Procedure for the authorisation of the same biocidal products

**Submission of the application**

1. **Validation**
   - A check is made that the proposed differences between the "same product" and the reference product simply concern information that can be the subject of an administrative change.
   - Validation*

2. **Acceptance**

3. **Decision taken by the Member State**

4. **Authorisation**

5. **Refusal of authorisation**

6. **Validation**

7. **Rejection of the application**

8. **Payment period for the fee**

9. **If non-payment**

10. **30 d**

11. **60 d**

*If the reference product is being assessed, the validation date corresponds to the date of adoption of the authorisation decision for the reference product.