

Vedolizumab, MLN0002, Ulcerative colitis, Crhon's disease, integrin inhibitor, Remicade, Humira, Simponi, Tysabri, Etrolizumab, Tofacitinib, GEMINI 1,2, STELARA, GSK-1605786, Vercirnon, Traficet-EN

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Abstracts

Takeda received priority review status from US FDA for its late stage candidate Vedolizumab (filed for UC and Crhon's disease (CD), $\alpha 4\beta 7$ integrin inhibitor) to treat ulcerative colitis indication. Vedolizumab targets the disease by preventing leukocyte extravasations through inhibiting $\alpha 4\beta 7$ integrin in the gut which is different from currently used second line options, mainly anti-TNFs (Remicade, Humira, Simponi). The key limitations with these drugs are their low response rate (~40% do not respond) and lost of response over time (~30%-40%). We understand that dose escalation and switch to a second anti-TNF recapture the response in 40-80% of patients but after 12 months the response rate drops significantly. This creates a huge unmet need for a drug which On pipeline frontThe report discuss in details unmet need, late stage pipeline drugs and competitive landscape of ulcerative colitis and crohn's disease treatment options.

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