



- [510\(k\)](#) | [Registration & Listing](#) | [Adverse Events](#) | [PMA](#) | [Classification](#) | [CLIA](#)
- [CFR Title 21](#) | [Advisory Committees](#) | [Assembler](#) | [Recalls](#) | [Guidance](#) | [Standards](#)



[New Search](#)

[Back To Search Results](#)

510(k) Premarket Notification Database

Device Classification Name	set, administration, intravascular
510(k) Number	K003854
Device Name	INTRAVASCULAR ADMINISTRATION SET INFUSION DEVICES, INC.
Applicant	6329 w. waterview ct. mccordsville, IN 46055
Contact	paul e dryden
Regulation Number	880.5440
Classification Product Code	FPA
Date Received	12/13/2000
Decision Date	02/05/2001
Decision	substantially equivalent (SE)
Classification Advisory Committee	General Hospital
Review Advisory Committee	General Hospital
Statement/Summary/Purged Status	Summary only
summary	summary
Type	Traditional
Reviewed by Third Party	No
Expedited Review	No

[CDRH Home Page](#) | [CDRH A-Z Index](#) | [Contact CDRH](#) | [Accessibility](#) | [Disclaimer](#)
[FDA Home Page](#) | [Search FDA Site](#) | [FDA A-Z Index](#) | [Contact FDA](#) | [HHS Home Page](#)

Center for Devices and Radiological Health / CDRH