

NATIONAL PROCEDURE

Opublikowany na Urząd Rejestracji Produktów Leczniczych, Wyrobów Medycznych i Produktów Biobójczych (<https://archiwum.urpl.gov.pl>)

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If at least one active substance of the biocidal product is still in the review program in relevant PT and is not approved yet, then the biocidal product should be authorised in Poland according to the national procedure (during the transitional period). The fees for the registration under the national procedure are presented in the table below.

| Application Type | Total fee [PLN] |
|---|-----------------|
| application for authorisation for placing of a biocidal product on the market | 1 000 |
| application for data modification of the authorisation | 500 |
| application for change of the authorisation holder | 100 |

Źródłowy URL: <https://archiwum.urpl.gov.pl/pl/node/4286>