

Бренд ученого

*Как сделать так, чтобы
нас цитировали*

Сергей Парамонов, к.х.н.

Специалист по информационным ресурсам, научные исследования
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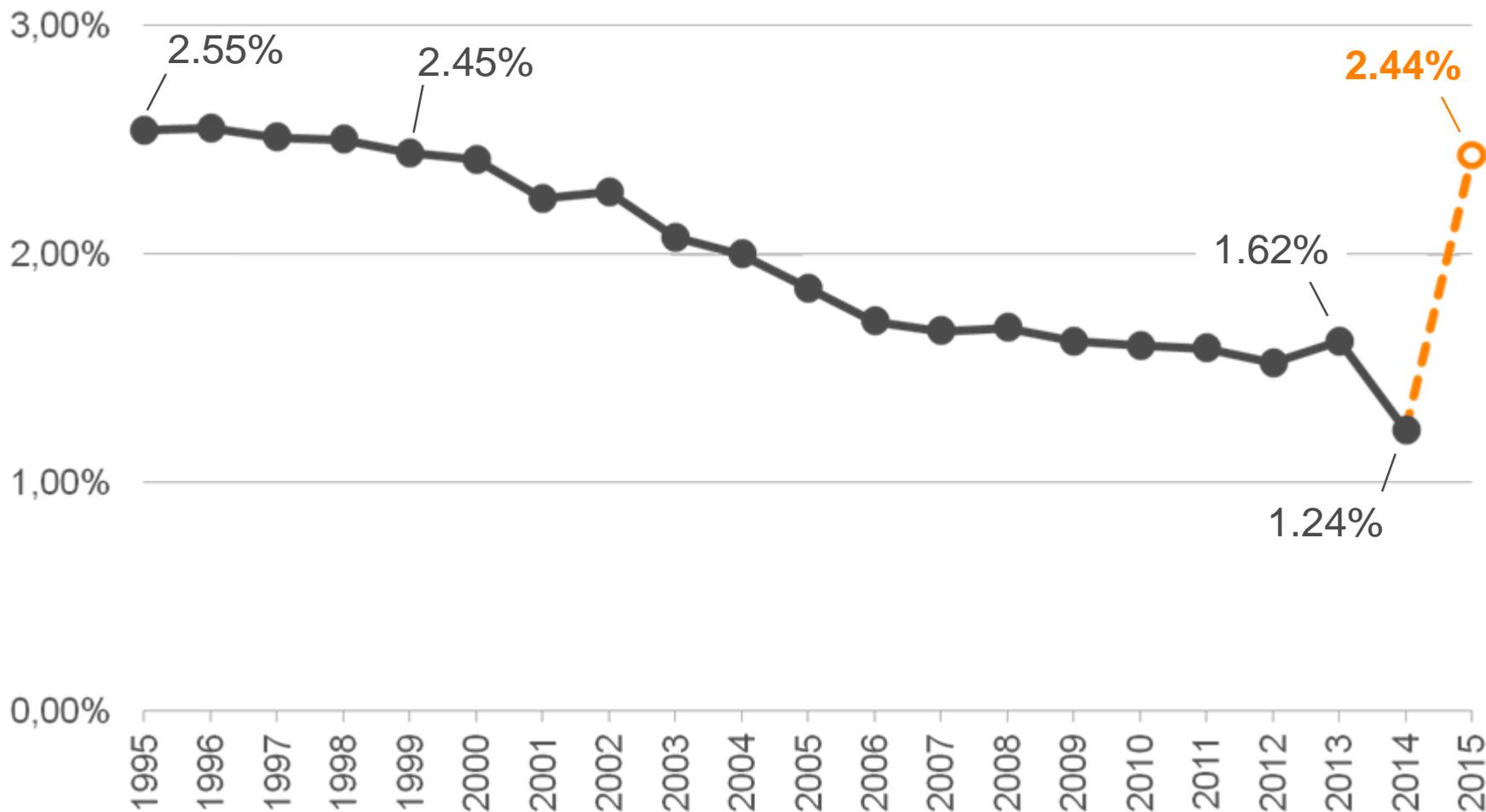
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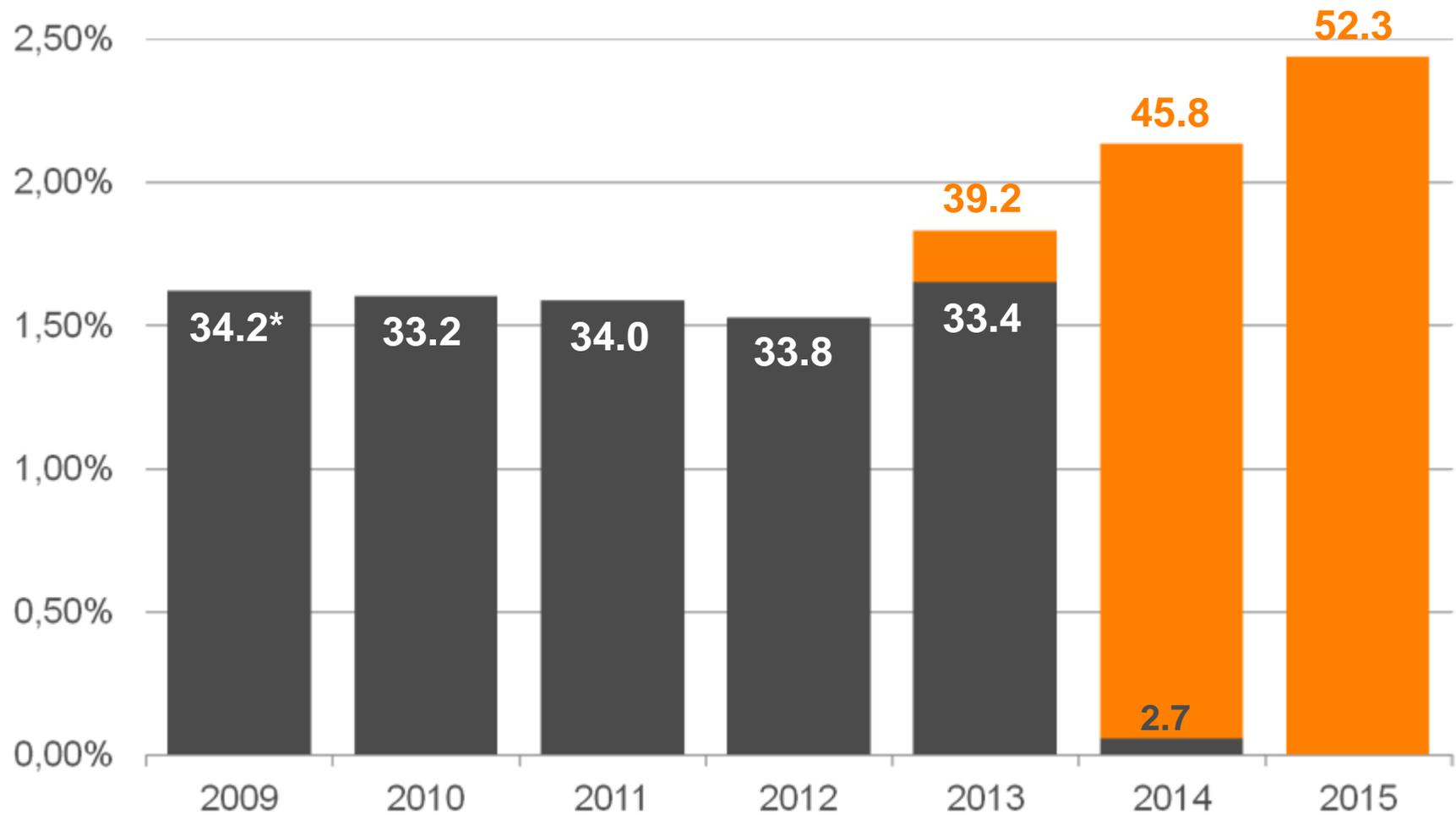
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- «вхождение к 2020 году не менее пяти российских университетов в первую сотню ведущих мировых университетов согласно мировому рейтингу университетов» (программа 5-100-2020)

2.44% – это сколько?

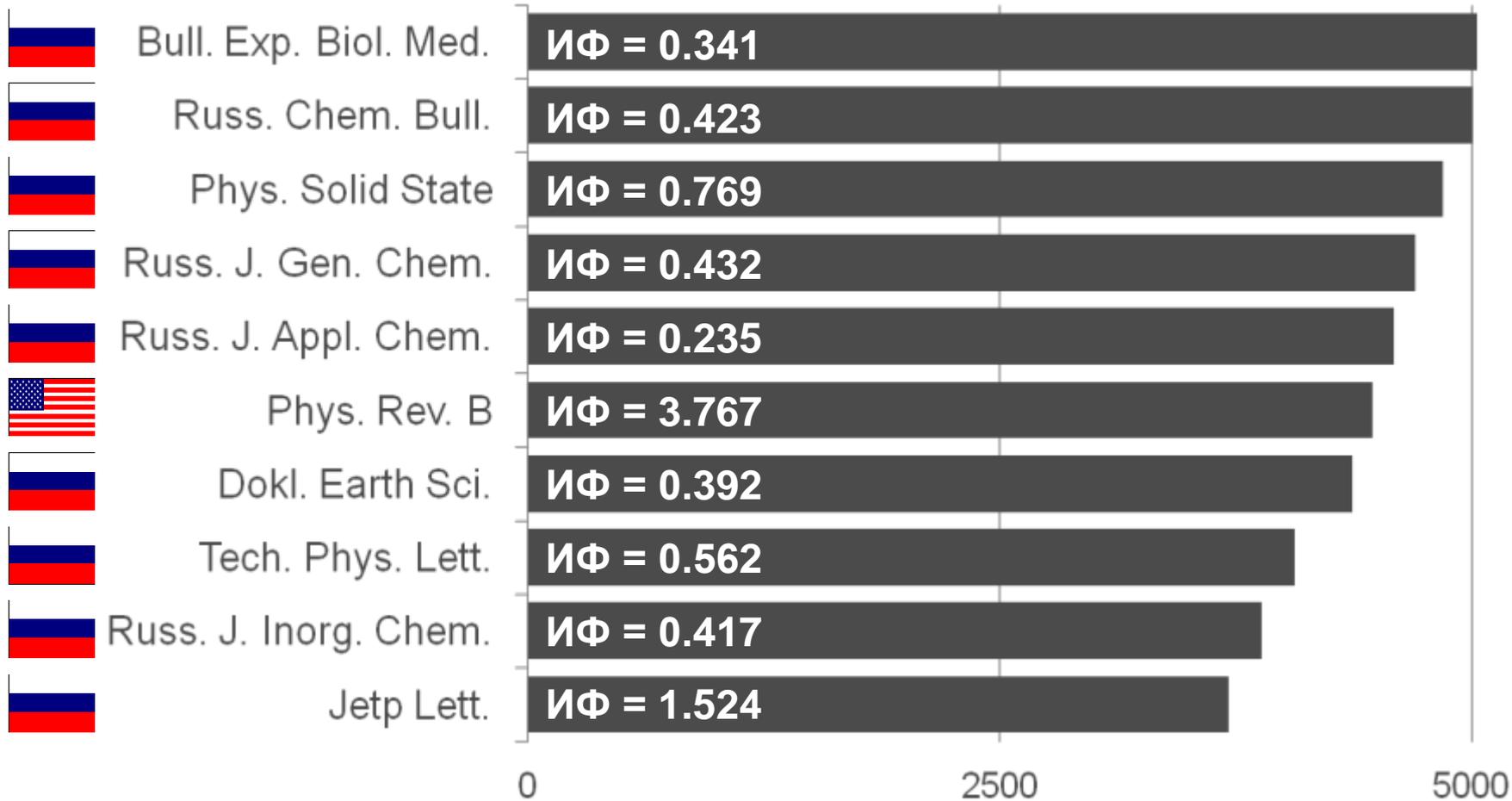


...сколько?



* ТЫСЯЧИ

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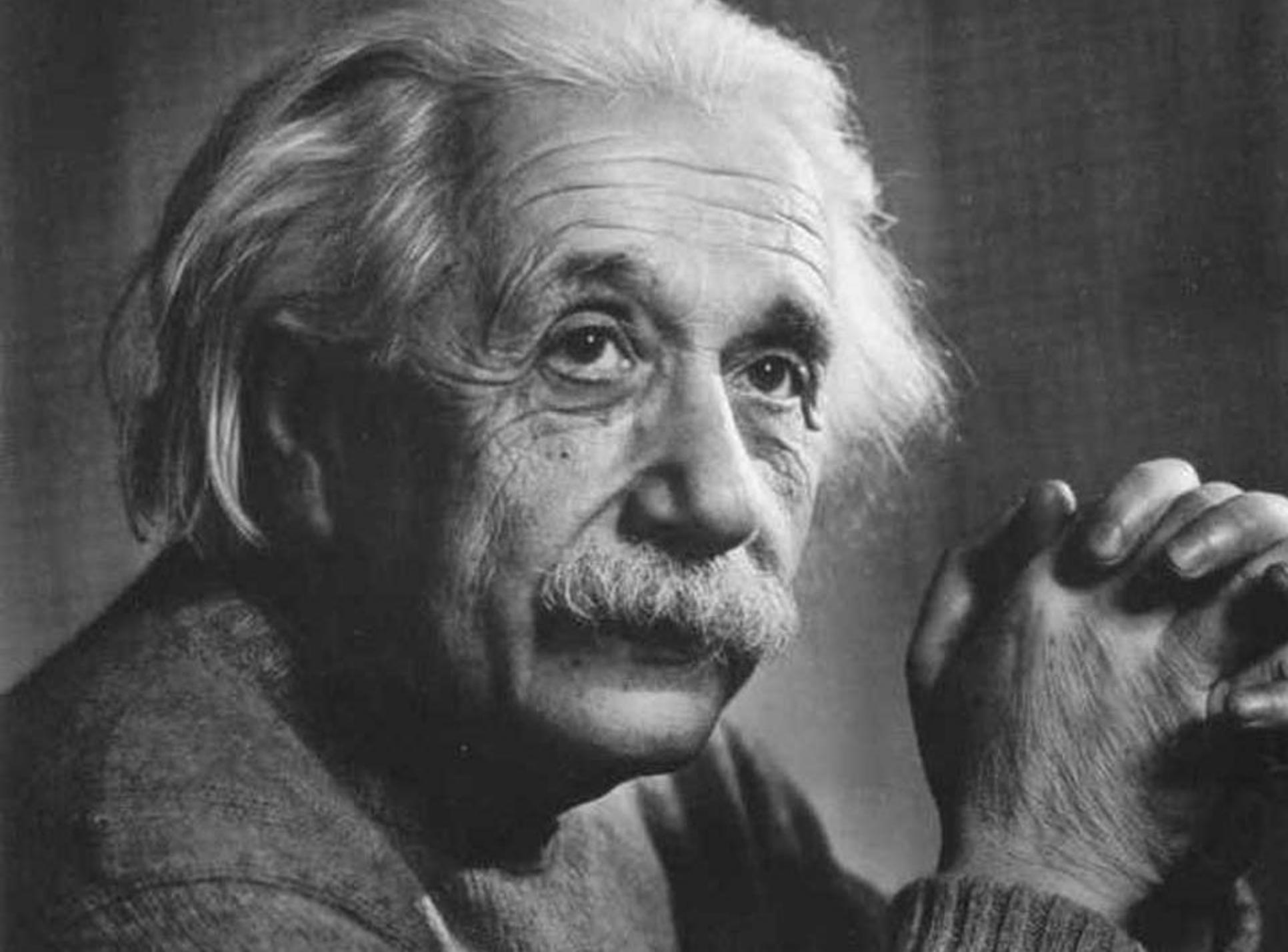
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 *Профессионалы есть в каждой области*



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*Каждый ученый –
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Бренд ученого



*Основа бренда –
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Актуалност и новизна



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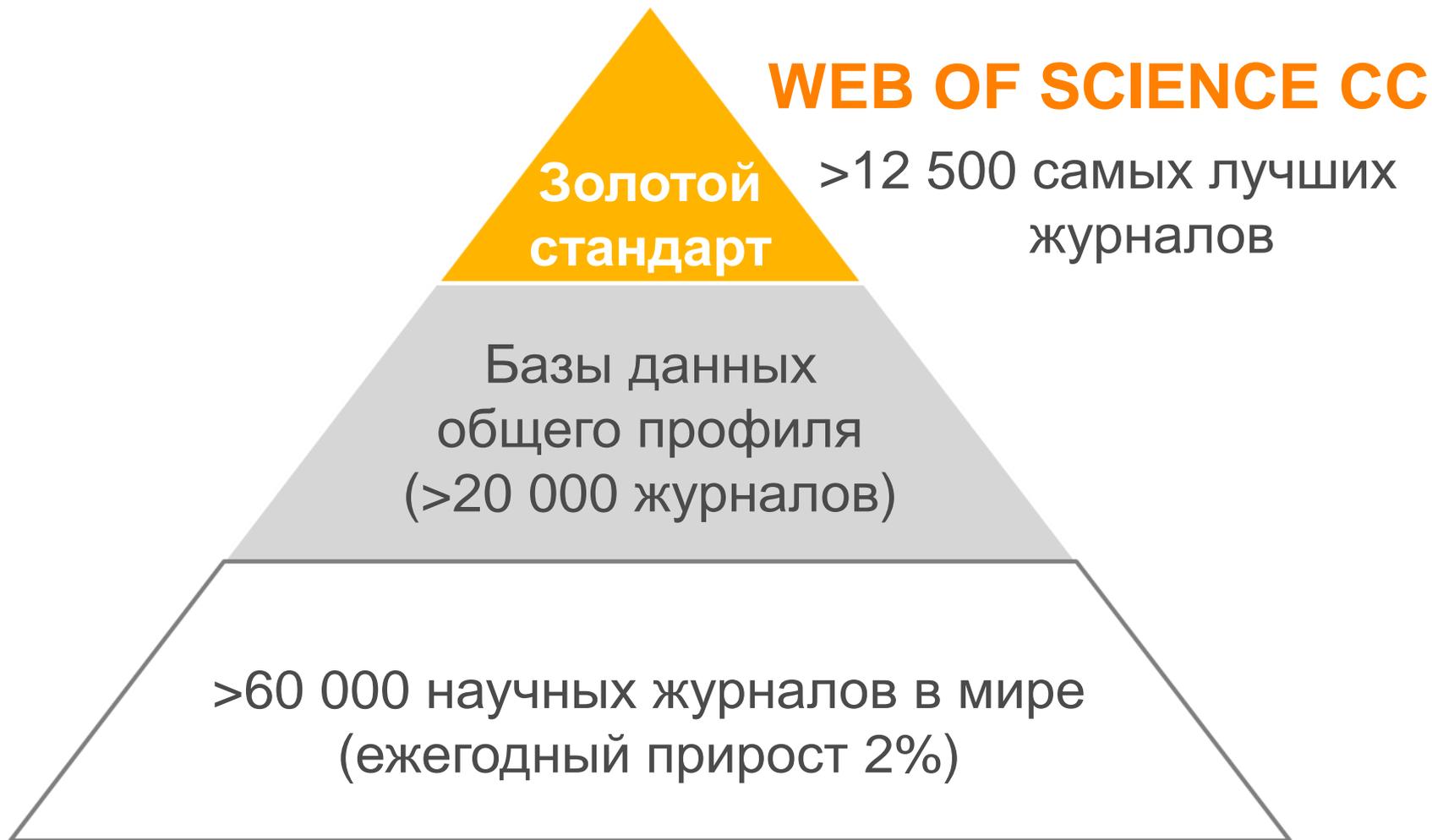
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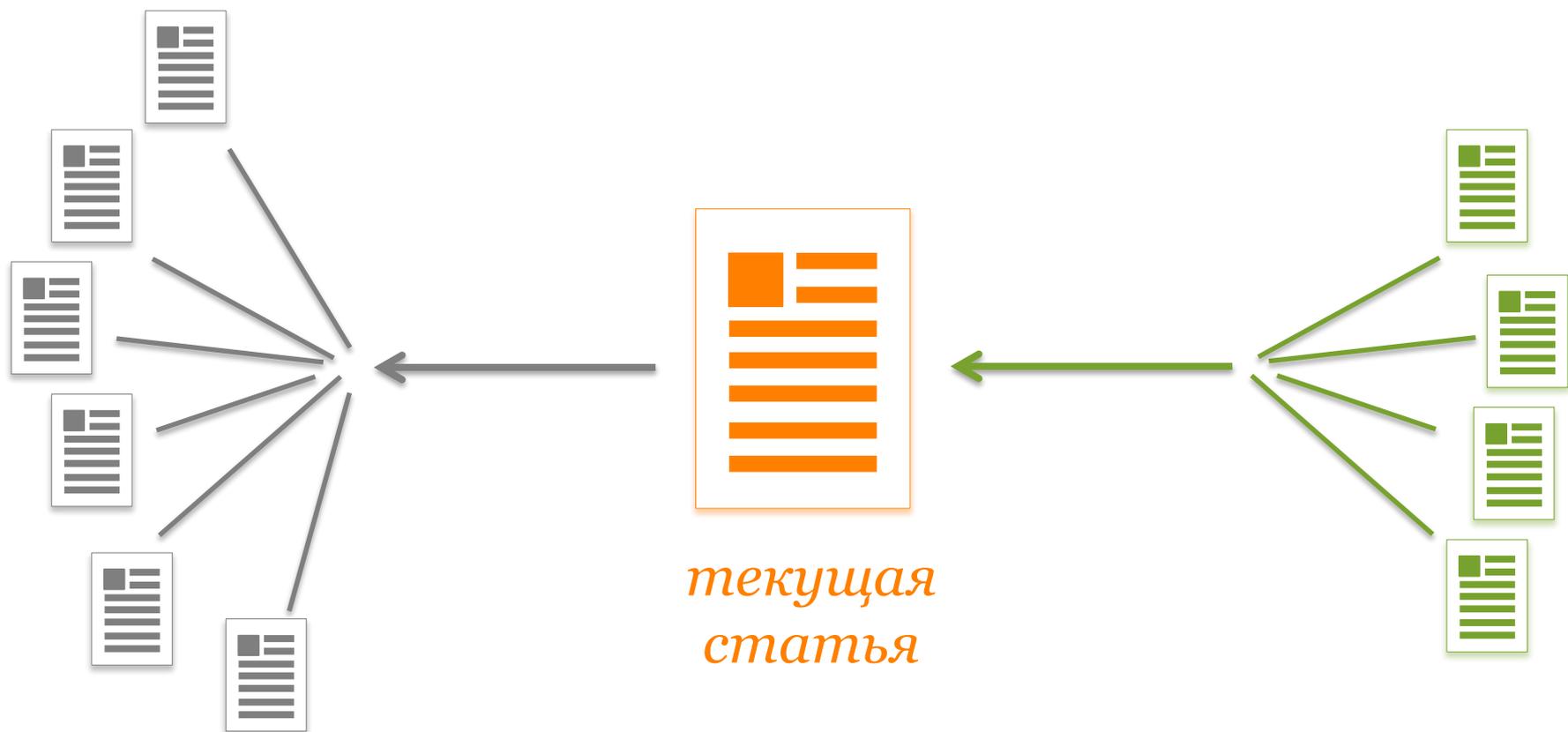
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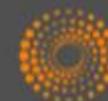
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JOURNAL OF CHROMATOGRAPHY A Volume: 1217 Issue: 16

Pages: 2674-2684 Published: APR 16 2010

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2. **Temperature controlled ionic liquid-dispersive liquid phase microextraction for determination of trace lead level in blood samples prior to analysis by flame atomic absorption spectrometry with multivariate optimization**

By: Shah, Faheem; Kazi, Tasneem Gul; Naeemullah; et al.

MICROCHEMICAL JOURNAL Volume: 101 Pages: 5-10

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Temperature controlled ionic liquid-dispersive liquid phase microextraction for determination of trace lead level in blood samples prior to analysis by flame atomic absorption spectrometry with multivariate optimization

By: Shah, F (Shah, Faheem)^[1,2]; Kazi, TG (Kazi, Tasneem Gul)^[1]; Naeemullah (Naeemullah)^[1]; Afridi, HI (Afridi, Hassan Imran)^[1]; Soy lak, M (Soy lak, Mustafa)^[2]

MICROCHEMICAL JOURNAL

Volume: 101 Pages: 5-10

DOI: 10.1016/j.microc.2011.09.009

Published: MAR 2012

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Abstract

This paper described a new approach for the preconcentration of lead (Pb²⁺) by temperature controlled ionic liquid-dispersive liquid phase microextraction (TIL-DLME) prior to analyzing by flame atomic absorption spectrometry (FAAS). An ionic liquid (IL) 1-Butyl-3-methylimidazolium hexafluorophosphate [C4MIM][PF6] was used as an extractant solvent. The Pb²⁺ was complexed with ammonium pyrrolidinedithiocarbamate (APDC) and then entered into the infinite IL drops at high temperature (>70 degrees C). Important variables affecting the microextraction efficiency such as pH, ligand concentration, amount of IL temperature and incubation time were investigated. The results showed that the coexistent ions had no obvious negative effect on the determination of Pb²⁺. In the optimum experimental conditions

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ions had no obvious negative effect on the determination of Pb²⁺. In the optimum experimental conditions, the limit of detection (LOD) and the enhancement factor (EF) were 0.13 µg L⁻¹ and 93, respectively. The relative standard deviation (RSD) of 10 µg L⁻¹. Pb²⁺ was 4.3%. The developed method was validated by determining Pb²⁺ in certified reference material (CRM) and the results showed that the determined values of Pb²⁺ were in good agreement with the certified value. The proposed method was applied satisfactorily for the preconcentration of Pb²⁺ in acid digested blood samples of children with different respiratory disorders. (C) 2011 Elsevier B.V. All rights reserved.

Keywords

Author Keywords: 1-Butyl-3-methylimidazolium; hexafluorophosphate; Lead; Ammonium pyrrolidinedithiocarbamate; Microextraction; Blood samples

KeyWords Plus: SINGLE DROP MICROEXTRACTION; CLOUD POINT EXTRACTION; AIR-POLLUTION; ENVIRONMENTAL-SAMPLES; WATER SAMPLES; PRECONCENTRATION; EXPOSURE; CHILDREN; MANGANESE(II); IMIDAZOLIUM

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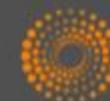
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By: Woolfenden, E (Woolfenden, Elizabeth)

JOURNAL OF CHROMATOGRAPHY A

Volume: 1217 Issue: 16 Pages: 2674-2684

DOI: 10.1016/j.chroma.2009.12.042

Published: APR 16 2010

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Abstract

Sorbent tubes/traps are widely used in combination with gas chromatographic (GC) analytical methods to monitor the vapour-phase fraction of organic compounds in **air**. Target compounds range in volatility from acetylene and freons to phthalates and PCBs and include apolar, polar and reactive species. Airborne vapour concentrations will vary depending on the nature of the location, nearby **pollution** sources, weather conditions, etc. Levels can range from low percent concentrations in stack and vent emissions to low part per trillion (ppt) levels in ultra-clean outdoor locations. Hundreds, even thousands of different compounds may be present in any given atmosphere. GC is commonly used in combination with mass spectrometry (MS) detection especially for environmental monitoring or for screening uncharacterised workplace atmospheres. Given the complexity and variability of organic vapours in **air**, no one sampling approach suits every monitoring scenario. A variety of different sampling strategies and sorbent media have been

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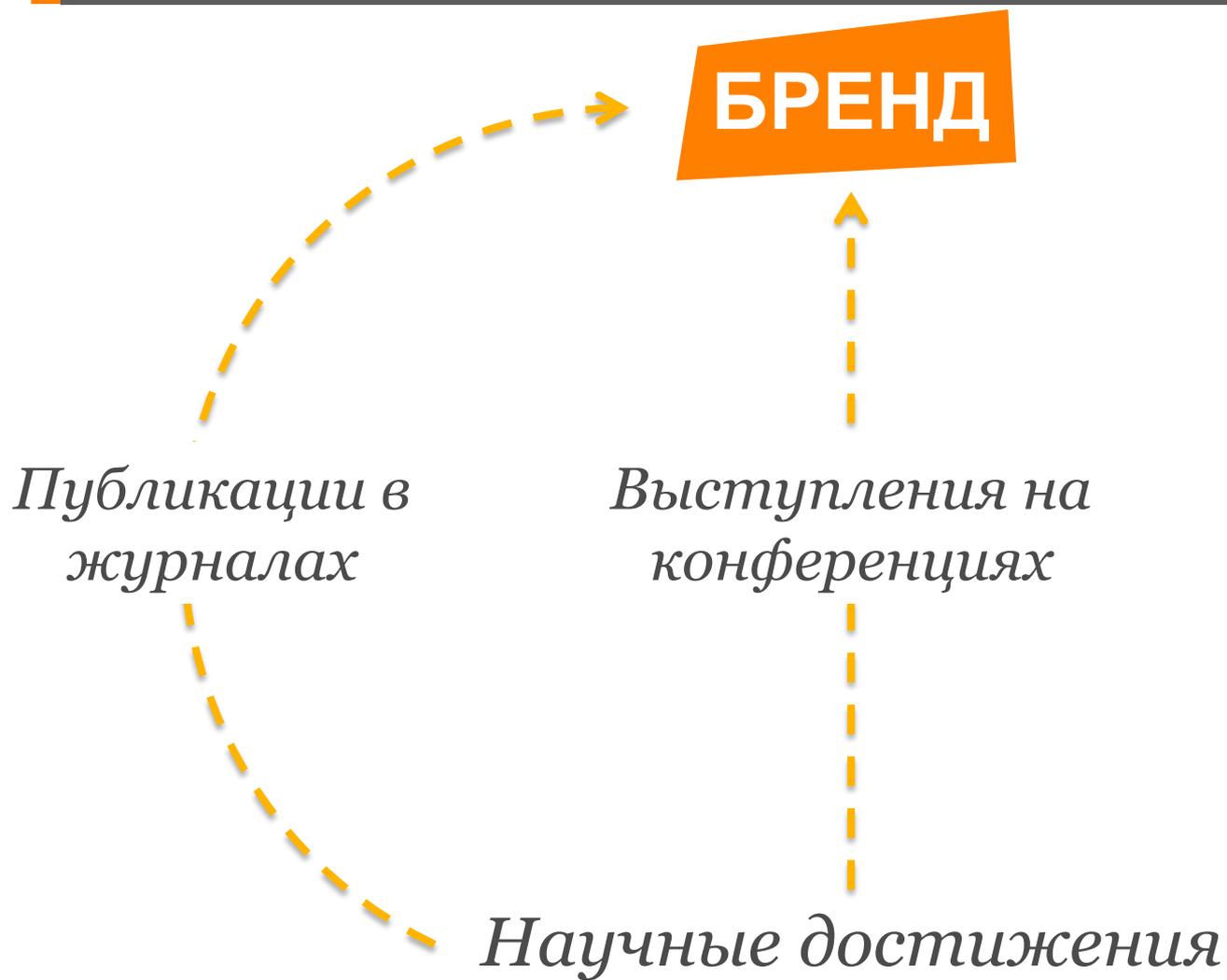
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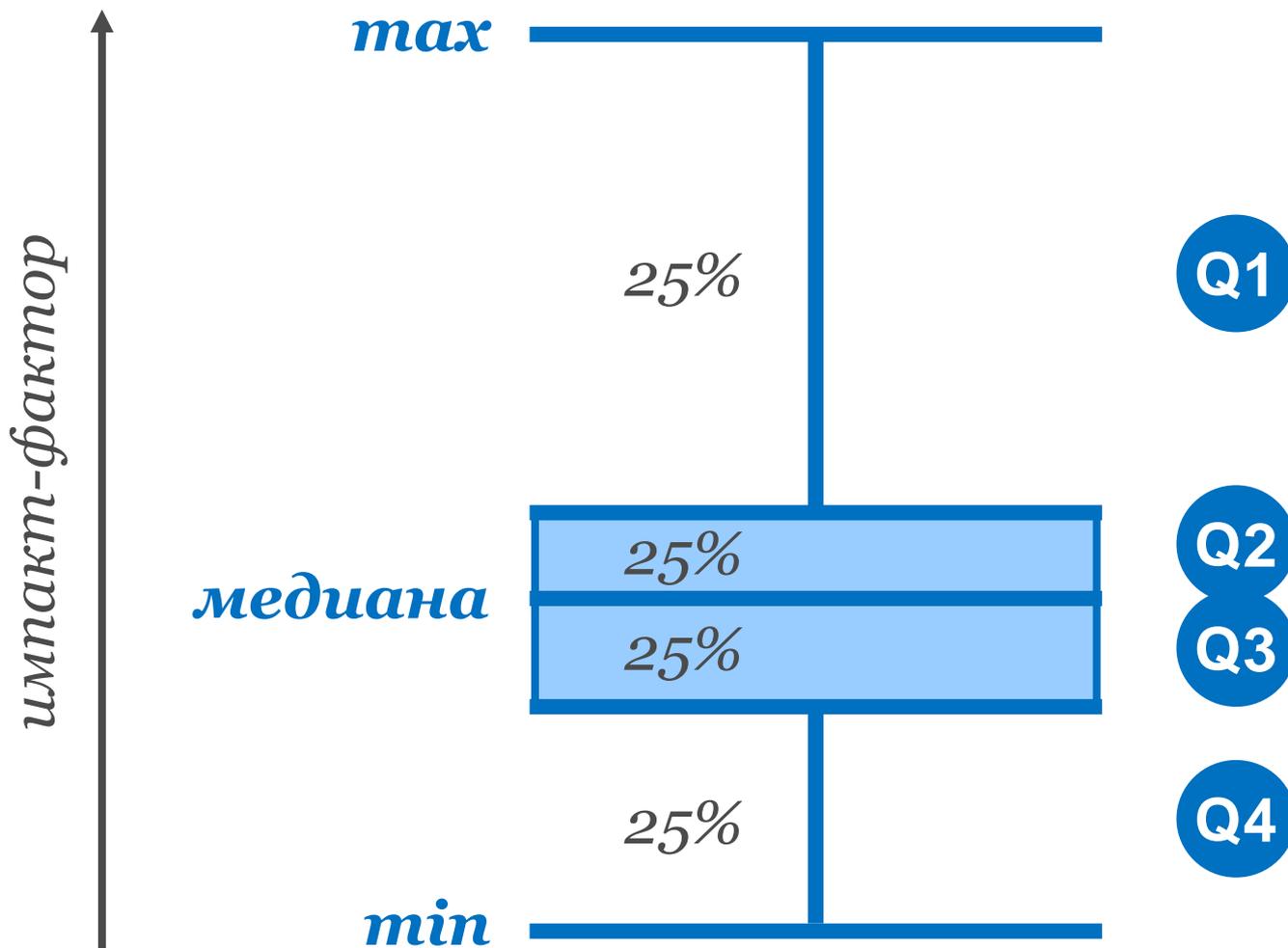


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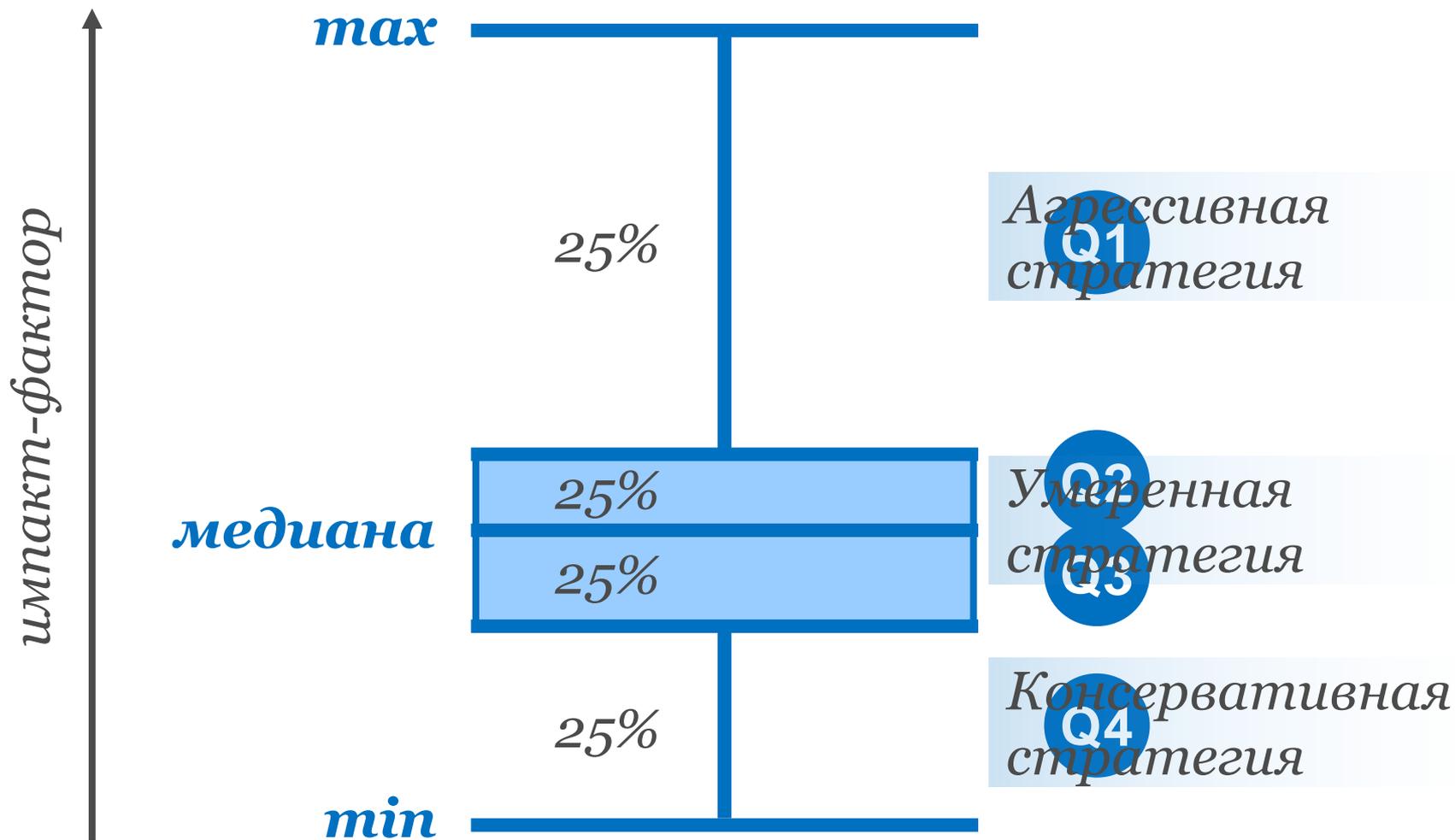
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Sorbent-based sampling methods for volatile and semi-volatile organic compounds in **air** Part 1: Sorbent-based **air** monitoring options

By: Woolfenden, E (Woolfenden, Elizabeth)

JOURNAL OF CHROMATOGRAPHY A

Volume: 1217 Issue: 16 Pages: 2674-2684

DOI: 10.1016/j.chroma.2009.12.042

Published: APR 16 2010

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Abstract

Sorbent tubes/traps are widely used in combination with gas chromatographic (GC) analytical methods to monitor the vapour-phase fraction of organic compounds in **air**. Target compounds range in volatility from acetylene and freons to phthalates and PCBs and include apolar, polar and reactive species. Airborne vapour concentrations will vary depending on the nature of the location, nearby **pollution** sources, weather conditions, etc. Levels can range from low percent concentrations in stack and vent emissions to low part per trillion (ppt) levels in ultra-clean outdoor locations. Hundreds, even thousands of different compounds may be present in any given atmosphere. GC is commonly used in combination with mass spectrometry (MS) detection especially for environmental monitoring or for screening uncharacterised workplace atmospheres. Given the complexity and variability of organic vapours in **air**, no one sampling approach suits every monitoring scenario. A variety of different sampling strategies and sorbent media have been

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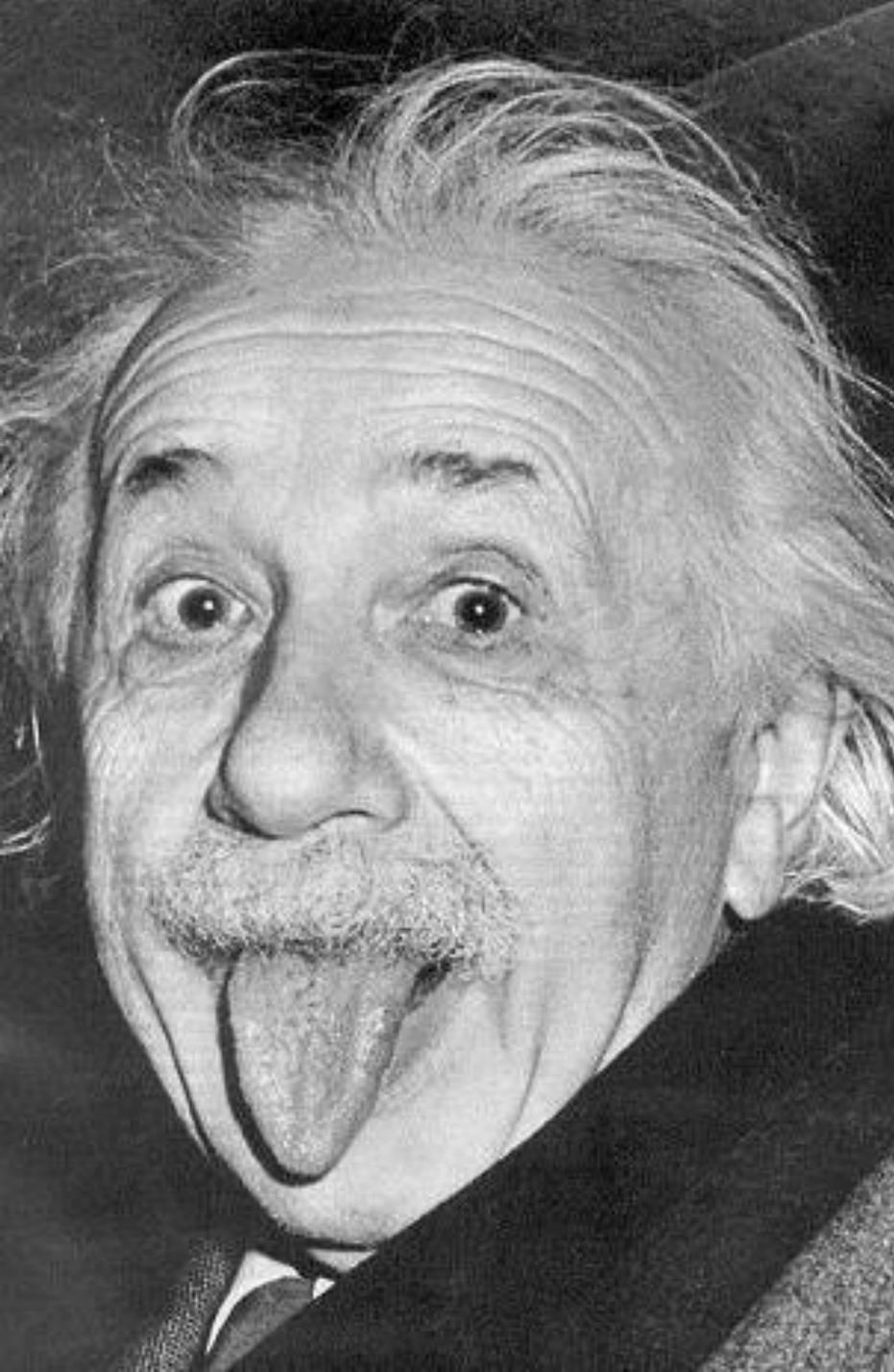
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3. H. K. Liu and P. J. Sadler, *Accounts of Chemical Research*, 2011, **44**, 349-359.
4. H. Mansouri-Torshizi, M. Eslami-Moghadam, A. Divsalar and A. A. Saboury, *Acta Chimica Slovenica*. 2011. **58**. 811-822.

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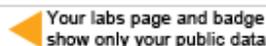


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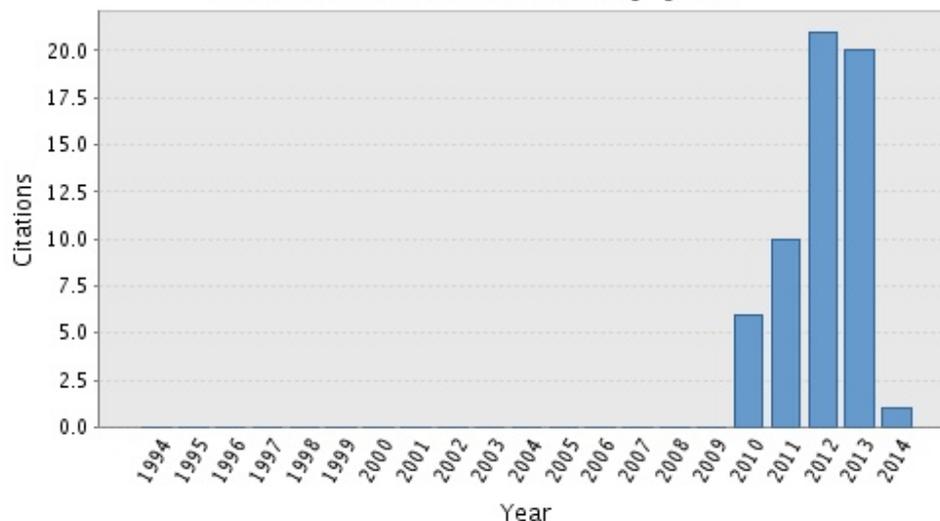
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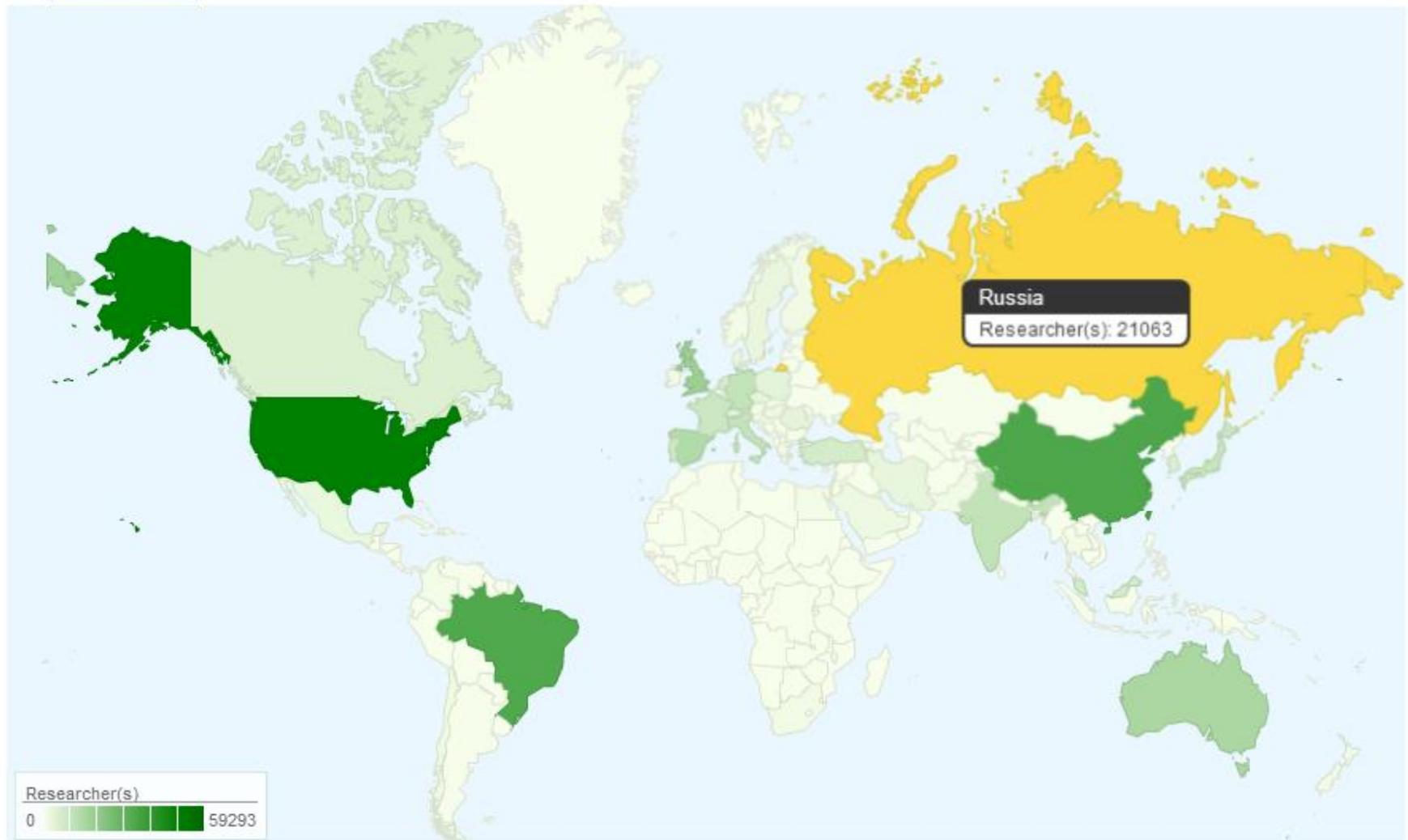
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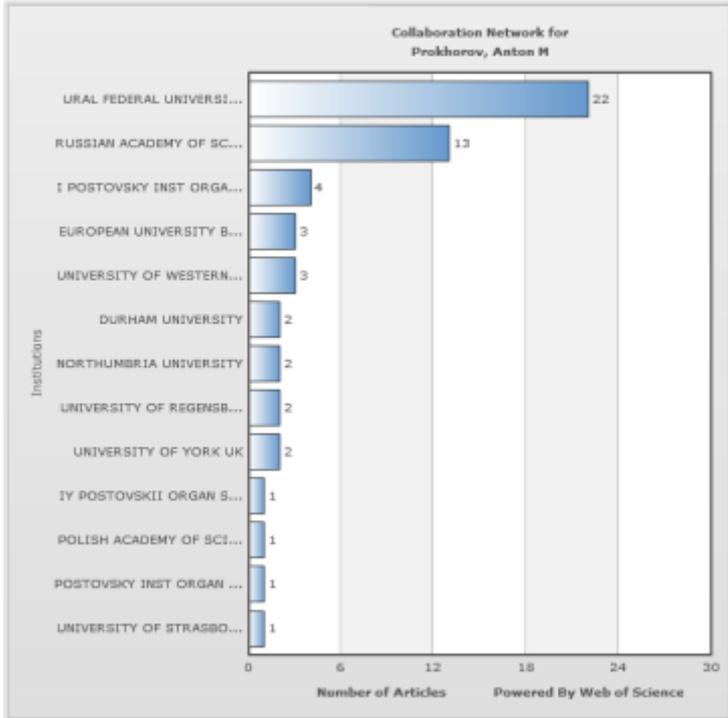
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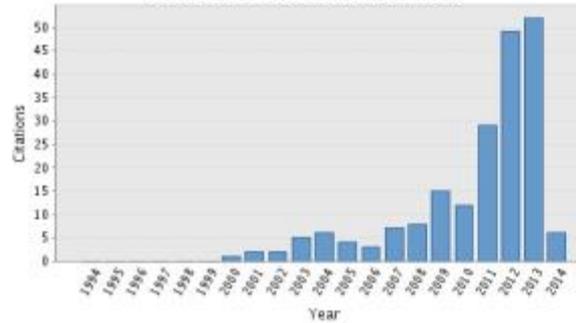
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