

Formularul ofertei (F3.1)

[Ofertantul va completa acest formular în conformitate cu instrucțiunile de mai jos. Nu se vor permite modificări în formatul formularului, precum și nu se vor accepta înlocuiri în textul acestuia.]

Data depunerii ofertei: **25.11.2019**

Către: **IMSP Institutul Mamei și Copilului**

SA "M-INTER-FARMA" declară că:

- a) Au fost examinate și nu există rezervări față de documentele de atribuire, inclusiv modificările nr.
 - a) SA "M-INTER-FARMA" se angajează să furnizeze/presteze, în conformitate cu documentele de atribuire și condițiile stipulate în specificațiile tehnice și preț, următoarele bunuri și/sau servicii **Filme radiologice.**
 - a) Suma totală a ofertei **fără TVA** constituie **133 500,00 (o sută treizeci și trei mii cinci sute lei 00 bani)**
 - b) Suma totală a ofertei **cu TVA** constituie **160 200,00 (o sută șaiszeci mii două sute 00 bani)**
- c) Prezenta ofertă va rămâne valabilă pentru perioada de timp specificată în **FDA4.8.**, începînd cu data-limită pentru depunerea ofertei, în conformitate cu **FDA5.2.**, va rămîne obligatorie și va putea fi acceptată în orice moment pînă la expirarea acestei perioade;
În cazul acceptării prezentei oferte, SA "M-INTER-FARMA" se angajează să obțină o Garanție de bună execuție în conformitate cu **FDA7**, pentru executarea corespunzătoare a contractului de achiziție publică.
- d) Nu sîntem în nici un conflict de interese, în conformitate cu punctul **IPO5.4.**
- e) Compania semnatară, afiliații sau sucursalele sale, inclusiv fiecare partener sau subcontractor ce face parte din contract, nu au fost declarate neeligibile în baza prevederilor legislației în vigoare sau a regulamentelor cu incidență în domeniul achizițiilor publice, în conformitate cu punctul **IPO5.5.**

Semnat:

[semnătura persoanei autorizate pentru semnarea ofertei]

L.Ș.

Nume: Palii Nina

În calitate de: Specialist achiziții publice

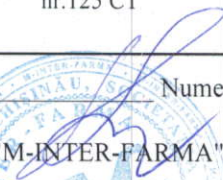
Ofertantul: SA "M-INTER-FARMA"

Adresa: mun. Chișinău, str. Grenoble 23

Data: 25.11.2019

Specificații tehnice (F4.1)

Numărul procedurii de achiziție ocds-b3wdp1-MD-1573824659435 din 25.11.2019							
Denumirea procedurii de achiziție: Filme radiologice							
Co d CP V	Denumirea bunurilor	Modelul articolului	Țara de origine	Produ- cătorul	Specificarea tehnică deplină solicitată de către autoritatea contractantă	Specificarea tehnică deplină propusă de către oferant	Standa rde de referin ță
1	2	3	4	5	6	7	8
3310000 0-1	Film radiologic digital 35x43 nr.125 CT	35x43 nr.125 CT	SUA	Carestream Health, INC	Film radiologic digital 35x43 nr.125 CT	Film radiografic pentru imprimare laser DRYVIEW DVB+, 35X43 cm, № 125 sheet	CE, ISO, înreg. AMED

Semnat:  Numele, Prenumele: Palina Nina În calitate de: specialist achizitii publice

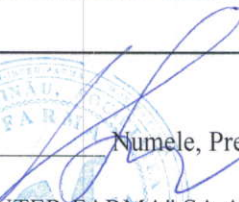
Oferantul: "M-INTER-FARMA" SA Adresa: mun. Chișinău, str. Grenoble 23



Specificații de preț (F4.2)

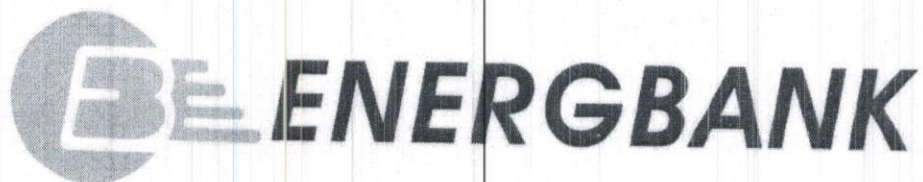
[Acest tabel va fi completat de către ofertant în coloanele 5,6,7,8, iar de către autoritatea contractantă – în coloanele 1,2,3,4,9,10]

Numărul procedurii de achiziție ocds-b3wdp1-MD-1573824659435 din 25.11.2019								
Denumirea procedurii de achiziție: Filme radiologice								
Cod CPV	Denumirea bunurilor	Cantitatea	Unitatea de măsură	Preț unitar (fără TVA)	Preț unitar (cu TVA)	Suma	Suma	Termenul de Livrare
Bunuri								
1.	Film radiologic digital 35x43 nr.125 CT	buc	18	7416,67	8900,00	133500,00	160200,00	În decurs de 10 zile de la comanda cumpărătorului
Total						133500,00	160200,00	

Semnat:  Numele, Prenumele: Pali Nina În calitate de: specialist achizitii publice

Ofertantul: "M-INTER-FARMA" SA Adresa: mun. Chișinău, str. Grenoble 23





Banca Comercială „ENERGBANK” Societate pe Acțiuni
Republica Moldova, MD-2001, mun. Chișinău, str. Tighina 23/3
tel.: +(373 22) 544-377, 858-000; fax: +(373 22) 858-080
www.energbank.com; e-mail: office@energbank.com
IDNO 1003600008150, cod TVA 0202040
codul băncii: ENEGMD22
cod IBAN: MD10NB00000000035215845 la CD BNM
Capital social - 100 000 000 lei MD

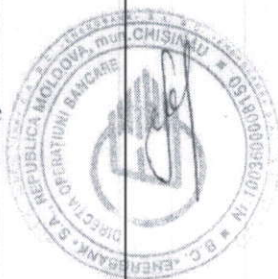
Nr. 13/01-08/1342
din «21» ianuarie 2019

S.A. “M-Inter-Farma”
mun. Chișinău, str. Grenoble, 23

Prin prezenta B.C. „ENERGBANK” S.A. (codul băncii ENEGMD22) confirmă că S.A. “M-Inter-Farma”, cod fiscal 1003600005263 deține contul IBAN MD37EN00000022245246845 în MDL, USD, EUR, RUB, UAH, GBP, CHF. .

Prezentul certificat este eliberat pentru a fi prezentat la solicitare.

Șef Direcției Operațiuni Bancare



Culiuc Svetlana





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: Non-X-ray Film, Sheet

Product List: DRYVIEW DVB Laser Imaging Film
DRYVIEW DVB+ Laser Imaging Film
DRYVIEW DVB+ Premium Laser Imaging Film
DRYVIEW DVC Laser Imaging Film
DRYVIEW DVE Laser Imaging Film
DRYVIEW DVM Mammography Laser Imaging Film
DRYVIEW DVM+ Mammography Laser Imaging Film
—End of List—

Device Classification: Europe - Class I, ANNEX IX, Rule I
Australia - Class I, Schedule 2, Part 2, Rule 2. 1

GMDN Code and Term: 40980. Medical x-ray film, non-screen

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.

Quality-Management-System: Certified to EN ISO 13485 by:
BSI No. FM 701584
BSI No. FM 46141
TUV No. Q2N 17 11 61500 005
BSI No. FM 507315

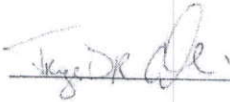


European Authorized Representative:

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):

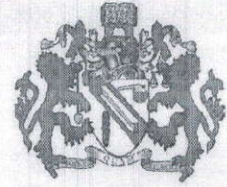
EN ISO 14971
EN 1041
EN ISO 15223-1
EN 62366



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



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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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Page 1 of 1

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

This certificate was issued electronically and is bound by the conditions of the contract.

Information and Contact: BSI, Bay Building, John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands Tel: + 31 20 346 0780
BSI Group The Netherlands B.V. registered in The Netherlands under 33264284.
A member of BSI Group of Companies.



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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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EC Certificate - Full Quality Assurance System

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

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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 Gallée 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

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

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Rochester
New York
14608
USA**

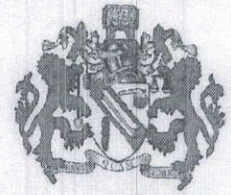
Subcontractor:	Service(s) supplied
Carestream Health, Inc. 8124 Pacific Avenue White City Oregon 97503 USA	Manufacture
Carestream Health, Inc. Global R & D Center (Shanghai) No. 27 Xinjinqiao Road Shanghai 201206 China	Design Development
Communication & Power Industries Canada Inc. 45 River Drive Georgetown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

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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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List of Significant Subcontractors

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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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S.A. M-INTER-FARMA

IDNO 1003600005263 Cod OKPO 37020536 Cod TVA 0207934
MD 2025, Chișinău, strada Grenoble 23
Tel 00 373 22/904-005, 904-006, 904-007

Nr. 179 din 25 noiembrie 2019

IMSP Institutul Mamei și Copilului
Către grupul de lucru la

LD nr. ocds-b3wdp1-MD-1573824659435 din 25.11.2019

DEMERS

Prin prezenta "M-INTER-FARMA" SA confirma ca, va prezenta mostrele produselor, oferite în cadrul LD nr. ocds-b3wdp1-MD-1573824659435 din 25.11.2019 privind achiziționare de "Filme radiologice" la solicitare autorității contractante.

Cu respect,
Director SA "M- INTER-FARMA"



Maria Morozov



S.A. M-INTER-FARMA

IDNO 1003600005263 Cod OKPO 37020536 Cod TVA 0207934
MD 2025, Chișinău, strada Grenoble 23
Tel 00 373 22/904-005, 904-006, 904-007

Nr. 178 din 25 noiembrie 2019

IMSP Institutul Mamei și Copilului
Către grupul de lucru la
LD nr. ocds-b3wdp1-MD-1573824659435 din 25.11.2019

DEMERS

Prin prezenta "M-INTER-FARMA" SA confirma ca, la solicitarea autorității contractante va prezenta actele, conform DUAЕ.

Cu respect,
Director SA "M- INTER-FARMA"



Maria Morozov