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**GOVERNMENT NOTICE**

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**DEPARTMENT OF HEALTH****No. R. 389****12 May 2010****MEDICINES AND RELATED SUBSTANCES ACT, 1965 (ACT 101 OF 1965)****GENERAL REGULATIONS MADE IN TERMS OF THE MEDICINES AND RELATED  
SUBSTANCES ACT, 1965: AMENDMENT**

The Minister for Health has, in consultation with the Medicines Control Council, in terms of section 22C (1) (a) of the Medicines and Related Substances Act, 1965 (Act 101 of 1965), made the regulations in the Schedule.

**SCHEDULE**

1. In these regulations “the Regulations” means the General Regulations, made in terms of the Medicines and Related Substances Act, 1965 (Act 101 of 1965) published under Government Notice No. R. 510, *Gazette* 24727 of 10 April 2003 as amended.

**Amendment of regulation 18 of the Regulations**

2. Regulation 18 of the Regulations is hereby amended by—

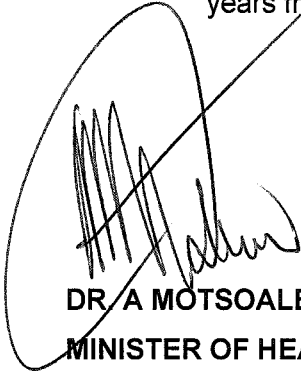
- (a) the deletion of subregulation (3)(f);
- (b) the deletion of subregulation (3)(g);
- (c) the deletion of subregulations (5(a), (c), (d) and (e);
- (d) the deletion of subregulation (6); and
- (e) the substitution for subregulation (7) of the following subregulation:

- “(7) Any person may support or oppose an application referred to in subregulation (1) by making written representations to the Director-General.”.

**Amendment of regulation 20 of the Regulations**

3. Regulation 20 of the Regulations is hereby amended by substitution for subregulation (1) of the following subregulation:

“(1) A licence issued in terms of regulation 18 or 19 shall be valid for a period of 5 years from the date of issue.”.



**DR. A MOTSOLEDI, MP**  
**MINISTER OF HEALTH**

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