A Study to Learn About How Well Riociguat Works, How Safe it is and How it is Used Under Real World Conditions in Patients in the United States Who Are Receiving Riociguat for High Blood Pressure in the Arteries That Carry Blood From the Heart to the Lungs (Pulmonary Arterial Hypertension, PAH)

Status: Recruiting

Eligibility Criteria

Age: 18 years and over

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

- Patients aged ≥18 years at the time of riociguat treatment initiation
- Diagnosis of PAH per National Institute for Health and Care Excellence (NICE) 2018 classification
- · Decision to initiate treatment with riociguat as per investigator's routine treatment practice made prior to enrollment in the study
- Initiation of riociguat, as per the FDA-approved US label:
- · At enrollment OR
- ≤90 days prior to enrollment, with a documented titration regimen (defined as all documented dose changes including, but not limited to: starting dose and dates and highest tolerated dose and dates)
- · Signed informed consent

Exclusion Criteria:

- Previously treated with and discontinued use of riociguat for any reason prior to study enrollment (discontinuation defined as an interruption of therapy ≥30 days)
- Participating in any of the following: 1. Blinded clinical trial 2. Clinical trial involving an unapproved drug 3. Investigational program with interventions outside of routine clinical practice
- · Life expectancy <12 months
- · Contraindicated to receive riociguat per the FDA approved US label
- · Use of nitrates or NO donors in any form
- Use of PDE5 inhibitors
- · PH associated with idiopathic interstitial pneumonias
- Unable or unwilling to provide informed consent

Conditions & Interventions

Interventions:

Drug: Riociguat (Adempas, BAY63-2521)

Conditions:

Pulmonary Arterial Hypertension

More Information

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Phase: IV IRB

Number: HM20022707 **System ID:** NCT04813926

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