

## Amgen picks its moment to strike for Chemocentryx



[Elizabeth Cairns](#)



**The \$3.7bn price is cheaper than it would have been last year, but Tavneos must still fulfill its pipeline-in-a-product potential.**

Some investors had been concerned that the launch of Tavneos, Chemocentryx's vasculitis therapy, would prove a damp squib. Amgen, it seems, has no such worries, today buying the company for \$3.7bn, more than twice the target's market value.

This is perhaps not as expensive as it looks. The \$52 a share Amgen paid may be a 116% premium to Chemocentryx's closing price yesterday, but the smaller group had been trading at this level just over a year ago.

Amgen investors will therefore hope the company has managed to bag a bargain, but much will depend on whether Tavneos can expand beyond ANCA-associated vasculitis, an autoimmune disease that causes inflammation of small blood vessels.

Notably, Chemocentryx has also been evaluating Tavneos in hidradenitis suppurativa, C3 glomerulopathy and lupus nephritis; however, results in the former two indications [have been mixed at best](#). Indeed, it was somewhat surprising that the drug [got the nod in October](#) for ANCA-associated vasculitis, given a [split adcom vote](#) and concerns about liver toxicity.

Tavneos sold \$5.4m in the US in the first quarter of 2022, surpassing consensus sellside expectations of \$3.8m, although it is too soon to label the launch a success. The drug is also approved in the EU and Japan, but Amgen will only gain the US sales; Kissei Pharmaceutical has the Japanese rights and Otsuka the Canadian, with all other territories being mopped up by Vifor.

### Growth driver

It will now be up to Amgen to make Tavneos a success in the world's biggest drug market and justify Chemocentryx's price.

The product's NPV, as calculated by *Evaluate Pharma* and based on current sellside consensus, stands at just under \$2bn. SVB analysts, meanwhile, see sales potential of over \$2bn by 2030.

As for the rest of Chemocentryx's pipeline, CCX140, which had been in phase 2 for kidney disease, seems to have been quietly dropped as it does not appear in [the company's latest update](#). There are also four other

projects, including a PD-(L)1 inhibitor, listed.

Chemocentryx's pipeline		
Product	Target	Status
CCX140	CCR2	Ph2 in nephrotic syndrome and diabetic nephropathy; since abandoned?
CCX507	CCR9	Ph1 in ulcerative colitis
CCX559	PD-(L)1	Ph1 in unspecified cancer
CCX111	CCR6	Preclinical in TH17-driven disease
CCXTBD	CCR4	Preclinical in "other dermatological"

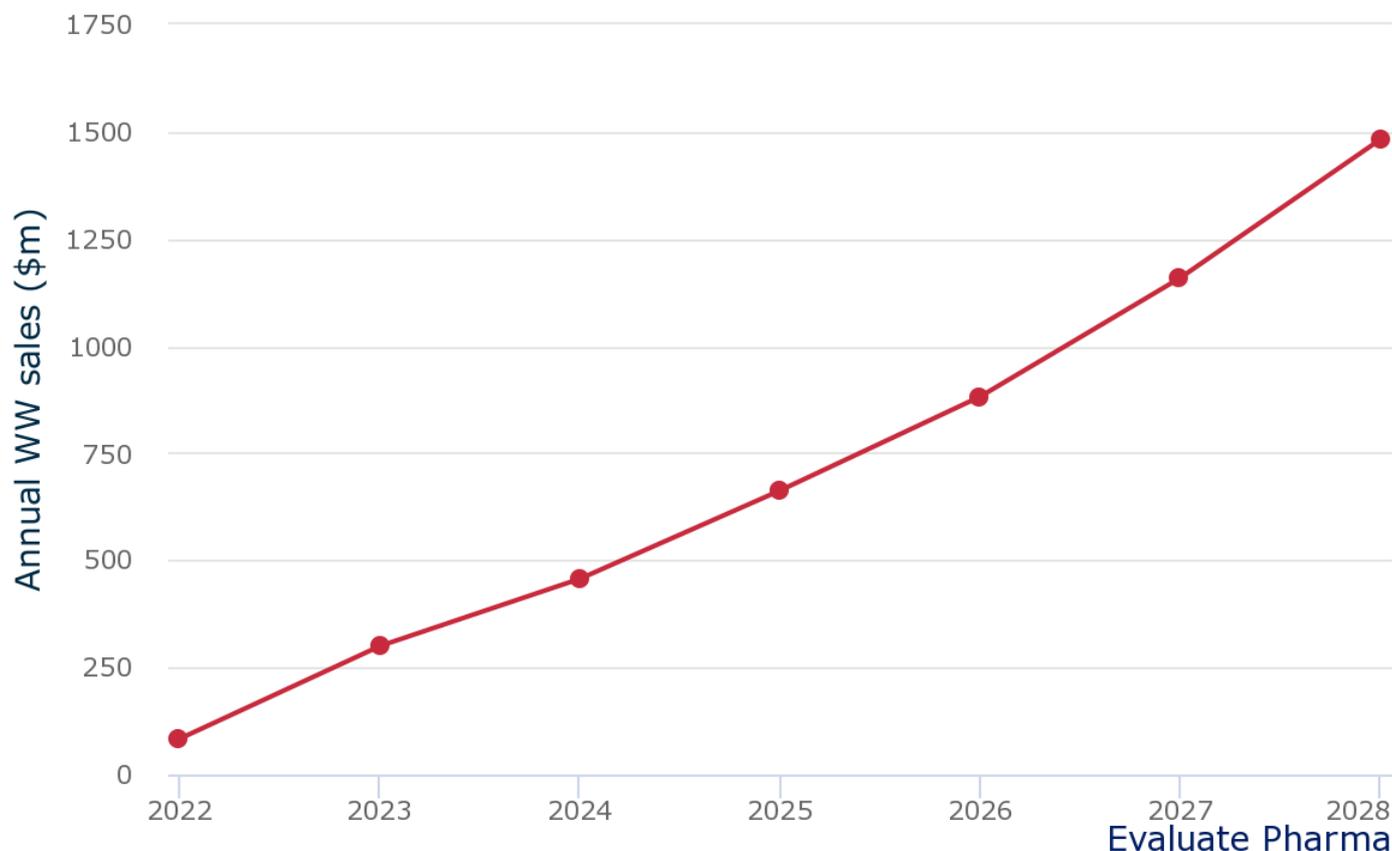
Source: [company presentation](#).

In any case, Tavneos is the main driver behind the deal. Chemocentryx previously said it would start two studies of avacopan in the second half of this year: a phase 3 in hidradenitis, despite the earlier miss, and a study in lupus nephritis, inspired by the renal function improvement seen in the phase 2 glomerulopathy trial.

In Tavneos, Amgen has got its hands on a much-needed growth driver. The group's existing portfolio is facing a patent cliff, with several key products, including Enbrel and Otezla, due to lose exclusivity in 2025-30.

Tavneos's US exclusivity could last until 2036 if patent extension is granted, according to SVB. Wringing the highest possible sales out of a drug has become an Amgen speciality, and the group will need all that expertise to justify today's outlay.

## Forecast sales for Tavneos



[More from Evaluate Vantage](#)

Evaluate HQ  
[44-\(0\)20-7377-0800](tel:44-020-7377-0800)

Evaluate Americas  
[+1-617-573-9450](tel:+16175739450)

Evaluate APAC  
[+81-\(0\)80-1164-4754](tel:+8108011644754)

© Copyright 2022 Evaluate Ltd.