

# Factor XIa Inhibitors After Heart Attack: A New Antithrombotic Path?

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Preventing recurrent thrombotic events after an acute myocardial infarction (AMI) remains a delicate balance between efficacy and bleeding risk. Current antithrombotic regimens, while effective, carry a non-trivial burden of haemorrhagic complications, particularly in high-risk patients.<sup>1</sup> Factor XIa inhibitors present a potential alternative, targeting a pathway thought to be less critical for physiological haemostasis.

## KEY TAKEAWAYS

- **The Pivot:** Factor XIa inhibition offers a novel antithrombotic mechanism, potentially separating thrombotic protection from bleeding risk.
- **The Data:** Early phase trials suggest a reduction in thrombotic events without a commensurate increase in major bleeding.
- **The Action:** Clinicians should monitor ongoing Phase III trials for definitive efficacy and safety data in post-AMI patients.

Patients surviving an acute myocardial infarction face a persistent risk of recurrent ischaemic events, including subsequent MI, stroke, and cardiovascular death. Dual antiplatelet therapy (DAPT) with aspirin and a P2Y<sub>12</sub> inhibitor forms the cornerstone of secondary prevention, but this regimen, especially when extended, significantly increases the risk of major bleeding. This inherent trade-off has driven the search for antithrombotic agents that can maintain or improve ischaemic protection while mitigating haemorrhagic complications.

The coagulation cascade, a complex series of enzymatic reactions, culminates in fibrin formation and clot stabilisation. Traditional anticoagulants, such as warfarin and direct oral anticoagulants (DOACs), broadly inhibit multiple factors within this cascade, leading to effective thrombosis prevention but also a heightened risk of bleeding. Factor XIa (FXIa) has emerged as a target of interest because of its unique role in pathological thrombus formation, while appearing to be less critical for primary haemostasis, the process that stops bleeding from an injured vessel. This distinction suggests that inhibiting FXIa might offer a wider therapeutic window, providing antithrombotic benefits with a lower propensity for bleeding.

## The Rationale for Factor XIa Inhibition

Factor XIa sits upstream in the intrinsic pathway of coagulation. Its activation by thrombin or Factor XIIa amplifies thrombin generation, particularly under conditions of high shear stress, such as those found in arterial thrombi. But, unlike other factors like Factor Xa or thrombin itself, Factor XIa's contribution to stopping bleeding from minor injuries appears to be less pronounced. Individuals with congenital Factor XI deficiency, for example, often exhibit only mild

bleeding phenotypes, or even none at all, unless challenged by surgery or trauma. This observation provides the biological basis for the hypothesis that FXIa inhibition could selectively target pathological thrombosis without unduly impairing physiological haemostasis.

The LIBREXIA ACS program, while not yet fully reported, represents a significant investment in exploring this hypothesis in the high-stakes setting of acute coronary syndromes. These trials typically enrol patients recently stabilised after an MI, a population at elevated risk for both recurrent ischaemic events and bleeding from intensive antithrombotic therapy. The design of such trials often involves comparing the investigational FXIa inhibitor against placebo, or against an active comparator, on top of standard antiplatelet therapy, usually DAPT. Primary endpoints generally focus on major adverse cardiovascular events (MACE), a composite of cardiovascular death, non-fatal MI, and non-fatal stroke, alongside key safety endpoints, particularly major bleeding events as defined by criteria such as TIMI or BARC.

### Early Clinical Signals and Design Considerations

Prior to the LIBREXIA ACS program, several Factor XIa inhibitors have entered clinical development, providing initial insights into the class. Early phase studies, often in populations undergoing elective knee or hip replacement surgery, demonstrated that these agents could reduce venous thromboembolism (VTE) with a favourable bleeding profile compared to enoxaparin. These initial signals, while not directly translatable to arterial thrombosis in ACS patients, supported the concept of a dissociation between antithrombotic efficacy and bleeding risk.

For patients with acute coronary syndromes, the challenge is more complex. The thrombotic milieu is highly procoagulant, and patients often have multiple comorbidities that increase both ischaemic and bleeding risks. A typical LIBREXIA ACS trial would randomise several thousand patients, perhaps 5,000 to 10,000, across hundreds of sites globally. Patients would receive either the FXIa inhibitor or placebo, in addition to their prescribed DAPT. Follow-up periods usually extend for 12 to 24 months to capture a sufficient number of MACE and bleeding events. Key inclusion criteria would involve a confirmed MI within a specific timeframe, often within 7 to 14 days, and a stable clinical course. Exclusion criteria would typically include a high bleeding risk, severe renal or hepatic impairment, or other conditions that might confound safety assessments.

### Measuring Efficacy and Safety

The primary efficacy endpoint in a trial like LIBREXIA ACS would be a composite of MACE. Secondary efficacy endpoints might include individual components of MACE, or other ischaemic events such as urgent revascularisation. The critical safety endpoint is major bleeding, often defined by BARC type 3 or 5, or TIMI major bleeding. Other safety outcomes include clinically relevant non-major bleeding (CRNMB) and all-cause mortality. The statistical powering of such trials is designed to detect a clinically meaningful reduction in MACE, typically around 15-20%, with a P-value threshold of  $P < .05$ . Non-inferiority designs for bleeding are also common, aiming to show that the FXIa inhibitor does not significantly increase bleeding compared to placebo, or that it is superior to an active comparator.

One of the key challenges in interpreting data from this class of drugs is the precise definition and adjudication of bleeding events. Different bleeding scales (TIMI, BARC, GUSTO) can yield varying rates and classifications of haemorrhage, making direct comparisons across trials difficult. Central adjudication committees, blinded to treatment assignment, are essential for maintaining the integrity of these safety endpoints. Furthermore, the duration of follow-up is crucial.

Bleeding risk often accumulates over time, and a shorter follow-up might underestimate the true haemorrhagic burden of a new agent.

### The Catch: Potential Limitations and Unanswered Questions

While the theoretical advantages of FXIa inhibition are compelling, several practical considerations remain. The optimal dose of an FXIa inhibitor in the ACS setting is not yet fully established. Dose-finding studies are critical to identify the sweet spot where antithrombotic efficacy is maximised and bleeding risk is minimised. Subgroup analyses, though often exploratory, will be vital to understand if certain patient populations, such as those with renal impairment, diabetes, or a history of prior bleeding, derive differential benefits or risks. For example, patients with chronic kidney disease are at a higher risk of both thrombotic and bleeding events, making them a particularly challenging group to manage with antithrombotic agents. Whether FXIa inhibitors can offer a safer alternative in this population is a key unanswered question.

The interaction with existing antiplatelet therapies also warrants careful consideration. Most FXIa inhibitor trials in ACS patients administer the investigational drug on top of DAPT. This raises questions about potential additive bleeding risks, even if the FXIa pathway is theoretically distinct. The duration of DAPT is also a variable; some patients may be on short-term DAPT (e.g., 6 months), while others may continue for 12 months or longer. How FXIa inhibitors integrate into these evolving DAPT strategies will be important for clinical practice. The Oxford Handbook of Cardiology offers a concise guide to these complex antithrombotic regimens.

Another limitation often seen in cardiovascular trials is the generalisability of results. Patients enrolled in clinical trials are typically younger, healthier, and have fewer comorbidities than the real-world population. Whether the benefits observed in a highly selected trial population extend to the broader spectrum of ACS patients, including the frail elderly or those with multiple complex conditions, remains to be seen. Post-marketing surveillance and real-world evidence studies will be essential to fully characterise the safety and effectiveness of FXIa inhibitors once they become available. The cost-effectiveness of a new antithrombotic agent is also a significant factor for widespread adoption. If FXIa inhibitors offer only a modest reduction in MACE or bleeding, their higher cost compared to generic antiplatelet agents might limit their use. Health economic analyses, often conducted alongside Phase III trials, will be crucial for informing reimbursement decisions and guideline recommendations. The long-term adherence to these new agents, particularly given the chronic nature of secondary prevention after MI, is another practical consideration. Complex dosing regimens or significant side effects could undermine the benefits observed in controlled trial settings.

The promise of Factor XIa inhibition lies in its potential to uncouple antithrombotic efficacy from the bleeding burden that has plagued current regimens. If LIBREXIA ACS delivers on this, it will redefine secondary prevention. Sarah Gellar, Clinical Trials EditorFinally, the specific mechanism of action of FXIa inhibitors, while theoretically safer, still requires comprehensive evaluation for rare but serious adverse events. While major bleeding is the primary safety concern, other potential off-target effects or idiosyncratic reactions must be meticulously monitored. The development of specific reversal agents for FXIa inhibitors, similar to those available for DOACs, would also be a significant advantage, enhancing their safety profile in emergency situations. Without such agents, managing bleeding complications could be more challenging.

### CLINICAL IMPLICATIONS

The prospect of a Factor XIa inhibitor for post-MI patients is genuinely intriguing. Current antithrombotic strategies, while effective, force clinicians to constantly weigh ischaemic protection against bleeding risk. A drug that can shift this balance, offering comparable or superior thrombotic prevention with a demonstrably lower bleeding rate, would be a significant advance.

For prescribing clinicians, this means a potential new tool in the armamentarium, particularly for those high-risk patients where extended DAPT is often curtailed due to bleeding concerns. If LIBREXIA ACS shows a clear benefit, it could allow for more aggressive, longer-term antithrombotic therapy without the current associated haemorrhagic burden. This would be a welcome relief for managing complex ACS patients.

But the devil will be in the details. The magnitude of benefit, the specific patient populations most likely to gain, and the real-world bleeding rates will all dictate uptake. Industry will need to demonstrate not just efficacy, but a clear safety advantage that justifies the likely premium price point. Without a compelling reduction in major bleeding, the clinical utility will be limited.

Patients, of course, stand to gain the most from a safer, more effective antithrombotic. Fewer major bleeding events mean fewer hospitalisations, fewer transfusions, and ultimately, a better quality of life. The hope is that Factor XIa inhibitors can deliver on this promise, moving beyond theoretical advantages to tangible clinical improvements.

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