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:

TISS FCP, 5-1/2IN(140MM), STD PAT, 1X2 TTH SER HDL

UDI Codes cross-matching summary table

HS REF	GTIN NUMBER (01)	LOT NUMBER (10)	MANUFACT. DATE (11)
101-5132	00304040023006	AF2003	2020-03
101-5132	00304040023006	AF2201	2022-01
101-5132	00304040023006	AF2206	2022-06