

Legislative Tracking



If you would like advice with regard to any of these issues, please do not hesitate to contact the Tax & Legal Department of Deloitte CIS at +7 (495) 787 06 00 (Moscow) or + 7 (812) 703 71 06 (St. Petersburg).

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18 July 2016

Official website of the OECD

<http://www.oecd.org/tax/panama-decides-to-sign-multilateral-tax-information-sharing-convention.htm>

Panama plans to sign Multilateral Convention on Mutual Administrative Assistance in Tax Matters

The Panamanian authorities plan to sign the OECD's Multilateral Convention on Mutual Administrative Assistance in Tax Matters.

This decision was communicated in a letter published on the official website of Panama's Ministry of Foreign Affairs on 15 July 2016.

To date, 98 countries and jurisdictions have signed the Convention, including Russia.

16 July 2016

Official website of the Russian Ministry of Economic Development

<http://economy.gov.ru/minec/about/structure/depino/2016160701>

Russia puts forward new cluster initiatives

The Russian Ministry of Economic Development came forward with new proposals and approaches envisaging comprehensive government support for leading innovation clusters.

The Ministry intends to identify promising innovative territorial clusters in at least five federal subjects. These clusters will receive support from the Ministry of Economic Development and other public agencies and development institutes to achieve levels of investment attractiveness and economic yield from government-backed business support consistent with international standards.

This initiative has not yet been officially released.

18 July 2016

ConsultantPlus

http://www.consultant.ru/document/cons_doc_LAW_201811/

New procedure for registration and inspection of pharmaceuticals within Eurasian Economic Union

The Eurasian Economic Council published Resolution No. 46 of 12 February 2016 "On the Rules for Registering and Assessing the Safety, Quality and Efficacy of Pharmaceuticals".

These rules outline the procedure for registering and assessing the safety, quality and efficacy of pharmaceuticals, making amendments to register entries, issuing copies of marketing authorisation documents, rejecting the registration of pharmaceuticals, as well as suspending and cancelling marketing authorisation.

Appendices to the rules include application templates and lists of documents required for the registration of a pharmaceutical product.

The Resolution will come into effect on 22 July 2016.

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