

APYRAL<sup>®</sup> ACTILOX<sup>®</sup> B



## Metal hydrates for cables

Product	D50 [µm]			Main application			
APYRAL <sup>®</sup> – Ground							
APYRAL® 8	15	1.3	24	• Bedding / Filling compounds			
APYRAL® 16	16	1.8	17	· Bedding / Fining compounds			
APYRAL® – Fine-pr	ecipitated	·					
APYRAL® 40CD	1.5	3.5	22	Thermoplastic			
APYRAL® 60CD	1	6	28	<ul><li>Insulation compounds</li><li>Sheathing compounds</li></ul>			
APYRAL <sup>®</sup> 120E	0.9	11	37	Elastomers <ul> <li>Sheathing compounds</li> </ul>			
APYRAL® 40 VS1	1.5	3.5	33	Crosslinked • Insulation compounds			
APYRAL <sup>®</sup> 60 VS1	1.3	6	45	<ul> <li>Silicone compounds</li> </ul>			
ACTILOX® B - Boeh	mite	·					
ACTILOX <sup>®</sup> B30	2.3	3	28				
ACTILOX® B60	1.2	5	30	<ul> <li>Insulation compounds</li> <li>Sheathing compounds</li> </ul>			
ACTILOX <sup>®</sup> 200SM	0.3	17	36				
APYRAL® – Submicron co-flame retardant							
APYRAL <sup>®</sup> 200SM	0.4	15	48	<ul> <li>Highly flame retardant compounds</li> </ul>			

All data listed in this brochure are reference values and subject to production tolerance. These values are exclusive to the product description and no guarantee is placed on the properties. It remains the responsibility of the users to test the suitability of the product for their application.

## List of abbreviations

BETSpecific surface area [m²/g]DS0Commonly used for average particle sizeE@BElongation at BreakEPDMEthylene-Propylene-Diene-RubberEVAPolyethylene-co-VinylacetateEVMPolyethylene-co-Vinylacetate with a high VA-contentFRLS PVCFire Retardant Low Smoke PVCHFFRHalogen Free Flame RetardantL/DLength versus diameter ratio (extruder parameter)LLDPELinear Low Density PolyethyleneLOILimiting Oxygen IndexLSFOHLow Smoke Free Of HalogenMFIMelt Flow IndexMPaMegapascalMVRMelt Volume RateNBRNitrile-Butadiene-RubberPAPolyethylenePEPolyothylenePFPolyothylenePFPolyothylenePFPolyothylenePFSilicone elastomers (Silicone Rubber)PFThermoplastic elastomerTPOThermoplastic polyolefinTPUThermoplastic polyolefinTPUThermoplastic polyolefinTPUUnderwriters LaboratoriesVAVinylacetateXIPPECosslinked polyethylene						
E@BElongation at BreakEPDMEthylene-Propylene-Diene-RubberEVAPolyethylene-co-VinylacetateEVMPolyethylene-co-Vinylacetate with a high VA-contentFRLS PVCFire Retardant Low Smoke PVCHFFRHalogen Free Flame RetardantL/DLength versus diameter ratio (extruder parameter)LLIDPELinear Low Density PolyethyleneLOILimiting Oxygen IndexLSFOHLow Smoke Free Of HalogenMFIMelt Flow IndexMPaMegapascalMVRMelt Volume RateNBRNitrile-Butadiene-RubberPAPolybutylene TerephthalatePEPolybutylenePEgMAMaleic acid anhydride grafted PEPPPolypopylenePVCPolypopylenePVCPolypopyleneSIRSilicone elastomers (Silicone Rubber)TFEThermoplastic polyoefinTPUThermoplastic polyuerthaneTSTensile StrengthULUnderwriters LaboratoriesVAVinylacetate	BET	Specific surface area [m²/g]				
EPDMEthylene-Propylene-Diene-RubberEVAPolyethylene-co-VinylacetateEVMPolyethylene-co-Vinylacetate with a high VA-contentFRLS PVCFire Retardant Low Smoke PVCHFFRHalogen Free Flame RetardantL/DLength versus diameter ratio (extruder parameter)LLDPELinear Low Density PolyethyleneL0Limiting Oxygen IndexLSFOHLow Smoke Free Of HalogenMFIMelt Flow IndexMPaMegapascalMVRMelt Volume RateNBRNitrile-Butadiene-RubberPAPolyathylenePBTPolyathylenePEPolyathylenePEgMAMaleic acid anhydride grafted PEPPPolyopoplenePVCPolyinylchlotiderpmRevolutions per minSb20, / ATOAntimony trioxideSIRSilicone elastomer (Silicone Rubber)TPUThermoplastic polyoefinTPUThermoplastic polyuerthaneTSTensile StrengthULUnderwriters LaboratoriesVAVinylacetate	D50	Commonly used for average particle size				
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SIR     Silicone elastomers (Silicone Rubber)       TPE     Thermoplastic elastomer       TPO     Thermoplastic polyolefin       TPU     Thermoplastic polyurethane       TS     Tensile Strength       UL     Underwriters Laboratories       VA     Vinylacetate	rpm	Revolutions per min				
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TPOThermoplastic polyolefinTPUThermoplastic polyurethaneTSTensile StrengthULUnderwriters LaboratoriesVAVinylacetate	SIR	Silicone elastomers (Silicone Rubber)				
TPU     Thermoplastic polyurethane       TS     Tensile Strength       UL     Underwriters Laboratories       VA     Vinylacetate	ТРЕ	Thermoplastic elastomer				
TS     Tensile Strength       UL     Underwriters Laboratories       VA     Vinylacetate	ТРО	Thermoplastic polyolefin				
UL     Underwriters Laboratories       VA     Vinylacetate	ТРИ	Thermoplastic polyurethane				
VA Vinylacetate	TS	Tensile Strength				
	UL	Underwriters Laboratories				
XLPE Crosslinked polyethylene	VA	Vinylacetate				
	XLPE	Crosslinked polyethylene				

# Content

## 01 Flame retardancy of cables

## 02 APYRAL®

Compounding 0	9
LSFOH compounds 1	1
Compounds for insulation and sheating1	1
Influence of BET-surface area 1	2
Mechanical properties, MVR and LOI 1	3
Water uptake 1	4
Coupling agents 1	
Exemplary formulations 1	6
PE / EVA and vinylsilane / peroxide1	
PE / EVA with PEgMA 1	8
PE / TPO with PEgMA 1	
EVA / EVM with aminosilane (non crosslinked) 2	0
Bedding / Filling compounds 2	1
Elastomer compounds (crosslinked) 2	1
Process aid ACTILOX® PA-14	3

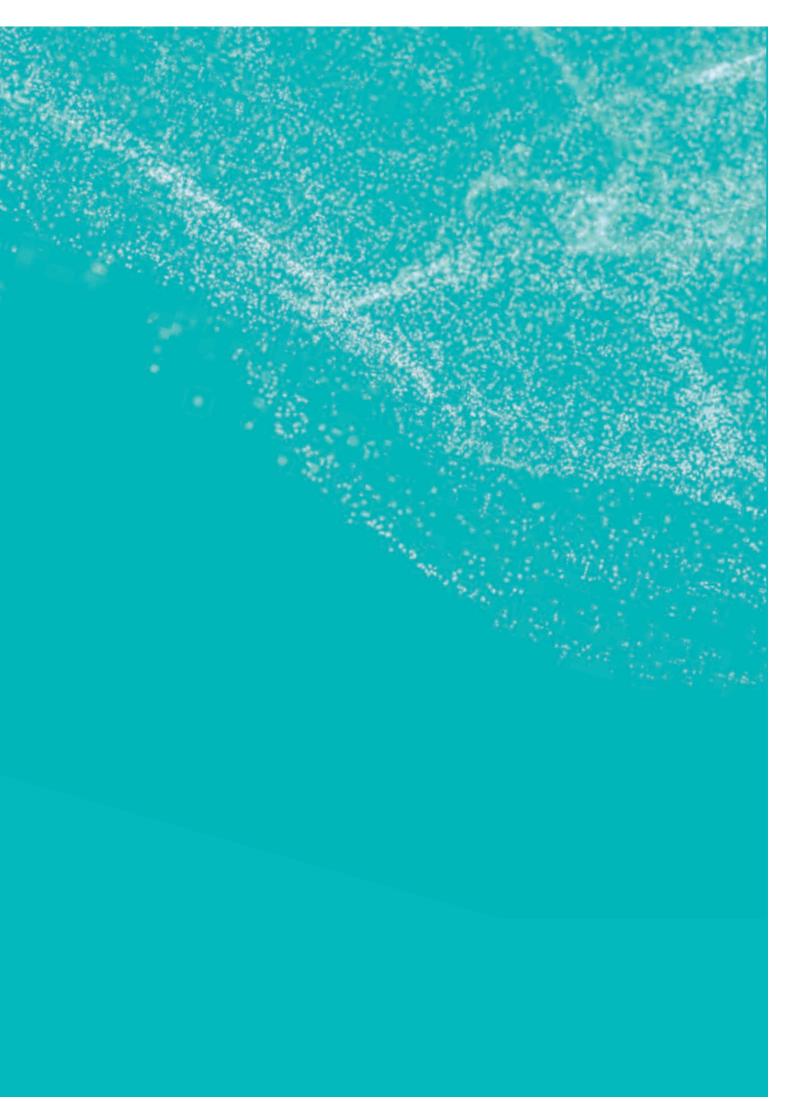
## 03 ACTILOX<sup>®</sup> B (Boehmite)

EVA with aminosilanes	28
Synergistic effects with ACTILOX <sup>®</sup> 200SM	28
ACTILOX <sup>®</sup> 200SM as char forming agent	28
Increased LOI / Improved flame retardancy	29

## 04 APYRAL<sup>®</sup> 200SM – A submicron scale synergists

## 05 FRLS PVC compounds

Smoke suppressed PVC compounds
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### Flame retardancy of cables

Wires and cables are present in our everyday life. Cables are found in buildings, cars and public transport vehicles as well as in all electric appliances.

Preventive fire protection of cables is a topic of ever increasing importance since forty years now.

Especially in buildings, spread of fire by cables is an important issue. Electrical, machinery and data cables found in buildings can be several kilometers long. They are installed horizontally within individual levels and penetrate multistorey buildings vertically.

Flame retarded cables can prevent a small cause, such as an electrical short circuit, from becoming a major fire catastrophe which can result in major material damages and even loss of lifes.

Over the years, metal hydrates, especially aluminium hydroxide and magnesium hydroxide have been established as the most important flame retardants for the wire and cable industry.

In particular their environmental friendliness and their favourable priceperformance ratio make **APYRAL**<sup>®</sup>, aluminium hydroxide and **ACTILOX**<sup>®</sup> **B**, aluminium oxide hydroxide (boehmite) important and sustainable flame retardants.

<b>APYRAL®</b>	AI(OH) <sub>3</sub>		
Chemical	Aluminium hydroxide		
Mineral	Gibbsite		
Common name	Aluminium trihydrate		
Abbreviation	ATH		
APYRAL <sup>®</sup> AOH ACTILOX <sup>®</sup> B	АЮОН		
	AlooH Aluminium oxide hydroxide		
ACTILOX® B			
ACTILOX® B Chemical	Aluminium oxide hydroxide		

## **APYRAL**<sup>®</sup>



## **APYRAL®**

Our **APYRAL**<sup>®</sup> products for wire and cable applications can be divided into two classes on the basis of property profiles:

- · fine precipitated grades
- ground grades

Fine precipitated **APYRAL**<sup>®</sup> products are broadly used to manufacture flame retardant insulation and especially sheathing compounds.

Of highest importance are **APYRAL® 40CD**, **APYRAL® 60CD** and **APYRAL® 120E** used in halogen free flame retardant (HFFR) cable compounds, also called LSFOH (Low Smoke, Free Of Halogen) cable compounds.

**APYRAL® 40 VS1** and **APYRAL® 60 VS1** are vinylsilane treated products, especially used in crosslinked LSFOH compounds based on EPDM, EVM and SIR (silicone rubber).

Additionally, increase of flame retardancy of PVC cable compounds and especially their smoke reduction is another important wire and cable application for fine precipitated **APYRAL®** products.

The ground grades **APYRAL® 8** and **APYRAL® 16** find widespread use in highly filled flame retardant bedding compounds.

**APYRAL**<sup>®</sup> products have a very high chemical purity of at least 99.5 %. The remaining constituents consist mainly of sodium oxide, which is part of the crystal lattice and partly adhering to the **APYRAL**<sup>®</sup> surface in hydrated form, as well as traces of iron and silica. Due to its whiteness **APYRAL**<sup>®</sup> behaves neutrally to the colouring of polymers. The Mohs hardness of 2.5 − 3 causes no problem even in highly filled molten masses. The relatively high heat capacity c<sub>p</sub> of 1.65 J/g\*K at 400 K (127 °C) has a beneficial effect on the dimensional stability under heat for **APYRAL**<sup>®</sup> filled polymers.

With a specific density of 2.4 g/cm<sup>3</sup>, **APYRAL**<sup>®</sup> is a medium dense mineral filler. As a result of the required filling ratios, the density of the flame retarded plastics is increased compared to the unfilled polymer. However, **APYRAL**<sup>®</sup> is one of the most attractive flame retardants, even on the basis of a volume-specific cost balance.

Application	APYRAL®					Polymers		
	8	16	40CD	60CD	120E	40 VS1	60 VS1	
Bedding	•	•						EPDM, EVM, EVA, TPO
Insulation			•	•		•	•	PE, XLPE, EVA, PVC
Sheathing			•	•	•	•	•	PE, EVA, TPO, TPE, TPU, PVC, EVM, EPDM, NBR, SIR

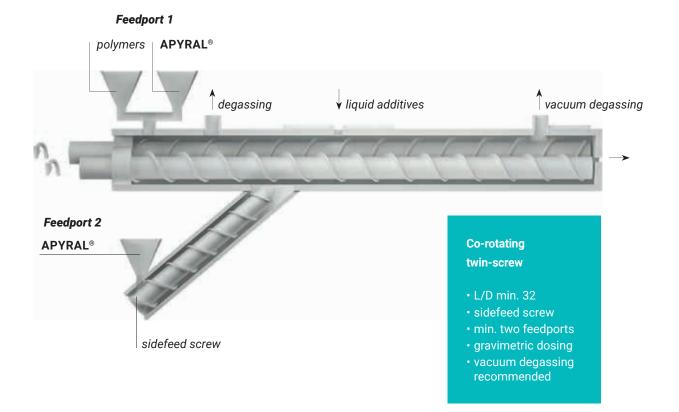
### Compounding

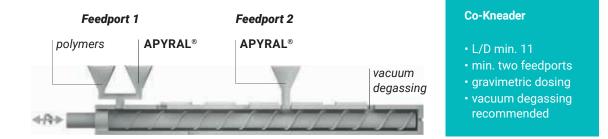
When producing **APYRAL**<sup>®</sup>-filled thermoplastics via melt compounding, special attention must be given to the process temperature limit of 200 °C.

The temperature limit may vary in a marginal way depending on the residence time and the applied shear force during compounding in different compounding machines (see our brochure "Mineral Based Flame Retardancy with Metal Hydrates"). Therefore, it is recommended that the temperature of the molten mass is kept below 190 °C. In the case of continuous processing, higher temperatures are possible for short residence times. However, this option must be tested depending on the individual compound formulation and machinery settings. In general, temperature control must be as exact as possible.

In the manufacture of filled elastomers as well as thermoplastics, the internal mixer (kneader) has proved most flexible. The continuous processes of the co-kneader and the corotating twin screw extruder offer better results in terms of dispersion and quality consistency. These compounding machines exhibit via their special designed screw geometry and special kneading elements excellent dispersing abilities, even in highly filled compounds.

The twin screw extruder should have an L/D ratio of at least 32 and two feeding ports. In order to ensure constant filling levels, gravimetric dosing of **APYRAL**<sup>®</sup> is recommended. The bulk of the filler should be applied via the second feeding port (preferred via a sidefeed screw) into the polymer melt. Vacuum degassing combined with a number of ventilation openings ensure a good ventilation of the molten mass and the removal of volatile organic substances.





Contrary to the co-rotating twin screw extruder, the co-kneader features one single screw which is put into an axial oscillation additional to the screw rotation. The compound is kneaded and transported through intermitting kneading elements and hence dispersed in a smooth oscillating manner. The co-kneader should have an L/D ratio of min. 11 and two feeding ports.

2/3 of the **APYRAL**<sup>®</sup> is generally applied via the first port along with the polymer. For the dosing of fluid additives the injecting between the first and the second feed port through a hollow kneading bolt has proved most effective.

Vacuum degassing before entry of the molten mass into the discharge screw is recommended, particularly when using silanes. In principle, screw geometries with low compression should be used for both continuous compounding processes. Specific screw designs are based on the compound formula, degree of filling and the **APYRAL**<sup>®</sup> grade used.

Recently, it became more common to use co-kneaders with higher L/D ratios (up to 22), especially for the production of silane crosslinkable HFFR compounds in a onestep procedure.

### LSFOH compounds

Low smoke free of halogen (LSFOH) compounds are sometimes based on elastomers such as EPDM or NBR, but much more frequently on polyolefines, in particular PE and its copolymers.

In Europe permanently installed cables are classified according to CPR (Construction Products Regulation, EU 305/2011). For cables, EN 50575 defines the different classifications. The most important test method is EN 50399, which is a vertical cable bundle test with integrated measurement of temperature, heat and smoke evolution. Reaction to fire, resistance to fire and release of dangerous substances are the main characteristics in the European as well as in international standards for flame retardant cables.

In order to pass these tests, compounds filled with 60 – 65 wt.-% of **APYRAL**<sup>®</sup> must be used.

Alongside the material properties, determined by the polymer and flame retardant used, the combustion behaviour in these cable tests, however, depends to a great extent on the respective cable geometry and cable design. Thus, the quoted filling levels are only guidelines.

### Compounds for insulation and sheathing

Because of its excellent insulation properties, crosslinked low density polyethylene (XLPE) is by far the most important coating material used to prevent short circuiting of electrical conductors. XLPE is also the most used insulation material for flame retardant cable constructions. Anyhow, LSFOH compounds based on fine precipitated **APYRAL**<sup>®</sup> are also applied in wire insulation. **APYRAL**<sup>®</sup> grades have a very low electrolyte content which guarantees for the lowest possible impact on compound insulation properties.

Most important is the use of LSFOH compounds as outer sheathing of cable constructions. This enables very effective fire prevention against external ignition sources.



### Influence of BET-surface area

The table shows the formulation as well as the main characteristic data for three fine precipitated **APYRAL**<sup>®</sup> grades in a test compound based on an EVA type with a VA-content of 26 %.

The most important parameter to distinguish between the fine precipitated **APYRAL**<sup>®</sup> grades is the specific surface area according to BET. Ranging from 3.5 m<sup>2</sup>/g for **APYRAL**<sup>®</sup> **40CD** and 6 m<sup>2</sup>/g for **APYRAL<sup>®</sup> 60CD** up to 11 m<sup>2</sup>/g for **APYRAL<sup>®</sup> 120E**, the BET surface area is covering a broad span.

In addition to the table, the influence of the **APYRAL**<sup>®</sup> grade and its BET surface on mechanical properties (tensile strength and elongation at break), flame retardancy according to LOI and processability (MVR) is visualized on the following page. The values displayed in the two graphs are taken from a LSFOH-compound based on an EVA grade with a VA-content of 19 %.

With increasing specific surface area of the mineral flame retardant a decrease in elongation at break but an increase in tensile strength is observed.

The melt index, cited in the table as Melt Volume Rate (MVR), is an important value to estimate the processing behavior of compounds during extrusion of insulated conductors or cables. The high value for **APYRAL® 60CD** and especially **APYRAL® 40CD** is remarkable and allows the design of fast extrudable LSFOH compounds.

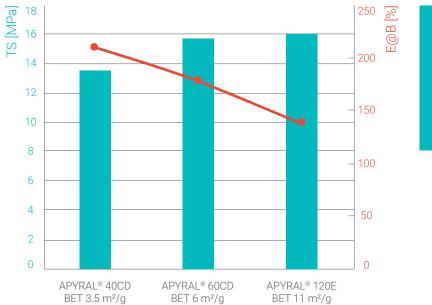
Component [wt%]		APYRAL® 40CD	APYRAL® 60CD	APYRAL® 120E			
EVA, Escorene™ UL 00226		38.3	38.3	38.3			
Aminosilane, Dynasylan® AM	/IEO	0.4	0.4	0.4			
APYRAL® 40CD		61.3	-	-			
APYRAL® 60CD		-	61.3	-			
APYRAL® 120E				61.3			
Characteristic data	Characteristic data						
Tensile Strength (TS) [MPa]		12.6 15		17.6			
Elongation at Break (E@B)	[%]	243	243 206				
MVR at 190 °C / 21.6 kg [cm³/10 min]		10.6	7.3	2.3			
LOI	[% O <sub>2</sub> ]	37	42	45			

### EVA (26 % VA-content) and Aminosilane

Compounding on internal lab kneader. Samples made of compression moulded plaques

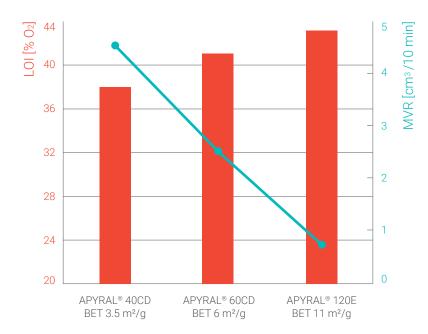
### Mechanical properties, MVR and LOI (EVA with 19 % VA-content)

EVA, Escorene UL 00119 – 38.3 wt.-% Aminosilane, Dynasylan (AMEO) – 0.4 wt.-% **APYRAL® 40CD** / **60CD** / **120E** – 61.3 wt.-%



## Mechanical properties with increasing BET

- increase in tensile strength
- decrease in elongation at break

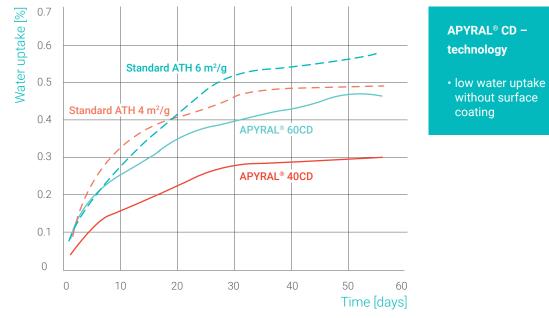


### Flame retardancy and processability with increasing BET

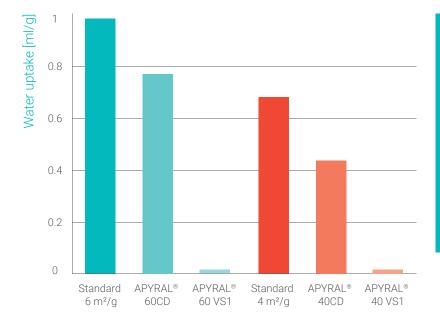
- increase in LOI
- decrease in MVR

### Water uptake

Water uptake of LSFOH compounds is essential for insulated wires. But also when used as sheathing material, a low water uptake is crucial to pass critical ageing tests performed under water bath conditions. The diagram on the left displays the water uptake for an EVA compound based on **APYRAL® CD** grades and standard ATH grades of similar fineness.



Water uptake of EVA, Escorene UL00119 filled with 61.3 wt.-% ATH / APYRAL<sup>®</sup>; T = 23 °C



With decreasing BET, decrease of filler water uptake

By hydrophobic coating further decrease possible

#### **APYRAL® VS1**

 lowest water uptake by vinylsilane coating

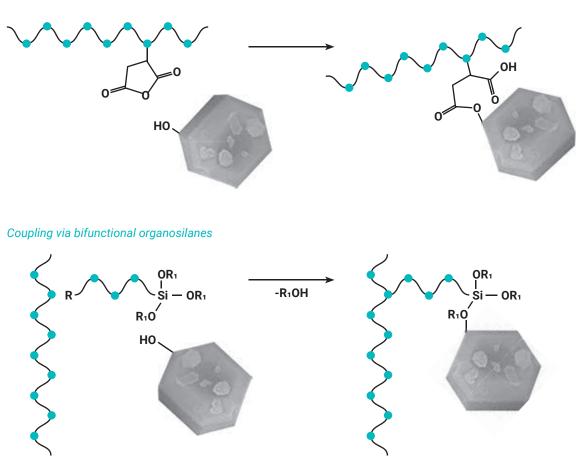
Water uptake of ATH / **APYRAL®** powder according to Baumann values of saturation after 30 min

**APYRAL® CD** results in significantly reduced water uptake compared to standard ATH grades of comparable BET surface. The reason for this is related to the comparably low water uptake of the **APYRAL**<sup>®</sup> **CD** powders, which is shown in the second graph.

### **Coupling agents**

The use of coupling or bonding agents to achieve good physical properties in highly filled LSFOH compounds is elementary. Coupling agents enable a binding between inorganic **APYRAL**<sup>®</sup> filler and polymer. The most preferred way of binding is by chemical reaction, but strong enough physico-chemical reaction is also feasible.

Coupling with maleic-acid-anhydride grafted polymers



 $R = NH_2$  (H-bridging and covalent bonding on polar groups within polymer) R = - (covalent bonding on polymer, peroxide initiated)  $R_1 = CH_{3_2} C_2 H_5$  Alongside the use of polyolefines grafted with maleic anhydride (in case of PE abbreviated as PEgMA), the use of bifunctional organosilanes, in particular with amino or vinyl end group, is the most widely used technology.

Both product groups function as chemical mediators between the mineral with a polar, hydroxide groups carrying surface and the polymer. The illustration on previous page shows the basic chemical mechanisms.

The reaction of the silane with the hydroxide surface releases ethyl- and/or methyl alcohol which must be removed.

Organosilanes are usually used in an amount of 1 % of the overall filler mass. In the case of

### **Exemplary formulations**

### PE / EVA and vinylsilane / peroxide

The compound formulation listed on the next page shows a simple thermoplastic recipe based on PE / EVA and vinylsilane. Peroxide concentration is high enough to graft the vinyl groups onto the polymer backbone, but small enough to avoid crosslinking of polymer chains.

### **APYRAL® CD** technology

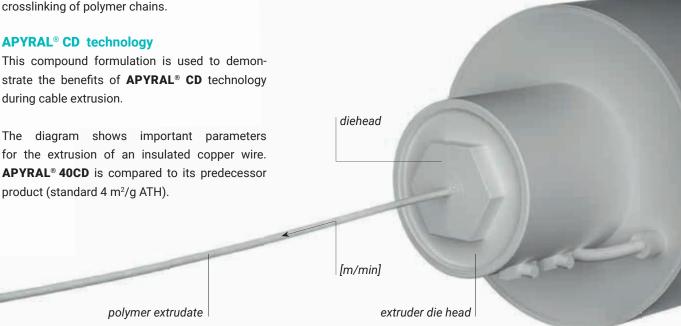
for the extrusion of an insulated copper wire. APYRAL® 40CD is compared to its predecessor product (standard 4 m<sup>2</sup>/g ATH).

PEgMA the dose is a few percent of the overall composition.

While PEgMA products are generally supplied in pellet form, organosilanes are liquid. If dosing of very small quantities of fluids is non favourable, they can also be used as a master batch on a porous polymer carrier.

Alternatively, compounders may use precoated fillers, like APYRAL® 40 VS1 and APYRAL® 60 VS1. In this case the silane is already bound to the metal hydrate surface.

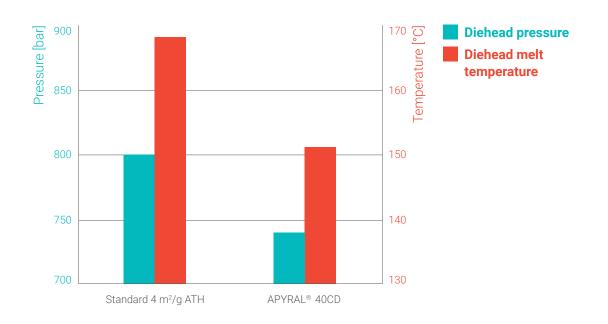
The significantly reduced values for diehead pressure and diehead melt temperature for APYRAL® 40CD enable an approximate 30 % increase in extrusion speed.



Component	[wt%]	Trade name	
LLDPE	9.66	ExxonMobile™ LL 1004 YB	
EVA	29 Escorene <sup>™</sup> UL 00226		
Vinylsilane	0.8	Silquest™ FR-693	
Process aid	0.3	Silquest™ PA-826	
Peroxide	0.04	Interox <sup>®</sup> TMCH-75-AI	
Stabiliser	0.2	Irganox® 1010	
Aluminium hydroxide	60	APYRAL <sup>®</sup> 40CD	

### Characteristic data

Tensile Strength	[MPa]	11
Elongation at Break	[%]	260
MVR at 160 °C / 21.6 kg	[cm <sup>3</sup> /10 min]	9.4
LOI	[% O <sub>2</sub> ]	37



Diehead pressure and diehead melt temperature at:

- 0.5 mm<sup>2</sup> copper wire (single, round)
- die diameter 1.4 mm
- insulation thickness approx. 0.35 mm
- line speed 650 m/min

### PE / EVA with PEgMA

Compounding in Werner & Pfleiderer dispersion kneader (kneading chamber); Specimen made of compression moulded plaques

Component		[wt%]	Trade name
LLDPE		10	ExxonMobile™ LL 1004 YB
EVA		24.8	Escorene™ UL 00226
PEgMA		5	Compoline CO/LL
Stabiliser		0.2 Irganox® 1010	
Aluminium hydroxide		60	APYRAL <sup>®</sup> 40CD / 60CD
Characteristic data		APYRAL® 40CD	APYRAL® 60CD
Tensile Strength	[MPa]	12.6	13.6
Elongation at Break	[%]	227	214
MVR at 160 °C / 21.6 kg	[cm <sup>3</sup> /10 min]	3.8	2.8
LOI	[% O <sub>2</sub> ]	33	34

Just like the previous exemplary LSFOH compound, the above listed formulation is based on a blend of PE and EVA. In this case, maleic anhydride grafted PE (PEgMA) was used as coupling agent.

This compound results in very good balanced physical properties for both **APYRAL® CD** grades. The LOI of the PEgMA coupled compound is slightly reduced in comparison to the vinylsilane coupled formulation on page 17.

Furthermore, differences in the melt viscosity (given as MVR) of the compounds are observable using the same ATH grade but different coupling agents. Besides the fundamental difference in coupling agent and the comparison of the characteristic compound properties, one has to consider that the above values have been examined on compounds produced on a discontinuous lab kneader and specimen, which have been cut out of compression moulded plaques instead of extruded tapes. This important influence of the compounding technology and specimen preparation is demonstrated for the following basic formulation on page 19.

A polymer blend of PE and a PE-copolymer (TPO) was used, comprising a filler loading level of 65 wt.-% and a PEgMA coupling agent. The characteristic data generated on specimen made of compression moulded plaques and compounds produced on the lab kneader are excellent for both **APYRAL® CD** grades.

For **APYRAL® 40CD** it is demonstrated that compounding equipment and technology used for specimen preparation have a significant influence on the final properties. The extruded tapes result in further improved tensile and elongation balance and compound pellets produced by co-kneader give higher MVR values.

## PE / TPO with PEgMA

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Influence of compounding	equipment on	meenumour		Lor orr compoundo

Component		Trade name		[wt%]
LLDPE		ExxonMobile™ LL 1004 YB		13.3
TPO (PE-co-octene)		Engage <sup>™</sup> 8452		17.5
PEgMA		Fusabond® E MB 226 D		4
Stabiliser		Irganox <sup>®</sup> 1010		0.2
Aluminium hydroxide		APYRAL <sup>®</sup> 40CD / 60CD		65
Compounding		Buss-Co-Kneader	Internal mixer	Internal mixer
Specimen		Extruded tape	Plaques	Plaques
Characteristic data		APYRAL <sup>®</sup> 40CD		APYRAL® 60CD
Tensile Strength	[MPa]	16.8	13.8	14
Elongation at Break	Iongation at Break [%]		244	188
MVR at 160 °C / 21.6 kg	MVR at 160 °C / 21.6 kg [cm <sup>3</sup> /10 min]		2.6	2.5
LOI [% 0 <sub>2</sub> ]		34	33	33

The compound below is based on an EVA with 26 % VA-content blended with an EVM grade. EVM are also copolymers of VA and ethylene, but comprising very high VA-contents (in this case 60 %).

These copolymers show rubber like properties. EVM accepts very high filler loads, but has a plasticizing effect.

Component		Trade	<b>[wt%]</b>	
EVA		Escorene™ UL 00226		29
EVM		Levamelt <sup>®</sup> 600		5.6
Aminosilane		Dynasylan® AMEO		0.4
Aluminium hydroxide		APYRAL <sup>®</sup> 40CI	60	
Characteristic data		APYRAL <sup>®</sup> 40CD	APYRAL® 60CD	APYRAL® 120E
Tensile Strength [MPa]		6.1	10.3	14.3
Elongation at Break [%]		289	226	172
MVR at 160 °C / 21.6 kg [cm <sup>3</sup> /10 min]		8.7	4.7	1.7
LOI	[% 0 <sub>2</sub> ]	40	46	47

Compounding on lab dispersion kneader. Specimen made of compression moulded plaques



**APYRAL® 120E** most preferred for rubers and rubber-like compounds

Consequently, compounds containing high contents of EVM have to be crosslinked to achieve sufficiently good tensile properties. Therefore, mostly silica and/ or carbon black are added as reinforcing fillers.

With a moderate level of EVM it is sufficient to counterbalance by the right choice of **APYRAL**<sup>®</sup> grade.

When using **APYRAL**<sup>®</sup> **40CD** the tensile strength is far too low for the non-crosslinked compound.

In contrast, for **APYRAL® 60CD** and especially **APYRAL® 120E** the higher specific surface areas pay out for such types of LSFOH compounds.

**APYRAL® 60CD** and **APYRAL® 120E** are the most preferred grades for elastomer compounds on the basis of NBR, EPDM, and EVM as well as extremely soft thermoplastic polymer blends e.g. based on copolymers of acrylic esters and ethylene.

### Bedding / Filling compounds

Electrical cables are usually designed with a filling mass surrounding the individual insulated conductors and thus filling the empty space between the insulated conductors and the cable sheath. These bedding compounds are also mineral filled. In order to design flame retarded cables with lowest possible fire load, **APYRAL**<sup>®</sup> is also used here, partially or fully substituting calcium carbonate.

The mechanical requirements for a filling mass are very low so that such compounds are designed with extremely high loadings of ground **APYRAL**®



grades. The following is a basic example based on **APYRAL**<sup>®</sup> **8** and a PE-co-butene.

Filling compounds are generally based on PE-coalkenes, EPDM or EVA with a high VA-content.

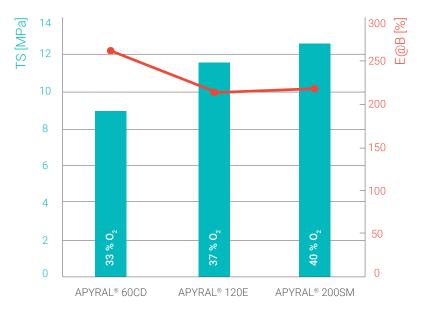
Component	[wt%]	MVR (190 °C/21.6 kg)	LOI
PE-co-butene (Engage™ ENR 7380)	15		(1) <sup>W</sup> 0
PE-Wax (Licowax® PE 520)	4	10.6 cm³/10 min	
Stearic acid	1		62 % 0 <sub>2</sub>
APYRAL® 8	80		

Compounding on lab dispersion kneader. Specimen made of compression moulded plaques

### Elastomer compounds (crosslinked)

On the next page EVM based elastomer compounds with a VA-content of 70 % are exemplarily shown. Compared to the commonly used **APYRAL® 60CD** and **APYRAL® 120E** grades, submicron sized **APYRAL® 200SM** shows an advantageous balance of elongation at break and tensile strength combined with a high flame retardant efficiency given as LOI (see graph p. 22). The submicron sized particles and the corresponding high specific surface area of **APYRAL® 200SM** is pivotal to a flame retardant boost effect reflected in the remarkable high LOI of 40 %.

Additional information regarding elastomers and representative compound formulations are given in our brochure "Metal Hydrates in Elastomers".



Mechanical properties and LOI of EVM 700 compounds with 160 phr  ${\bf APYRAL}^{\circ}$ 

### Elastomer Compound / EVM with 70 % VA-content

Component [phr]		APYRAL® 60CD	APYRAL® 120E	APYRAL® 200SM
EVM, LEVAPRENE <sup>®</sup> 700		100	100	100
Zinc Borate	Zinc Borate		10	10
Plasticiser (mix)		9.9	9.9	9.9
Stabiliser (mix)		4	4	4
Crosslinker (mix)		7	7	7
Adhesive	Adhesive		1.6	1.6
APYRAL® 60CD		160	-	-
APYRAL® 120E		-	160	-
APYRAL <sup>®</sup> 200SM		-	-	160
Characteristic data				
Tensile Strength	[MPa]	8.9	11.6	12.6
Elongation at Break [%]		264	214	220
MVR at 160 °C / 21.6 kg	MVR at 160 °C / 21.6 kg [cm³/10 min]		36	45
Shore A (at 23 °C)	Shore A (at 23 °C) (dimensionless)		77	81
LOI	[% O <sub>2</sub> ]	33	37	40

Technical data given after crosslinking, except MVR viscosity

### **Process aid ACTILOX® PA-14**

The benefits of **APYRAL® CD** technology on processability of LSFOH compounds, especially the tremendous increase in the line speed compared to standard fine precipitated ATH grades, has already been discussed on page 14. To enable even better processing of LSFOH compounds Nabaltec AG has developed a processing aid designed to ease extrusion, but avoiding an increase in diehead temperature.

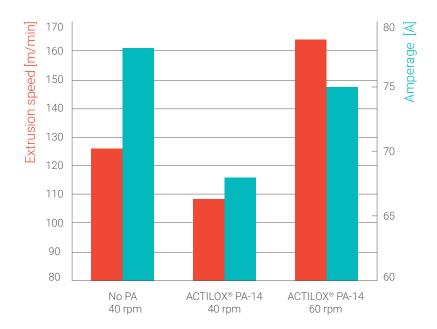
**ACTILOX**<sup>®</sup> **PA-14** is a powdery process aid based on mineral carriers and silicones added at 0.5 – 2 wt.-% during compounding.

The formation of "die drool" is a principle problem of highly filled compounds. It describes the leakage of resin around the nozzle area of the steel block through which the wire is extruded.

The diagram displays extrusion speed and amperage of the pilot plant extruder used.

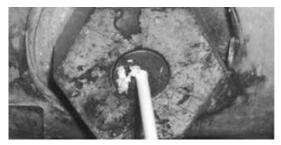
When using **ACTILOX**<sup>®</sup> **PA-14** at 1.2 wt.-% and keeping extruder revolution at 40 rpm, amperage is significantly reduced, but unfortunately also the line speed (insulated copper with 3 mm diameter).

At increased extruder revolution (60 rpm) the potential of **ACTILOX® PA-14** is fully utilized. The line speed increases by 30 % while amperage remains at the same level as for the reference compound not containing any process aid.

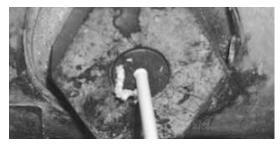


*Extrusion speed and amperage during extrusion of a LSFOH compound; Left without, middle and right with* **ACTILOX® PA-14** 

### After 3 min.



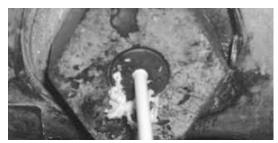
### After 6 min.



### After 9 min.

When the "die drool" has achieved a critical size and mass, it can be drawn away by the insulated wire and often causes failures within the insulation. Out of specification production and extruder shut downs are the unavoidable consequence.

The addition of **ACTILOX® PA-14** very effectively retards "die drool" formation. Additionally the resin residue formed is not conically closed around the nozzle, but forms on one side of the die only. When growing it slowly drifts away from the nozzle and finally drops. A potential pick up by the insulated wire can be avoided. This effect is displayed by the three images.



Drifting and dropping "die drool" during extrusion of a LSFOH-compound containing **ACTILOX® PA-14** 

# ACTILOX® B (Boehmite)



## **ACTILOX® B (Boehmite)**

ACTILOX® **B**, boehmite, is finding an ever-increasing range of new applications as functional filler, especially when higher temperature stability is required. Hence, highmelting compounds with process temperatures of up to 340 °C can be processed using **ACTILOX® B**.

All our **ACTILOX**<sup>®</sup> **B** products are extremely pure, crystalline boehmites with a very low ATH residue (purity min. 99 %). This guarantees an extraordinarily high temperature stability.

**ACTILOX**<sup>®</sup> **B** can be processed up to 340 °C. Therefore it can be used in TPOs and TPEs.

Furthermore **ACTILOX**<sup>®</sup> grades are applicable in technical compounds based on PA, PBT etc. In such polycondensates strongly alkaline fillers such as magnesium hydroxide can lead to hydrolytic decomposition of the polymer chains.

Therefore **ACTILOX**<sup>®</sup> **B** is much more favourable for polycondensates due to its relatively low basicity and moisture content.

#### Boehmite application in cable compounds

Application		ACTILOX	B	Delumere
Application	B30	B60	200SM	Polymers
Insulation	•	•		PP, PP-copolymers, PE, XLPE, EVA
Sheathing	•	•	•	PP, PP-copolymers, PE, XLPE, EVA, TPO, TPE, EVM, EPDM, NBR

Its high heat capacity ( $c_p=1.54$  J/gK at 500 K, 227 °C) combined with high temperature stability make **ACTILOX**<sup>®</sup> boehmites ideal for use in electrical insulation heat sinks.

Our boehmites show very low electrolyte contents. This predestines **ACTILOX® B** for electrical insulation applications.

Its Mohs hardness of 3 causes no tool abrasion problems even in highly filled molten masses.With a specific density of 3 g/cm<sup>3</sup>, **ACTILOX® B** is a medium dense mineral filler.

Additionally, **ACTILOX**<sup>®</sup> **B** shows a very good chemical resistance, especially a very high acid resistance.

### Solubility in battery acid

Mineral filler	dissolved fraction in 34 % sulphuric acid (H <sub>2</sub> SO <sub>4</sub> )
<b>APYRAL® 40CD</b> Aluminium hydroxide	8 %
ACTILOX® B30 Boehmite	0.2 %

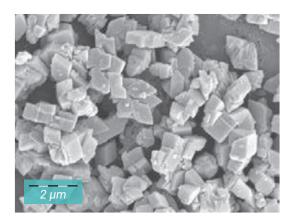
The **ACTILOX**<sup>®</sup> **B** grades differ in their specific BET surface area and median particle size D50.

The fineness increases from **ACTILOX**<sup>®</sup> **B30** (BET= 3 m<sup>2</sup>/g, D50 = 2.3  $\mu$ m) via **ACTILOX**<sup>®</sup> **B60** (BET= 5 m<sup>2</sup>/g, D50 = 1.2  $\mu$ m) to **ACTILOX**<sup>®</sup> **200SM** (BET= 17 m<sup>2</sup>/g, D50 = 0.3  $\mu$ m).

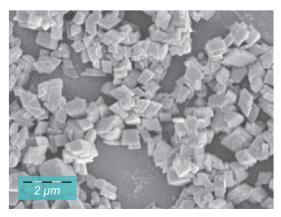
ACTILOX® B30 and ACTILOX® B60 are characterized by an excellent processability. This is due to the unique cubic like morphology of the boehmite crystals these two ACTILOX® B grades consist of. This morphology leads to low viscosities and causes a very good dispersability in the polymer matrix respectively in the cable compound. Therefore ACTILOX® B30 and ACTILOX® B60 are especially destinated for fast extrudable LSFOH compounds, whenever high line speeds and high extrusion temperatures are required.

**ACTILOX® 200SM** has proved being effective in increasing flame retardancy of LSFOH compounds when used together with **APYRAL®** and magnesium hydroxide. This combination of mineral fillers improves the ash or char stability as well as the LOI in a given compound (see pages 28 – 31).

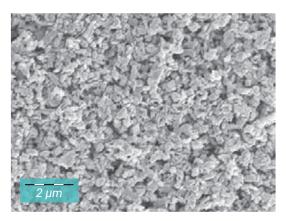
The table on page 28 shows a simple compound based on an EVA grade with a VA-content of 19 % and the most important compound data for the three different **ACTILOX**<sup>®</sup> boehmite grades.



SEM image of ACTILOX® B30



SEM image of ACTILOX® B60



SEM image of ACTILOX® 200SM

### **EVA with aminosilanes**

Component [wt%]		ACTILOX <sup>®</sup> B30	ACTILOX <sup>®</sup> B60	ACTILOX® 200SM		
EVA, Escorene™ UL 00119		38.3	38.3	38.3		
Aminosilane, Dynasylan®	AMEO	0.4	0.4	0.4		
ACTILOX <sup>®</sup> B30		61.3	-	-		
ACTILOX® B60		-	61.3	-		
ACTILOX <sup>®</sup> 200SM		-	-	61.3		
Characteristic data	Characteristic data					
Tensile Strength	[MPa]	12.5	16.8	18		
Elongation at Break [%]		219	221	165		
MVR at 190 °C / 21.6 kg	MVR at 190 °C / 21.6 kg [cm³/10 min]		3.5	0.5		
LOI [% 0 <sub>2</sub> ]		26	27	34		

Compounding on lab dispersion kneader. Specimen made of compression moulded plaques

The trends observed for mechanical properties and processability (MVR) are comparable to the BET dependant relationships found for fine precipitated **APYRAL**<sup>®</sup> (see pages 12 - 13).

For flame retardancy according to LOI, only a small influence of BET surface difference between **ACTILOX® B30** and **ACTILOX® B60** is detectable. Anyhow, the explicitely higher BET surface of

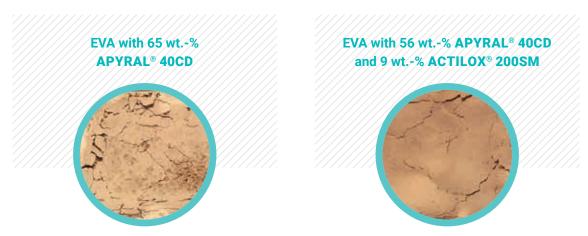
ACTILOX<sup>®</sup> 200SM results in a significant LOI increase.

Generally, boehmites of the **ACTILOX**<sup>®</sup> product range result in lower LOI values compared to fine precipitated **APYRAL**<sup>®</sup> at identical loadings. For more detailed information of such general considerations on flame retardancy, we kindly request to refer to our brochure "Mineral Flame Retardancy with Metal Hydrates".

### Synergistic effects with ACTILOX<sup>®</sup> 200SM

ACTILOX® 200SM as char forming agent

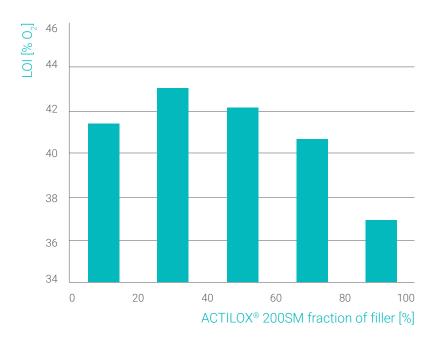
When using **ACTILOX**<sup>®</sup> **200SM** as a co-additive to **APYRAL**<sup>®</sup>, the remaining char after combustion forms a very homogenous, nearly fully closed surface as the below comparison of the residues of two plaques tested in the cone calorimeter shows. During the start of a fire, chars serve as a protective shield against heat and retard the release of volatile and flammable decomposition products on the incipient thermal breakdown of the polymer matrix. This shielding effect and especially transport inhibition is the more successful the more closed and stable the char layer is. Char / ash formation after Cone-Calorimeter



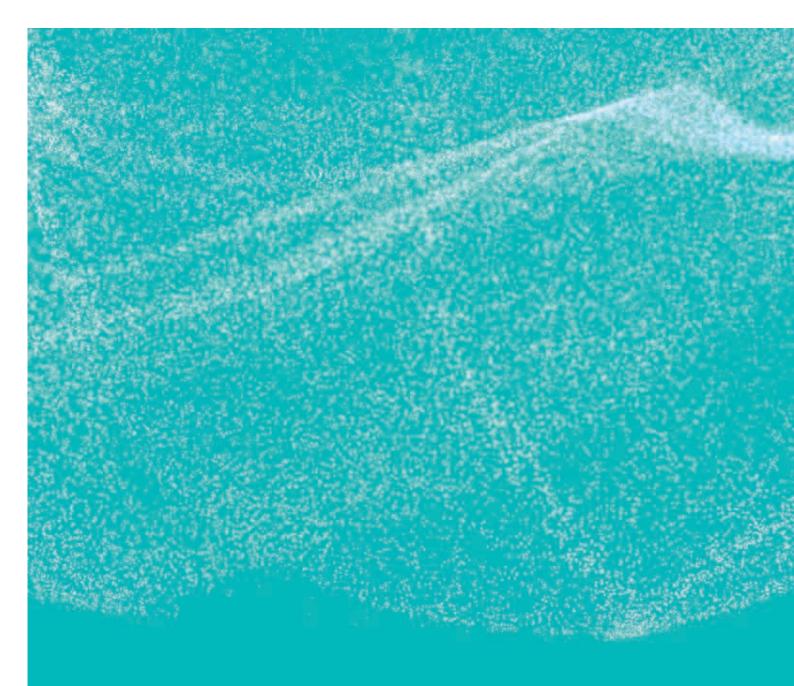
Increased LOI / Improved flame retardancy

As mentioned before, combinations of **ACTILOX® 200SM** and **APYRAL®** exhibit a flame retardant efficiency that each individual component will not achieve at comparable filling levels. This synergistic effect is given in the graph below, showing a successive substitution of **APYRAL®** by **ACTILOX® 200SM**. Obviously this

combination passes an effective maximum at a ratio of 30 % + 70 % (related to an overall filling level of 61.3 wt.-% in an EVA compound). At this ratio, the LOI increases by 3.5 % compared to the sole use of **APYRAL® 60CD**. Only when the boehmite content exeeds 80 %, the LOI drops below the reference values (ATH or MDH without boehmite).



LOI as a function of the mineral filler composition (total filling level 61.3 wt.-%); LOI development by a successive exchange of **APYRAL® 60CD** with **ACTILOX® 200SM** in an EVA compound



# APYRAL® 200SM

### APYRAL<sup>®</sup> 200SM – A submicron scale synergist

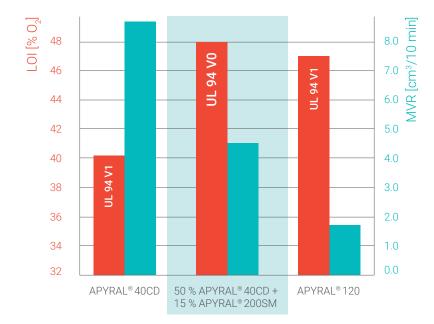
**APYRAL® 200SM** is an ultrafine precipitated ATH. This product can be used in combination with standard ATH products to boost flame retardancy at same filling levels or to reduce the total filler load of a compound at same level of flame resistivity.

In the following, the advantages of **APYRAL® 200SM** are demonstrated in an EVA/EVM compound. Compared to a common **APYRAL®** filled compound, a mixture of **APYRAL® 40CD** and **APYRAL® 200SM** shows best balance of LOI and processability / viscosity.

LOI and MVR for three compounds based on standard **APYRAL**<sup>®</sup> grades only and a com-

pound based on a blend of **APYRAL® 40CD** and **APYRAL® 200SM** are compared in the graph.

The blend of **APYRAL® 40CD** and **APYRAL® 200SM** shows an increase of 8 % O<sub>2</sub> in LOI compared to **APYRAL® 40CD** and a MVR comparable to the compound based on **APYRAL® 60CD** (see table p. 20) and is the only compound in this series to be classified according to UL94 V0 at only 1.6 mm specimen thickness. A combination of **APYRAL® 200SM** and **APYRAL® 40CD** also increases the tensile strength in an EVA compound.



High fire resistance combined with good processability. LOI, MVR and UL94-V0 at 1.6 mm for EVA/EVM filled with 65 wt.-% **APYRAL**®

Combination of APYRAL® 40CD and APYRAL® 200SM • UL94 V0 at 1.6 mm • good processability

### APYRAL® 200SM

- at same loading LOI boost of 8 % 0<sub>2</sub>
- very good MVR
- UL94 V0 classification at only 1.6 mm thickness

# FRLS PVC compounds

## FRLS PVC compounds

Component		[phr]	[phr]	[phr]	
PVC, K = 70 (Evipol <sup>®</sup> SH 7020)		100	100	100	
Plasticizer, DINP		50	50	50	
Ca/Zn-Stabilisator (Bäropan R 8850KA-5)		5	5	5	
PE-Wax (A-C 6A)		0.5	0.5	0.5	
Flame retardant		5	50	50	
		ΑΤΟ	APYRAL® 40CD	APYRAL® 60CD	
Characteristic data					
Tensile Strength [MPa]		20	17	17.4	
Elongation at Break [%]		310	310	310	
LOI [% 0 <sub>2</sub> ]		20	30	30	

#### Examples for flame retardant and smoke suppressed PVC compounds

Compounding on lab dispersion kneader. Specimen made of compression moulded plaques

### Smoke suppressed PVC compounds

Plasticized PVC is still by far the most widespread material in the cable sector and is used for the production of insulations but mainly for cable sheaths. PVC intrinsically has relatively low flammability and is self extinguishing when a small flame is applied. A major disadvantage with almost all halogenated materials is the generated black smoke in the case of fire.

Plasticised PVC filled with **APYRAL**<sup>®</sup> develops considerably less smoke than conventional flame retarded PVC with antimony trioxide (ATO) at the same ignitability (LOI values).

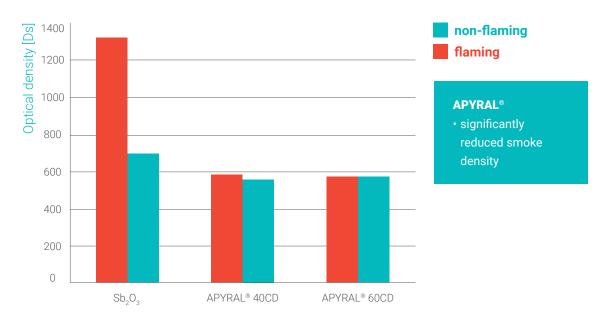
The table above shows an example of such a Fire Retardant Low Smoke PVC (FRLS-PVC). For **APYRAL® 40CD** and **APYRAL® 60CD** the characteristic values received are compared with a compound traditionally flame retarded using ATO. The mechanical properties of all compounds are very

good and easily fulfill all the requirements by common industry standards, easily.

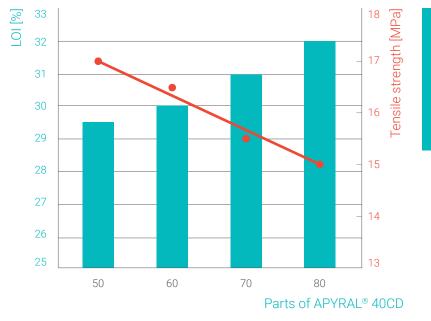
The diagram next page shows the maximum optical density of the four compounds listed in the table, received by smoke density measurements in accordance with ASTM E662.

**APYRAL**<sup>®</sup> filled compounds show reduced smoke density compared to ATO  $(Sb_2O_3)$  filled PVC at conditions resembling a smouldering fire (non flaming mode).

Much more considerable differences are recorded when a pilot flame is used (flaming mode, simulating an open fire). The compounds filled with **APYRAL**<sup>®</sup> metal hydrates have considerably reduced smoke values when an open fire breaks out (See more in our brochoure "Metal Hydrates for PVC").



Maximum smoke density according to ASTM E662 of plasticized PVC using different flame retardants



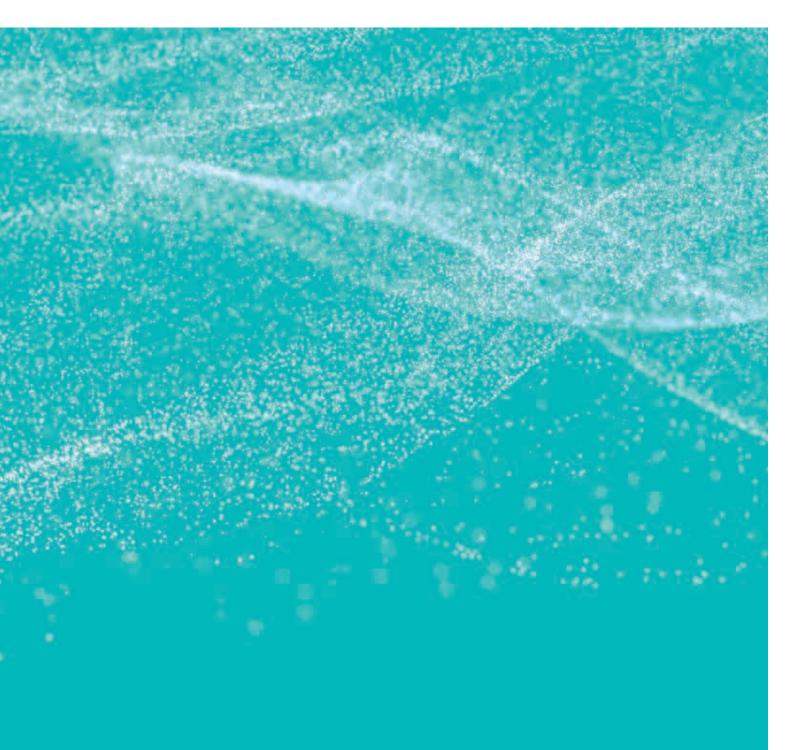
### APYRAL<sup>®</sup> 40CD

- $\boldsymbol{\cdot}$  LOI increases with
- loading level
- good tensile
- strength even up to 80 parts loading

LOI and tensile strength of plasticized PVC at increasing  $\textbf{APYRAL}^{\circledast}~\textbf{40CD}$  loading

The second diagram displays LOI and tensile strength for a plasticized PVC compound as a function of **APYRAL® 40CD** loading. LOI increases with increasing **APYRAL® 40CD** load.

The tensile strength of the compound drops, but remains at a very good level even up to 80 parts on 100 parts PVC and easily fulfils values required by common industry standards.



# Service for our customers

## Service for our customers

### Technical service development / production

Nabaltec AG develops new products and refines innovative products in close cooperation with our customers and raw material suppliers.

Here we use our own lab facilities as well as our excellent contacts to external test institutes and laboratories to offer our customers a wide range of service to support them in formulation development and test procedures.

The successful implementation of this development and the intensive customer consultations enable Nabaltec AG an interaction with our customers in a cooperative, responsible and innovative manner. This culminates in the development of high performance products at the customer as well as in our facility.

Additionally, we have the capacity to fashion tailor made products for special customer requirements and their highly sophisticated and demanding markets.

### **Laboratory services**

Our analysis centre is responsible for independent production control and quality offers laboratory services for customers intending to use our large analytical equipment.

With this excellent equipment we are able to execute analytic tests in the area of inorganic solids, trace elements and water quality.

The certification in accordance with DIN EN ISO 17025 confirms the high service standards of our lab.

We will gladly inform you about our capabilities.

## Nabaltec product portfolio

### **NABALOX®**

Aluminium oxides, for the production of ceramic, refractory and polishing products

### **APYRAL® AOH**

Boehmite, as flame retardant filler and functional filler

### **APYRAL®**

Aluminium hydroxides, as flame retardant and functional filler

### **GRANALOX®**

Ceramic bodies, for the production of engineering ceramics

### **NABACAST®**

Hydraulic, cement-free binder, based on α-alumina

### **SYMULOX®**

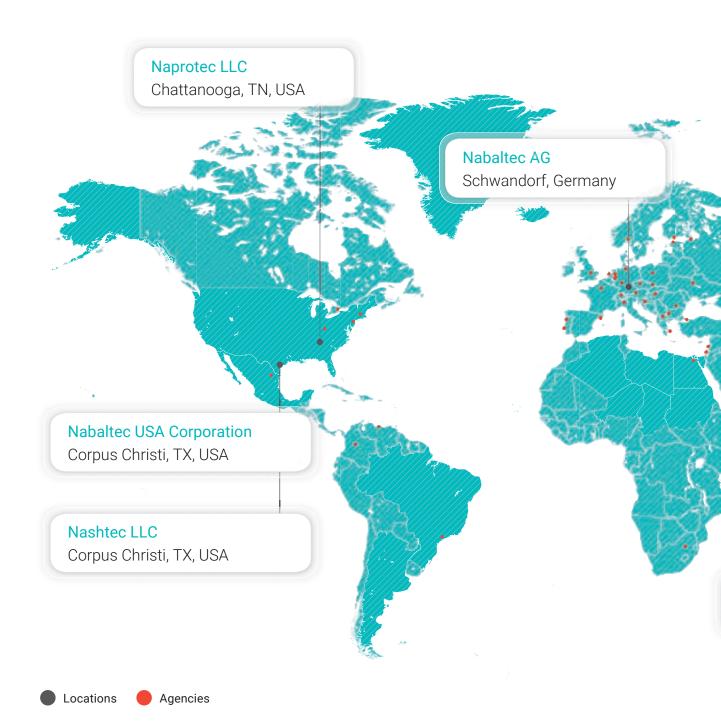
Synthetic sintered Mullite, for the production of e.g. refractory products

### **ACTILOX®**

Boehmite, as flame retardant filler and catalyst carrier



Visit us at our website www.nabaltec.de where you will find the latest company updates and recent versions of all available certificates free for download as PDF-documents.





1

Nabaltec (Shanghai) Trading Co., Ltd. Shanghai, China

### **Further information:**

### Nabaltec AG

P.O. Box 1860 · 92409 Schwandorf Phone +49 9431 53-0 www.nabaltec.de info@nabaltec.de

### **Customer Service**

Phone +49 9431 53 910 sales@nabaltec.de

### **Technical Service**

Phone +49 9431 53 920 tec-service@nabaltec.de

Nabaltec AG P.O. Box 1860 · 92409 Schwandorf Tel +49 9431 53-0 Fax +49 9431 61 557

www.nabaltec.de info@nabaltec.de

All data listed in this brochure are reference values and subject to production tolerance. These values are exclusive to the product description and no guarantee is placed on the properties. It remains the responsibility of the users to test the suitability of the product for their application.

02 / 2020