental knee joint component: Prosthesis designed to replace the bearing surfaces of both condyles of either the femur or the tibia (see figure 1, positions A and B, or C and D).

NOTE 1 In surgical practice unicondylar components may be used either for unicondylar replacements, for unicompartamental replacements, or for total replacements, whereas generally bicondylar components are used only for total knee joint replacement.

2.3 unicompartamental partial knee joint replacement: Procedure of replacing the bearing surfaces of the contiguous condyles of the femur and tibia in one

compartment of the knee (see figure 1, positions A and C, or B and D).

2.4 total replacement of the knee joint: Procedure of replacing the femoral and tibial bearing surfaces in both compartments of the knee (see figure 1, positions A, B, C and D).

NOTE 2 Provision for articulation with the patella may or may not be provided.

2.5 non-constrained total knee joint prosthesis: Total knee joint prosthesis in which there is no mechanical attachment between the tibial and femoral components and which allows movement in all three planes (see figures 2, 3 and 4).

2.6 partially-constrained total knee joint prosthesis: Total knee joint prosthesis having some mechanical constraint between the tibial and femoral components and which allows movement in more than one plane (see figure 5).

2.7 fully-constrained total knee joint prosthesis: Total knee joint prosthesis in which the two parts are mechanically articulated and which allows movement principally in one plane (see figure 6).

2.8 effective bone resection distance: Minimum distance between resected surfaces of the femur and/or tibia in contact with the plateau(x) of the implant.

2.9 stem: Part of bicondylar component designed to enter the medullary cavity (see figures 2, 5 and 6, dimensions t and f).