





CZECH REPUBLIC

Recent and planned changes in pharmaceutical pricing and reimbursement and overview of the medical devices system

	Changes in pricing 2013-2014 O1/07/2013: Croatia has been added to the reference basket (price procedure) O1/01/2015 plan: Decrease of VAT rate on medicines from 15% to 10% (will affect price automatically)	 Changes in reimbursement 2013-2014 01/07/2013: Croatia has been added to the reference basket (reimbursement price procedure) 01/01/2015 plan: Decrease of VAT rate on medicines from 15% to 10% (will affect reimbursement price automatically)
	 Other changes related to medicines 25/07/2013: Generic prescription enabled (indicative) 01/01/2014: Hospital stay fee cancelled (was approx. €3.6 per day). Plans on cancelation of other fees (dispensing fee, clinical examination fee, currently approx. €1.1). 	
M E D D E V I C E S	 Price: prices of medical devices are not regulated. No competent authority for pricing is established. List of medical devices is available at www.RZPRO.cz. Reimbursement price: based on Act No. 48/1997 Coll., on Public Health Insurance and on Amendments to Some Related Acts. Reimbursement prices are published by insurance fund: http://www.vzp.cz/poskytovatele/ciselniky/zdravotnicke-prostredky IN-PATIENT CARE – represents approx. 2/3 of all medical devices. Generally 100 % reimbursement rate (based on the least expensive variant in the cluster) is applied. Common hospital devices (e.g. needles, bandages) are financed as part of the health service (DRG reimbursement). Extra financing of expensive medical (hardware) devices (CTs, X-ray etc.) is established. OUT-PATIENT CARE - represents approx. 1/3 of all medical devices. Positive (185 clusters) and negative list (23 clusters) are defined by Act No. 48/1997 Coll. Reimbursement rate is 100 % of the least expensive variant in the cluster of medical devices defined by the positive list. Medical devices not listed on the positive or negative list are reimbursed at 75 % rate (Act No. 48/1997 Coll.) Currently new Act on Medical Devices is in the legislation process: Should lead to unifying the agenda of "registration" of medical devices by establishing single registry of all safe and effective MDs in the Czech Republic (currently 7 competent institutions operate in the area). In the future, reimbursement of MDs in out-patient sector will be decided by the State Institute for Drug Control (SÚKL) also with the use of HTA 	