



Федеральная служба
по надзору в сфере
здравоохранения

MINISTRY OF HEALTH
ROSZDRAVNADZOR

Administration in the sphere of circulation of medical devices



Starting from January 1st 2013

In accordance with Art. 38 / Federal law № 323:

**Pre-market approval
and post-market
surveillance of
medical devices**

**State registration
of medical devices**

**Issuance of import permits for medical devices
for the purpose of state registration**

**Control over the entire life cycle
of medical devices**

**Monitoring of safety
of medical devices**

**Licensing the manufacturing
of medical devices**

Regulation of the Circulation of MD

In accordance with the Federal law «On the fundamental principles of healthcare in the Russian Federation»

RESOLUTIONS OF THE GOVERNMENT OF THE RUSSIAN FEDERATION

27.12.2012 №
1416

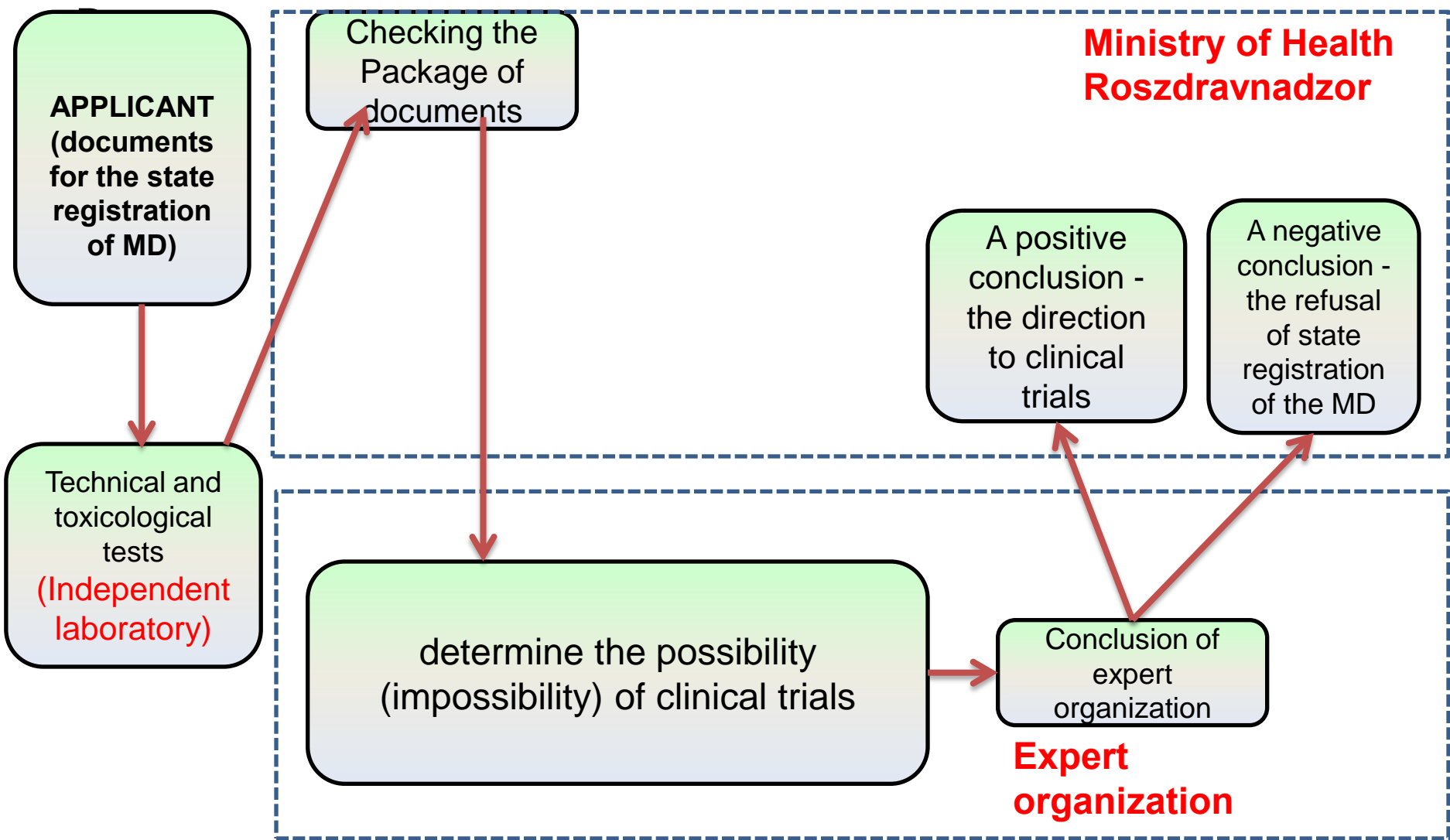
19.06.2012
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26.09.2012
№ 970



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The scheme of registration (I level)





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The scheme of registration of MD (2 level)

