ForPatients
*by Roche*

Neurodegenerative Disorder

# A Study of Epoetin Beta Treatment in Anemic Participants With Myelodysplastic Syndrome (MDS)

| Trial Status | Trial Runs In | Trial Identifier |
|---|---|---|
| Completed | 1 Countries | NCT02145026 ML29005 |

*The source of the below information is the publicly available website ClinicalTrials.gov. It has been summarised and edited into simpler language.*

## *Trial Summary:*

This is a Phase IV, prospective, multi-center, open-label study to assess the effectiveness and safety profile of epoetin beta (Recormon®) for treatment of symptomatic anemia in adult participants associated with low/intermediate-1-risk MDS. After screening, eligible participants will be treated with epoetin beta as recommended in the approved label and international guidelines for the use of epoetin in MDS participants and the dosage will be adjusted on the basis of erythroid response.

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|---|---|
| **Hoffmann-La Roche**<br>Sponsor | **Phase 4**<br>Phase |

**NCT02145026 ML29005**
Trial Identifiers

## *Eligibility Criteria:*

| Gender | Age | Healthy Volunteers |
|---|---|---|
| All | >= 18 Years | No |