1. For devices without the direct mark, the user may obtain the UDI from the table below by referencing the corresponding information marked on the physical device.
    a. Unique Device Identification: Policy Regarding Compliance Dates for Class I and Unclassified Devices, Direct Marking, and Global Unique Device Identification Database Requirements for Certain Devices (July 25, 2022)

2. This information can be reprinted from the product web page on the Henry Schein website at www.HenrySchein.com

Medical Device description:

EDNA TOWEL FCP, 3-7/8 INCH

UDI Codes cross-matching summary table

HS REF	GTIN NUMBER (01)	LOT NUMBER (10)	MANUFACT. DATE (11)
104-8244	00304040016541	145050	2016-08
104-8244	00304040016541	166701	2017-01
104-8244	00304040016541	222588	2017-10
104-8244	00304040016541	222651	2017-10
104-8244	00304040016541	229703	2017-11
104-8244	00304040016541	229704	2017-11
104-8244	00304040016541	229763	2017-11
104-8244	00304040016541	243934	2018-02
104-8244	00304040016541	243935	2018-02
104-8244	00304040016541	AF2003	2020-03
104-8244	00304040016541	AF2005	2020-05
104-8244	00304040016541	AF2111	2021-11