

These clinical study results are supplied for informational purposes only in the interests of scientific disclosure. They are not intended to substitute for the FDA-approved package insert or other approved labeling.

Drug Details

| Company Name | Business Partner | Drug Name | Generic Name | Unique ID | Studied Indications or Disease | Phase |
|------------------------|--|-----------|--------------|-----------|--------------------------------|-----------|
| Abbott Laboratories | | Humira | Adalimumab | M02-570 | Psoriatic Arthritis | Phase III |
| Clinical Study Summary | Protocol No. M02-570: A Phase III Multicenter Study of the Safety and Efficacy of Human Anti-TNF Monoclonal Antibody Adalimumab (D2E7) in Moderate to Severely Active Psoriatic Arthritis (PsA) Subjects with Inadequate Response to Disease Modifying Anti-Rheumatic Drug Therapy | | | | | |

Abbott Laboratories (Abbott) will use reasonable efforts to include accurate and up-to-date information, consistent with Abbott policies and procedures, on the ClinicalStudyResults.org web site. However, because, among other reasons, the status of studies often changes, Abbott can make no, and makes no, warranties or representations of any kind as to the currency or completeness of the information contained therein. Persons accessing and using information posted by Abbott on the ClinicalStudyResults.org web site do so at their own risk. Abbott disclaims all warranties, express or implied, including warranties of merchantability of fitness for a particular purpose. Abbott shall not be liable for any damages, including without limitation, direct, incidental, consequential, indirect or punitive damages, arising out of access to, use of, or inability to use information posted by Abbott on the ClinicalStudyResults.org web site, or any errors or omissions in the content thereof.