KYORIN Co., Ltd. 5, Kanda Surugadai 2-chome, Chiyoda-ku, Tokyo Securities Code: 4569; TSE 1st Section

RELEASE OF "MUCODYNE® DS50%"

KYORIN Pharmaceutical Co., Ltd. (Headquarters: Tokyo, President: Keiji Hirai), a subsidiary of KYORIN Co., Ltd. announces the release of Mucodyne® DS50% (Nonproprietary name: L-Carbocisteine), a mucoregulating drug (for adults and children) today.

Since the release of Mucodyne® Tablets (for adults) in January 1981, Kyorin Pharmaceutical Co., Ltd. has made planned and consistent efforts to meet the needs of patients and medical workers, by developing new formulations and obtaining new indications. (Please refer to the Summary of Product Profile in the attachment for the present formulation and indications.)

The Company hopes that the new Mucodyne® DS50% will improve convenience for adult and pediatric patients, and contribute to treatment in such fields as pulmonology and otology.

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< Product Profile for MUCODYNE $^{\circledR}$ DS 50% >

Product Profile for MIUC	DE 30/02			
PRODUCT NAME	MUCODYNE® DS50%			
NONPROPRIETARY	L-Carbocisteine (JAN)			
NAME				
INDICATIONS	For adults			
	Expectoration for the following diseases: Upper respiratory inflammation			
	(pharyngitis, laryngitis), acute bronchitis, bronchial asthma, chronic bronchitis,			
	bronchiectasis and pulmonary tuberculosis. Drainage in chronic sinusitis.			
	For children			
	Expectoration for the following diseases: Upper respiratory inflammation			
	(pharyngitis, laryngitis), acute bronchitis, bronchial asthma, chronic bronchitis,			
	bronchiectasis and pulmonary tuberculosis. Drainage in chronic sinusitis. Drainage in			
	otitis media with effusion.			
DOSAGE AND	For adults			
ADMINISTRATION	For oral use, the usual dose for adults is 500 mg of L-carbocisteine (<u>1.0g</u> of MUCODYNE® DS50%) three times daily. MUCODYNE® DS50% should be suspended immediately before use. The dose may be adjusted according to age and symptoms.			
	For children			
	For oral use, the usual dose for children is 10 mg/kg of L-carbocisteine (0.02g/kg of			
	MUCODYNE® DS50%) three times daily. MUCODYNE® DS50% should be			
	suspended immediately before use. The dose may be adjusted according to age and			
	symptoms.			
NHI PRICE	¥37.70/g			
PACKAGING	Boxes of 120 packets (<u>1.0g</u> ×120 packets)			
	Bottles of 100 g, and bottles of 500 g.			

Underlined sections correspond to changes from MUCODYNE® DS 33.3%.

Reference: Indications and marketing launch for each MUCODYNE formulation

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Formulations	For adults	For children	Marketing launch	
MUCODYNE® Tablets	0	_	Date of initial marketing in Japan: July 2001	
250mg			* Marketed in January 1981 under the name of	
			MUCODYNE [®] Tablets	
MUCODYNE® Tablets	0	_	Date of initial marketing in Japan: October 1996	
500mg				
MUCODYNE [®] Fine	0	_	Date of initial marketing in Japan: November	
Granules 50%			1987	
MUCODYNE [®] Syrup 5%	_	\circ	Date of initial marketing in Japan: October 1987	
MUCODYNE® DS 33.3%	0	\circ	Date of initial marketing in Japan: October 2001	
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MUCODYNE® DS 50%		\circ	Date of initial marketing in Japan: May 2010	
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