| | Finished product | Qualifying process to be performed within the Area |
|---|---|---|
| ex 30.04 | Plasters, capable of adhering to themselves or to the skin, impregnated or coated with pharmaceutical substances or put up in retail packings for medical or surgical purposes | Manufacture from materials not falling in 30.04 or 40.06 |
| ex 30.05 | Sterile surgical catgut and similar sterile suture materials; sterile absorbable surgical haemostatics; dental cements and fillings | Manufacture from materials not falling in 30.05 |
| ex 30.05 | Opacifying preparations for X-ray examination and diagnostics reagents | Manufacture from materials not falling in 30.05, provided that all the active ingredients[1]), other than any listed in the Basic Materials List, have been made in the Area by chemical transformation[2]) or are of Area origin |
| | *Chapter 31. Fertilisers.* | |
| 31.02 | Mineral or chemical fertilisers, nitrogenous | Manufacture from natural sodium nitrate (ex 28.39 or ex 31.02) or from materials not falling in 28.30, 28.39, 29.25, 31.02 or 38.19 |
| 31.03 | Mineral or chemical fertilisers, phosphatic | Manufacture from materials not falling in 28.10, 28.40 or 31.03 |
| 31.04 | Mineral or chemical fertilisers, potassic | Manufacture from potassium chloride (ex 31.04) or crude natural potassium salts (ex 31.04) or from materials not falling in 31.04 |
| ex 31.05 | Other fertilisers; goods of the present Chapter in tablets, lozenges and similar prepared forms or in packings of a gross weight not exceeding ten kilogrammes (other than the fertilisers covered by the following item) | Manufacture from ammonium phosphates (ex 31.05) of Area origin or from materials not falling in 31.05, provided that any materials falling in 31.02 (other than natural sodium nitrate), 31.03, 38.11, 38.19 or 39.01 or in Chapter 28 or 29 are of Area origin |

[1]) An "active ingredient" means any substance which forms part of the finished product and in respect of which any therapeutic or prophylactic claim is made by the producer or exporter. The documentary evidence of origin relating to the goods must include a statement of the ingredients which are claimed to be active.

[2]) As defined in the Introductory Notes to the respective Chapters.