

CASE STUDY

AES Shortens Timelines by up to 20 Months for a Uterine Fibroid Program

AT A GLANCE

Indication: **Uterine Fibroids**



Enrollment
time saved:

20 months
(Protocol 1)

8 months
(Protocol 2)

THE CHALLENGE

This was an important Phase III, **2-protocol** study for a **first-in-class treatment** of abdominal uterine bleeding in women with uterine fibroids.

The sponsor was facing significant enrollment challenges including:

- Delayed site start-up, as sites were unable to meet their overly optimistic commitments to deliver enough patients;
- Patients who were unwilling to wash out their current birth control medications to participate in the study; and
- Women who had irregular menstrual cycles or other uncontrolled chronic conditions that excluded them from participating

THE PLAN

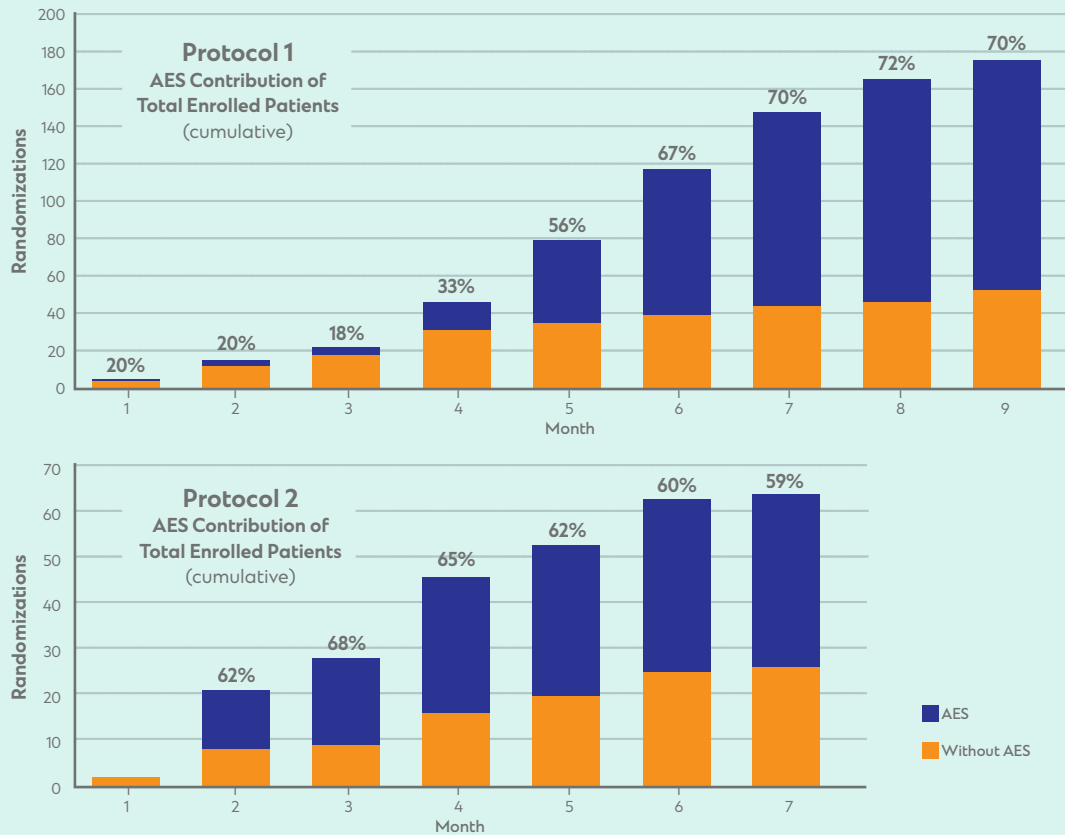
The clinical trial team needed a highly reliable solution to avoid an enrollment delay of almost 2 years. Accelerated Enrollment Solutions (AES) was hired and quickly launched a media and online marketing campaign.

Additionally, our Global Site Relations team worked with monitors to encourage site follow-up of patient referrals, and supplemented monitor conversations with delinquency reporting and email reminders.



THE RESULTS

Results were exceptional. AES contributed 70% and 59% of total patient enrollment for Protocols 1 and 2, respectively, which increased the site enrollment rate by 206% overall.



THE BENEFITS

By significantly increasing Site Enrollment Rates, AES saved 20 months of enrollment for Protocol 1, and 8 months of enrollment for Protocol 2.

About AES

Accelerated Enrollment Solutions (AES) is the new standard in clinical trial productivity. Our innovative and integrated site and patient services secure the success of clinical trials by providing enrollment, timing, and budget certainty.

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