

**Rochtain agus Imeascadh**

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**MEMORANDUM**

To	Pharmacy Executive Manager, Children's Health Ireland
From	Access and Integration Drug Management Programme
Subject	New HSE-Approved Medicinal Product(s) – Dolutegravir sodium, abacavir, lamivudine (Triumeq®)
Date	18/02/2025

Dear Colleagues,

The following medicinal product has received HSE approval for pricing and reimbursement.

<b>Medicinal Product(s)</b>		<b>Price to Wholesaler (excluding VAT)</b>
Dolutegravir sodium, abacavir, lamivudine (Triumeq®) 5 mg/60 mg/30 mg dispersible tablets x 90		€283.38
<b>Active Ingredient(s)</b>	Dolutegravir sodium, abacavir, lamivudine	
<b>Date of Approval</b>	06/02/2025	
<b>Manufacturer</b>	ViiV Healthcare BV	
<b>ATC Code and Pharmacotherapeutic Group</b>	J05AR13; Antivirals for systemic use, antivirals for treatment of HIV infections, combinations.	
<b>HSE-Approval Details</b>	For the treatment of Human Immunodeficiency Virus type 1 (HIV-1) infected children of at least 3 months of age and weighing at least 6kg to less than 25kg.	
<b>Licensed Indication(s)</b>	As listed in the Summary of Product Characteristics (SmPC).	
<b>Details of HSE Price Agreements (if applicable)</b>	Centrally agreed Commercial in Confidence Patient Access Scheme in place that ViiV Healthcare will share with hospitals directly. The product will be reimbursed in the same manner that drugs of this category are currently being reimbursed. Under the current IPHA Agreement, hospitals can negotiate revised arrangements with individual manufacturers.	
<b>Additional Information</b>		

Best wishes,

Access and Integration Drug Management Programme