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Perivasc drug is suspended

Fábrica presented nonconformities during inspection to verify Good Manufacturing Practices.


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Anvisa ordered the suspension of the importation, distribution, marketing and use of the drug Perivasc (diosmin + hesperidin). The formulation is used to treat manifestations of chronic venous insufficiency, such as varicose veins, sequelae of thrombophlebitis and varicose ulcers.


The decision, contained in Resolution RE 1,592 / 2017 (<http://pesquisa.in.gov.br/imprensa/jsp/visualiza/index.jsp?data=16/06/2017&jornal=1&pagina=34&totalArquivos=72>) , published on Friday (16/6) in the Official Gazette of the Union, was based on non-conformities detected during inspection to verify Good Manufacturing Practices in the Spanish company Kern Pharma, The manufacturer of the medicinal product. The product is imported into Brazil by Eurofarma Laboratórios SA Therefore, the company must collect the products that still exist in the market.

Medicines (http://portal.anvisa.gov.br/noticias?p_p_id=101_INSTANCE_FXrpx9qY7FbU&p_p_lifecycle=0&p_p_state=normal&p_p_mode=view&p_p_col_id=column-2&p_p_col_count=1&p_r_p_564233524_tag=medicamentos) suspension (http://portal.anvisa.gov.br/noticias?p_p_id=101_INSTANCE_FXrpx9qY7FbU&p_p_lifecycle=0&p_p_state=normal&p_p_mode=view&p_p_col_id=column-2&p_p_col_count=1&p_r_p_564233524_tag=suspens%C3%A3o)

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