



Volume number

248

Layer number

18

DOMESTIC MEDICAL PREPARATION LICENSE

Analysis charge receipt

License charge receipt

 Date
 Number

 25/01/2013
 /
 65

DateNumber21.02.2012F0419305.09.2012F00223

It is allowed that the product formulated by VEM İLAÇ SAN. VE TİC. LTD. ŞTİ./ANKARA, manufactured by MEFAR İLAÇ SAN. A.Ş. Pendik/İSTANBUL and called "MILRICOR 10 mg/10 ml I.V. Solution for Injection/Infusion Ampoule (Milrinone)" can be sold as prescription drug by VEM İLAÇ SAN. VE TİC. LTD. ŞTİ / ANKARA.

Signed by Dr. Hakkı GÜRSÖZ Vice President of Instutiton (Seal and Signature)

11 February 2013

Shelf life of drug product is 24 months. (At room temperature below 25°C) After it is diluted with %0.45 and 0.9 NaCl for infusion and %5 Glucose solutions, it is stable for 24 hours at room temperature below 25°C. 11 February 2013 (Seal and Signature)

Packaging: In box, Type I glass ampoule, 1 unit. 11 February 2013 (Seal and Signature)

Starting date of five-year scientific examination period is 11 February 2013 and it is confirmed. (Seal and Signature)

Shelf life of drug product is 48 months and storage condition is room temperature below 25°C. 04 August 2016 (Seal and Signature)

It was approved to change company title of registered product hereby with this license as "Vem İlaç San. ve Tic. A.Ş." 18 August 2016 (Seal and Signature) Results of the scientific examination results are confirmed and license remains valid. 06 June 2018 (Seal and Signature)

It was approved to add "Vem İlaç San. ve Tic. A.Ş. Kapaklı/TEKİRDAĞ" as alternative to the current release site including manufacture, primary/secondary packaging and batch control/analysis of the product registered hereby with this certificate. 16 July 2018 (Seal and Signature)

It is suitable to remove "Mefar İlaç San. A.Ş., Pendik/İSTANBUL" from batch release sites including current approved bulk manufacturing, primary-secondary packaging, batch control site of the preparate registered in this certificate. 12 July 2019 (Seal and Signature)