ESTIMATION OF THE POSSIBILITY OF EXPANDING THE INSTRUMENT BASE FOR THE RAPID DETECTION OF FALSIFIED MEDICINAL PRODUCTS IN THE RIVNE REGION

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1. Introduction
One of the main tasks facing the State Service of Ukraine on Medicines and Drug Control is to take measures to ensure state quality control of medicines, which must include the selection and laboratory analysis of drug samples. It is well known that the threat of an increase in the number of counterfeits in pharmaceutical markets increases during epidemics, pandemics, natural disasters, catastrophes, when drug consumption increases significantly. During these periods, there is usually an increase in the workload of territorial laboratories for quality control of medicines, as it is observed today with the spread of acute respiratory disease COVID-19 caused by the coronavirus SARS-CoV-2. Increasing the workload can be solved in several ways – either by increasing the number of laboratories and instrumentation for drug control, or by significantly reducing the time for sample preparation and analysis of each sample.

At present, it is impossible to imagine the quality control of drugs without the use of physico-chemical methods that are optimal in terms of "price-quality" in terms of the accuracy of determining the identity of drugs in relation to the cost of the equipment. These methods are increasingly being implemented in basic pharmaceu-
tical research and in the practice of pharmaceutical analysis for the identification and quantification of various groups of drugs, their standard samples and dosage forms [1, 2]. The most optimal for use in pharmaceutical analysis, including for the detection of falsified drugs (FD) and API, today are photometric methods, namely: fluorimetry, atomic emission spectrometry, atomic absorption spectrometry, absorption spectrophotometric spectrophotometry absorption spectrophotometry in the ultraviolet (UV) and visible regions, Raman spectrometry, mass spectrometry, near-infrared spectrometry (NIRS). Photometric methods are widely used both in scientific research and to identify various counterfeits, for example, IR spectrophotometry is used to identify APIs, including broad-spectrum antibiotics (ampicillin, chloramphenicol, indomethacin) [3, 4]; the method of IR spectroscopy is used in the pharmacopeial analysis of ranitidine hydrochloride tablets [5], there are studies of the application of the method in the analysis of medicinal plant raw materials [6]. Thus, photometric methods are widely used in establishing the authenticity of drugs and are used as a reliable method of detecting their falsification.

The aim of the article was to assess ways to expand the instrument base of photometric equipment for quality control of medicines, substantiation of prospects for their implementation and application in the activities of the territorial service on medicines and drug control in Rivne region.

2. Research planning (methodology)

To achieve this goal, the following tasks were identified: analysis and justification of the choice of photometric method; advantages of using IR and Raman spectrometry; conducting experimental research involving IR spectrophotometers of higher educational institutions in Rivne; identify the possibility of involving the instrument base of higher education institutions for the analysis of drugs in cases of heavy workload of laboratories of territorial services for medicines and drug control; summarizing the analysis and prospects for further research.

Experimental studies of drugs by IR spectrophotometry were planned on the basis of the National University of Water Management of Rivne and Rivne State University for the Humanities. The list of drugs and APIs was selected according to the lists of counterfeit drugs detected in Ukraine [7], in particular, the drug “Sumamed” manufactured by “Pliva” (Croatia) containing azithromycin [8, 9] and other drugs in demand.

3. Materials and methods of research

The research materials are literature data, scientific publications on the application of photometric methods in drug quality control and own research on drug quality control. The objects of experimental studies were azithromycin (“Sumamed”, Croatia, series 424129), erythromycin (“Zinerit”, the Netherlands, series 19A05 / 74), ibuprofen (“Ilbuprom max”, Poland, series U1904351), complex preparation of ibuprofen and paracetamol (“Nurofen Intensive, Great Britain, series HJ580”), clarithromycin (“Clerimed 500, Cyprus, series A2J053”), paracetamol (“Panadol”, Spain, series, F76W), cefuroxime sodium (“Axetin”, Cyprus, series C722A1). The following methods were used in the work: system analysis, bibliosemantic, data generalization, the method of absorption spectrophotometry in the infrared region using the instrument base was used in experimental research: IR Fourier spectrophotometers “IRAfinity-1S” manufactured by Shimadzu and “Nicolet iS5” manufactured by Fisher Scientific.

4. Research results

Infrared spectrometers such as UR-10, UR-20, Specord75 IR, Specord M40 (Carl Zeiss Jena, Germany), IR-Fourier spectrometers Nicolet 6700 (Thermo Fisher Scientific, USA) are most often used in the practice of pharmaceutical research [10] JASCO FT-IR 460 PLUS spectrometer (Pike Technologies, Madison, USA), as well as spectrometers IRPrestige-21, FTIR-8000S (Shimadzu, Japan). As a rule, helium-neon laser is used as a source of infrared radiation in such devices, the device also includes an optical system with mirrors and light dividers, a Michelson interferometer, an IR detector, and a processor. According to the technical characteristics, modern IR spectrometers have characteristics not less than: spectral range 7800–375 cm⁻¹, resolution 1 cm⁻¹, linearity of the ordinate (ASTM E1421) 0.1 % T, accuracy on the wave number 0.02 cm⁻¹, scan speed 3 scans / s.

These characteristics meet the requirements of SPPhU and are sufficient for pharmaceutical analysis.

Currently, IR spectrometers are very widely used, created and available databases of IR spectra for technical and food additives, drugs, poly- and monomers, surfactants, plasticizers, pesticides, solvents, petroleum products, toxic substances, steroids and other compounds, in the composition which includes one component. The literature is increasingly devoted to the publication of IR spectroscopy for the detection of FD, for example, samples of drugs of the steroid group [11], to identify derivatives of 1,4-benzodiazepines [12], qualitative and quantitative analysis of antitumor drugs derived from bis-β- chloroethylylamine [13], drugs based on benzensulfamide derivatives in the presence of degradation products [14], nicotinic acid and its derivatives [15], pyrazole and salicylic acid derivatives [16], sildenafil [17], oxcarbazepine [18]. In the presence of an azido group having a characteristic band in the range of 4.69–4.71 μm, it is possible to identify azidothymidine [19], by a set of absorption bands in the high-, medium- and low-frequency (less than 7.69 μm) regions – synthetic peptide compounds: thymogen, thymodepressin, neogene [20]. The IR spectrum of cordonum (adrenolytic substance) allows to identify it in biological material when performing forensic chemical studies [21].

A variant of IR spectrometry is the method of Raman spectrometry [22]. Raman spectrometry or Raman scattering obtains optical spectra when a substance or material is irradiated with near-infrared monochromatic laser radiation, in which the molecules of the substance are polarized and scatter light in the range from 2.5 to 5000 μm [23, 24]. With such irradiation, the frequency of scattered radiation changes compared to the frequency of radiation incident on the object. Due to this, the analysis can take only 30 seconds. This method allows you to conduct research in addition to the identification of
chemical compounds, as well as on-line monitoring of the reactions of organic synthesis and polymerization in real time; biochemical studies, such as cells in the wild; mineralogical research. That is why this method is increasingly used for drug analysis, including the detection of FD as described in [25]. A comprehensive study using a portable Raman IR spectrometer in the analysis of counterfeit solid dosage forms is given in the publication Dégardin K, Guillemin A, Roggo Y [26].

The main advantages of the method of Raman spectroscopy in comparison with IR spectrophotometry is that it:
- is non-destructive;
- can be made contactless and without opening the original packaging;
- usually does not require sample preparation;
- provides an opportunity to analyze samples in different physical states (solid materials, liquids, and in some cases also gases);
- the analysis is express (from seconds to minutes);
- has the possibility of remote non-contact analysis (for systems with optical fiber);
- has the ability to work with aqueous solutions (no overlap of the H2O signal as in IR spectrometry);
- has the ability to obtain a signal from the depth of the sample, transparent in the selected range, with a penetration depth of from 0.1 to 10 μm (depending on the frequency of the radiation source). Portable Raman spectrometers, which are used both for input API control in pharmaceutical companies and for drug analysis, are becoming more and more widespread. Comparative characteristics of portable Raman spectrometers are given in Table 1.

These devices and the method of Raman spectrometry have proven themselves in the work of the Food and Drug Administration (FDA) and the US customs. The spectral information obtained by this method can be used to identify powders, liquids, gases, aqueous solutions and mixtures thereof. The device on the basis of KR can simultaneously detect up to 5 chemicals in the test sample. The method of Raman spectrometry is especially important for the detection of FD, as it allows you to quickly and quickly detect deviations from the parameters set by the manufacturer directly at the test site.

In order to confirm the possibility of involving the instrument base of higher education institutions, we conducted research on IR spectrometers of a certain list of drugs, including antipyretics and antibiotics that can be used to treat complications of COVID-19. Experimental studies were performed on spectrophotometers of the National University of Water Management in Rivne and Rivne State University for the Humanities. Samples of drugs were purchased in pharmacies in Rivne in the period from 13.03.2020–20.03.2020. The results are presented in Table 2. The table shows the analysis of the drugs determination, including the correlation coefficient R, the determination limit (DL) and the limit of quantification (QDL).

### Table 1

| Model                  | Manufacturer      | Excitation wavelength | Spectral range          | Resolution | Detector                      |
|------------------------|-------------------|-----------------------|-------------------------|------------|-------------------------------|
| NanoRam                | B&W Tek Taiwan    | 785 nm                | 176–2900 cm⁻¹           | 9 cm⁻¹     | TE-Cooled Linear CCD Array    |
| Progeny                | Rigaku            | 1064 nm               | 200–2500 cm⁻¹           | 8–11 cm⁻¹  | 512 pixels, TE cooled InGaAs  |
| Mira DS Advanced       | Metrohm           | 785 nm                | 400–2300 cm⁻¹           | 8–10 cm⁻¹  | CCD-matrix                   |
| Vaya Handheld Raman Spectrometer | Agilent        | 830 nm                | 300–2000 cm⁻¹           | 12–20 cm⁻¹ | CCD-matrix                   |
| TruScan                | Thermo Scientific | 785 nm                | 250–2875 cm⁻¹           | 8–10.5     | liquid nitrogen cooled Germanium |
| BRAVO                  | Bruker            | 1064 nm               | 300–3200 cm⁻¹           | 8–10 cm⁻¹  | liuid nitrogen cooled Germanium |

### Table 2

| Name of the substance | Solvent, wave number, cm⁻¹ | Characteristics |
|-----------------------|-----------------------------|-----------------|
|                       |                             | $R^2$ | DL, mg | QDL, mg | $D$, % |
| Azithromycin          | Toluene, 1700–1750          | 0.999 | 1.0    | 0.2    | 98    |
| Erythromycin          | KBr, 1680–1750              | 0.998 | 0.006  | 0.02   | 99    |
| Ibuprofen             | Chloroform, 1465–1814       | 0.998 | 0.78 μg/ml | 2.6 μg/ml        | 98    |
| Ibuprofen and paracetamol | KBr, 1680–1785 for ibuprofen and 1531–1631 for paracetamol | 0.999 | 0.001 mg/g | 0.013 mg/g | 98    |
| Clarithromycin        | KBr, 1660–1780              | 0.996 | 0.01   | 0.03   | 98    |
| Paracetamol           | KBr, 1001–1802              | 0.999 | 0.005  | 0.019  | 99    |
| Cefuroxime sodium     | KBr, 1470–1610              | 0.999 | 0.16   | 0.6    | 99    |
5. Discussion of research results

According to the results of research of drug analysis by IR spectrophotometry in the transmission mode, it was found that the spectrometers of educational institutions in Rivne meet the requirements of SPbU and the indicator of the limit of determination significantly outweighs the chemical methods of determination. The main disadvantage of IR spectrophotometry of drugs, in our opinion, is that they require sample preparation (deformation, destruction and dissolution, especially for the solid phase). This requires additional legal insurance when transferring samples from territorial services for medicines and drug control to another laboratory (sealing of samples of seized drugs, photoregistration of sample preparation methods, spectrum sampling process, its comparison with the reference, etc.). Comparisons of the obtained results with the results of other researchers indicate that some of the studied substances with the help of our equipment are determined with greater accuracy [27–29]. Deviations of API content correspond to content tolerances in drugs (5–10 %). According to the results of experimental studies by IR spectrophotometry, FD were not detected, which indicates the quality of drugs sold in pharmacies in Rivne. Since these studies were the first to use the equipment of Rivne institutions of higher education for this task, we see further prospects in improving the methods of IR spectrometry to use them for such specific tasks.

6. Conclusions

Today, IR spectrophotometry in terms of reliability, cost of one test and accuracy is one of the first places in the analysis of API and drugs. From the experience of the State Service for Medicines and Drug Control in Rivne region it is known that it is necessary to introduce new, express and mobile methods of detecting counterfeit drugs without any damage to their packaging. That is why in Ukraine it is necessary to implement two significant projects that can minimize and even prevent the entry of counterfeits on the Ukrainian pharmaceutical market, which are associated with the use of new technologies in research. Namely, the introduction of codification of 2D drug packaging – bar code and equipping territorial bodies of state quality control of medicines with portable devices, for example, as illustrated in the article – Raman spectrometers to detect substandard and counterfeit drugs. The use of Raman spectroscopy will not only significantly reduce the time of inspections, but also significantly increase their efficiency. In addition, since this analysis is performed without damaging the original packaging of the medicine, the use of such a device will significantly save money of businesses, because for a normal laboratory analysis, as a rule, requires several packages of drugs.

According to the results of experimental research on the basis of the National University of Water Management of Rivne and Rivne State University for the Humanities, it is expedient and possible to involve the equipment of regional educational institutions and laboratories to detect counterfeits and prevent their use by the health care system.

Conflict of interest

Authors declare no conflict of interest.

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Received date 20.05.2020
Accepted date 18.06.2020
Published date 30.06.2020

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