authority is contingent on Respondent being a practitioner with a valid DEA registration, *see* 21 U.S.C. 823(g)(2)(A); 21 CFR 1301.28(a), I will revoke his DATA-Waiver authority as well.

Order

Pursuant to the authority vested in me by 21 U.S.C. 823(f) and 824(a), as well as 28 CFR 0.100(b), I order that DEA Certificate of Registration No. BZ5641419 and DATA-Waiver Identification Number XZ5641419, issued to Witold Marek Zajewski, M.D., be, and they hereby are, revoked. I further order that any pending application of Witold Marek Zajewski to renew or modify the above registration, or any pending application of Witold Marek Zajewski for any other registration in the State of Illinois, be, and it hereby is, denied. This Order is effective immediately.[4]

Dated: April 4, 2018.

Robert W. Patterson,

Acting Administrator.

[FR Doc. 2018–07454 Filed 4–10–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various classes of schedule I or II controlled substances.

SUPPLEMENTARY INFORMATION:

The companies listed below applied to be registered as importers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

Company	FR docket	Published
Sharp (Bethlehem), LLC	83 FR 539	January 4, 2018.
Catalent Pharma Solutions, LLC	83 FR 2215	January 16, 2018.
Janssen Pharmaceuticals, Inc	83 FR 2214	January 16, 2018.
Mylan Pharmaceuticals, Inc	83 FR 5809	February 9, 2018.
Meridian Medical Technologies, Inc	83 FR 5810	February 9, 2018.
Noramco, Inc	83 FR 5810	February 9, 2018.
Johnson Matthey, Inc	83 FR 5811	February 9, 2018.
Mylan Technologies, Inc	83 FR 5811	February 9, 2018.
Mylan Pharmaceuticals, Inc	83 FR 8107	February 23, 2018.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable basic classes of schedule I or II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I or II controlled substances to the above listed companies.

Dated: April 4, 2018.

Susan A. Gibson,

Deputy Assistant Administrator.

[FR Doc. 2018–07444 Filed 4–10–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as bulk manufacturers of various classes of schedule I and II controlled substances.

SUPPLEMENTARY INFORMATION:

The companies listed below applied to be registered as bulk manufacturers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted for these notices.

Company	FR docket	Published
Nanosyn, Inc	82 FR 56993	December 1, 2017.
Janssen Pharmaceutical, Inc	82 FR 58027	December 8, 2017.
Cambrex High Point, Inc	82 FR 61795	December 29, 2017.
AMPAC Fine Chemicals LLC	82 FR 61795	December 29, 2017.
Organix, Inc	83 FR 150	January 2, 2018.
Johnson Matthey Inc	83 FR 2215	January 16, 2018.
Chemtos, LLC	83 FR 2671	January 18, 2018.
Alcami Wisconsin Corporation	83 FR 2675	January 18, 2018.

[4] For the same reasons which led the IDFPR to revoke Respondent's controlled substance license, I conclude that the public interest necessitates that this Order be effective immediately. 21 CFR 1316.67.