PIHAK BERKUASA PERANTI PERUBATAN
Medical Device Authority
KEMENTERIAN KESIHATAN MALAYSIA
Ministry of Health Malaysia
Aras 5 & 6, Prima 9, Prima Avenue II,
Blok 3547, Persiaran Apec,
63000 Cyberjaya, Selangor
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Tel: (+603)8230 0300
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Portal Rasmi: www.mdb.gov.m
Email: mdb@mdb.gov.my

MEDICAL DEVICE RECALL NOTICE

MDA REFERENCE NUMBER:	MDA/PMSV/R2018-067
MEDICAL DEVICE NAME:	**Non Latex Durex Real Feel Condoms** (Poly Isopropene Condom) Device Registration Number : GC38774698818
AFFECTED BATCH NUMBERS:	• 1000433144 • 1000438055 • 1000422259
INTENDED USE:	Contraception and the prevention of sexually transmitted infections
RECALLING ESTABLISHMENT:	**Reckitt Benckiser (Malaysia) Sdn. Bhd.** Licensing Number: KP5915634715
REASON FOR RECALL/ALERT:	*-Please Refer Attachment-*
RECALL CLASS:	Recall Class III
ACTION & RECOMMENDATION:	**Any individuals and premises possessing the above mentioned device are advised to respond to this communication immediately.**
REFERENCES (if available):	
CONTACT:	For further information, please contact: Authorised Representative : **RECKITT BENCKISER (MALAYSIA) SDN. BHD.** **LEVEL 5 MENARA UAC, NO 12 JALAN PJU 7/5,** **MUTIARA DAMANSARA,** **PETALING JAYA,** **47800 SELANGOR.** www.durex.com.my **Durex Consumer Careline: - 1-800-888-839.** **Media Enquiries: +603 2276 0990** Atiq Safirah Dau Ming Seling

MEDICAL DEVICE AUTHORITY
MINISTRY OF HEALTH MALAYSIA
1ST AUGUST 2018