Dexcom product Declarations of Conformity
EU DECLARATION OF CONFORMITY
RoHS Directive

Manufacturer's Name: DexCom, Inc.
Manufacturer's Address: 6340 Sequence Drive
San Diego, CA 92121
United States of America

Product Model/Name(s):
- STP- GS -(001 to 007)/ mg/dL, Dexcom G6 CGM System
- STP- GS -(100 to 112)/ mmol/L, Dexcom G6 CGM System
- STK- GS -(001 to 099)/ mg/dL, Dexcom G6 Receiver Kit
- STK- GS -(101 to 199)/ mmol/L, Dexcom G6 Receiver Kit
- STR- GS -(002 to 099)/ mg/dL, Dexcom G6 Replacement Receiver Kit
- STR- GS -(101 to 199)/ mmol/L, Dexcom G6 Replacement Receiver Kit
- STT- GS -(001 to 010)/ Dexcom G6 Transmitter Kit
- STS- GS -(002 to 144)/ Dexcom G6 Sensor
- STSGS002 and STSGS003/ Dexcom G6 Sensor

Directive: Restriction on the use of certain Hazardous Substances (RoHS)

Devices Comply with:
- EN IEC 63000:2018 Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances.

This document is valid until the date of the certificate expiration of May 26th, 2024 unless superseded by a more recent revision.

We, the undersigned, hereby declare that the equipment specified above conforms to the above Standards.

Signature: [Signature]
Date: 09 DEC 2021

Full Name: Chris Cuddy Position: Director Quality Assurance, Dexcom Inc.
EU DECLARATION OF CONFORMITY
Radio Equipment Directive

Manufacturer's Name: DexCom, Inc.
Manufacturer's Address: 6340 Sequence Drive
                        San Diego, CA 92121
                        United States of America

Product Model/Name(s):
  o STP- GS -(001 to 007)/ mg/dL, Dexcom G6 CGM System
  o STP- GS -(100 to 112)/ mmol/L, Dexcom G6 CGM System
  o STK- GS -(001 to 099)/ mg/dL, Dexcom G6 Receiver Kit
  o STK- GS -(101 to 199)/ mmol/L, Dexcom G6 Receiver Kit
  o STR- GS -(002 to 099)/ mg/dL, Dexcom G6 Replacement Receiver Kit
  o STR- GS -(101 to 199)/ mmol/L, Dexcom G6 Replacement Receiver Kit
  o STT- GS -(001 to 010)/ Dexcom G6 Transmitter Kit
  o STS- GS -(002 to 144)/ Dexcom G6 Sensor
  o STSGS002 and STSGS003/ Dexcom G6 Sensor

Devices Comply with:

  Frequency Band: 2402 – 2480 MHz
  Output Power: -1.7dBm

The equipment described above is in conformity with the Directive 2014/53/EU, in accordance with
the following harmonized standards:

<table>
<thead>
<tr>
<th>Essential Requirements</th>
<th>Compliance Standard</th>
</tr>
</thead>
<tbody>
<tr>
<td>Article 3(1)(a) Safety</td>
<td>EN 60601-1.2006/AMD1.2013</td>
</tr>
<tr>
<td>Article 3(1)(b) EMC</td>
<td>ETSI EN 301 489-1 V2.2.3 (2019-11), ETSI EN 301 489-17 (2017-03), EN 60601-1-2:2015</td>
</tr>
<tr>
<td>Article 3(2) Radio Spectrum Usage</td>
<td>ETSI EN 300 328 V2.2.2 (2019-07)</td>
</tr>
<tr>
<td>Article 3(3)</td>
<td>N/A, No delegated acts currently in force for radio equipment listed in this Declaration.</td>
</tr>
</tbody>
</table>

This document is valid until the date of the certificate expiration of May 26th, 2024 unless
superseded by a more recent revision.

We, the undersigned, hereby declare that the equipment specified above conforms to the above Standards.

Signature: [Signature]  Date: 09/03/2021

Full Name: Chris Cuddy  Position: Director Quality Assurance, Dexcom Inc.
# EC DECLARATION OF CONFORMITY

<table>
<thead>
<tr>
<th>Manufacturer Name:</th>
<th>Dexcom, Inc.</th>
</tr>
</thead>
</table>
| Manufacturer Address: | 6340 Sequence Drive  
San Diego, CA 92121  
United States of America |
| Manufacturer SRN: | US-MF-000010694 |
| Authorised Representative Name: | MDSS GmbH |
| Authorised Representative Address: | Schiffgraben 41  
30175 Hannover, Germany |
| Authorised Representative’s SRN: | DE-AR-000005430 |

<table>
<thead>
<tr>
<th>Product Name(s):</th>
<th>Dexcom G7 Continuous Glucose Monitoring System</th>
</tr>
</thead>
<tbody>
<tr>
<td>Product Type:</td>
<td>Active Medical Device, Continuous Glucose Monitor</td>
</tr>
<tr>
<td>Intended Purpose:</td>
<td>The Dexcom G7 Continuous Glucose Monitoring System is a continuous glucose monitoring system intended to continuously measure the glucose in the interstitial fluid and is designed to replace fingerstick blood glucose (BG) testing for treatment decisions.</td>
</tr>
<tr>
<td>Basic UDI-DI:</td>
<td>038627G7CGM2Y</td>
</tr>
<tr>
<td>EMDN:</td>
<td>Z12040115</td>
</tr>
<tr>
<td>GMDN:</td>
<td>44611</td>
</tr>
</tbody>
</table>

### Applicable Regulation/Directive(s):

The above-mentioned product is in conformity with the following legislative acts:
- Regulation (EU) 2017/745 on medical devices
- Directive 2014/53/EU on the harmonisation of the laws of the Member States relating to the making available on the market of radio equipment
- Directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment

### Regulation (EU) 2017/745 on medical devices

<table>
<thead>
<tr>
<th>Classification:</th>
<th>Class IIb, Rule 8 according to Annex VIII of Regulation (EU) 2017/745</th>
</tr>
</thead>
<tbody>
<tr>
<td>Notified Body:</td>
<td>BSI Group The Netherlands B.V.</td>
</tr>
</tbody>
</table>
| Notified Body Address: | Say Building, John M. Keynesplein 9  
1066 EP Amsterdam, Netherlands |
| Notified Body ID#: | 2797 |
| Conformity Assessment: | Annex IX Chapters I and III |
| EC Certificate: | MDR 727974 |
### Radio Equipment Directive 2014/53/EU

<table>
<thead>
<tr>
<th>Harmonised Standard</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>IEC 60601-1:/AMD1 :2012</td>
<td></td>
</tr>
<tr>
<td>ETSI EN 301 489-1 V.2.2.3 (2019-11)</td>
<td></td>
</tr>
<tr>
<td>ETSI EN 301 489-17 (2017-03)</td>
<td></td>
</tr>
<tr>
<td>IEC 60601-1-2 (2014)</td>
<td></td>
</tr>
<tr>
<td>ETSI EN 300 328 V2.2.2 (2019-07)</td>
<td></td>
</tr>
</tbody>
</table>

### Restriction of Hazardous Substances Directive 2011/65/EU

<table>
<thead>
<tr>
<th>Harmonised Standard</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>EN IEC 63000:2018</td>
<td></td>
</tr>
</tbody>
</table>

**Applicable Common Specifications:**
There are no relevant common specifications for the above-mentioned products.

**We, the undersigned, hereby declare that the products specified above conforms to the above Standards.**

The EU declaration of conformity is issued under the sole responsibility of Dexcom Inc. as manufacturer and the device is in conformity with Regulation (EU) 2017/745 on medical devices.

**Manufacturer:** Dexcom, Inc.

I hereby declare that the products named above has been designed to comply with the relevant sections of Regulation (EU) 2017/745 on medical devices and the applicable Union legislative acts specified above. I hereby declare that the products are manufactured in conformity with the technical documentation referred to in Annex II and Annex III and meet the requirements of this Regulation which apply to them. The product complies with all general safety and performance requirements of the Regulation. All supporting documentation is retained under the premises of the manufacturer.

**Signature:** Digitally signed by Neeta Sharma  
**Date:** 2022.04.06 15:34:09 -07'00'  
**for and on behalf of**  
Dexcom Inc.

**Date:** ________________

**Full Name:** Neeta Sharma  
**Position:** Vice President, Regulatory Affairs  
**Place, Date of Issue:** San Diego, April 6, 2022

**Start of CE-Marking:** March 11, 2022
EU DECLARATION OF CONFORMITY
Radio Equipment Directive

Manufacturer's Name: DexCom, Inc.
Manufacturer's Address: 6340 Sequence Drive
San Diego, CA 92121
United States of America

Product Model/Name(s):
- STP-DO-(001 to 015)/ mg/dL, Dexcom ONE Starter Kit
- STP-DO-(101 to 115)/ mmol/L, Dexcom ONE Starter Kit
- STT-DO-(001 to 010)/ Dexcom ONE Transmitter
- STS-DO-(001 to 015)/ Dexcom ONE Sensor
- STK-DO-(001 to 015)/ mg/dL, Dexcom ONE Receiver
- STK-DO-(101 to 115)/ mmol/L, Dexcom ONE Receiver

Devices Comply with:

<table>
<thead>
<tr>
<th>Frequency Band:</th>
<th>2402 – 2480 MHz</th>
</tr>
</thead>
<tbody>
<tr>
<td>Output Power:</td>
<td>-1.7dBm</td>
</tr>
</tbody>
</table>

The equipment described above is in conformity with the Directive 2014/53/EU, in accordance with the following harmonized standards:

<table>
<thead>
<tr>
<th>Essential Requirements</th>
<th>Compliance Standard</th>
</tr>
</thead>
<tbody>
<tr>
<td>Article 3(1)(a) Safety</td>
<td>EN 60601-1:2006/AMD1:2013</td>
</tr>
<tr>
<td>Article 3(1)(b) EMC</td>
<td>ETSI EN 301 489-1 V2.2.3 (2019-11), ETSI EN 301 489-17 (2017-03), EN 60601-1-2:2015</td>
</tr>
<tr>
<td>Article 3(2) Radio Spectrum Usage</td>
<td>ETSI EN 300 328 V2.2.2 (2019-07)</td>
</tr>
<tr>
<td>Article 3(3)</td>
<td>N/A, No delegated acts currently in force for radio equipment listed in this Declaration.</td>
</tr>
</tbody>
</table>

This document is valid until the date of the certificate expiration of May 26th, 2024 unless superseded by a more recent revision.

We, the undersigned, hereby declare that the equipment specified above conforms to the above Standards.

Signature: [Signature]
Date: 14 FEB 2022

Full Name: Chris Cuddy  Position: Director Quality Assurance, Dexcom Inc.
EU DECLARATION OF CONFORMITY
RoHS Directive

Manufacturer’s Name: DexCom, Inc.
Manufacturer’s Address: 6340 Sequence Drive
San Diego, CA 92121
United States of America

Product Model/Name(s):
- STP-DO-(001 to 015)/ mg/dL, Dexcom ONE Starter Kit
- STP-DO-(101 to 115)/ mmol/L, Dexcom ONE Starter Kit
- STT-DO-(001 to 010)/ Dexcom ONE Transmitter
- STS-DO-(001 to 015)/ Dexcom ONE Sensor
- STK-DO-(001 to 015)/ mg/dL, Dexcom ONE Receiver
- STK-DO-(101 to 115)/ mmol/L, Dexcom ONE Receiver

Directive: Restriction on the use of certain Hazardous Substances (RoHS)

Devices Comply with:
- EN IEC 63000:2018 Technical documentation for the assessment of electrical and electronic products with respect to the restriction of hazardous substances.

This document is valid until the date of the certificate expiration of May 26th, 2024 unless superseded by a more recent revision.

We, the undersigned, hereby declare that the equipment specified above conforms to the above Standards.

Signature: [Signature]
Date: 16 FEB 2022

Full Name: Chris Cuddy
Position: Director Quality Assurance, Dexcom Inc.