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1. Procedural advice for users of the Centralised Procedure for Similar Biological Medicinal Products applications (http://www.ema.europa.eu/docs/en_GB/document_library/Regulatory_and_procedural_guideline/2012/04/WC500125166.pdf)
2. European Commission:What you Need to Know about Biosimilar Medicinal Products (http://ec.europa.eu/DocsRoom/documents/8242/attachments

/1/translations/en/renditions/native)

3. EMA:Improving Understanding of Biosimilars in the EU (https://www.ema.europa.eu/en/news/improving-understanding-biosimilars-eu)

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