DECLARATION OF CONFORMITY
according to ISO/IEC 17050-1 and EN 17050-1

DoC #: HSD-0002-Q- R1 Original/en

Manufacturer's Name: **HP Inc.**
Manufacturer's Address: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

declare, under its sole responsibility that the product

Product Name and Model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulatory Model Number:[1)] HSD-0002-Q
Product Options: None

conforms to the following Product Specifications and Regulations:

Safety
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ecodesign
Regulation (EC) No. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

The product herewith complies with the requirements of the Low Voltage Directive 2014/35/EU, the EMC Directive 2014/30/EU, the Ecodesign Directive 2009/125/EC, the RoHS Directive 2011/65/EU and carries the CE-marking accordingly.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Additional Information:

1) This product is assigned a Regulatory Model Number which stays with the regulatory aspects of the design. The Regulatory Model Number is the main product identifier in the regulatory documentation and test reports, this number should not be confused with the marketing name or the product numbers.
2) This product was tested in a typical HP environment.
3) The compliance statement for EU/EFTA/Turkey is legally binding in the Union from the date when the product is placed in the market. Supporting documentation and declarations of compliance distributed before such a date are shared on confibential basis and may contain business critical information. Please do not redistribute without explicit consent of HP Inc.

Palo Alto, CA
07-09-2018

HP Inc.
1501 Page Mill Road, Palo Alto, CA
Gilles Soulard, Manager
Product Compliance Center

Gilles Soulard, Manager
Product Compliance Center

Local contact for regulatory topics only:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
http://www.hp.eu/certificates

Click here to go to the translation(s)

KONFORMITÄTSERKLÄRUNG

nach ISO / IEC 17050-1 und EN 17050-1

DoC #: HSD-0002-Q-R1Übersetzung/de

Herstellername: **HP Inc.**
Adresse des Herstellers: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

erklärt, dass das Produkt

Produktname und Modell:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulatorische Model Nummer:[1)] HSD-0002-Q
Produktoptionen: Keine Optionen

entspricht den folgenden Produktspezifikationen und Vorschriften:

Sicherheit:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMV
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Öko-Design
Verordnung (EG) Nr. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Das Produkt erfüllt die Anforderungen der Niederspannungsrichtlinie 2014/35/EU, der EMV-Richtlinie 2014/30/EU, die Ökodesign-Richtlinie 2009/125/EG, die RoHS-Richtlinie 2011/65/EU und trägt das CE-Kennzeichnung entsprechend.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Zusätzliche Information:

1) Für die regulatorischen Aspekte zum Design wurde diesem Produkt eine Regulatorische Model Nummer zugeordnet. Zur Produktidentifizierung in der regulatorischen Dokumentation und den Prüfberichten wird diese Regulatorische Model Nummer verwendet und sollte nicht zu Verwechslungen von Marketingnamen oder Produktnummern führen.
2) Dieses Produkt wurde in einer typischen HP Konfiguration getestet.

Palo Alto, CA
07-09-2018

überprüfen Sie die Unterschrift auf der ursprünglichen Erklärung beigefügt
Gilles Soulard, Manager
Product Compliance Center

Lokale Ansprechpartner für Richtlinien und Bestimmungen:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DÉCLARATION DE CONFORMITÉ

selon la norme ISO / IEC 17050-1 et EN 17050-1

DoC #: HSD-0002-Q-R1Traduction/fr

Nom du fabricant: **HP Inc.**
Adresse du fabricant: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

déclare que le produit

Nom du produit et le modèle:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Numéro de modèle réglementaire:[1)] HSD-0002-Q
Options du produit: Pas d'options

est conforme aux normes et règlements de produit suivantes:

Sécurité:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

écoconception
Règlement (UE) n ° 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Ce produit est conforme aux exigences de la Directive Basse Tension 2014/35/UE, la directive CEM 2014/30/UE, la directive sur l'écoconception 2009/125/CE, la directive RoHS 2011/65/UE et porte la marque CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Informations complémentaires:

1) A ce produit est assigné un numéro de modèle réglementaire qui reste avec les aspects réglementaires de la conception. Le numéro de modèle réglementaire est le principal identificateur de produit dans la documentation réglementaire et les rapports d'essais, ce nombre ne doit pas être confondu avec le nom commercial ou les numéros de produit.
2) Ce produit a été testé dans un environnement typique HP en conjonction avec un système hôte HP.

Palo Alto, CA
07-09-2018

vérifier la signature sur la déclaration d'origine annexé
Gilles Soulard, Manager
Product Compliance Center

Contact local pour les réglementations uniquement:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DICHIARAZIONE DI CONFORMITA'
secondo la norma ISO / IEC 17050-1 e EN 17050-1

DoC #: HSD-0002-Q-R1Traduzione/it

Nome del Fornitore: **HP Inc.**
Indirizzo del Fornitore: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

dichiara che il prodotto

Nome del prodotto e modello:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Numero di modello normativo:[1)] HSD-0002-Q
Opzioni del prodotto: Nessuna opzione

è conforme alle seguenti specifiche e regolamenti di prodotto:

Sicurezza
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ecodesign
Regolamento (CE) No. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Questo prodotto è conforme ai requisiti della direttiva Bassa Tensione 2014/35/UE, la direttiva EMC 2014/30/UE, la direttiva sulla Progettazione Ecocompatibile 2009/125/CE, la direttiva RoHS 2011/65/UE e porta il marchio CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Informazioni aggiuntive:

1) A questo prodotto è assegnato un Numero di modello normativo (RMN) che rimane con gli aspetti normativi della progettazione. Il numero di modello normativo è l'identificativo principale del prodotto nella documentazione normativa e rapporti di prova, questo numero non deve essere confuso con il nome commerciale o il numero di prodotto.
2) Questo prodotto è stato testato in un tipico ambiente HP.

Palo Alto, CA
07-09-2018

controllare la firma sulla dichiarazione originale allegata
Gilles Soulard, Manager
Product Compliance Center

Contatto locale solo per informazioni sulla conformità al marchio CE:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501

www.hp.eu/certificates

DECLARACIÓN DE CONFORMIDAD

según la norma ISO / IEC 17050-1 y EN 17050-1

DoC #: HSD-0002-Q-R1Traducción/es

Nombre del fabricante: **HP Inc.**
Dirección del fabricante: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

declara que el producto

Nombre del producto y modelo:[2] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Número de modelo reglamentario:[1] HSD-0002-Q
Opciones del producto: Sin opciones

cumple con las siguientes especificaciones y normas de productos:

Seguridad:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Diseño ecológico
Reglamento (CE) No. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Este producto cumple con los requisitos de la Directiva de baja tensión 2014/35/UE, la Directiva EMC 2014/30/UE, la Directiva sobre diseño ecológico 2009/125/CE, la Directiva RoHS 2011/65/UE y lleva la marca CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Información Adicional:

1) A este producto se asigna un número de modelo regulatorio que cumple con los aspectos regulatorios del diseño. El número de modelo normativo es el identificador principal del producto en la documentación reglamentaria e informes de ensayo, este número no se debe confundir con el nombre comercial o los números del producto.
2) Este producto se ha probado en un entorno típico de HP.

Palo Alto, CA
07-09-2018

comprobar la firma de la primera declaración anexa
Gilles Soulard, Manager
Product Compliance Center

Contacto local únicamente para temas de normativa:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DEKLARACJA ZGODNOŚCI

zgodnie z normą ISO / IEC 17050-1 i EN 17050-1

DoC #: HSD-0002-Q-R1Tłumaczenie/pl

Nazwa producenta: **HP Inc.**
Adres producenta: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

oświadcza, że produkt

Nazwa produktu i modelu:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Numer modelu:[1)] HSD-0002-Q
Opcje produktu: Brak opcji

jest zgodny z następującymi specyfikacjami produktu i rozporządzeń:

Bezpieczeństwo:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ekoprojektu
Rozporządzenie (WE) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Niniejszy produkt jest zgodny z wymaganiami Dyrektywa o niskim napięciu 2014/35/WE, Dyrektywa EMC 2014/30/WE, Dyrektywa w sprawie ekoprojektu 2009/125/WE, Dyrektywa RoHS 2011/65/WE i posiada odpowiednio oznakowanie CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Dodatkowe informacje:

1) Produkt ten został przypisany numer modelu z aspektów regulacyjnych projekt. Numer modelu jest głównym identyfikatorem produktu w dokumentacji normatywnej i raporty z badań, liczba ta nie powinna być mylona z nazwą handlową lub też liczby produktów.
2) Produkt był testowany w typowym środowisku HP, w połączeniu z systemem gospodarza HP.

Palo Alto, CA
07-09-2018

sprawdzić podpis na oryginale deklaracji załączonej
Gilles Soulard, Manager
Product Compliance Center

Lokalny kontakt na temat przepisów:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DECLARATIE DE CONFORMITATE

în conformitate cu ISO / IEC 17050-1 și EN 17050-1

DoC #: HSD-0002-Q-R1Traducere/ro

producătorului Nume: **HP Inc.**
Adresa producătorului: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

declară că produsul

Nume produs și model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Numărul de model de reglementare:[1)] HSD-0002-Q
Opțiuni de produse: Nu există opțiuni

în conformitate cu următoarele specificații și regulamentelor de produse:

Siguranță:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ecodesign
Regulamentul (CE) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Produsul prezentat aici, corespunde cu cerintele Directiva de joasă tensiune 2014/35/UE, la Directiva EMC 2014/30/UE, Directiva privind proiectarea ecologică 2009/125/CE, la Directiva RoHS 2011/65/UE și poartă marcajul CE în consecință.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Informații suplimentare:

1) Acest produs este atribuit un număr de model de reglementare care rămâne cu aspectele de reglementare ale proiectare. Numărul de model de reglementare este identificatorul principal produs în documentația de reglementare și rapoarte de încercare, acest număr nu trebuie confundat cu numele de marketing sau umerele de produs.
2) Acest produs a fost testat într-un mediu tipic HP împreună cu un sistem gazdă HP.

Palo Alto, CA
07-09-2018

verifica semnătura de pe declarația inițială anexat
Gilles Soulard, Manager
Product Compliance Center

Local de contact pentru subiecte de reglementare numai:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

CONFORMITEITSVERKLARING
volgens ISO / IEC 17050-1 en EN 17050-1

DoC #: HSD-0002-Q-R1Vertaling/nl

Naam van de fabrikant: **HP Inc.**
Adres van de fabrikant: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

verklaart, dat het product

Naam van het product en model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulatory Model Number:[1)] HSD-0002-Q
Product opties: Geen opties

voldoet aan de volgende productspecificaties en wetgeving:

Veiligheid:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ecodesign
Verordening (EG) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Het product voldoet aan de eisen van de LVD-richtlijn 2014/35/EU, de EMV-Richtlijn 2014/30/EU, de Ökodesign-Richtlijn 2009/125/EG, de RoHS-Richtlijn 2011/65/EU en draagt het k CE-keurmer.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Extra informatie:

1) Dit product is voorzien van een wettelijk modelnummer die blijft bij de regelgevende aspecten van de ontwerp. Het voorgeschreven modelnummer is het belangrijkste product-id in de regelgeving documentatie en testrapporten, moet dit nummer niet verwarren met de merknaam of de productnummers.
2) Dit product werd getest in het HP-omgeving in combinatie met een HP hostsysteem.

Palo Alto, CA
07-09-2018

controleren van een handtekening op het oorspronkelijke verklaring gehecht
Gilles Soulard, Manager
Product Compliance Center

Lokale contactpersoon voor mbtvoorschriften:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

MEGFELELŐSÉGI NYILATKOZAT

az ISO / IEC 17050-1 és EN 17050-1

DoC #: HSD-0002-Q-R1Fordítás/hu

A gyártó neve: **HP Inc.**
Gyártó címe: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

kijelenti, hogy a termék

A termék neve és a modell:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Szabályozási Modell száma:[1)] HSD-0002-Q
Termék opciók: Nincs lehetőség

megfelel az alábbi termék dokumentáció és rendeletek:

Biztonság:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Környezetbarát tervezési
Rendelet (EC) No. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

A termék eleget tesz az alacsony feszültségre vonatkozó 2014/35/EU irányelv, az EMÖ 2014/30/EU irányelv, a környezetbarát tervezésről szóló 2009/125/EK irányelv, az RoHS 2011/65/EU irányelv viseli a CE jelzést viseli.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

További információ:

1) Ez a termék tartozik egy hatósági azonosítási típusszám ami marad szabályozási szempontból a design. A szabályozási modellszámmal a fő termék azonosítóját a szabályozási dokumentációban és vizsgálati jelentések, ez a szám nem keverendő össze a kereskedelmi névvel vagy a termék számokat.
2) Ez a termék tesztelése egy tipikus HP környezetben együtt egy HP gazda rendszert.

Palo Alto, CA
07-09-2018

ellenőrzi az aláírást az eredeti nyilatkozatot mellékelt
Gilles Soulard, Manager
Product Compliance Center

Helyi kapcsolattartó a jogi tudnivalókkal:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DECLARAÇÃO DE CONFORMIDADE

de acordo com a ISO / IEC 17050-1 e EN 17050-1

DoC #: HSD-0002-Q-R1Tradução/pt

Nome do fabricante: **HP Inc.**
Endereço do fabricante: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

declara, que o produto

Nome do produto e modelo:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Número de Modelo de Regulamentação:[1)] HSD-0002-Q
Opções do produto: Não há opções

está em conformidade com as seguintes especificações e regulamentos do produto:

Segurança:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ecodesign
Regulamento (CE) nº 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

O produto está em conformidade com os requisitos da a Directiva de Baixa Tensão 2014/35/UE, a Directiva EMC 2014/30/UE, a Directiva Ecodesign 2009/125/CE, a Directiva RoHS 2011/65/UE e leva a marca CE de acordo.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Informações adicionais:

1) Este produto recebeu um Número de modelo de regulamentação que fica com os aspectos regulatórios do design. O Número de Modelo de Regulamentação é o identificador principal do produto na documentação regulamentar e relatórios de ensaio, este número não deve ser confundido com o nome comercial ou os números dos produtos.
2) Este produto foi testado em um ambiente típico de HP.

Palo Alto, CA
07-09-2018

verificar a assinatura na declaração original anexado
Gilles Soulard, Manager
Product Compliance Center

O contato local para tópicos regulamentares apenas:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

ΔΉΛΩΣΗ ΣΥΜΜΌΡΦΩΣΗΣ
σύμφωνα με το πρότυπο ISO / IEC 17050-1 και EN 17050-1

DoC #: HSD-0002-Q-R1Μετάφραση**/el**

Όνομα κατασκευαστή: **HP Inc.**
Διεύθυνση του κατασκευαστή: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

δηλώνει ότι το προϊόν

Όνομα προϊόντος και μοντέλου:[2] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Αριθμός Μοντέλο κατά τους κανονισμούς:[1] HSD-0002-Q
Επιλογές Προϊόντος: Δεν υπάρχουν επιλογές

συμμορφώνεται προς τις εξής προδιαγραφές και κανονισμοί:

Ασφάλεια:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

οικολογικού σχεδιασμού
Κανονισμός (ΕΚ) αριθ 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Το παρόν προϊόν συμμορφώνεται με τις απαιτήσεις του η Οδηγία Χαμηλής Τάσης 2014/35/ΕΕ, η Οδηγία EMC 2014/30/ΕΕ, η Οδηγία για τον οικολογικό σχεδιασμό 2009/125/ΕΚ, η Οδηγία RoHS 2011/65/ΕΕ και φέρει τη σήμανση CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Πρόσθετες πληροφορίες:

1) Αυτό το προϊόν έχει εκχωρηθεί ένας ρυθμιστικός αριθμός μοντέλου που μένει με τις ρυθμιστικές πτυχές των σχεδιασμό. Ο αριθμός Ρυθμιστική μοντέλου είναι το κύριο αναγνωριστικό του προϊόντος με την κανονιστική τεκμηρίωση και εκθέσεις δοκιμών, ο αριθμός αυτός δεν πρέπει να συγχέεται με την εμπορική ονομασία ή τους αριθμούς προϊόντος.
2) Αυτό το προϊόν έχει δοκιμαστεί σε ένα τυπικό περιβάλλον HP.

Palo Alto, CA
07-09-2018

ελέγξτε την υπογραφή για την αρχική δήλωση που προσαρτάται
Gilles Soulard, Manager
Product Compliance Center

Τοπική επαφής για θέματα κανονισμών μόνο:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

FÖRKLARING OM ÖVERENSSTÄMMELSE

i enlighet med ISO / IEC 17050-1 och EN 17050-1

DoC #: HSD-0002-Q-R1Översättning/sv

Tillverkarens namn: **HP Inc.**
Tillverkarens adress: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

förklarar att produkten

Namn och modell:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regleringsmodellnummer:[1)] HSD-0002-Q
Produktalternativ: Inga optioner

överensstämmer med följande produktspecifikationer och förordningar:

Säkerhet:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ekodesign
Förordning (EG) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Produkten uppfyller därmed kraven i Lågspänningsdirektivet 2014/35/UE, EMC-direktivet 2014/30/UE, Ekodesigndirektivet 2009/125/CE, RoHS-direktivet 2011/65/UE och är CE-märkning i enlighet därmed.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Ytterligare information:

1) Den här produkten har tilldelats ett kontrollmodellnummer som stannar med de regulatoriska aspekterna av design. Kontrollmodellnumret är den viktigaste produktbeteckning i lagstiftnings dokumentation och provningsrapporter, detta nummer ska inte förväxlas med produktnamn eller produktnummer.
2) Produkten testades i en typisk HP-miljö tillsammans med en HP-värdsystem.

Palo Alto, CA
07-09-2018

kontrollera signaturen på den ursprungliga deklarationen som bifogas
Gilles Soulard, Manager
Product Compliance Center

Lokal kontakt ENDAST med frågor:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

PROHLÁŠENÍ O SHODĚ

podle normy ISO / IEC 17050-1 a ČSN EN 17050-1

DoC #: HSD-0002-Q-R1Překlad/cs

Jméno výrobce: **HP Inc.**
Adresa výrobce: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

prohlašuje, že výrobek

Název produktu a model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulační číslo modelu:[1)] HSD-0002-Q
Možnosti výrobku: Žádné volby

vyhovuje následujícím specifikacím a nařízení produktu:

Bezpečnost:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ekodesignu
Nařízení Rady (ES) č 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Tento výrobek splňuje požadavky Směrnice o nízkém napětí 2014/35/UE, Směrnice EMC 2014/30/UE, Směrnice o ekodesignu 2009/125/CE, Směrnice RoHS 2011/65/UE a nese označení CE v souladu s CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Další informace:

1) Tento produkt je přiděleno regulatorní číslo modelu, který zůstává s regulačními aspekty konstrukce. Počet Regulační model je hlavním identifikátorem produktu v regulační dokumentaci a protokoly o zkouškách, toto číslo by nemělo být zaměňováno s marketingovým názvem nebo čísla produktu.
2) Tento výrobek byl testován v typické prostředí HP ve spolupráci s hostitelským systémem HP.

Palo Alto, CA
07-09-2018

zkontrolovat podpis na původní prohlášení připojeném
Gilles Soulard, Manager
Product Compliance Center

Lokální kontakt pro informace pouze o směrnicích:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

ДЕКЛАРАЦИЯ ЗА СЪОТВЕТСТВИЕ
в съответствие с ISO / IEC 17050-1 и EN 17050-1

DoC #: HSD-0002-Q-R1Превод/bg

Наименование на производителя: **HP Inc.**
Производителя Адрес: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

заявява, че продуктът

Име на продукта и модел:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Нормативен номер на модела:[1)] HSD-0002-Q
Опции на продукта: Няма опции

съответства на следните спецификации на продукта и регламенти:

Безопасност:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

екодизайна
Регламент (ЕС) № 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Този апарат в съответствие с изискванията на Директивата за ниско напрежение 2014/35/EC, Директивата EMC 2014/30/EC, Директивата за екодизайна 2009/125/ОТ, Директивата RoHS 2011/65/EC и носи маркировката съответно CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Допълнителна информация:

1) Този продукт има нормативен номер на модела, който остава с регулаторните аспекти на дизайн. Номер на Нормативен модел е идентификатора основен продукт в нормативната документация и протоколи от изпитвания, този брой не трябва да се бърка с пазарното име или номера на продукта.
2) Този продукт е изпробван в типична HP среда във връзка с хост система HP.

Palo Alto, CA
07-09-2018

проверка на подписа върху оригинала на декларацията, приложена
Gilles Soulard, Manager
Product Compliance Center

Local контакт за регулаторни теми само:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

VYHLÁSENIE O ZHODE

podľa normy ISO / IEC 17050-1 a STN EN 17050-1

DoC #: HSD-0002-Q-R1Preklad/sk

Meno výrobcu: **HP Inc.**
adresa výrobcu: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

prehlasuje, že výrobok

Názov produktu a model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulačné číslo modelu:[1)] HSD-0002-Q
Možnosti výrobku: Žiadne voľby

spĺňa nasledujúce špecifikácie a nariadenia produktu:

Bezpečnosť:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ekodizajne
Nariadenie Rady (ES) č 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Tento výrobok spĺňa požiadavky Smernice o nízkom napätí 2014/35/UE, Smernice EMC 2014/30/UE, Smernice o ekodizajne 2009/125/CE, Smernice RoHS 2011/65/UE a nesie označenie CE v súlade s CE.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Ďalšie informácie:

1) Tento produkt je pridelené regulačné číslo modelu, ktorý zostáva s regulačnými aspektmi konštrukcie. Počet Regulačný model je hlavným identifikátorom produktu v regulačnej dokumentácii a protokoly o skúškach, toto číslo by sa nemalo zamieňať s marketingovým názvom alebo s výrobnými číslami.
2) Tento výrobok bol testovaný v typickej prostredí HP v spolupráci s hostiteľským systémom HP .

Palo Alto, CA
07-09-2018

skontrolovať podpis na pôvodnú vyhlásení pripojenom
Gilles Soulard, Manager
Product Compliance Center

Lokálne kontakt pre informácie iba o smerniciach:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

OVERENSSTEMMELSESERKLÆRING
i henhold til ISO / IEC 17050-1 og EN 17050-1

DoC #: HSD-0002-Q-R10versættelse/da

Producentens navn: **HP Inc.**
Producentens adresse: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

erklærer, at produktet

Produktnavn og Model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Lovpligtigt modelnummer:[1)] HSD-0002-Q
Produkt Valg: Ingen optioner

opfylder følgende produktspecifikationer og forordninger:

Sikkerhed:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

miljøvenligt design
Forordning (EF) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Produktet overholder kravene ilavspændingsdirektivet 2014/35/EU, EMC-direktivet 2014/30/EU, direktivet om miljøvenligt design 2009/125/EC, RoHS-direktivet 2011/65/EU og bærer CE-mærket i overensstemmelse hermed.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Yderligere information:

1) Dette produkt er tildelt et lovpligtigt modelnummer, som forbliver de lovgivningsmæssige aspekter af den design. Det lovpligtige modelnummer er det vigtigste produkt-id i den lovgivningsmæssige dokumentation og testrapporter, bør dette antal ikke forveksles med navn markedsføringen eller varenumre.
2) Dette produkt blev testet i en typisk HP-miljø i forbindelse med en HP host system.

Palo Alto, CA
07-09-2018

kontrollere signaturen på den oprindelige erklæring, der er knyttet
Gilles Soulard, Manager
Product Compliance Center

Lokal kontaktperson for regulative emner:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

VAATIMUSTENMUKAISUUSVAKUUTUS
ISO / IEC 17050-1: n ja EN 17050-1

DoC #: HSD-0002-Q-R1Käännös/fi

Valmistajan nimi: **HP Inc.**
Valmistajan osoite: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

ilmoittaa, että tuote

Tuotteen nimi ja malli:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Säädösmallinumero:[1)] HSD-0002-Q
Tuotevaihtoehdot: No vaihtoehdot

täyttää seuraavat tuotevaatimukset ja asetukset:

Turvallisuus:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ecodesign
Asetuksen (EY) N: o 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Tuote täyttää vaatimukset Pienjännitedirektiivin 2014/35/EU, EMC-direktiivin 2014/30/EU, Ekologista suunnittelua koskevan direktiivin 2009/125/EY, RoHS-direktiivin 2011/65/EU ja siinä on vastaava CE-merkintä.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Lisätiedot:

1) Tämä tuote on annettu virallinen numero, joka pysyy hallinnolliset näkökohdat suunnittelu. Säädöksiä koskeva mallinumero on tärkein tuotteen tunniste sääntely asiakirjat ja testausselosteet, tätä numeroa ei pidä sekoittaa tuotteen nimeen tai tuotteen numeroita.
2) Tämä tuote testattu tyypillisessä HP ympäristöministeriö yhdessä HP isäntä järjestelmä.

Palo Alto, CA
07-09-2018

Tarkista allekirjoitus alkuperäisen ilmoituksen liitteenä
Gilles Soulard, Manager
Product Compliance Center

Paikallinen yhteyshenkilö säännöksistä antavat:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

ATITIKTIES DEKLARACIJA

pagal ISO / IEC 17050-1 ir EN 17050-1

DoC #: HSD-0002-Q-R1Vertimas/lt

Gamintojo pavadinimas: **HP Inc.**
Gamintojo adresas: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

pareiškia, kad produktas

Produkto pavadinimas ir modelis:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Normatyvinis modelio numeris:[1)] HSD-0002-Q
Prekės pasirinkimai: Nėra variantai

atitinka šias produkto specifikacijas ir reglamentų:

Sauga:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMS
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ekologinio
Reglamentas (EB) Nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Šis gaminys atitinka reikalavimus, Žemų įtampų direktyva 2014/35/ES, EMS direktyva 2014/30/ES, Ekologinio projektavimo direktyva 2009/125/EB, RoHS direktyva 2011/65/ES ir yra pažymėtas CE ženklu atitinkamai.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Papildoma informacija:

1) Šis produktas pažymėtas normatyviniu modelio numeriu, kuris lieka su reglamentavimo aspektų dizainas. Normatyvinis modelio numeris yra pagrindinė produkto identifikatorius reguliavimo dokumentus ir bandymų ataskaitos, šis skaičius neturėtų būti painiojamas su prekės pavadinimu arba gaminio numeriais.
2) Ši prekė buvo išbandytas standartinės HP aplinkoje kartu su HP priimančioji sistema.

Palo Alto, CA
07-09-2018

patikrinti paraša, pridėto pradinio deklaracijos
Gilles Soulard, Manager
Product Compliance Center

Vietinis kontaktas normatyviniais klausimais tik:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

SAMSVARSERKLÆRING
i henhold til ISO / IEC 17050-1 og EN 17050-1

DoC #: HSD-0002-Q-R10versettelse/no

Produsentens navn: **HP Inc.**
Produsentens adresse: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

erklærer at dette produktet

Produktnavn og modell:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Forskriftsmessig modellnummer:[1)] HSD-0002-Q
Produktvalg: Ingen opsjoner

er i samsvar med følgende produktspesifikasjoner og forskrift:

Sikkerhet:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

økodesign
Forordning (EF) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Produktet er i samsvar med kravene i lavspenningsdirektiv 2014/35/EU, EMC-direktiv 2014/30/EU, den økodesign direktiv 2009/125/EF, RoHS-direktiv 2011/65/EU og bærer derfor CE-merket.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Tilleggsinformasjon:

1) Dette produktet er tildelt et modellnummer som holder med de regulatoriske aspekter av design. Forskriftsmodellnummeret er den viktigste produkt-ID i regulatoriske dokumentasjon og testrapporter, bør dette nummeret ikke forveksles med markedsføringsnavnet eller produktnumrene.
2) Produktet ble testet i et typisk miljø HP i forbindelse med en HP-vertssystemet.

Palo Alto, CA
07-09-2018

sjekk signaturen på originalerklæringen annektert
Gilles Soulard, Manager
Product Compliance Center

Lokal kontakt for spørsmål om forskrifter:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

IZJAVA O SKLADNOSTI

v skladu z ISO / IEC 17050-1 in EN 17050-1

DoC #: HSD-0002-Q-R1Prevajanje/sl

Ime proizvajalca: **HP Inc.**
Naslov proizvajalca: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

izjavlja, da izdelek

Ime izdelka in model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Upravna številka modela:[1)] HSD-0002-Q
Možnosti izdelka: Ni opcije

v skladu z naslednjimi specifikacijami in predpisi o izdelku:

Siguranţă:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

okoljsko primerni zasnovi
Regulamentul (CE) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Izdelek je skladen z zahtevami Direktiva o nizki napetosti 2014/35/EU, Direktiva EMC 2014/30/EU, Direktiva o okoljsko primerni zasnovi 2009/125/ES, Direktiva RoHS 2011/65/EU in nosi oznako CE v skladu s tem.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Dodatne informacije:

1) Ta izdelek je dodeljena upravna številka modela, ki ostane pri regulativnih vidikih zasnova. Število Regulativni model je glavni identifikator izdelka v regulativnem dokumentacije in poročila o preskusih, to število ne sme zamenjevati s tržnim imenom ali številkami izdelka.
2) Acest produs fost testat intr-un Mediu tipic HP împreună cu un sistem Gazda HP.

Palo Alto, CA
07-09-2018

verifica semnătura de pe declaraţia iniţială anexat
Gilles Soulard, Manager
Product Compliance Center

Lokalni kontakt za upravnimi temami:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

VASTAVUSDEKLARATSIOON

vastavalt ISO / IEC 17050-1 ja EN 17050-1

DoC #: HSD-0002-Q-R1Tõlkimine/et

Tootja nimi: **HP Inc.**
Tootja aadress: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

kinnitab, et toode

Toote nimi ja mudel:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Normatiivne mudelinumber:[1)] HSD-0002-Q
Toote lisaseadmed: No valikud

vastab järgmistele tootespetsifikatsioonidele ja määrused:

Ohutus:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ökodisaini
Määruse (EÜ) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Käesolev toode vastab nõuetele Madalpinge direktiivi 2014/35/EL, EMC direktiivi 2014/30/EL, Ökodisaini direktiivi 2009/125/EÜ, RoHS direktiivi 2011/65/EL ning kannab CE-märgistusega.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Lisainfo:

1) See toode on määratud normatiivne mudelinumber mis jääb regulatiivsed aspektid disain. Normatiivne mudelinumber on peamine toode tunnuse reguleerivad dokumendid ja katseprotokollid, see number ei tohiks segamini ajada toote nime või toode numbrid.
2) See toode on testitud tüüpiliste HP keskkonnas koos HP vastuvõtva süsteemi .

Palo Alto, CA
07-09-2018

kontrollida allkirja esialgse deklaratsiooni lisatud
Gilles Soulard, Manager
Product Compliance Center

Kohalik kontakt Ainult normatiivsete küsimuste korral:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

IZJAVA O SUKLADNOSTI

u skladu s ISO / IEC 17050-1 i EN 17050-1

DoC #: HSD-0002-Q-R1Prijevod/hr

Naziv proizvođača: **HP Inc.**
Proizvođača adresa: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

izjavljuje da je proizvod

Naziv proizvoda i Model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Broj modela prema propisima:[1)] HSD-0002-Q
Mogućnosti proizvoda: Nema opcija

sukladan sljedećim specifikacijama proizvoda i propisa:

Sigurnost:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ekološki dizajn
Uredba (EZ) br 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Navedeni proizvod sukladan sa zahtjevima Direktiva o niskom naponu 2014/35/EU, EMC Direktivom 2014/30/EU, Ekološki dizajn Direktivom 2009/125/EC, RoHS Direktivom 2011/65/EU i nosi CE oznaka u skladu s tim.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Dodatne informacije:

1) Ovaj proizvod je dodijeljen broj modela prema propisima koji ostaje s regulatornim aspektima Dizajn. Broj modela prema propisima je glavni identifikator proizvod u regulatornom dokumentacije i izvješća o ispitivanju, ovaj broj ne treba brkati s marketinškim nazivom ili brojeve proizvoda.
2) Ovaj proizvod je ispitan u tipičnoj HP okoliš u suradnji s HP-host sustava.

Palo Alto, CA
07-09-2018

provjeriti potpis na izvorne izjave priložen
Gilles Soulard, Manager
Product Compliance Center

Lokalno kontakt za pravna pitanja samo:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

ATBILSTĪBAS DEKLARĀCIJA

saskaņā ar ISO / IEC 17050-1 un EN 17050-1

DoC #: HSD-0002-Q-R1Tulkojums/lv

Ražotāja nosaukums: **HP Inc.**
Ražotāja adrese: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

paziņo, ka produkts

Produkta nosaukums un modelis:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulējošais modeļa numurs:[1)] HSD-0002-Q
Izstrādājuma iespējas: Nav iespējas

atbilst šādām iekārtas specifikācijām un regulām:

Sábháilteacht:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

ekodizains
Rialachán (CE) Uimh 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Produkts ar šo atbilst prasībām Zemsprieguma direktīva 2014/35/ES, EMC direktīva 2014/30/ES, Ekodizaina direktīva 2009/125/EK, RoHS direktīva 2011/65/ES un veic CE marķējums atbilstoši.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Papildus informācija:

1) Šis produkts ir piešķirts regulējošais modeļa numurs, kas paliek ar normatīvajiem aspektiem dizainu. Regulējošais modeļa numurs ir galvenais produkts identifikators pārvaldes dokumentācijā un testēšanas pārskati, šis skaits nedrīkst sajaukt ar mārketinga nosaukumu vai produktu numuriem.
2) Rinneadh tástáil ar táirge SEO i dtimpeallacht HP tipiciúil i gcomhar le CORAS óstach HP.

Palo Alto, CA
07-09-2018

seiceáil par síniú ar dearbhú bunaidh i gceangal
Gilles Soulard, Manager
Product Compliance Center

Vietējā kontaktpersona jautājumos par reglamentāciju tikai:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

LEYFISYFIRLÝSING
samkvæmt ISO / IEC 17050-1 og EN 17050-1

DoC #: HSD-0002-Q-R1Þýðing/is

Framleiðanda Heiti: **HP Inc.**
Heimilisfang framleiðanda: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

því yfir, að varan

Nafn vöru og Model:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Regulatory Model Number:[1)] HSD-0002-Q
Valmöguleikar vöru: Engar valkosti

í samræmi við eftirfarandi forskriftir vara og reglugerðir:

Öryggi:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

visthönnun
Reglugerð (EB) nr 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Varan hér með í samræmi við kröfur Lágspennu tilskipun 2014/35/EU, EMC tilskipun 2014/30/EU, Tilskipun um vistvæna hönnun 2009/125/EB, RoHS tilskipun 2011/65/EU og ber CE-merkið í samræmi.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Viðbótarupplýsingar:

1) Þessi vara er úthlutað Regulatory tegundarnúmer sem dvelst hjá reglugerðum þætti í hönnun. The Regulatory Model Number er helsta vara auðkenni í reglugerðum skjöl og prófunarskýrslur, þetta númer ætti ekki að rugla saman við nafn markaðssetningu eða vörunúmerum.
2) Þessi vara var prófað í dæmigerðum HP umhverfi í tengslum við HP móti kerfinu.

Palo Alto, CA
07-09-2018

athuga undirskrift á upprunalegu yfirlýsingu fylgir
Gilles Soulard, Manager
Product Compliance Center

Upplýsingaskrifstofa fyrir reglusetningu efni eingöngu:
EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

DIKJARAZZJONI TA 'KONFORMITÀ

skond ISO / IEC 17050-1 u EN 17050-1

DoC #: HSD-0002-Q-R1Traduzzjoni/mt

Isem tal-manifattur: **HP Inc.**
Indirizz tal-manifattur: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

jiddikjara, li l-prodott

Isem tal-prodott u Mudell:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Mudell regolatorju Numru:[1)] HSD-0002-Q
Għażliet tal-prodott: ebda għażliet

jikkonforma mal-Prodott Speċifikazzjonijiet li ġejjin u Regolamenti:

Sigurtà:
EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC
EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ekodisinn
Regolament (KE) Nru 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS
EN 50581:2012

Il-prodott hawnhekk jikkonforma mar-rekwiżiti tal-Direttiva Vultaġġ Baxx 2014/35/UE, Direttiva EMC 2014/30/UE, Direttiva tal-Ekodisinn 2009/125/KE, Direttiva RoHS 2011/65/UE u jkollu l-marka CE kif xieraq

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Informazzjoni addizzjonali:

1) Dan il-prodott huwa assenjat għal mudell regolatorju Numru li jibqa 'mal-aspetti regolatorji ta' l- disinn. Il-Numru tal-Mudell regolatorja hija l-identifikatur tal-prodott ewlieni fid-dokumentazzjoni regolatorju u rapporti tat-test, dan in-numru ma għandux jiġi mfixkel ma 'isem marketing jew in-numri tal-prodott.
2) Dan il-prodott ġie ttestjata f'ambjent HP tipiku flimkien ma 'sistema ospitanti HP.

Palo Alto, CA
07-09-2018

jivverifika l-firma fuq id-dikjarazzjoni oriġinali annessa
Gilles Soulard, Manager
Product Compliance Center

Kuntatt lokali għal suġġetti regolatorji biss:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

UYGUNLUK BEYANI

ISO / IEC 17050-1 ve EN 17050-1 uygun olarak

DoC #: HSD-0002-Q-R1Çeviri/tr

Üretici Adı: **HP Inc.**
Üretici Adresi: 1501 Page Mill Road, Palo Alto, CA 94303-1112 USA

beyan eder, ürün

Ürün Adı ve Modeli:[2)] HP Healthcare Edition HC271 Clinical Review Monitor, HP Healthcare Edition HC271p Clinical Review Monitor
Yasal Model Numarası:[1)] HSD-0002-Q
Ürün Opsiyonları: Opsiyon yok

aşağıdaki Ürün spesifikasyonlarına ve Yönetmeliklere uygun olduğunu beyan eder:

Güvenlik

EN 60950-1:2006 + A11:2009 +A1:2010 +A12:2011 +A2:2013
IEC 60950-1:2005 +A1:2009 +A2:2013
EN 62479:2010

EMC

EN 55032:2012 Class B
EN 55032:2015 Class B
EN 61000-3-2:2014
EN 61000-3-3:2013
EN 55024:2010
FCC CFR 47 Part 15
ICES-003, Issue 6

Ekotasarım

Yönetmelik (EC) No. 1275/2008
EN 50564:2011
IEC 62301:2011

RoHS

EN 50581:2012

Ürün, bu belgeyle birlikte, şu şartlarına uygundur Düşük Voltaj Direktifi 2014/35/AB, EMC Direktifi EMC 2014/30/AB, Ekotasarım Direktifi 2009/125/EC, RoHS Direktifi 2011/65/AB ve uygun CE işaretini taşır.

This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions:
(1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

Ek Bilgiler:

1) Bu ürüne tasarım sırasında, teknik mevzutlar gözetilerek bir Yasal Model Numarası atanmıştır. Yasal Model Numarası, ana ürün için düzenleyici belgelerinde ve test raporlaronda tanımlayıcıdır. Bu numara pazarlama adı veya ürün numaraları ile karıştırılmamalıdır.
2) Bu ürün, tipik bir HP ortamında test edilmiştir .

Palo Alto, CA
07-09-2018

imza için orginal DoC ye bakınız
Gilles Soulard, yetkili
Product Compliance Center

Sadece teknik yasal konular için kontak:

EU: HP Deutschland GmbH, HP HQ-TRE, 71025 Boeblingen, Germany
U.S.: HP Inc., 1501 Page Mill Road, Palo Alto 94304, U.S.A. 650-857-1501
www.hp.eu/certificates

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