

**Programme  
IQWiG in Dialogue  
21 June 2013**

**„Importance of approval status for the benefit assessment“**

**Presentation: Professor Dr Ralf Bender**

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| <b>10.00 - 10.10</b> | <b>Introduction to the topic</b><br>Dr Beate Wieseler, IQWiG   |
| <b>10.10 - 10.30</b> | <b>Criteria for checking the applicability of study results</b><br>PD Dr Ulrich Grouven, IQWiG   |
| <b>10.30 - 11.00</b> | <b>Approval studies and therapeutic indication - experience from the early benefit assessment</b><br>Dr Thomas Kaiser, IQWiG   |
| <b>11.00 - 12.00</b> | <b>Discussion</b>  |
| <b>12.00 - 13.00</b> | <b>Lunch break</b>   |
| <b>13.00 - 13.30</b> | <b>From clinical data to therapeutic indication – a compass for the benefit assessment?</b><br>Dr Karl Broich, Federal Institute for Drugs and Medical Devices, Bonn |
| <b>13.30 - 14.00</b> | <b>New drugs between approval and benefit assessment – how do manufacturers manage?</b><br>Professor Dr Torsten Strohmeyer, GlaxoSmithKline, Munich                  |
| <b>14.00 - 14.30</b> | <b>Importance of approval status for G-BA decisions</b><br>Max Grüne, Federal Joint Committee, Berlin  |
| <b>14.30 - 15.30</b> | <b>Discussion</b>  |
| <b>15.30 - 16.00</b> | <b>Conclusion of the event with coffee and biscuits</b>  |

**Venue: KOMED, Im Mediapark 6, 50670 Cologne**