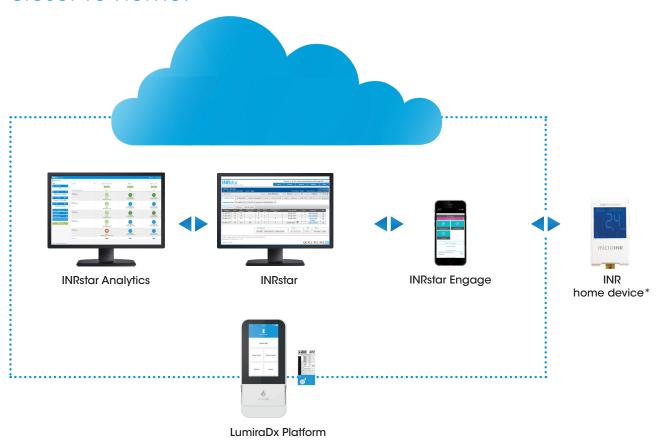
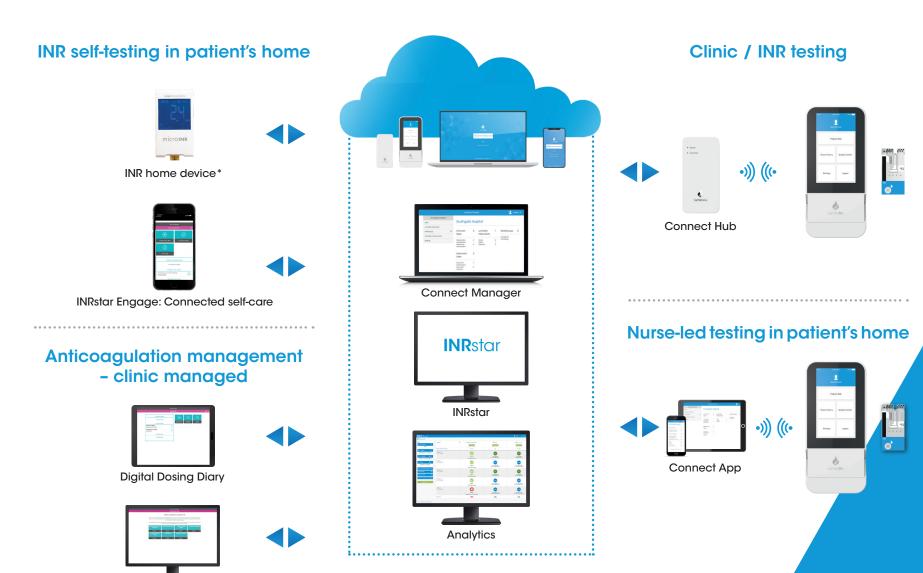


Our complete digital anticoagulation management solution is especially attractive to healthcare organizations, allowing them to manage all anticoagulation patients in one place, display multiple anticoagulation services across an area, gain instant access to data allowing monitoring of best practice and achievement of improved clinical outcomes. Ultimately, offering greater patient choice and bringing care closer to home.



LumiraDx complete anticoagulation management solution



DOAC care programme



INRstar – Award-winning anticoagulation software

INRstar provides complete anticoagulation support for the induction, dosing, review and connected self-care of patients on both warfarin and direct oral anticoagulant drugs (DOACs/NOACs), helping care teams to manage patients safely and effectively.

INRstar is used in over 2,700 locations with 284,000 active patients making it the most used anticoagulation clinical decision support software in the UK.

INRstar integrates with the LumiraDx Platform enabling complete digital anticoagulation management with diagnostics and diagnostic-

led care.

- Clinical decision support software, INRstar for healthcare professionals (81.2 % of VKA patients managed in INRstar with a TTR (Time in Therapeutic Range) of greater than 65 %)
- Ability to integrate with PAS, LIS, EHR, Clinical Systems, PT-INR point-of-care meters (clinics and self-testing)
- Central reporting at population level, INRstar analytics for commissioners
- Connected self-care, INRstar Engage app for patients



INRstar Engage – Stay connected with the INRstar Engage app

One single patient app: INRstar Engage our easy-to-use app supports you to deliver tailored care for all your anticoagulation patients remotely including patients on direct or novel oral anticoagulants (DOACs / NOACs).

INRstar Engage transforms the way you and your patients connect – securely integrating patient data and devices with their INRstar record and clinical system.



INRstar, INRstar analytics and INRstar Engage specifications

INRstar

| Safety | Clinically risk assessed, IVD complies with all UK / EU medical device regulations, CE marked, ISO 13845:2016 certified, 9001:2015 certified, 27001: 2013 certified | |
|-------------|---|--|
| Auditing | Audit capabilities / quarterly audit | |
| Reports | Location specific reporting | |
| TTR | Monitoring at patient and location level | |
| Integration | Clinical systems (TPP, EMIS & INPS) and HL7 integration | |
| Permissions | Role-based permissions | |
| Drugs | Supports warfarin, DOACs, acenocoumarol and LMWH | |

INRstar analytics

| Drugs | Supports warfarin, DOACs, acenocoumarol and LMWH | | |
|-----------------|--|--|--|
| Benchmarking | National benchmarking against all INRstar locations | | |
| Reports | Monitoring clinical outcomes and for auditing purposes | | |
| Anonymised Data | Allows commissioners to view location specific data without identifying patients | | |

INRstar Engage app

| Care Programmes | Digital Dosing Diary, Warfarin self-testing, DOAC support | |
|-----------------|---|--|
| Platforms | Available on smartphones, tablets and web | |
| Safety | Clinically risk assessed | |





LumiraDx INR Test – Convincing assay performance

The LumiraDx INR test is an enzyme kinetic assay based on Thrombin activity.

The coagulation cascade is activated by the test reagents and INR is calculated from the measured PT-time. Haematocrit is also measured on the strip to ensure patients are within the 25-55% range. The accuracy of the LumiraDx INR test is comparable to the ACL Elite lab reference method.

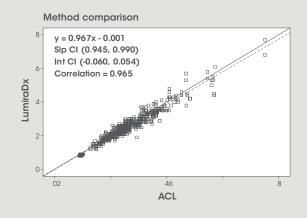
INR Test Strip specifications

| Displayed results | INR | | |
|--|--|--|--|
| —————————————————————————————————————— | 11/1/2 | | |
| Storage temperature | Between 5 °C and 32 °C (41 °F and 89 °F) | | |
| Operating temperature | Between 15 °C and 30 °C (59 °F and 86 °F) | | |
| Measuring range | 0.8–7.5 INR | | |
| Sample size | Capillary: 8 µL | | |
| Sample type | Fresh capillary whole blood – taken by finger-stick or via non-anticoagulated transfer tub | | |
| Time to result | INR results in 60-90 seconds* | | |
| ISI | Approx. 1.0 | | |
| Interferences | See INR Test Product Insert for details | | |
| | • | | |

^{*}Supratherapeutic INR > 4.0 test result could take 90-180 seconds

Accuracy

Direct Capillary Blood (596 samples from 326 patients) was collected from multiple sites. Reference method was ACL Elite Pro Coagulation Analyser.



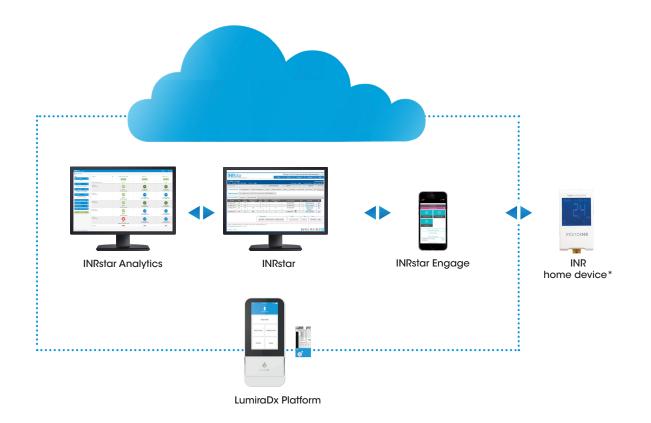
Precision

Determined for capillary blood from sample duplicates collected at multiple sites. The following results represent the mean paired rep %CV for both direct and transfer tube application.

| Sample | n | Mean INR | Mean % CV |
|--------------------|-----|----------|-----------|
| Direct application | 284 | 2.54 | 3.46 |
| Transfer tube | 291 | 2.53 | 3.73 |



LumiraDx Cloud Services



The cloud services components of the LumiraDx solution in the UK are hosted in secure data centres, certified to UK Government standards. The data and the encrypted back-ups taken to enable data recovery in the unlikely event that it would be required, will not leave the UK/EEA unless appropriate data transfer safeguards are in place. LumiraDx work closely with the data centre provider to ensure all aspects of the cloud infrastructure are managed and operated to required standards, the data centre is also certified to meet information security and data protection requirements for other countries.

- Primary data centre locations in the UK: Reading and Harrogate
- Comply with GDPR requirements

For more information on the LumiraDx Platform, LumiraDx Connect, INRstar and INRstar Engage refer to related content pages on lumiradx.com

The LumiraDx Platform is for use by healthcare professionals only. LumiraDx Platform not available in all countries. Not available in the USA. Products subject to product specifications and terms and conditions. LumiraDx and INRstar are trademarks of the LumiraDx Group.

* iLine home testing device available through LumiraDx channels in UK and Ireland only.



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