DATE DESPRE PARTICIPANT

Denumirea firmei	**SA "M-INTER-FARMA"**
Adresa firmei	**Strada Grenoble 23, mun. Chişinău, Moldova MD-2025**
IDNO - Codul fiscal	**1003600005263**
Codul TVA	**O207934**
Oficiul de înregistrare	**Camera Înregistrării de Stat, Chişinău**
Anul întemeierii	**12 mai 1994**
Denumirile precedente	**SA "M-INTER-FARMA"**
Telefoane de contact	**904-005, 904-006**
Fax	**904-007,**
E-mail	**anatolie@minterfarma.md /nina@minterfarma.md**
web pagină	**www.minterfarma.md**
Tipul firmei	**societate pe acţiuni, privată**
Director	**Morozov Maria**
Firme subsidiare	**nu**
Numarul angajaţilor	**16**
Numărul depozitelor	**9**
Baza tehnico materiala	**înzestrată**
Volumul de producţie	**-**
IBAN COD	**MD37EN000000022245246845**
Denumirea băncii în care se deserveşte	**BC "ENERGBANK" SA**
Codul băncii	**ENEGMD22**
Adresa băncii	**Str.Tighina 23/3, mun. Chişinău, Republica Moldova**

L.Ş.

AGENŢIA SERVICII PUBLICE
DEPARTAMENTUL ÎNREGISTRARE ŞI LICENŢIERE A UNITĂŢILOR DE DREPT

EXTRAS

din Registrul de stat al persoanelor juridice
Nr. **413433** data **27.05.2019**

Denumirea completă: **SOCIETATEA PE ACŢIUNI "M-INTER-FARMA".**
Denumirea prescurtată: **"M-INTER-FARMA" S.A.**
Forma juridică de organizare: **Societate pe acţiuni**
Numărul de identificare de stat şi codul fiscal (IDNO): **1003600005263**
Data înregistrării de stat: **12.05.1994**
Modul de constituire: **nou creată.**
Sediul: MD-2028, str. Grenoble, 23, mun. CHIŞINĂU, R. MOLDOVA.
Obiectul principal de activitate:
1. Practica medicală
2. Comerţul cu ridicata al produselor agricole brute şi animalelor vii
3. Comerţul cu amănuntul al produselor farmaceutice
4. Comerţul cu ridicata al produselor farmaceutice
5. Activitatea farmaceutică
6. Alte activităţi de asistenţă medicală
7. Comerţul cu amănuntul în magazine nespecializate, cu vînzare predominantă de produse alimentare, băuturi şi produse din tutun
8. Practica stomatologică
9. Comerţul cu ridicata al parfumurilor şi produselor cosmetice
10. Importul, fabricarea, comercializarea, asistenţa tehnică şi (sau) reparaţia dispozitivelor medicale şi (sau) a opticii
11. Importul şi (sau) fabricarea, depozitarea, comercializarea angro a substanţelor şi materialelor chimice, articolelor şi produselor chimice de menaj.

Capitalul social: 5160000 lei,
Administrator: MOROZOV MARIA, IDNP 0970208486488, tel. tel. 069148966.
Actionari, conform registrului actionarilor "M-INTER-FARMA" S.A.

Prezentul extras este eliberat în temeiul art.34 al Legii nr.220-XVI din 19 octombrie 2007 privind înregistrarea de stat a persoanelor juridice şi a întreprinzătorilor individuali şi confirmă datele din Registrul de stat la data de: **27.05.2019**.

Registrator **Ion MERLICI**

INSTITUŢIA PUBLICĂ "AGENŢIA SERVICII PUBLICE" IDNO 1002600024700 REPUBLICA MOLDOVA CHIŞINĂU 006

EB 0255498

REPUBLICA MOLDOVA

CERTIFICAT DE ÎNREGISTRARE

PRIN PREZENTUL SE CERTIFICĂ , CĂ **SOCIETATEA PE ACŢIUNI "M-INTER-FARMA"** ESTE ÎNREGISTRATĂ LA CAMERA ÎNREGISTRĂRII DE STAT

Numărul de indentificare de stat - codul fiscal

1003600005263

Data înregistrării **12.05.1994**

Data eliberării **17.12.2004**

Iovu Galina, registrator de stat
Funcţia, numele, prenumele persoanei care a eliberat certificatul

semnătura

CAMERA ÎNREGISTRĂRII DE STAT L.Ş. 103

MD 0006726

Anexa nr.7.2 la Instrucţiunea
aprobată prin ordinul IFPS
nr. 400 din 14 martie 2014

CC 04 AE

CERTIFICAT
privind lipsa sau existenţa restanţelor faţă de bugetul public naţional

Nr. № A1947398 **din** от 06.12.2019

1. Destinaţia / Назначение

AGENŢIA SERVICII PUBLICE

2. Date despre contribuabil / Информация о налогоплательщике

Denumirea Наименование	**Codul fiscal / Numărul de identificare** Фискальный код / Идентификационный номер
M-INTER-FARMA S.A.	1003600005263
Adresa sediului de bază (strada, numărul) Адрес основного месторасположения (улица, номер)	**Codul - Denumirea localităţii** Код - Наименование населенного пункта
Grenoble nr.23	0130-SEC.CENTRU

3. Atestarea lipsei sau existenţei restanţelor conform datelor Sistemului Informaţional Automatizat / Подтверждение отсутствия или наличия недоимки согласно данных Информационной автоматизированной системы

La data emiterii prezentului certificat restanţa faţă de bugetul public naţional constituie/ На дату выдачи данной справки недоимка перед национальным публичным бюджетом составляет: **0,00** lei/лей.

4. Valabil pînă la / Действителен до 21.12.2019

5. Autentificarea Serviciului Fiscal de Stat / Подтверждение Государственной налоговой службы

Sef DDF Centru
Funcţia/Должность

Semnătura/Подпись

Viorica CĂUŞ
Numele şi prenumele/Фамилия и имя

L.Ş/ М.П.

Executor: A. Nemerenco
Numele şi prenumele/Фамилия и имя

Este extras din Sistemul Informaţional al SFS SIA „Contul curent al contribuabilului"// 06.12.2019 ora 13:07:11 cu aplicarea prevederilor pct. 82-83 Ordin IFPS nr.400 din 14.03.2014 (Monitorul Oficial 72-77/399, 28.03.2014)

NOTA (0,00)

Declaraţia privind conduita etică şi

neimplicarea în practici frauduloase şi de corupere (F3.4)

[Ofertantul va semna şi va include această declaraţie în oferta depusă. Declaraţia semnată va fi de asemenea inclusă în contractele ofertanţilor cîştigători. Nu se vor permite modificări în formatul formularului, precum şi nu se vor accepta înlocuiri în textul acestuia.]

Data: 19.12.2019

LD nr. ocds-b3wdp1-MD-1576248013641

Către: **IMSP Spitalul de Psihiatrie Bălţi**

"M-INTER-FARMA"SA confirmă prin prezenta că:

1. Nici unul dintre angajaţii, companionii, agenţii, acţionarii, consultanţii, partenerii noştri sau rudele sau asociaţi ai lor nu este în relaţii care ar fi putut considerate ca un conflict de interese, conform prevederilor din documentele de atribuire.
2. În cazul în care vom afla despre faptul unui conflict potenţial, vom raporta imediat informaţia respectivă către autoritatea contractantă.
3. Nici unul dintre angajaţii, companionii, agenţii, acţionarii, consultanţii, partenerii noştri sau rudele sau asociaţii ai lor nu a fost angajat în practici de corupere, escrocherie, complotare, constrîngere sau alte practici anticoncurenţiale în procesul pregătirii ofertei din cadrul prezentei licitaţii, conform prevederilor din documentele de atribuire, punctul IPO10.
4. În legătură cu procedura respectivă de licitaţie şi cu orice contract care, eventual, ne va fi adjudecat ca rezultat al acesteia, nu au fost, nici nu vor fi efectuate nici un fel de plăţi către angajaţii, companionii, agenţii, acţionarii, consultanţii, partenerii noştri sau rudele lor, care sînt implicaţi în achiziţia publică, implementarea contractului şi aprobarea plăţilor contractuale în numele autorităţii contractante.

Semnat:_____

Nume: Palii Nina

Funcţia în cadrul companiei: jurist

Denumirea companiei: "M-INTER-FARMA" SA

L.Ş.

Carestream

DECLARATION OF CONFORMITY

Carestream Health, Inc., hereby declares that the product(s) listed are made in conformity with: Medical Device Directive [Directive 93/42/EEC], ANNEX II Conformity Assessment Procedure and the Australian Therapeutic Goods (Medical Devices) Regulations 2002, Clause 1.8 of Schedule 3.

Manufacturer's Name and Address:	Carestream Health, Inc. 150 Verona Street Rochester, New York, USA 14608
Medical Device:	X-ray Film, Sheet
Product List:	X-OMAT BT Film Medical X-ray Blue / MXB Film Medical X-ray Blue / MXBE Film Medical X-ray Green / MXG Film T-MAT G/RA Film T-MAT L/RA Film INSIGHT Pediatric Film INSIGHT Thoracic Film X-SIGHT G/RA Film MIN-R EV Film MIN-R S Film MIN-R 2000 Plus Film L Green X-ray Film WB HG Green X-ray Film WB Green X-ray Film WB Full Blue X-ray Film WB Mammo Film WB —End of List—
Device Classification:	Europe - Class IIa, ANNEX IX , Rule 16 Australia - Class IIa, Schedule 2, Part 5, Rule 5.4
GMDN Code and Term:	40979. Medical x-ray film, screen
Scope of Application	All declared products

Each kind of medical device to which the quality-management system has been applied complies with the applicable provisions of the essential principles, the classification rules, and the full-quality assurance procedures at each stage, from the design of the device until its final inspection before being supplied.

Quality-Management-System	Certified to EN ISO 13485 by BSI No. FM 701584 BSI No. FM 507315 DNV No. 245266-2017-AQ-USA-NA-PS Rev. 1 BSI No. FM 46141 TUV No. Q2N 17 11 61500 005
European Notified Body:	British Standards Institute, BSI (2797)
Full-Quality-Assurance-System Certificate (CE):	BSI Certificate Number CE 01233
European Authorized Representative:	Carestream Health France 1, rue Galilée 93192 NOISY-LE-GRAND CEDEX FRANCE

The relevant sections of the following European Union Harmonized standards apply to the listed product(s):

EN ISO 14971
EN 1041
EN ISO 15223-1
EN 62366

Treya D. R. Collins
Director, Global Quality Systems
Carestream Health, Inc.
150 Verona Street
Rochester, New York 14608, USA
Telephone: +1-970-304-4654

Carestream

DECLARATION OF CONFORMITY

Carestream Health, Inc., hereby declares that the product(s) listed are made in conformity with:

Medical Device Directive [Directive 93/42/EEC], ANNEX VII Conformity Assessment Procedure and the Australian Therapeutic Goods (Medical Devices) Regulations 2002, Clause 6.6 of Schedule 3.

Manufacturer's Name and Address:

Carestream Health, Inc.
150 Verona Street
Rochester, New York, USA 14608

Medical Device:

Medical Imaging Photoprocessing Devices - Photochemicals

Product List:

Chemistry used to develop or fix the image on medical and dental X-ray films:
GBX Developer and Replenisher
GBX Fixer and Replenisher
GBX Twin Pack
X-OMAT MX Fixer and Replenisher
X-OMAT MX Developer and Replenisher
READYMATIC Developer and Replenisher
READYMATIC Fixer and Replenisher
READYMATIC Chem Pack
RP X-OMAT Developer and Replenisher
RP X-OMAT LO Fixer and Replenisher
X-OMAT EX II Developer and Replenisher
X-OMAT Developer Starter
X-OMAT LE+ Developer and Replenisher
X-OMAT LE+ Fixer and Replenisher
CARESTREAM DENTAL X-ray Monobath
CARESTREAM DENTAL X-ray Developer
CARESTREAM DENTAL X-ray Fixer
Rapid Access Twin Pack
Rapid Access Developer
Rapid Access Fixer
XCE Developer and Replenisher
XCF Fixer and Replenisher
XPE Developer
XPF Fixer
—End of List—

Device Classification:

Europe - Class I, ANNEX IX, Rule 1
Australia - Class I, Schedule 2, Part 2, Rule 2.1

GMDN Code and Term:

41009 Radiographic film processing chemical, automated
41008 Radiographic film processing chemical, manual

Scope of Application All declared products

Each kind of medical device to which the Declaration of Conformity (not requiring assessment by the Secretary) procedures have been applied complies with the applicable provisions of the essential principles and the classification rules before being supplied

European Authorized Representative: Carestream Health France
1, rue Galilée
93192 NOISY-LE-GRAND CEDEX
FRANCE

The relevant sections of the following European Union Harmonized standards apply to the listed product(s):

EN 1041
EN ISO 15223-1
EN ISO 14971
EN 62366

Kevin C. Wright
Senior Director, Worldwide Regulatory Affairs
Carestream Health, Inc.
150 Verona Street
Rochester, New York 14608, USA
Telephone: +1-585-627-6878

bsi.

By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Carestream Health, Inc
150 Verona Street
Rochester
New York
14608
USA

Holds Certificate No: **FM 701584**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

The design, manufacture, distribution, (integration, installation and servicing excluding film products) of diagnostic image recording devices, photo chemicals, medical and dental imaging systems, cone beam computed tomography, information technology software for healthcare information systems and medical imaging and detection. Manufacture, service, installation and distribution of Dry View Printers. Storage, Handling, Packaging and Distribution of Pharmaceutical Products.

For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk - Medical Devices

Original Registration Date: 2008-12-20
Latest Revision Date: 2019-09-24

Effective Date: 2019-10-03
Expiry Date: 2022-10-02

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By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. **CE 01233**

Issued To: **Carestream Health, Inc.**
150 Verona Street
Rochester
New York
14608
USA

In respect of:

The design, development and manufacture of diagnostic image recording devices including storage phosphor screens and reader systems, medical x-ray films, direct digital radiography systems, dental x-ray systems including film, dental and medical imaging equipment, and medical imaging and PACS Software. Those aspects of metrology related to the design and manufacture of dimensional measuring PACS software.

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 2797):

Albert Roossien, Regulatory Lead

First Issued: **1996-03-06** Date: **2019-02-27** Expiry Date: **2021-03-05**

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EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No: **CE 01233**

Date: **2019-02-27**

Issued To: **Carestream Health, Inc.**
150 Verona Street
Rochester
New York
14608
USA

Subcontractor:	Service(s) supplied
Agfa-Gevaert HealthCare GmbH Bürgermeister-Götz-Str. 10 Schrobenhausen 86529 Germany	**Manufacture**
Algotec Systems Ltd 2 Hapnina Street PO BOX 46 43107 Ra'anana Israel	**Design** **Development** **Software**
Analogic Corporation 8 Centennial Drive Peabody Massachusetts 01960 USA	**Design** **Manufacture**

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By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 01233**
Date:	**2019-02-27**
Issued To:	**Carestream Health, Inc.** **150 Verona Street** **Rochester** **New York** **14608** **USA**

Subcontractor:	**Service(s) supplied**
Carestream Health France 1, rue Galilée 93192 NOISY-LE-GRAND CEDEX France	**EU Representative**
Carestream Health, Inc 5450 Campus Drive Canandaigua New York 14424 USA	**Manufacture**
Carestream Health, Inc. 1049 West Ridge Road Rochester New York 14615 USA	**Design** **Development** **Manufacture**

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EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 01233**
Date:	**2019-02-27**
Issued To:	**Carestream Health, Inc.** **150 Verona Street** **Rochester** **New York** **14608** **USA**

Subcontractor:	**Service(s) supplied**
Carestream Health, Inc. 1669 Lake Avenue Rochester New York 14652 USA	**Design** **Development** **Manufacture**
Carestream Health, Inc. 1964 Lake Ave Rochester New York 14615 USA	**Design**
Carestream Health, Inc. 2000 Howard Smith Avenue West Windsor Colorado 80550 USA	**Manufacture**

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EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 01233**
Date:	**2019-02-27**
Issued To:	**Carestream Health, Inc.** **150 Verona Street** **Rochester** **New York** **14608** **USA**

Subcontractor:	**Service(s) supplied**
Carestream Health, Inc. 8124 Pacific Avenue White City Oregon 97503 USA	**Manufacture**
Carestream Health, Inc. Global R & D Center (Shanghai) No. 27 Xinjinqiao Road Shanghai 201206 China	**Design** **Development**
Communication & Power Industries Canada Inc. 45 River Drive Georgetown Ontario L7G 2J4 Canada	**Manufacture**

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By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No: **CE 01233**

Date: **2019-02-27**

Issued To: **Carestream Health, Inc.**
150 Verona Street
Rochester
New York
14608
USA

Subcontractor:	Service(s) supplied
Micro-X Ltd 1284 South Road Clovelly Park South Australia 5042 Australia	**Design** **Manufacture**
Rayco (Shanghai) Medical Products Company Limited Building 7, No. 1510 Chuanqiao Road China (Shanghai) Pilot Free Trade Z Shanghai 201206 China	**Manufacture**

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By Royal Charter

EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 01233**
Date:	**2019-02-27**
Issued To:	**Carestream Health, Inc.** **150 Verona Street** **Rochester** **New York** **14608** **USA**

Subcontractor:	**Service(s) supplied**
Rayco (Xiamen) Medical Products Company Limited 308 Wengjiao Road Haicang District Xiamen Fujian 361022 China	**Manufacture**
ScImage, Inc. 4916 El Camino Real, Suite 200 Los Altos California 94022 USA	**Design** **Development** **Software**

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EC Certificate - Full Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No:	**CE 01233**
Date:	**2019-02-27**
Issued To:	**Carestream Health, Inc.** **150 Verona Street** **Rochester** **New York** **14608** **USA**

Subcontractor:	**Service(s) supplied**
Soluciones Médicas Exportación S de RL de CV Calle Anillo Periferico Poniente No. 3100 Colonia Paraísos Del Colli Zapopan Jalisco 45069 Mexico	**Manufacture**
Varian Medical Systems, Inc. X-Ray Products 1678 South Pioneer Road Salt Lake City Utah 84104 USA	**Manufacture**

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By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 9001:2008

This is to certify that:

Carestream Health, Inc
150 Verona Street
Rochester
New York
14608
USA

Holds Certificate No: **FM 537916**

and operates a Quality Management System which complies with the requirements of ISO 9001:2008 for the following scope:

The design, development, manufacture, and service of digital radiography imaging systems (such as INDUSTREX digital systems) and accessories for the non-destructive testing industry.

For and on behalf of BSI:

Carlos Pitanga, SVP, System Certification and Compliance

Original Registration Date: 07/17/2008
Latest Revision Date: 10/03/2016

Effective Date: 10/21/2016
Expiry Date: 09/14/2018

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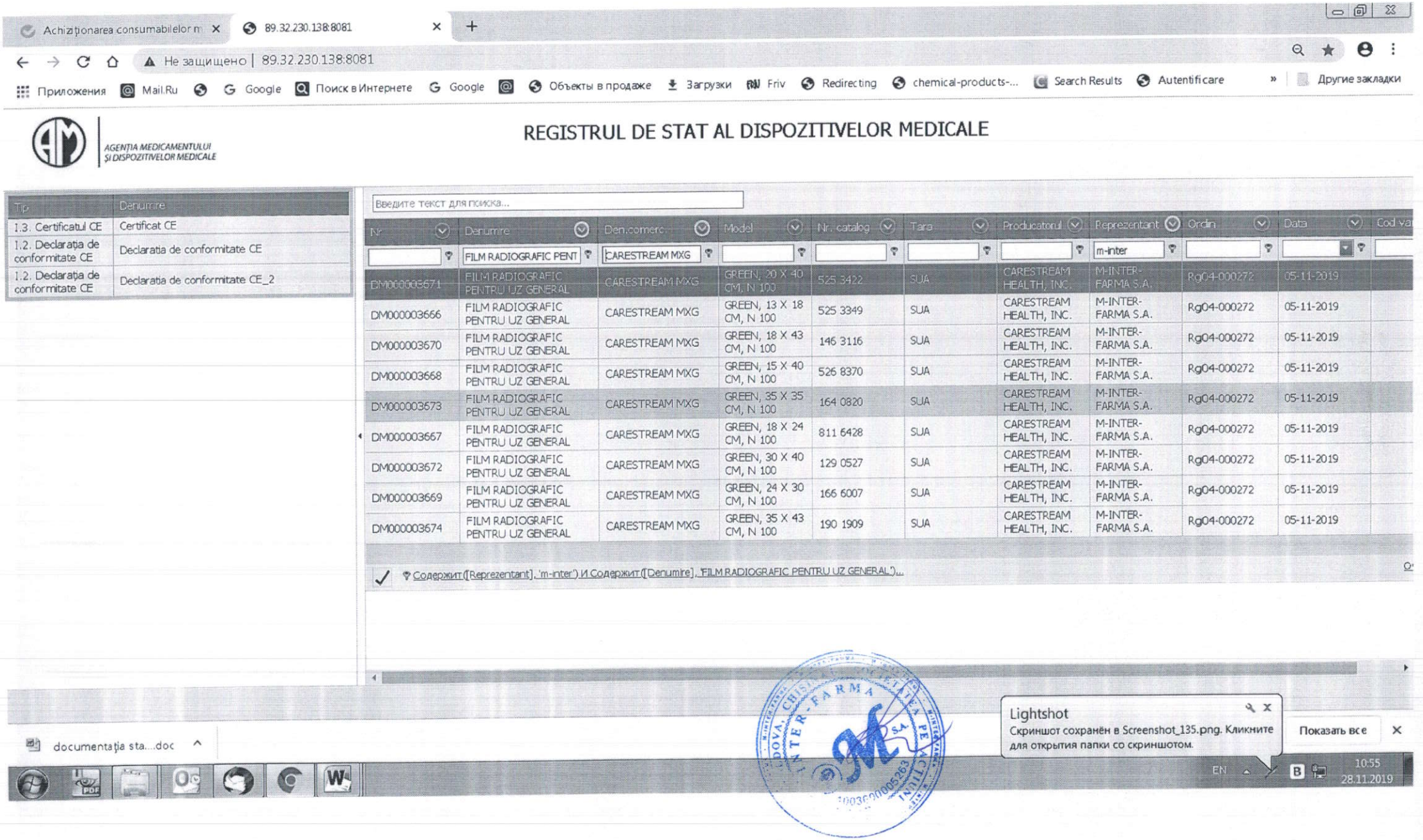
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FILM RADIOGRAFIC PENT
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DM000003671 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 20 X 40 CM, N 100 525 3422 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003666 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 13 X 18 CM, N 100 525 3349 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003670 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 18 X 43 CM, N 100 146 3116 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003668 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 15 X 40 CM, N 100 526 8370 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003673 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 35 X 35 CM, N 100 164 0820 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003667 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 18 X 24 CM, N 100 811 6428 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003672 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 30 X 40 CM, N 100 129 0527 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003669 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 24 X 30 CM, N 100 166 6007 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
DM000003674 FILM RADIOGRAFIC PENTRU UZ GENERAL CARESTREAM MXG GREEN, 35 X 43 CM, N 100 190 1909 SUA CARESTREAM HEALTH, INC. M-INTER-FARMA S.A. Rg04-000272 05-11-2019
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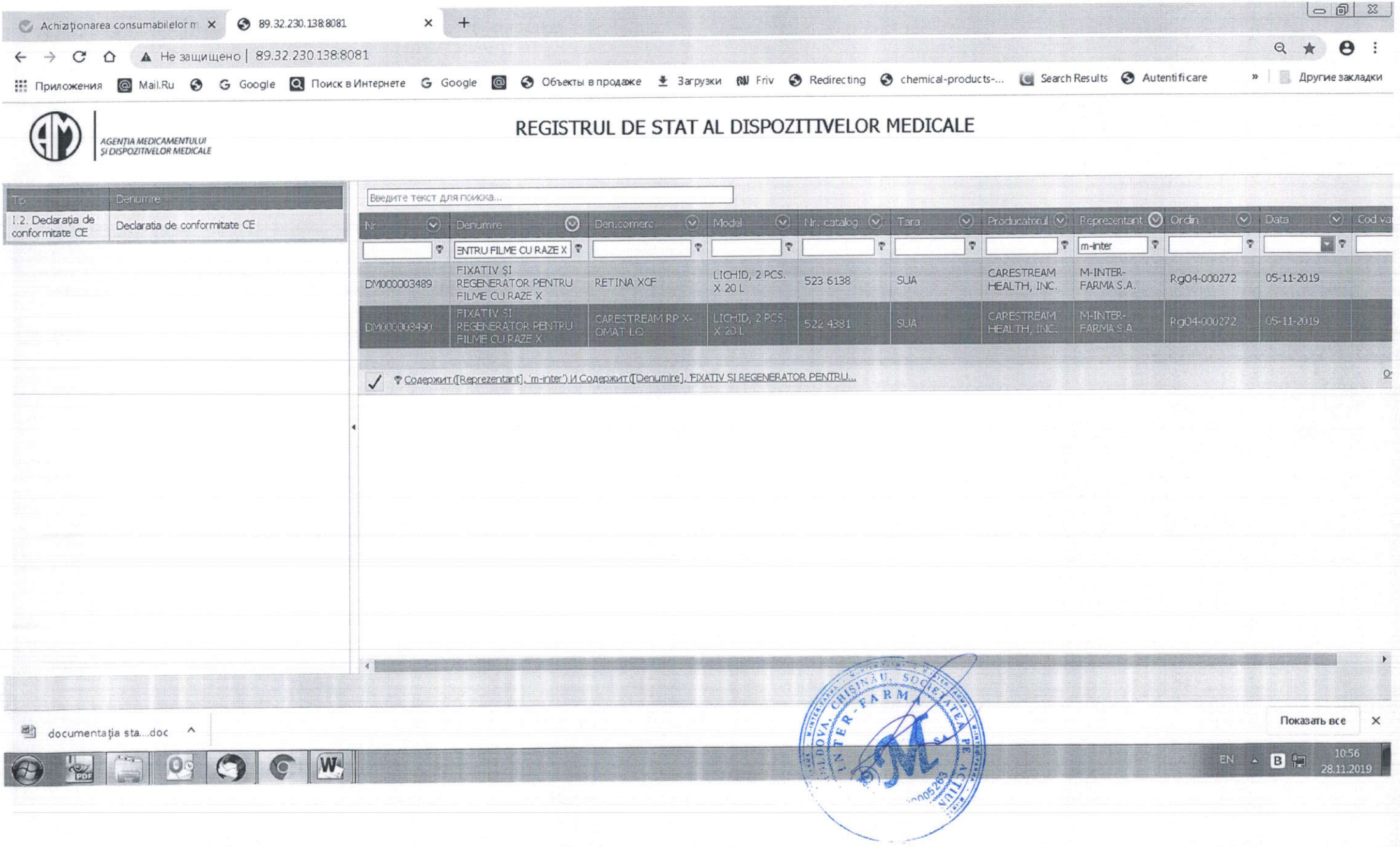
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RETINA XCF
LICHID, 2 PCS. X 20 L
523 6138
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
05-11-2019
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FIXATIV ȘI REGENERATOR PENTRU FILME CU RAZE X
CARESTREAM RP X-OMAT LO
LICHID, 2 PCS. X 20 L
522 4381
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
05-11-2019
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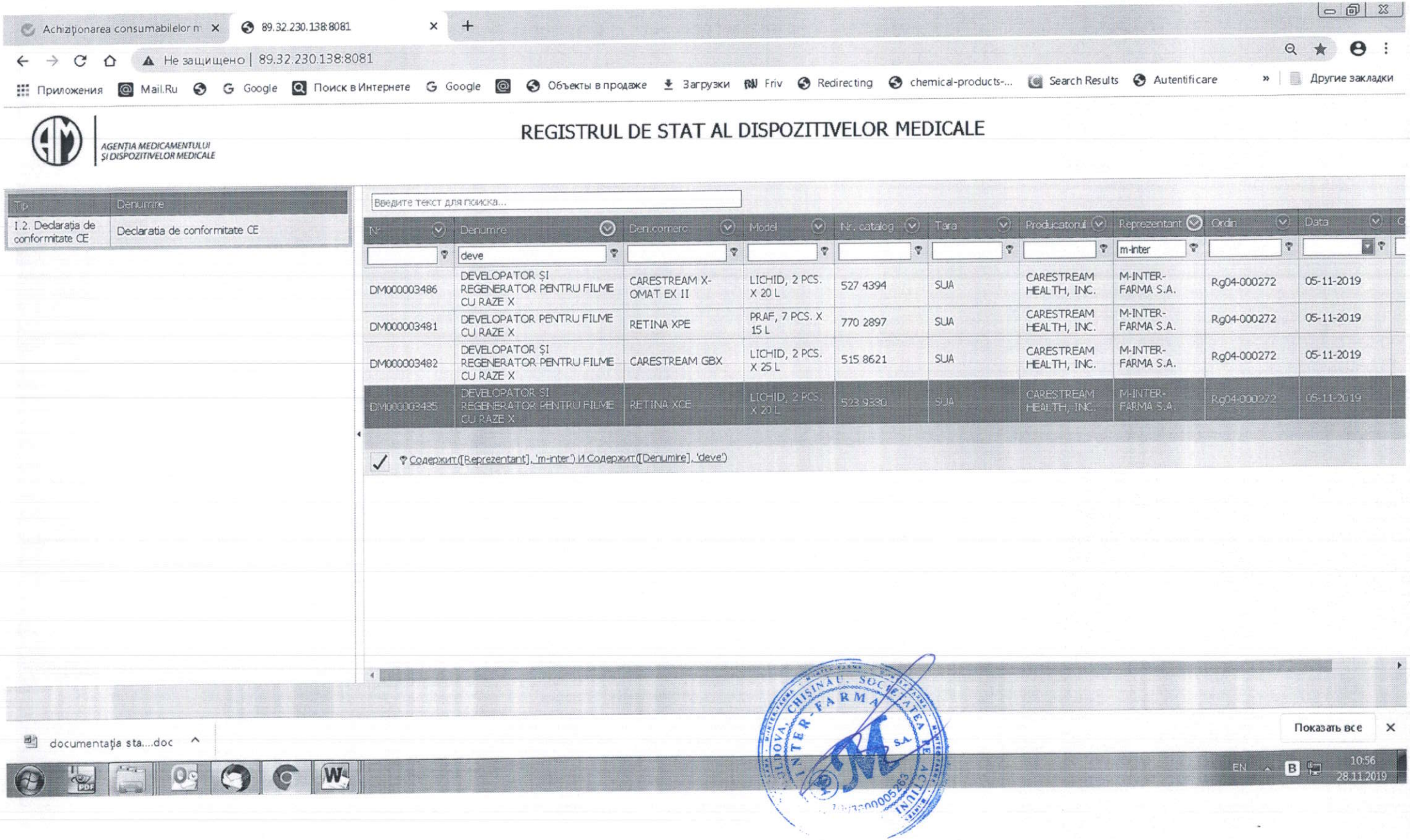
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DEVELOPATOR ȘI REGENERATOR PENTRU FILME CU RAZE X
CARESTREAM X-OMAT EX II
LICHID, 2 PCS. X 20 L
527 4394
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
05-11-2019
DM000003481
DEVELOPATOR PENTRU FILME CU RAZE X
RETINA XPE
PRAF, 7 PCS. X 15 L
770 2897
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
05-11-2019
DM000003482
DEVELOPATOR ȘI REGENERATOR PENTRU FILME CU RAZE X
CARESTREAM GBX
LICHID, 2 PCS. X 25 L
515 8621
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
05-11-2019
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DEVELOPATOR ȘI REGENERATOR PENTRU FILME CU RAZE X
RETINA XCE
LICHID, 2 PCS. X 20 L
523 9380
SUA
CARESTREAM HEALTH, INC.
M-INTER-FARMA S.A.
Rg04-000272
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