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| **Form** |
| **Declaration of radiopharmaceuticals** |
| **Identification number:** | ZL000\_00\_029 |
| **Version:** | 1.2 |
| **Valid from:** | 29.06.2023 |

# Basic information

|  |  |
| --- | --- |
| Name of the medicinal product: | …… |
| Authorisation no.:*If known* | …… |
| Application ID:*If known* | …… |
| Date: | …… |

# Further information

|  |  |
| --- | --- |
| Do you possess an FOPH licence for the handling of radioactive medicinal products? | [ ]  Yes [ ]  No |
| If yes, licence number:*(please enclose a copy of the licence)* | …… |

|  |  |
| --- | --- |
| Indications: | …… |
| Mode of action / operation: | …… |

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| --- |
| **Composition** |
| Radionuclide | Active substance to be labelled [[1]](#footnote-1) or labelled molecule [[2]](#footnote-2) /Quantity per dosage form / molecular weight | Specific activity | Activity per dosage form |
| …… | * ……
 | …… | …… |

Change history

| **Version** | **Change** | **sig** |
| --- | --- | --- |
| 1.2 | New layout, no content adjustments to the previous version. | dei |
| 1.1 | Formal adjustments to the header and footerNo content adjustments to the previous version. | dei |
| 1.0 | Implementation of TPO4 | dts |

1. for labelling kits [↑](#footnote-ref-1)
2. for ready-to-use preparations with labelled molecules [↑](#footnote-ref-2)